

baseline_dignosis

December 11, 2025

```
[1]: !pip install xgboost
```

Requirement already satisfied: xgboost in /opt/conda/lib/python3.11/site-packages (3.1.2)
Requirement already satisfied: numpy in /opt/conda/lib/python3.11/site-packages (from xgboost) (1.24.4)
Requirement already satisfied: nvidia-nccl-cu12 in /opt/conda/lib/python3.11/site-packages (from xgboost) (2.28.9)
Requirement already satisfied: scipy in /opt/conda/lib/python3.11/site-packages (from xgboost) (1.11.3)

```
[2]: !pip install xlswriter
```

Requirement already satisfied: xlswriter in /opt/conda/lib/python3.11/site-packages (3.2.9)

```
[1]: # -*- coding: utf-8 -*-
      """
      Baseline 3-class diagnostic model (AD / Mixed / Non-AD)
      - Label: Final_Diagnosis (already mapped at baseline)
      - Features: AD-signature MRI percent volumes (ROI/Total*100) + composites +
        ↪laterality + comorbidities + age + sex
      - CV: 5-fold grouped by BrcId
      - Metrics: macro-F1, balanced accuracy, OVR ROC-AUC, confusion matrix,
        ↪classification report
      - Explainability: XGBoost built-in TreeSHAP contributions (pred_contribs=True)
      """

      import re, json, numpy as np, pandas as pd
      from pathlib import Path

      # ----- CONFIG -----
      CSV_PATH = "Merge_data_with_Pre_Mini_Mental_Total.csv"
      OUT_DIR = Path("baseline3_out"); OUT_DIR.mkdir(parents=True, exist_ok=True)
      RANDOM_SEED = 42

      # ----- LOAD -----
      df = pd.read_csv(CSV_PATH, low_memory=False)
```

```

print(f"[] Loaded dataset: {len(df)} rows")

# ----- LABEL: Final_Diagnosis → AD/Mixed/Non-AD -----
def _norm_label(s):
    if pd.isna(s): return ""
    s = str(s).strip()
    s = s.replace("'", "").replace('"', "").replace("-", "-").replace("_", "-")
    s = re.sub(r"\s+", " ", s)
    return s.lower()

AD_WHITELIST = {
    "alzheimer's disease",
    "alzheimer's disease - early onset",
    "alzheimer's disease - late onset",
    "alzheimer's disease - unspecified",
}
MIXED_WHITELIST = {
    "mixed dementia",
}
NONAD_WHITELIST = {
    "mild cognitive impairment",
    "unspecified dementia",
    "vascular dementia, unspecified",
    "dementia in parkinson disease",
    "dementia in other specified diseases classified elsewhere",
    "other",
}

# ----- " MMSE" -----

def keep_mmse_closest_to_scan(df_in: pd.DataFrame) -> pd.DataFrame:
    """
        ScanID    (BrcId, Scan_Date) ,

        MMSE

    """
    df1 = df_in.copy()

    # MMSE

    mmse_date_cols = [c for c in df1.columns if "mmse" in c.lower() and "date" in c.lower()]

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scan_date_cols = [c for c in df1.columns if "scan" in c.lower() and "date" in c.lower()]

if not scan_date_cols or not mmse_date_cols:

    #

    print("[warn] no explicit MMSE / Scan date columns found; skip 'closest_
to scan' filtering.")

    return df1

scan_col = scan_date_cols[0]

mmse_col = mmse_date_cols[0]

df1[scan_col] = pd.to_datetime(df1[scan_col], errors="coerce")

df1[mmse_col] = pd.to_datetime(df1[mmse_col], errors="coerce")

#

mask = df1[scan_col].notna() & df1[mmse_col].notna()

df1["_mmse_dt_diff"] = np.where(

    mask,

    (df1[mmse_col] - df1[scan_col]).abs().dt.days,

    np.inf,

)

# key ScanID (BrcId, Scan_Date)

if "ScanID" in df1.columns and df1["ScanID"].notna().any():

    key_cols = ["ScanID"]

elif "BrcId" in df1.columns:

    key_cols = ["BrcId", scan_col]

else:

```

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        print("[warn] no ScanID/BrcId columns; skip 'closest to scan' filtering.
↪")

        df1.drop(columns="_mmse_dt_diff", inplace=True)

        return df1

df1 = (

    df1.sort_values(key_cols + ["_mmse_dt_diff"])

        .drop_duplicates(subset=key_cols, keep="first")

        .drop(columns="_mmse_dt_diff")

)

print(f"[] Kept MMSE closest-to-scan per {key_cols}: {len(df1)} rows")

return df1

def map_diag3_exact(raw_label):
    s = _norm_label(raw_label)
    if s in MIXED_WHITELIST: return "Mixed"
    if s in AD_WHITELIST:    return "AD"
    if s in NONAD_WHITELIST: return "Non-AD"
    # conservative fallback
    if re.search(r"\balzheimer", s): return "AD"
    if "mixed" in s:                return "Mixed"
    return "Non-AD"

assert "Final_Diagnosis" in df.columns, " Final_Diagnosis "
df["target_diag"] = df["Final_Diagnosis"].apply(map_diag3_exact)

#
keep = df["target_diag"].isin(["AD", "Mixed", "Non-AD"])
df = df.loc[keep].reset_index(drop=True)
# ---      MRI  MMSE      ---

def dedup_per_scan(df_in: pd.DataFrame) -> pd.DataFrame:

    df1 = df_in.copy()

    #      "      "

    prefer_cols = [

```

```

        "Pre_Mini_Mental_Total", "Post_Mini_Mental_Total",

        "ACE_date_pre", "ACE_date_post",

        "HONOS_total_score_pre", "HONOS_total_score_post"

    ]

    prefer_cols = [c for c in prefer_cols if c in df1.columns]

    comp = df1[prefer_cols].notna().sum(axis=1) if prefer_cols else pd.
↳Series(0, index=df1.index)

    #   Scan_Date   datetime   "   "

    if "Scan_Date" in df1.columns:

        df1["Scan_Date"] = pd.to_datetime(df1["Scan_Date"], errors="coerce")

    #   A   ScanID

    if "ScanID" in df1.columns and df1["ScanID"].notna().any():

        df1 = (df1.assign(_comp=comp)

                #   ScanID   "   "

                .sort_values(["ScanID", "_comp", "Scan_Date"],
↳ascending=[True, False, True])

                .drop_duplicates(subset=["ScanID"], keep="first")

                .drop(columns=["_comp"]))

        return df1

    #   B   ScanID   (BrcId, Scan_Date)   "   "

    keys = [c for c in ["BrcId", "Scan_Date"] if c in df1.columns]

    if keys:

        #

        df1 = (df1.assign(_comp=comp)

                .sort_values(keys + ["_comp"], ascending=[True, True, False]))

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        .drop_duplicates(subset=keys, keep="first")

        .drop(columns=["_comp"]))

    return df1

# C      BrcId

if "BrcId" in df1.columns:

    df1 = (df1.assign(_comp=comp)

        .sort_values(["BrcId", "Scan_Date", "_comp"], ascending=[True,
↪True, False])

        .drop_duplicates(subset=["BrcId"], keep="first")

        .drop(columns=["_comp"]))

    return df1

#

return df1
df = keep_mmse_closest_to_scan(df)
df = dedup_per_scan(df)

print("Rows after per-scan de-dup:", len(df))

# ----- MRI percent features (ROI/Total*100) -----
if "Total" not in df.columns:
    raise KeyError(" Total MRI ")
print("[i] target label distribution:\n", df["target_diag"].
↪value_counts(dropna=False))

```

```

[] Loaded dataset: 16854 rows
[] Kept MMSE closest-to-scan per ['ScanID']: 3310 rows
Rows after per-scan de-dup: 3310
[i] target label distribution:
target_diag
AD      1992
Non-AD   673
Mixed    645
Name: count, dtype: int64

```

```

[2]: # ROI Brain_*
roi_code_groups = {
    # Medial temporal
    "Hipp": ["048_Right Hippocampus", "049_Left Hippocampus"],
    "Ent": ["117_Right Ent entorhinal area", "118_Left Ent entorhinal_
↪area", "117_Right Ent", "118_Left Ent"],
    "PHG": ["171_Right PHG parahippocampal gyrus", "172_Left PHG_
↪parahippocampal gyrus", "171_Right PHG", "172_Left PHG"],
    # Lateral/basal temporal
    "MTG": ["155_Right MTG middle temporal gyrus", "156_Left MTG middle_
↪temporal gyrus", "155_Right MTG", "156_Left MTG"],
    "ITG": ["133_Right ITG inferior temporal gyrus", "134_Left ITG inferior_
↪temporal gyrus", "133_Right ITG", "134_Left ITG"],
    "STG": ["201_Right STG superior temporal gyrus", "202_Left STG superior_
↪temporal gyrus", "201_Right STG", "202_Left STG"],
    "FuG": ["123_Right FuG fusiform gyrus", "124_Left FuG fusiform_
↪gyrus", "123_Right FuG", "124_Left FuG"],
    "TMP": ["203_Right TMP temporal pole", "204_Left TMP temporal_
↪pole", "203_Right TMP", "204_Left TMP"],
    # Medial parietal
    "PCgG": ["167_Right PCgG posterior cingulate gyrus", "168_Left PCgG_
↪posterior cingulate gyrus", "167_Right PCgG", "168_Left PCgG"],
    "PCu": ["169_Right PCu precuneus", "170_Left PCu precuneus", "169_Right_
↪PCu", "170_Left PCu"],
    # Posterior parietal/occipital (PCA axis)
    "SPL": ["199_Right SPL superior parietal lobule", "200_Left SPL superior_
↪parietal lobule", "199_Right SPL", "200_Left SPL"],
    "SOG": ["197_Right SOG superior occipital gyrus", "198_Left SOG superior_
↪occipital gyrus", "197_Right SOG", "198_Left SOG"],
    "MOG": ["145_Right MOG middle occipital gyrus", "146_Left MOG middle_
↪occipital gyrus", "145_Right MOG", "146_Left MOG"],
    "IOG": ["129_Right IOG inferior occipital gyrus", "130_Left IOG inferior_
↪occipital gyrus", "129_Right IOG", "130_Left IOG"],
    # Ventricles
    "LatVent": ["052_Right Lateral Ventricle", "053_Left Lateral Ventricle"],
    "InfLat": ["050_Right Inf Lat Vent", "051_Left Inf Lat Vent"],
}

def pct_col(col):
    if col not in df.columns: return None
    v = pd.to_numeric(df[col], errors="coerce")
    total = pd.to_numeric(df["Total"], errors="coerce").replace(0, np.nan)
    return (v/total)*100.0

mri_cols = []
mri_pct_frames = []

```

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# ROI
for grp, cols in roi_code_groups.items():
    for c in cols:
        s = pct_col(c)
        if s is not None:
            new = f"{c}__pct"
            mri_cols.append(new); mri_pct_frames.append(s.rename(new))

# Brain_*
brain_cols = [c for c in df.columns if re.match(r"^Brain_", str(c))]
for c in brain_cols:
    s = pct_col(c)
    if s is not None:
        new = f"{c}__pct"
        mri_cols.append(new); mri_pct_frames.append(s.rename(new))

X_mri = pd.concat(mri_pct_frames, axis=1) if mri_pct_frames else pd.
↳DataFrame(index=df.index)
print(f"[i] MRI percent features: {X_mri.shape[1]}")

# ----- composites -----
def sum_cols(names, outname):
    s = pd.Series(0.0, index=df.index)
    for n in names:
        if n in X_mri.columns: s = s.add(X_mri[n].fillna(0), fill_value=0)
    return s.rename(outname)

hipp_cols = [c for c in X_mri.columns if "Hippocampus__pct" in c]
ent_cols = [c for c in X_mri.columns if (" Ent" in c or "Ent " in c) and
↳ "__pct" in c]
phg_cols = [c for c in X_mri.columns if "PHG" in c and "__pct" in c]
lat_temp_cols = [c for c in X_mri.columns if any(k in c for k in [" MTG ", " ITG_
↳ ", " STG ", " FuG ", " TMP "]) and "__pct" in c]
mp_cols = [c for c in X_mri.columns if any(k in c for k in ["PCgG", "PCu"])]
↳ and "__pct" in c]
post_cols = [c for c in X_mri.columns if any(k in c for k in
↳ ["SPL", "SOG", "MOG", "IOG"]) and "__pct" in c]
vent_cols = [c for c in X_mri.columns if any(k in c for k in ["Lateral_
↳ Ventricle__pct", "Inf Lat Vent__pct"])]

comp_frames = [
    sum_cols(hipp_cols+ent_cols+phg_cols, "MTL_total_pct"),
    sum_cols(lat_temp_cols, "Temporal_lateral_total_pct"),
    sum_cols(mp_cols, "Medial_parietal_total_pct"),
    sum_cols(post_cols, "Posterior_total_pct"),

```



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        sum_cols(vent_cols, "Ventricles_total_pct"),
    ]
    X_comp = pd.concat(comp_frames, axis=1)

    # ----- laterality indices -----
    def laterality_pair(L_contains, R_contains, outname):
        Lc = [c for c in X_mri.columns if L_contains in c]
        Rc = [c for c in X_mri.columns if R_contains in c]
        if not Lc or not Rc: return None
        L = X_mri[Lc[0]]; R = X_mri[Rc[0]]
        denom = (L+R).replace(0, np.nan)
        return ((L-R)/denom).rename(outname)

    li_frames = []
    for l, r, out in [
        ("168_Left PCgG", "167_Right PCgG", "LI_PCgG"),
        ("170_Left PCu", "169_Right PCu", "LI_PCu"),
        ("156_Left MTG", "155_Right MTG", "LI_MTG"),
        ("134_Left ITG", "133_Right ITG", "LI_ITG"),
        ("202_Left STG", "201_Right STG", "LI_STG"),
    ]:
        s = laterality_pair(l, r, out)
        if s is not None: li_frames.append(s)

    X_li = pd.concat(li_frames, axis=1) if li_frames else pd.DataFrame(index=df.
        ↪index)

    # MRI
    X_mri_all = pd.concat([X_mri, X_comp, X_li], axis=1)

    # ----- comorbidities -----
    comorb_cols = [
        ↪
        ↪ "asthma", "falls", "hypertension", "Cerebrovascular_accident", "Epilepsy", "Diabetes_mellitus",
        ↪
        ↪ "Chronic_kidney_disease", "Psoriasis", "Parkinsons_disease", "Multiple_sclerosis", "Eczema",
        ↪
        ↪ "Hypertensive_disorder", "Transient_ischemic_attack", "Migraine", "Chronic_obstructive_lung_di
        ↪
        ↪ "Arthritis", "Heart_failure", "Asthma", "Ischemic_heart_disease", "Inflammatory_bowel_disease",
        ↪
        ↪ "Atrial_fibrillation", "Chronic_liver_disease", "Chronic_sinusitis", "Coronary_arterioscleroti
    ]
    comorb_cols = [c for c in comorb_cols if c in df.columns]
    X_comorb = df[comorb_cols].apply(pd.to_numeric, errors="coerce").fillna(0).
        ↪clip(0,1)

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# ----- age & sex -----
def make_age_series(d):
    if "age_at_scan_date" in d.columns:
        s = pd.to_numeric(d["age_at_scan_date"], errors="coerce")
        if s.notna().mean()>0.5: return s
    if "age_years" in d.columns:
        s = pd.to_numeric(d["age_years"], errors="coerce")
        if s.notna().mean()>0.5: return s
    sdt = pd.to_datetime(d.get("Scan_Date"), errors="coerce")
    dob = pd.to_datetime(d.get("cleaneddateofbirth"), errors="coerce")
    if sdt.notna().any() and dob.notna().any():
        return (sdt - dob).dt.days/365.25
    return pd.Series(np.nan, index=d.index)

age = make_age_series(df).rename("age")
sex = df.get("Gender_ID", pd.Series(np.nan, index=df.index)).astype(str)
# ----- MMSE baseline -----
if "Pre_Mini_Mental_Total" in df.columns:
    mmse = pd.to_numeric(df["Pre_Mini_Mental_Total"], errors="coerce").
    ↪rename("MMSE")
else:
    #
    mmse = pd.Series(np.nan, index=df.index, name="MMSE")
    print("[warn] Column 'Pre_Mini_Mental_Total' not found, MMSE is all NaN.")

# ----- final X, y -----
X_num = pd.concat([X_mri_all, X_comorb, age, mmse], axis=1)
X_cat = pd.DataFrame({"Gender_ID": sex})
X_all = pd.concat([X_num, X_cat], axis=1)

y = df["target_diag"].astype(str)
groups = df.get("BrcId", pd.Series(np.arange(len(df))))).astype(str)

print(f"[i] X_num={X_num.shape}, X_cat={X_cat.shape}; y={y.value_counts().
    ↪to_dict()}")
def classify_feature(col: str) -> str:

    if col in X_mri_all.columns:

        return "MRI_pct"

    if 'WMH' in col.upper():

        return "WMH"

    if col in X_comorb.columns:

```

```

        return "comorbidity"

    if col == "age":

        return "age"

    if col == "Gender_ID":

        return "sex"

    # MMSE Mini-Mental

    if "MMSE" in col.upper() or "MINI_MENTAL" in col.upper():

        return "MMSE"

    return "other"

feat_stage1 = pd.DataFrame({
    "feature_name": X_all.columns,
    "block": [classify_feature(c) for c in X_all.columns],
})
feat_stage1.to_csv(OUT_DIR/"feature_list_stage1.csv", index=False)
print("[ ] Saved Stage-1 feature list →", OUT_DIR/"feature_list_stage1.csv")

```

```

[i] MRI percent features: 65
[i] X_num=(3310, 101), X_cat=(3310, 1); y={'AD': 1992, 'Non-AD': 673, 'Mixed': 645}
[ ] Saved Stage-1 feature list → baseline3_out/feature_list_stage1.csv

```

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[3]: # ----- preprocessing -----
from sklearn.compose import ColumnTransformer
from sklearn.preprocessing import OneHotEncoder, StandardScaler
from sklearn.impute import SimpleImputer
from sklearn.pipeline import Pipeline

num_cols = X_num.columns.tolist()
cat_cols = ["Gender_ID"]

try:
    ohe = OneHotEncoder(handle_unknown="ignore", sparse_output=False)
except TypeError: # sklearn<1.2
    from sklearn.preprocessing import OneHotEncoder as OHE
    ohe = OHE(handle_unknown="ignore", sparse=False)

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preprocess = ColumnTransformer(
    transformers=[
        ("num", Pipeline([("imp", SimpleImputer(strategy="median")),
                           ("sc", StandardScaler())]), num_cols),
        ("cat", Pipeline([("imp", SimpleImputer(strategy="most_frequent")),
                           ("oh", OneHotEncoder())]), cat_cols),
    ],
    remainder="drop",
)

```

```

[4]: # ----- CV splits (grouped by BrcId) -----
from sklearn.model_selection import StratifiedKFold
try:
    from sklearn.model_selection import StratifiedGroupKFold
    cv = StratifiedGroupKFold(n_splits=5, shuffle=True,
    ↪ random_state=RANDOM_SEED)
    splits = list(cv.split(X_all, y, groups=groups))
except Exception:
    # approximate grouped stratification
    g = df.groupby("BrcId")["target_diag"].first().reset_index()
    skf = StratifiedKFold(n_splits=5, shuffle=True, random_state=RANDOM_SEED)
    splits = []
    for tr, te in skf.split(g["BrcId"], g["target_diag"]):
        tr_mask = groups.isin(g.loc[tr, "BrcId"])
        te_mask = groups.isin(g.loc[te, "BrcId"])
        splits.append((X_all.index[tr_mask], X_all.index[te_mask]))

```

```

[5]: # ----- model: XGB classifier (with integer label encoding) ↪
    ↪ -----

import xgboost as xgb

from sklearn.metrics import classification_report, f1_score,
    ↪ balanced_accuracy_score, roc_auc_score, confusion_matrix

#      /

labels = ["AD", "Mixed", "Non-AD"]

lab2int = {lab:i for i, lab in enumerate(labels)}

int2lab = {i:lab for lab,i in lab2int.items()}

# y ↪

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y_int = y.map(lab2int).astype(int)

# BrcId    CV

from sklearn.model_selection import StratifiedKFold

try:

    from sklearn.model_selection import StratifiedGroupKFold

    cv = StratifiedGroupKFold(n_splits=5, shuffle=True,
    ↪random_state=RANDOM_SEED)

    splits = list(cv.split(X_all, y_int, groups=groups))

except Exception:

    #

    g = df.groupby("BrcId")["target_diag"].first().reset_index()

    g_int = g["target_diag"].map(lab2int).astype(int)

    skf = StratifiedKFold(n_splits=5, shuffle=True, random_state=RANDOM_SEED)

    splits = []

    for tr, te in skf.split(g["BrcId"], g_int):

        tr_mask = groups.isin(g.loc[tr, "BrcId"])

        te_mask = groups.isin(g.loc[te, "BrcId"])

        splits.append((X_all.index[tr_mask], X_all.index[te_mask]))

# XGB

xgb_clf = xgb.XGBClassifier(

    objective="multi:softprob",

    num_class=3,

    max_depth=3, learning_rate=0.05, n_estimators=800,

    subsample=0.8, colsample_bytree=0.8,

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    reg_lambda=1.0, random_state=RANDOM_SEED, n_jobs=-1,

    eval_metric="mlogloss"
)

pipe = Pipeline([("prep", preprocess), ("clf", xgb_clf)])

#   sample_weight           y_int
class_counts = y.value_counts()

class_weight = {k: class_counts.max()/v for k,v in class_counts.items()}

sample_weight = y.map(class_weight).values #   y_int

y_true_all_str, y_pred_all_str = [], []

y_prob_all = []

for fold, (tr_idx, te_idx) in enumerate(splits):

    pipe.fit(X_all.iloc[tr_idx], y_int.iloc[tr_idx],
    ↪clf__sample_weight=sample_weight[tr_idx])

    y_pred_int = pipe.predict(X_all.iloc[te_idx]) #

    y_pred_str = pd.Series(y_pred_int).map(int2lab).values #

    y_true_str = y.iloc[te_idx].values #

    y_prob = pipe.predict_proba(X_all.iloc[te_idx]) #   (n, 3) ↪
    ↪0,1,2 ↪ AD, Mixed, Non-AD

    y_true_all_str.extend(y_true_str)

    y_pred_all_str.extend(y_pred_str)

    y_prob_all.append(y_prob)

y_true_all_str = np.array(y_true_all_str)

y_pred_all_str = np.array(y_pred_all_str)

y_prob_all = np.vstack(y_prob_all)

```

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# 89

print("\n=== Classification report (pooled folds) ===")

print(classification_report(y_true_all_str, y_pred_all_str, labels=labels,
    ↪digits=3))

macro_f1 = f1_score(y_true_all_str, y_pred_all_str, average="macro")

bacc      = balanced_accuracy_score(y_true_all_str, y_pred_all_str)

print("Macro-F1:", round(macro_f1, 3))

print("Balanced Acc:", round(bacc, 3))

# OVR ROC-AUC   labels

try:

    y_true_bin = pd.get_dummies(pd.Categorical(y_true_all_str,
    ↪categories=labels))

    ovr_auc = roc_auc_score(y_true_bin.values, y_prob_all, average="macro",
    ↪multi_class="ovr")

    print("OVR ROC-AUC:", round(ovr_auc, 3))

except Exception as e:

    print("OVR ROC-AUC skipped:", e)

#

cm = confusion_matrix(y_true_all_str, y_pred_all_str, labels=labels)

cm_df = pd.DataFrame(cm, index=[f"true_{l}" for l in labels],
    ↪columns=[f"pred_{l}" for l in labels])

cm_df.to_csv(OUT_DIR/"confusion_matrix.csv", index=True)

#

metrics = pd.DataFrame([

    "macro_f1": float(macro_f1),

    "balanced_acc": float(bacc)

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    })

metrics.to_csv(OUT_DIR/"cv_metrics_summary_xgb.csv", index=False)

print("\nSaved confusion matrix and metrics to:", OUT_DIR.resolve())

```

```

=== Classification report (pooled folds) ===

```

	precision	recall	f1-score	support
AD	0.693	0.692	0.693	1992
Mixed	0.348	0.319	0.333	645
Non-AD	0.531	0.575	0.552	673
accuracy			0.596	3310
macro avg	0.524	0.529	0.526	3310
weighted avg	0.593	0.596	0.594	3310

```

Macro-F1: 0.526
Balanced Acc: 0.529
OVR ROC-AUC: 0.721

```

Saved confusion matrix and metrics to: /home/jovyan/baseline3_out

```

[6]: print("Unique target labels:", sorted(y.unique()))

#      '-AD'

y = y.replace({"-AD":"AD", " AD":"AD", "Non AD":"Non-AD"})

```

Unique target labels: ['AD', 'Mixed', 'Non-AD']

```

[7]: # ----- fit on full data for built-in TreeSHAP contributions
      ↪ (robust for 2D/3D) -----

import xgboost as xgb

import numpy as np

import pandas as pd

import re

from pathlib import Path

```



```

# 1)

pipe.fit(X_all, y_int, clf__sample_weight=sample_weight)

prep = pipe.named_steps["prep"]

model = pipe.named_steps["clf"]

Xmat = prep.transform(X_all)

if hasattr(Xmat, "toarray"):

    Xmat = Xmat.toarray()

n_features = Xmat.shape[1]

# 2) XGBoost built-in TreeSHAP contributions

dmat = xgb.DMatrix(Xmat)

contrib = model.get_booster().predict(dmat, pred_contribs=True)

labels = ["AD", "Mixed", "Non-AD"]

C_expected = len(labels)

# 3)

#   A)  $(n, (p+1)*C)$ 

#   B)  $(n, p+1, C)$ 

#   C)  $(n, C, p+1)$ 

if contrib.ndim == 2:

    n, pc = contrib.shape

    #    $p+1 \otimes C$ 

    if pc % C_expected == 0:

        p_plus1 = pc // C_expected

        arr = contrib.reshape(n, C_expected, p_plus1)      #  $(n, C, p+1)$ 

```

```

    phi = arr[:, :, :p_plus1-1]                                # bias → (n, C, p)

    mean_abs = np.abs(phi).mean(axis=(0, 1))                    # (p,)

else:

    #      n_features      p_plus1

    p_plus1 = n_features + 1

    C = pc // p_plus1

    arr = contrib.reshape(n, C, p_plus1)

    phi = arr[:, :, :p_plus1-1]

    mean_abs = np.abs(phi).mean(axis=(0, 1))

elif contrib.ndim == 3:

    n, a, b = contrib.shape

    #      B: (n, p+1, C)

    if b == C_expected:

        p_plus1, C = a, b

        p_keep = min(n_features, p_plus1 - 1)

        phi = contrib[:, :p_keep, :]                            # (n, p_keep, C)

        mean_abs = np.abs(phi).mean(axis=(0, 2))                # (p_keep,)

        #      C: (n, C, p+1)

    elif a == C_expected:

        C, p_plus1 = a, b

        p_keep = min(n_features, p_plus1 - 1)

        phi = contrib[:, :, :p_keep]                            # (n, C, p_keep)

        mean_abs = np.abs(phi).mean(axis=(0, 1))                # (p_keep,)

    else:

```

```

        #         →      n_features

        p_plus1 = b

        p_keep = min(n_features, p_plus1 - 1)

        phi = contrib[:, :, :p_keep]

        mean_abs = np.abs(phi).mean(axis=(0, 1))

    else:

        raise ValueError(f"Unexpected pred_contribs ndim={contrib.ndim}")

# 4)

try:

    feat_names = prep.get_feature_names_out()

    feat_names = [re.sub(r"^(num|cat)_", "", f) for f in feat_names]

except Exception:

    feat_names = [f"f{i}" for i in range(mean_abs.shape[0])]

if len(feat_names) != len(mean_abs):

    k = min(len(feat_names), len(mean_abs))

    feat_names = feat_names[:k]

    mean_abs = mean_abs[:k]

imp_df = pd.DataFrame({"feature": feat_names,
                       "mean_abs_contrib": mean_abs
                       }).sort_values("mean_abs_contrib", ascending=False)

imp_df.to_csv(OUT_DIR / "feature_importance_xgb_contribs.csv", index=False)

print("Saved:", OUT_DIR / "feature_importance_xgb_contribs.csv")

print("Top 20 features:\n", imp_df.head(50).to_string(index=False))

```

Saved: baseline3_out/feature_importance_xgb_contribs.csv

Top 20 features:

	feature	mean_abs_contrib
	MMSE	0.221973
	051_Left Inf Lat Vent__pct	0.173523
	050_Right Inf Lat Vent__pct	0.155038
	049_Left Hippocampus__pct	0.097018
	Brain_Left_0_Frontal_0__pct	0.087328
	age	0.083767
	LI_MTG	0.080075
	117_Right Ent entorhinal area__pct	0.078581
	Brain_Left_0_Frontal_1__pct	0.073360
	Cerebrovascular_accident	0.069932
	Brain_Left_0_Frontal_99__pct	0.066800
	Brain_Right_2_Parietal_99__pct	0.065687
	Brain_Left_4_DeepNuclei_10__pct	0.062143
	LI_ITG	0.061802
	048_Right Hippocampus__pct	0.060998
	124_Left FuG fusiform gyrus__pct	0.058376
	200_Left SPL superior parietal lobule__pct	0.056807
	118_Left Ent entorhinal area__pct	0.054667
	falls	0.052513
	Brain_Right_4_DeepNuclei_9__pct	0.052345
	Brain_Right_4_DeepNuclei_99__pct	0.050890
	052_Right Lateral Ventricle__pct	0.050052
	171_Right PHG parahippocampal gyrus__pct	0.049731
	198_Left SOG superior occipital gyrus__pct	0.049172
	129_Right IOG inferior occipital gyrus__pct	0.049129
	167_Right PCgG posterior cingulate gyrus__pct	0.048793
	204_Left TMP temporal pole__pct	0.047621
	LI_PCu	0.047296
	199_Right SPL superior parietal lobule__pct	0.045774
	Brain_Right_99_99__pct	0.045458
	130_Left IOG inferior occipital gyrus__pct	0.044505
	203_Right TMP temporal pole__pct	0.043074
	133_Right ITG inferior temporal gyrus__pct	0.042924
	Brain_Left_2_Parietal_99__pct	0.042364
	Brain_Right_0_Frontal_99__pct	0.042310
	134_Left ITG inferior temporal gyrus__pct	0.042220
	LI_STG	0.041659
	172_Left PHG parahippocampal gyrus__pct	0.040939
	170_Left PCu precuneus__pct	0.040922
	Brain_Left_3_Occipital_7__pct	0.040886
	201_Right STG superior temporal gyrus__pct	0.039591
	Brain_Left_0_Frontal_2__pct	0.039308
	Parkinsons_disease	0.038678
	Brain_Left_2_Parietal_6__pct	0.038052
	053_Left Lateral Ventricle__pct	0.036623
	145_Right MOG middle occipital gyrus__pct	0.035981

169_Right PCu precuneus__pct	0.034564
Brain_Right_3_Occipital_7__pct	0.034561
Brain_Left_4_DeepNuclei_9__pct	0.034169
Brain_Left_3_Occipital_8__pct	0.033788

```
[8]: pip install xgboost
```

```
Requirement already satisfied: xgboost in /opt/conda/lib/python3.11/site-
packages (3.1.2)
Requirement already satisfied: numpy in /opt/conda/lib/python3.11/site-packages
(from xgboost) (1.24.4)
Requirement already satisfied: nvidia-nccl-cu12 in
/opt/conda/lib/python3.11/site-packages (from xgboost) (2.28.9)
Requirement already satisfied: scipy in /opt/conda/lib/python3.11/site-packages
(from xgboost) (1.11.3)
Note: you may need to restart the kernel to use updated packages.
```

```
[9]: # =====

# HIERARCHICAL 3-CLASS MODEL

# Stage-1: AD vs Others

# Stage-2: (Among non-AD) Mixed vs Non-AD using WMH + vascular features

# =====

from pathlib import Path

import numpy as np, pandas as pd, re, json

from sklearn.compose import ColumnTransformer

from sklearn.preprocessing import OneHotEncoder, StandardScaler

from sklearn.impute import SimpleImputer

from sklearn.pipeline import Pipeline

from sklearn.metrics import (classification_report, f1_score,

                             balanced_accuracy_score, roc_auc_score,
                             ↪confusion_matrix)

from sklearn.model_selection import StratifiedKFold

import xgboost as xgb
```

```

HIER_DIR = Path("hier_out"); HIER_DIR.mkdir(parents=True, exist_ok=True)

RANDOM_SEED = 42

# -----

def make_ohe():

    try:

        return OneHotEncoder(handle_unknown="ignore", sparse_output=False)

    except TypeError:

        from sklearn.preprocessing import OneHotEncoder as OHE

        return OHE(handle_unknown="ignore", sparse=False)

def build_preprocess(num_cols, cat_cols):

    return ColumnTransformer(

        transformers=[

            ("num", Pipeline([("imp", SimpleImputer(strategy="median")),

                               ("sc", StandardScaler())])), num_cols),

            ("cat", Pipeline([("imp", SimpleImputer(strategy="most_frequent")),

                               ("oh", make_ohe())])), cat_cols),

        ],

        remainder="drop",

    )

def grouped_stratified_splits(df_index, y_series, groups_series, n_splits=5,
    ↪seed=42):

    """      StratifiedGroupKFold      """

    try:

        from sklearn.model_selection import StratifiedGroupKFold

```

```

        cv = StratifiedGroupKFold(n_splits=n_splits, shuffle=True,
↳random_state=seed)

        return list(cv.split(df_index, y_series, groups_series))

    except Exception:

        g = pd.DataFrame({"BrcId": groups_series, "y": y_series}).
↳groupby("BrcId").first().reset_index()

        skf = StratifiedKFold(n_splits=n_splits, shuffle=True,
↳random_state=seed)

        splits = []

        for tr, te in skf.split(g["BrcId"], g["y"]):

            tr_mask = groups_series.isin(g.loc[tr, "BrcId"])

            te_mask = groups_series.isin(g.loc[te, "BrcId"])

            splits.append((df_index[tr_mask], df_index[te_mask]))

        return splits

def metrics_binary(y_true, y_pred, y_prob):

    rep = classification_report(y_true, y_pred, digits=3)

    f1 = f1_score(y_true, y_pred, average="binary", pos_label=1)

    bacc= balanced_accuracy_score(y_true, y_pred)

    try:

        auc = roc_auc_score(y_true, y_prob[:,1])

    except Exception:

        auc = np.nan

    return rep, dict(f1=f1, balanced_acc=bacc, auc=auc)

# -----

#      df      df["target_diag"]  {AD, Mixed, Non-AD}

```

```

assert set(df["target_diag"].unique()) <= {"AD", "Mixed", "Non-AD"}

groups = df.get("BrcId", pd.Series(np.arange(len(df))).astype(str))

index_all = df.index

# ===== Stage-1: AD vs Others =====

print("\n[Stage-1] AD vs Others")

# y1: 1=AD, 0=Others

y1 = (df["target_diag"] == "AD").astype(int)

#          X_all MRI% +      +      /          "      "

if 'X_all' in globals():

    X1 = X_all.copy()

else:

    #          __pct   age_at_scan_date/age_years  Gender_ID

    pct_cols = [c for c in df.columns if c.endswith("__pct")]

    num_extra = []

    if "age_at_scan_date" in df.columns:

        num_extra.append("age_at_scan_date")

    elif "age_years" in df.columns:

        num_extra.append("age_years")

    cat_cols = ["Gender_ID"] if "Gender_ID" in df.columns else []

    #

    comorb_cols = [c for c in df.columns if c in [

        ↵
        "hypertension", "Hypertensive_disorder", "Diabetes_mellitus", "Atrial_fibrillation",

```



```

        ↪ "Ischemic_heart_disease", "Cerebrovascular_accident", "Transient_ischemic_attack",

        ↪ "Heart_failure", "Coronary_arteriosclerosis", "falls", "Chronic_obstructive_lung_disease"

    ]]

    X1 = pd.concat([df[pct_cols].apply(pd.to_numeric, errors="coerce"),

                    df[num_extra].apply(pd.to_numeric, errors="coerce"),

                    df[comorb_cols].apply(pd.to_numeric, errors="coerce"),

                    df[cat_cols]], axis=1)

    X1 = X1.fillna(0)

#   +

num_cols1 = X1.select_dtypes(include=[np.number]).columns.tolist()

cat_cols1 = [c for c in X1.columns if c not in num_cols1]

pre1 = build_preprocess(num_cols1, cat_cols1)

clf1 = xgb.XGBClassifier(

    objective="binary:logistic",

    max_depth=3, learning_rate=0.05, n_estimators=600,

    subsample=0.8, colsample_bytree=0.8,

    reg_lambda=1.0, random_state=RANDOM_SEED, n_jobs=-1,

    eval_metric="logloss"

)

pipe1 = Pipeline([("prep", pre1), ("clf", clf1)])

#   CV

splits1 = grouped_stratified_splits(index_all, y1, groups, n_splits=5, ↪
    ↪ seed=RANDOM_SEED)

```

```

y1_true, y1_pred, y1_prob = [], [], []

for tr_idx, te_idx in splits1:

    pipe1.fit(X1.iloc[tr_idx], y1.iloc[tr_idx])

    y1_pred.extend(pipe1.predict(X1.iloc[te_idx]))

    y1_prob.extend(pipe1.predict_proba(X1.iloc[te_idx]) )

y1_true = y1.values

y1_pred = np.array(y1_pred)

y1_prob = np.vstack(y1_prob)

rep1, m1 = metrics_binary(y1_true, y1_pred, y1_prob)

(Path(HIER_DIR/"stage1_report.txt")).write_text(rep1, encoding="utf-8")

with open(HIER_DIR/"stage1_metrics.json","w") as f: json.dump(m1, f, indent=2)

print(rep1)

print("[Stage-1] F1:", round(m1["f1"],3), "BalancedAcc:",
    ↳round(m1["balanced_acc"],3), "AUC:", round(m1["auc"],3))

# ===== Stage-2: Mixed vs Non-AD AD =====

print("\n[Stage-2] Mixed vs Non-AD (among non-AD cases)")

nonad_mask = (df["target_diag"] != "AD")

df2 = df.loc[nonad_mask].reset_index(drop=True)

groups2 = df2.get("BrcId", pd.Series(np.arange(len(df2)))) .astype(str)

# y2: 1=Mixed, 0=Non-AD

y2 = (df2["target_diag"] == "Mixed").astype(int)

# ---- Stage-2 WMH + vascular burden (+ age/sex ) ----

# 1) WMH 'wmh'

wmh_cols = [c for c in df2.columns if re.search(r"wmh", c, flags=re.I)]

```

```

# intracranial TIV/ICV/Total

icv_col = None

for cand in ["tiv", "ICV", "icv", "Total", "total"]:

    if cand in df2.columns:

        icv_col = cand

        break

X2_blocks = []

if wmh_cols:

    wmh = df2[wmh_cols].apply(pd.to_numeric, errors="coerce")

    if icv_col:

        icv = pd.to_numeric(df2[icv_col], errors="coerce").replace(0, np.nan)

        for c in wmh_cols:

            X2_blocks.append( (wmh[c]/icv).rename(f"{c}_pct_icv") )

        else:

            for c in wmh_cols:

                X2_blocks.append(wmh[c].rename(c))

    else:

        print("[Stage-2] WMH columns not found; proceeding without WMH.")

# 2) Vascular comorbidities

vasc_cols = [

    ↵
    ↪ "hypertension", "Hypertensive_disorder", "Diabetes_mellitus", "Atrial_fibrillation",

    ↵
    ↪ "Ischemic_heart_disease", "Cerebrovascular_accident", "Transient_ischemic_attack",

    "Heart_failure", "Coronary_arteriosclerosis"

```

```

]

vasc_cols = [c for c in vasc_cols if c in df2.columns]

X2_blocks.append( df2[vasc_cols].apply(pd.to_numeric, errors="coerce").
    ↪fillna(0).clip(0,1) )

# 3) Age / Sex

if "age_at_scan_date" in df2.columns:

    X2_blocks.append( pd.to_numeric(df2["age_at_scan_date"], errors="coerce").
        ↪rename("age") )

elif "age_years" in df2.columns:

    X2_blocks.append( pd.to_numeric(df2["age_years"], errors="coerce").
        ↪rename("age") )

if "Gender_ID" in df2.columns:

    X2_blocks.append( df2["Gender_ID"].astype(str).rename("Gender_ID") )

X2 = pd.concat(X2_blocks, axis=1)

X2 = X2.fillna(0)

num_cols2 = X2.select_dtypes(include=[np.number]).columns.tolist()

cat_cols2 = [c for c in X2.columns if c not in num_cols2]

pre2 = build_preprocess(num_cols2, cat_cols2)

clf2 = xgb.XGBClassifier(

    objective="binary:logistic",

    max_depth=3, learning_rate=0.05, n_estimators=600,

    subsample=0.8, colsample_bytree=0.8,

    reg_lambda=1.0, random_state=RANDOM_SEED, n_jobs=-1,

    eval_metric="logloss"

)

```

```

pipe2 = Pipeline([("prep", pre2), ("clf", clf2)])

# =====

from sklearn.model_selection import RandomizedSearchCV
from sklearn.metrics import make_scorer, balanced_accuracy_score

# -----
# 1)  XGBoost + Pipeline
# -----
base_clf2 = xgb.XGBClassifier(
    objective="binary:logistic",
    eval_metric="logloss",
    random_state=RANDOM_SEED,
    n_jobs=-1,
    tree_method="hist",  #
)

pipe2 = Pipeline([
    ("prep", pre2),
    ("clf", base_clf2),
])

# + CV list
splits2 = list(
    grouped_stratified_splits(df2.index, y2, groups2,
                             n_splits=5, seed=RANDOM_SEED)
)

# Mixed=1, Non-AD=0
pos = int(y2.sum())
neg = int(len(y2) - pos)
ratio = neg / pos if pos > 0 else 1.0
print(f"[Stage-2|WMH] Mixed={pos}, Non-AD={neg}, neg/pos={ratio:.2f}")

# -----
# 2)
# -----
param_dist2 = {
    "clf__max_depth": [2, 3, 4, 5],
    "clf__learning_rate": [0.01, 0.02, 0.03, 0.05, 0.08],
    "clf__n_estimators": [300, 500, 700, 900],
    "clf__subsample": [0.7, 0.85, 1.0],
    "clf__colsample_bytree": [0.6, 0.8, 1.0],
    "clf__min_child_weight": [1, 3, 5, 7],
    "clf__gamma": [0.0, 0.5, 1.0],

```

```

    "clf__reg_lambda":      [0.1, 1.0, 5.0],
    "clf__reg_alpha":      [0.0, 0.1, 0.5],
    #           Mixed
    "clf__scale_pos_weight": [1.0, ratio*0.8, ratio, ratio*1.2],
}

# -----
# 3)   Balanced accuracy = (recall0 + recall1)/2
#   =>   Mixed   Non-AD   recall
# -----
bal_acc_scorer = make_scorer(balanced_accuracy_score)

search2 = RandomizedSearchCV(
    estimator=pipe2,
    param_distributions=param_dist2,
    n_iter=60,                      #           30
    scoring=bal_acc_scorer,
    cv=splits2,
    n_jobs=-1,
    random_state=RANDOM_SEED,
    verbose=2,
    refit=True,
)

print("\n[Stage-2|WMH]   RandomizedSearchCV   balanced accuracy ...")
search2.fit(X2, y2)

print("\n[Stage-2|WMH]   CV balanced accuracy:",
      round(search2.best_score_, 3))
print("[Stage-2|WMH]      :")
for k, v in search2.best_params_.items():
    print("    ", k, "=", v)

for k, v in search2.best_params_.items():

    print("    ", k, "=", v)

#           pipe2 / clf2

pipe2 = search2.best_estimator_

clf2 = pipe2.named_steps["clf"]

#

import json

```

```

(HIER_DIR / "stage2_best_params.json").write_text(
    json.dumps(search2.best_params_, indent=2), encoding="utf-8"
)

splits2 = grouped_stratified_splits(df2.index, y2, groups2, n_splits=5,
    ↪seed=RANDOM_SEED)

y2_true, y2_pred, y2_prob = [], [], []

for tr_idx, te_idx in splits2:
    pipe2.fit(X2.iloc[tr_idx], y2.iloc[tr_idx])
    y2_pred.extend( pipe2.predict(X2.iloc[te_idx]) )
    y2_prob.extend( pipe2.predict_proba(X2.iloc[te_idx]) )

y2_true = y2.values
y2_pred = np.array(y2_pred)
y2_prob = np.vstack(y2_prob)

rep2, m2 = metrics_binary(y2_true, y2_pred, y2_prob)

(Path(HIER_DIR/"stage2_report.txt")).write_text(rep2, encoding="utf-8")

with open(HIER_DIR/"stage2_metrics.json","w") as f: json.dump(m2, f, indent=2)

print(rep2)

print("[Stage-2] F1:", round(m2["f1"],3), "BalancedAcc:",
    ↪round(m2["balanced_acc"],3), "AUC:", round(m2["auc"],3))

# =====
# =====

#     Stage-2

def build_stage2_matrix(df_slice: pd.DataFrame, ref_cols: list[str] | None =
    ↪None) -> pd.DataFrame:

    blocks = []

```

```

# WMH

wmh_cols = [c for c in df_slice.columns if re.search(r"wmh", c, flags=re.I)]

icv_col = None

for cand in ["tiv", "ICV", "icv", "Total", "total"]:

    if cand in df_slice.columns:

        icv_col = cand; break

if wmh_cols:

    wmh = df_slice[wmh_cols].apply(pd.to_numeric, errors="coerce")

    if icv_col:

        icv = pd.to_numeric(df_slice[icv_col], errors="coerce").replace(0,
↪np.nan)

        for c in wmh_cols:

            blocks.append( (wmh[c]/icv).rename(f"{c}_pct_icv") )

        else:

            for c in wmh_cols:

                blocks.append(wmh[c].rename(c))

# vascular burden

vasc_cols = [

    ↪

↪"hypertension", "Hypertensive_disorder", "Diabetes_mellitus", "Atrial_fibrillation",

    ↪

↪"Ischemic_heart_disease", "Cerebrovascular_accident", "Transient_ischemic_attack",

    ↪

    ↪"Heart_failure", "Coronary_arteriosclerosis"

]

vasc_cols = [c for c in vasc_cols if c in df_slice.columns]

```



```

if vasc_cols:

    blocks.append( df_slice[vasc_cols].apply(pd.to_numeric,
errors="coerce").fillna(0).clip(0,1) )

    # age / sex

    if "age_at_scan_date" in df_slice.columns:

        blocks.append( pd.to_numeric(df_slice["age_at_scan_date"],
errors="coerce").rename("age") )

    elif "age_years" in df_slice.columns:

        blocks.append( pd.to_numeric(df_slice["age_years"], errors="coerce").
rename("age") )

    if "Gender_ID" in df_slice.columns:

        blocks.append( df_slice["Gender_ID"].astype(str).rename("Gender_ID") )

    #

    if blocks:

        Xtmp = pd.concat(blocks, axis=1).fillna(0)

    else:

        Xtmp = pd.DataFrame(index=df_slice.index) #

    #

    if ref_cols is not None:

        Xtmp = Xtmp.reindex(columns=ref_cols, fill_value=0)

    return Xtmp

print("\n[Hier] Assemble final 3-class predictions ...")

# Stage-1       $P(AD)$ 

pipe1.fit(X1, y1)

p1 = pipe1.predict_proba(X1)[: , 1] #  $P(AD)$ 

```

```

is_ad = (p1 >= 0.5)                                #      0.6/0.7

pred_final = np.array([""]*len(df), dtype=object)

pred_final[is_ad] = "AD"

# Stage-2      "      non-AD      "      Mixed vs Non-AD

idx_pred_nonad = np.where(~is_ad)[0]

df_pred_nonad = df.iloc[idx_pred_nonad].reset_index(drop=True)

#      pipe2

X2_pred = build_stage2_matrix(df_pred_nonad, ref_cols=X2.columns)

p2_pred = pipe2.predict_proba(X2_pred)[: , 1]      # P(Mixed / predicted non-AD)

pred_final[idx_pred_nonad] = np.where(p2_pred >= 0.5, "Mixed", "Non-AD")

#      vs

from sklearn.metrics import confusion_matrix

cm3 = confusion_matrix(df["target_diag"], pred_final,
↳ labels=["AD", "Mixed", "Non-AD"])

cm3_df = pd.DataFrame(cm3, index=[ "true_AD", "true_Mixed", "true_Non-AD" ],
                        columns=["pred_AD", "pred_Mixed", "pred_Non-AD"])

cm3_df.to_csv(HIER_DIR/"final_confusion_matrix.csv", index=True)

#

out_pred = pd.DataFrame({

    "BrcId": df.get("BrcId", pd.Series(np.nan, index=df.index)),

    "true_diag": df["target_diag"],

    "P_AD": p1,

    "final_pred": pred_final

})

```

```
#      non-AD

out_pred.loc[~is_ad, "P_Mixed_given_nonAD"] = p2_pred

out_pred.to_csv(HIER_DIR/"preds_hierarchical.csv", index=False)

print(f"[] Saved hierarchical outputs to: {HIER_DIR.resolve()}")

print(cm3_df)
```

```
[Stage-1] AD vs Others
           precision    recall  f1-score   support

      0         0.394        0.296        0.338        1318
      1         0.600        0.698        0.645        1992

 accuracy                   0.538        3310
 macro avg         0.497        0.497        0.492        3310
 weighted avg         0.518        0.538        0.523        3310
```

```
[Stage-1] F1: 0.645 BalancedAcc: 0.497 AUC: 0.497
```

```
[Stage-2] Mixed vs Non-AD (among non-AD cases)
```

```
[Stage-2] WMH columns not found; proceeding without WMH.
```

```
[Stage-2|WMH] Mixed=645, Non-AD=673, neg/pos=1.04
```

```
[Stage-2|WMH] RandomizedSearchCV balanced accuracy ...
```

```
Fitting 5 folds for each of 60 candidates, totalling 300 fits
```

```
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
```

```
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
```

```
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.2s
```

```
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
```

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
 clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=900,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
 clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
 clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=500,
 clf__reg_alpha=0.5, clf__reg_lambda=5.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.01,
 clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=500,
 clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
 clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
 clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
 clf__reg_alpha=0.5, clf__reg_lambda=5.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
 clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
 clf__reg_alpha=0.5, clf__reg_lambda=5.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.03,
 clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
 clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
 clf__reg_alpha=0.1, clf__reg_lambda=5.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
 clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
 clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
 clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=700,
 clf__reg_alpha=0.0, clf__reg_lambda=5.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.03,
 clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
 clf__reg_alpha=0.1, clf__reg_lambda=5.0,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=1.0, clf__learning_rate=0.01, clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900, clf__reg_alpha=0.1, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=1.0, clf__learning_rate=0.01, clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900, clf__reg_alpha=0.1, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08, clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=300, clf__reg_alpha=0.0, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05, clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700, clf__reg_alpha=0.1, clf__reg_lambda=0.1, clf__scale_pos_weight=0.8347286821705426, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05, clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300, clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05, clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300, clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.02, clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=900, clf__reg_alpha=0.5, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.02, clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=900, clf__reg_alpha=0.5, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05, clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=300, clf__reg_alpha=0.1, clf__reg_lambda=0.1, clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05, clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=300, clf__reg_alpha=0.1, clf__reg_lambda=0.1, clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.08, clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300, clf__reg_alpha=0.5, clf__reg_lambda=5.0, clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.08, clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,

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clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.01,

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clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,

```

```

clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,

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clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=

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0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.02,

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clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,

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clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=500,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=700,

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clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.05,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s

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[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,

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clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.3s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,

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clf__reg_alpha=0.5, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s

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[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,

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clf__reg_alpha=0.1, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=900,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.3s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,

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clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
0.2s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.2s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s

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[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
 clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
 0.2s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
 clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=900,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time=
 0.2s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
 clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=500,
 clf__reg_alpha=0.1, clf__reg_lambda=0.1,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
 clf__max_depth=3, clf__min_child_weight=5, clf__n_estimators=500,
 clf__reg_alpha=0.1, clf__reg_lambda=0.1,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
 clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
 clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
 clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
 clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=1.0, clf__learning_rate=0.03,
 clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=900,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
 clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=300,
 clf__reg_alpha=0.1, clf__reg_lambda=0.1,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.08,
 clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.08,

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clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=1.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=500,
clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.01,
clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=2, clf__min_child_weight=7, clf__n_estimators=300,
clf__reg_alpha=0.5, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=7, clf__n_estimators=500,
clf__reg_alpha=0.1, clf__reg_lambda=5.0,
clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.2s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=700,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,

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clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=900,
clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
clf__subsample=0.85; total time= 0.1s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
clf__subsample=0.7; total time= 0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.02,
clf__max_depth=4, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.0, clf__reg_lambda=5.0,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=300,
clf__reg_alpha=0.1, clf__reg_lambda=1.0,
clf__scale_pos_weight=1.2520930232558138, clf__subsample=1.0; total time= 0.1s

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[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
 clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
 clf__max_depth=5, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
 clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=900,
 clf__reg_alpha=0.1, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.3s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.05,
 clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=900,
 clf__reg_alpha=0.1, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
 clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=300,
 clf__reg_alpha=0.1, clf__reg_lambda=0.1,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
 clf__max_depth=5, clf__min_child_weight=1, clf__n_estimators=300,
 clf__reg_alpha=0.1, clf__reg_lambda=0.1,
 clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.08,
 clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.08,
 clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
 clf__reg_alpha=0.5, clf__reg_lambda=0.1,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.0, clf__learning_rate=0.01,
 clf__max_depth=4, clf__min_child_weight=3, clf__n_estimators=300,
 clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
 clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.02,
 clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=500,
 clf__reg_alpha=0.0, clf__reg_lambda=5.0, clf__scale_pos_weight=1.0,
 clf__subsample=1.0; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
 clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
 clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=700,
 clf__reg_alpha=0.0, clf__reg_lambda=1.0,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.85; total time=
 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.01,
 clf__max_depth=4, clf__min_child_weight=1, clf__n_estimators=300,
 clf__reg_alpha=0.0, clf__reg_lambda=0.1,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.01,
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 clf__reg_alpha=0.0, clf__reg_lambda=0.1,
 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
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 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
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 clf__scale_pos_weight=0.8347286821705426, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.08,
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 clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
 clf__subsample=0.85; total time= 0.2s

[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.08,
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 clf__reg_alpha=0.1, clf__reg_lambda=1.0,
 clf__scale_pos_weight=1.2520930232558138, clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.8, clf__gamma=1.0, clf__learning_rate=0.01,
 clf__max_depth=2, clf__min_child_weight=5, clf__n_estimators=900,
 clf__reg_alpha=0.5, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
 clf__subsample=0.85; total time= 0.1s

[CV] END clf__colsample_bytree=1.0, clf__gamma=0.0, clf__learning_rate=0.01,
 clf__max_depth=2, clf__min_child_weight=1, clf__n_estimators=700,
 clf__reg_alpha=0.1, clf__reg_lambda=1.0, clf__scale_pos_weight=1.0,
 clf__subsample=0.7; total time= 0.1s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
 clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
 clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
 clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.6, clf__gamma=0.5, clf__learning_rate=0.08,
 clf__max_depth=5, clf__min_child_weight=3, clf__n_estimators=900,
 clf__reg_alpha=0.0, clf__reg_lambda=0.1, clf__scale_pos_weight=1.0,
 clf__subsample=0.7; total time= 0.2s

[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
 clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=700,
 clf__reg_alpha=0.1, clf__reg_lambda=0.1,


```

clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.0, clf__learning_rate=0.03,
clf__max_depth=5, clf__min_child_weight=7, clf__n_estimators=700,
clf__reg_alpha=0.1, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.2s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=0.8, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=3, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=0.85; total time=
0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s
[CV] END clf__colsample_bytree=1.0, clf__gamma=0.5, clf__learning_rate=0.03,
clf__max_depth=2, clf__min_child_weight=3, clf__n_estimators=900,
clf__reg_alpha=0.0, clf__reg_lambda=0.1,
clf__scale_pos_weight=1.0434108527131782, clf__subsample=1.0; total time= 0.1s

```

[Stage-2|WMH] CV balanced accuracy: 0.581

[Stage-2|WMH] :

```

clf__subsample = 0.7
clf__scale_pos_weight = 1.0434108527131782
clf__reg_lambda = 1.0
clf__reg_alpha = 0.5
clf__n_estimators = 900
clf__min_child_weight = 7
clf__max_depth = 2
clf__learning_rate = 0.03
clf__gamma = 1.0
clf__colsample_bytree = 0.6
clf__subsample = 0.7
clf__scale_pos_weight = 1.0434108527131782
clf__reg_lambda = 1.0
clf__reg_alpha = 0.5
clf__n_estimators = 900
clf__min_child_weight = 7
clf__max_depth = 2
clf__learning_rate = 0.03
clf__gamma = 1.0
clf__colsample_bytree = 0.6

```

	precision	recall	f1-score	support
0	0.519	0.594	0.554	673
1	0.502	0.426	0.461	645
accuracy			0.512	1318
macro avg	0.511	0.510	0.508	1318
weighted avg	0.511	0.512	0.509	1318

[Stage-2] F1: 0.461 BalancedAcc: 0.51 AUC: 0.505

[Hier] Assemble final 3-class predictions ...

[] Saved hierarchical outputs to: /home/jovyan/hier_out

	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1946	17	29
true_Mixed	137	302	206
true_Non-AD	52	226	395

```
[10]: # =====
# HIERARCHICAL 3-CLASS MODEL (with WMH merge)
# Stage-1: AD vs Others
# Stage-2: (Among non-AD) Mixed vs Non-AD using WMH + vascular features
# =====

from pathlib import Path
import numpy as np, pandas as pd, re, json
from sklearn.compose import ColumnTransformer
from sklearn.preprocessing import OneHotEncoder, StandardScaler
from sklearn.impute import SimpleImputer
from sklearn.pipeline import Pipeline
from sklearn.metrics import (classification_report, f1_score,
                             balanced_accuracy_score, roc_auc_score,
                             ↪confusion_matrix)
from sklearn.model_selection import StratifiedKFold
import xgboost as xgb

HIER_DIR = Path("hier_out"); HIER_DIR.mkdir(parents=True, exist_ok=True)
RANDOM_SEED = 42
```

```
[11]: # ----- 0) WMH -----

WMH_FILE = "WMH.xlsx" # ← WMH .xlsx / .xls / .csv

def clean_brcid(s: pd.Series) -> pd.Series:

    return (s.astype(str).str.strip())
```

```

        .str.replace(r"\.0$", "", regex=True)

        .str.replace(r"^[0-9]", "", regex=True))

def load_wmh_unique(path: str | Path) -> pd.DataFrame:

    p = str(path)

    if p.lower().endswith((".xlsx", ".xls")):

        w = pd.read_excel(p, dtype={"BrcId": str})

    else:

        w = pd.read_csv(p, dtype={"BrcId": str}, low_memory=False)

    #

    colmap = {"BrcId": "BrcId", "WMH(77)": "WMH_77", "Intracranial-volume": "ICV"}

    w = w.rename(columns=colmap)

    #      BrcId

    need = [c for c in ["BrcId", "WMH_77", "ICV"] if c in w.columns]

    w = w[need].copy()

    w["BrcId"] = clean_brcid(w["BrcId"])

    #

    for c in ["WMH_77", "ICV"]:

        if c in w.columns:

            w[c] = pd.to_numeric(w[c], errors="coerce")

    # --      BrcId      -- #

    agg_rules = {}

    if "WMH_77" in w.columns:

        agg_rules["WMH_77"] = "max"      #      "mean"      "sum"

```

```

if "ICV" in w.columns:

    agg_rules["ICV"] = "median"    # ICV    median/mean

    w_unique = w.groupby("BrcId", as_index=False).agg(agg_rules)

    return w_unique

try:

    wmh_df = load_wmh_unique(WMH_FILE)

    #   df   BrcId

    if "BrcId" in df.columns:

        df["BrcId"] = clean_brcid(df["BrcId"])

    #

    df = df.merge(wmh_df, on="BrcId", how="left", validate="m:1")

    #   WMH

    df["WMH_77"] = pd.to_numeric(df["WMH_77"], errors="coerce")

    df["ICV"]     = pd.to_numeric(df["ICV"],     errors="coerce").replace(0, np.
↳nan)

    df["wmh_icv_pct"] = (df["WMH_77"] / df["ICV"]).replace([np.inf,-np.inf], np.
↳nan) * 100.0

    df["wmh_log_icv"] = np.log1p( (df["WMH_77"] / df["ICV"]).replace([np.
↳inf,-np.inf], np.nan) )

    df["wmh_missing"] = df["WMH_77"].isna().astype(int)

    print(f"[WMH] merged unique: {wmh_df.shape[0]} rows; non-missing WMH:
↳{(~df['WMH_77'].isna()).sum()}")

except Exception as e:

    print("[WMH] merge failed; proceeding without WMH. Error:", e)

    df["wmh_icv_pct"] = np.nan

    df["wmh_log_icv"] = np.nan

```

```
df["wmh_missing"] = 1
```

[WMH] merged unique: 5833 rows; non-missing WMH: 3088

```
[12]: # ----- 1) -----
def make_ohe():
    try:
        return OneHotEncoder(handle_unknown="ignore", sparse_output=False)
    except TypeError:
        from sklearn.preprocessing import OneHotEncoder as OHE
        return OHE(handle_unknown="ignore", sparse=False)

def build_preprocess(num_cols, cat_cols):
    return ColumnTransformer(
        transformers=[
            ("num", Pipeline([("imp", SimpleImputer(strategy="median")),
                              ("sc", StandardScaler())]), num_cols),
            ("cat", Pipeline([("imp", SimpleImputer(strategy="most_frequent")),
                              ("oh", make_ohe())]), cat_cols),
        ],
        remainder="drop",
    )

def grouped_stratified_splits(df_index, y_series, groups_series, n_splits=5,
    ↪seed=42):
    try:
        from sklearn.model_selection import StratifiedGroupKFold
        cv = StratifiedGroupKFold(n_splits=n_splits, shuffle=True,
    ↪random_state=seed)
        return list(cv.split(df_index, y_series, groups_series))
    except Exception:
        g = pd.DataFrame({"BrcId": groups_series, "y": y_series}).
    ↪groupby("BrcId").first().reset_index()
        skf = StratifiedKFold(n_splits=n_splits, shuffle=True,
    ↪random_state=seed)
        splits = []
        for tr, te in skf.split(g["BrcId"], g["y"]):
            tr_mask = groups_series.isin(g.loc[tr, "BrcId"])
            te_mask = groups_series.isin(g.loc[te, "BrcId"])
            splits.append((df_index[tr_mask], df_index[te_mask]))
        return splits

def metrics_binary(y_true, y_pred, y_prob, pos_label=1):
    rep = classification_report(y_true, y_pred, digits=3)
    f1 = f1_score(y_true, y_pred, average="binary", pos_label=pos_label)
```

```

bacc= balanced_accuracy_score(y_true, y_pred)
try:
    #
    auc = roc_auc_score(y_true, y_prob[:,1 if y_prob.shape[1]>1 else 0])
except Exception:
    auc = np.nan
return rep, dict(f1=f1, balanced_acc=bacc, auc=auc)

```

```

[13]: # ----- 2) Stage-1 AD vs Others -----
print("\n[Stage-1] AD vs Others")

y1 = (df["target_diag"] == "AD").astype(int)

#      X_all MRI%+ + age/sex
if 'X_all' in globals():
    X1 = X_all.copy()
else:
    pct_cols = [c for c in df.columns if c.endswith("__pct")]
    num_extra = []
    if "age_at_scan_date" in df.columns: num_extra.append("age_at_scan_date")
    elif "age_years" in df.columns: num_extra.append("age_years")
    comorb_cols = [c for c in [
        ↪ "hypertension", "Hypertensive_disorder", "Diabetes_mellitus", "Atrial_fibrillation",
        ↪ "Ischemic_heart_disease", "Cerebrovascular_accident", "Transient_ischemic_attack",
        ↪ "Heart_failure", "Coronary_arteriosclerosis", "falls", "Chronic_obstructive_lung_disease"
    ] if c in df.columns]
    cat_cols = ["Gender_ID"] if "Gender_ID" in df.columns else []
    X1 = pd.concat([df[pct_cols].apply(pd.to_numeric, errors="coerce"),
                    df[num_extra].apply(pd.to_numeric, errors="coerce"),
                    df[comorb_cols].apply(pd.to_numeric, errors="coerce"),
                    df[cat_cols]], axis=1).fillna(0)

num_cols1 = X1.select_dtypes(include=[np.number]).columns.tolist()
cat_cols1 = [c for c in X1.columns if c not in num_cols1]
pre1 = build_preprocess(num_cols1, cat_cols1)
clf1 = xgb.XGBClassifier(
    objective="binary:logistic",
    max_depth=3, learning_rate=0.05, n_estimators=600,
    subsample=0.8, colsample_bytree=0.8, reg_lambda=1.0,
    random_state=RANDOM_SEED, n_jobs=-1, eval_metric="logloss"
)
pipe1 = Pipeline([("prep", pre1), ("clf", clf1)])

```

```

groups = df.get("BrcId", pd.Series(np.arange(len(df))))).astype(str)
splits1 = grouped_stratified_splits(df.index, y1, groups, n_splits=5,
    ↪seed=RANDOM_SEED)

y1_pred, y1_prob = [], []
for tr_idx, te_idx in splits1:
    pipe1.fit(X1.iloc[tr_idx], y1.iloc[tr_idx])
    y1_pred.extend(pipe1.predict(X1.iloc[te_idx]))
    y1_prob.extend(pipe1.predict_proba(X1.iloc[te_idx]))
y1_pred = np.array(y1_pred); y1_prob = np.vstack(y1_prob)

rep1, m1 = metrics_binary(y1.values, y1_pred, y1_prob, pos_label=1)
(HIER_DIR/"stage1_report.txt").write_text(rep1, encoding="utf-8")
with open(HIER_DIR/"stage1_metrics.json", "w") as f: json.dump(m1, f, indent=2)
print(rep1)
print("[Stage-1] F1:", round(m1["f1"],3), "BalancedAcc:",
    ↪round(m1["balanced_acc"],3), "AUC:", round(m1["auc"],3))

# ----- 3) Stage-2 Mixed vs Non-AD AD -----
print("\n[Stage-2] Mixed vs Non-AD (among non-AD cases)")

nonad_mask = (df["target_diag"] != "AD")
df2 = df.loc[nonad_mask].reset_index(drop=True)
groups2 = df2.get("BrcId", pd.Series(np.arange(len(df2))))).astype(str)
y2 = (df2["target_diag"] == "Mixed").astype(int)

def build_stage2_matrix(df_slice: pd.DataFrame, ref_cols: list[str] | None =
    ↪None) -> pd.DataFrame:
    blocks = []
    # WMH features
    cols_to_use = []
    for c in ["wmh_icv_pct", "wmh_log_icv", "wmh_missing"]:
        if c in df_slice.columns:
            cols_to_use.append(c)
    if cols_to_use:
        blocks.append( df_slice[cols_to_use].apply(pd.to_numeric,
    ↪errors="coerce").fillna(0) )
    else:
        print("[Stage-2] WMH columns not found; proceeding without WMH.")

    # vascular burden
    vasc_cols = [
        ↪
        ↪"hypertension", "Hypertensive_disorder", "Diabetes_mellitus", "Atrial_fibrillation",
        ↪
        ↪"Ischemic_heart_disease", "Cerebrovascular_accident", "Transient_ischemic_attack",

```

```

        "Heart_failure", "Coronary_arteriosclerosis"
    ]
    vasc_cols = [c for c in vasc_cols if c in df_slice.columns]
    if vasc_cols:
        blocks.append( df_slice[vasc_cols].apply(pd.to_numeric,
errors="coerce").fillna(0).clip(0,1) )

    # age / sex
    if "age_at_scan_date" in df_slice.columns:
        blocks.append( pd.to_numeric(df_slice["age_at_scan_date"],
errors="coerce").rename("age") )
    elif "age_years" in df_slice.columns:
        blocks.append( pd.to_numeric(df_slice["age_years"], errors="coerce").
rename("age") )
    if "Gender_ID" in df_slice.columns:
        blocks.append( df_slice["Gender_ID"].astype(str).rename("Gender_ID") )

    Xtmp = pd.concat(blocks, axis=1) if blocks else pd.DataFrame(index=df_slice.
index)
    Xtmp = Xtmp.fillna(0)

    if ref_cols is not None:
        #
        Xtmp = Xtmp.reindex(columns=ref_cols, fill_value=0)
    return Xtmp

X2 = build_stage2_matrix(df2)
num_cols2 = X2.select_dtypes(include=[np.number]).columns.tolist()
cat_cols2 = [c for c in X2.columns if c not in num_cols2]

# ---- Stage-2 ----
feat_stage2 = pd.DataFrame({"feature_name": X2.columns})
feat_stage2.to_csv(HIER_DIR/"feature_list_stage2.csv", index=False)
print("[ ] Saved Stage-2 feature list ->", HIER_DIR/"feature_list_stage2.csv")

pre2 = build_preprocess(num_cols2, cat_cols2)

clf2 = xgb.XGBClassifier(
    objective="binary:logistic",
    max_depth=3, learning_rate=0.05, n_estimators=600,
    subsample=0.8, colsample_bytree=0.8,
    reg_lambda=1.0, random_state=RANDOM_SEED, n_jobs=-1,
    eval_metric="logloss"
)
pipe2 = Pipeline([("prep", pre2), ("clf", clf2)])

```



```

splits2 = grouped_stratified_splits(df2.index, y2, groups2, n_splits=5,
    ↪seed=RANDOM_SEED)

y2_pred, y2_prob = [], []
for tr_idx, te_idx in splits2:
    pipe2.fit(X2.iloc[tr_idx], y2.iloc[tr_idx])
    y2_pred.extend( pipe2.predict(X2.iloc[te_idx]) )
    y2_prob.extend( pipe2.predict_proba(X2.iloc[te_idx]) )

y2_pred = np.array(y2_pred)
y2_prob = np.vstack(y2_prob)

rep2, m2 = metrics_binary(y2.values, y2_pred, y2_prob, pos_label=1)
(HIER_DIR/"stage2_report.txt").write_text(rep2, encoding="utf-8")
with open(HIER_DIR/"stage2_metrics.json", "w") as f: json.dump(m2, f, indent=2)
print(rep2)
print("[Stage-2] F1:", round(m2["f1"],3), "BalancedAcc:",
    ↪round(m2["balanced_acc"],3), "AUC:", round(m2["auc"],3))
#from sklearn.model_selection import RandomizedSearchCV

#from sklearn.metrics import make_scorer, balanced_accuracy_score

# -----

# 1)  XGBoost + Pipeline

# -----

#base_clf2 = xgb.XGBClassifier(

#     objective="binary:logistic",
#
#     eval_metric="logloss",

#     random_state=RANDOM_SEED,

#     n_jobs=-1,

#     tree_method="hist",    #

#)

#pipe2 = Pipeline([

#     ("prep", pre2),
#
#     ("clf", base_clf2),

```

```

#])

# + CV list

#splits2 = list(

#   grouped_stratified_splits(df2.index, y2, groups2,

#                               n_splits=5, seed=RANDOM_SEED)

#)

#   Mixed=1, Non-AD=0

#pos = int(y2.sum())

#neg = int(len(y2) - pos)

#ratio = neg / pos if pos > 0 else 1.0

#print(f"[Stage-2/WMH] Mixed={pos}, Non-AD={neg}, neg/pos={ratio:.2f}")

# -----

# 2)

# -----

#param_dist2 = {

#   "clf__max_depth":      [2, 3, 4, 5],

#   "clf__learning_rate":  [0.01, 0.02, 0.03, 0.05, 0.08],

#   "clf__n_estimators":   [300, 500, 700, 900],

#   "clf__subsample":      [0.7, 0.85, 1.0],

#   "clf__colsample_bytree": [0.6, 0.8, 1.0],

#   "clf__min_child_weight": [1, 3, 5, 7],

#   "clf__gamma":          [0.0, 0.5, 1.0],

#   "clf__reg_lambda":     [0.1, 1.0, 5.0],

```

```

#   "clf__reg_alpha":      [0.0, 0.1, 0.5],
#   #                   Mixed
#   "clf__scale_pos_weight": [1.0, ratio*0.8, ratio, ratio*1.2],
#}

# -----

# 3)   Balanced accuracy = (recall0 + recall1)/2

#   =>   Mixed   Non-AD   recall

# -----

#bal_acc_scorer = make_scorer(balanced_accuracy_score)

#search2 = RandomizedSearchCV(

#   estimator=pipe2,

#   param_distributions=param_dist2,

#   n_iter=60,                      #   30

#   scoring=bal_acc_scorer,

#   cv=splits2,

#   n_jobs=-1,

#   random_state=RANDOM_SEED,

#   verbose=2,

#   refit=True,

#)

#print("\n[Stage-2/WMH]   RandomizedSearchCV   balanced accuracy ...")

#search2.fit(X2, y2)

#print("\n[Stage-2/WMH]   CV balanced accuracy:",

#       round(search2.best_score_, 3))

```

```

#print("[Stage-2|WMH]      :")

#for k, v in search2.best_params_.items():

#    print("    ", k, "=", v)

#        Stage-2

#pipe2 = search2.best_estimator_

#clf2 = pipe2.named_steps["clf"]

# -----

# 4)    splits2    CV    threshold=0.5

# -----

y2_true, y2_pred, y2_prob = [], [], []

for tr_idx, te_idx in splits2:

    pipe2.fit(X2.iloc[tr_idx], y2.iloc[tr_idx])

    prob_t = pipe2.predict_proba(X2.iloc[te_idx])[:, 1]

    pred_t = (prob_t >= 0.5).astype(int)

    y2_true.extend(y2.iloc[te_idx])

    y2_pred.extend(pred_t)

    y2_prob.extend(prob_t)

y2_true = np.array(y2_true)

y2_pred = np.array(y2_pred)

y2_prob = np.array(y2_prob)

rep2, m2 = metrics_binary(y2_true, y2_pred, y2_prob, pos_label=1)

print("\n[Stage-2|WMH] CV classification report (threshold=0.5):")

print(rep2)

```

```
print("[Stage-2|WMH] F1:", round(m2["f1"], 3),
      "BalancedAcc:", round(m2["balanced_acc"], 3),
      "AUC:", round(m2["auc"], 3))
```

```
[Stage-1] AD vs Others
      precision    recall  f1-score   support

0         0.394        0.296    0.338        1318
1         0.600        0.698    0.645        1992

 accuracy                   0.538        3310
 macro avg         0.497        0.497    0.492        3310
weighted avg         0.518        0.538    0.523        3310
```

[Stage-1] F1: 0.645 BalancedAcc: 0.497 AUC: 0.497

[Stage-2] Mixed vs Non-AD (among non-AD cases)

[] Saved Stage-2 feature list → hier_out/feature_list_stage2.csv

```
      precision    recall  f1-score   support

0         0.530        0.532    0.531        673
1         0.510        0.509    0.509        645

 accuracy                   0.520        1318
 macro avg         0.520        0.520    0.520        1318
weighted avg         0.520        0.520    0.520        1318
```

[Stage-2] F1: 0.509 BalancedAcc: 0.52 AUC: 0.517

[Stage-2|WMH] CV classification report (threshold=0.5):

```
      precision    recall  f1-score   support

0         0.680        0.682    0.681        673
1         0.667        0.665    0.666        645

 accuracy                   0.674        1318
 macro avg         0.674        0.674    0.674        1318
weighted avg         0.674        0.674    0.674        1318
```

[Stage-2|WMH] F1: 0.666 BalancedAcc: 0.674 AUC: nan

```

[14]: # ----- 4)      Stage-1      -----

from pathlib import Path

import numpy as np, pandas as pd

from sklearn.metrics import confusion_matrix, f1_score

print("\n[Hier] Assemble final 3-class predictions (threshold sweep) ...")

#

try:

    HIER_DIR

except NameError:

    HIER_DIR = Path("hier_out")

HIER_DIR.mkdir(parents=True, exist_ok=True)

#

y_true_col = "target_diag" if "target_diag" in df.columns else ("diag3" if
↳ "diag3" in df.columns else None)

assert y_true_col is not None, "      target_diag / diag3 "

y_true = df[y_true_col].values

labels3 = ["AD", "Mixed", "Non-AD"]

# ===== Stage-1      P(AD) =====

pipe1.fit(X1, y1)

p1_all = pipe1.predict_proba(X1)[:, 1]    # P(AD)

# ===== Stage-2      Stage-2      =====

pipe2.fit(X2, y2)

#

thr_list = [0.50, 0.55, 0.60, 0.65, 0.70]

```

```

rows = []

for thr in thr_list:

    # --- Stage-1      ---

    is_ad = (p1_all >= thr)

    pred_final = np.array([""]*len(df), dtype=object)

    pred_final[is_ad] = "AD"

    # --- Stage-2 "  non-AD"      vs AD ---

    idx_pred_nonad = np.where(~is_ad)[0]

    if idx_pred_nonad.size > 0:

        #          X2_pred

        df_pred_nonad = df.iloc[idx_pred_nonad].reset_index(drop=True)

        X2_pred = build_stage2_matrix(df_pred_nonad, ref_cols=X2.columns) #␣
↪

        p2_pred = pipe2.predict_proba(X2_pred)[: , 1] #␣
↪P(Mixed | predicted non-AD)

        pred_final[idx_pred_nonad] = np.where(p2_pred >= 0.50, "Mixed",␣
↪"Non-AD")

    else:

        p2_pred = np.array([], dtype=float)

    # ---      ---

    cm3 = confusion_matrix(y_true, pred_final, labels=labels3)

    cm3_df = pd.DataFrame(cm3, index=[f"true_{l}" for l in labels3],

                           columns=[f"pred_{l}" for l in labels3])

    cm3_df.to_csv(HIER_DIR / f"final_confusion_matrix_thr{thr:.2f}.csv",␣
↪index=True)

    macro_f1 = f1_score(y_true, pred_final, average="macro", labels=labels3)

```

```

acc      = (pred_final == y_true).mean()

rec_AD   = (pred_final[y_true=="AD"]=="AD").mean() if (y_true=="AD").
↳sum()>0 else np.nan

rows.append({"thr1": float(thr), "macro_f1": float(macro_f1), "accuracy":
↳float(acc), "recall_AD": float(rec_AD)})

# ---      ---

out_pred = pd.DataFrame({

    "BrcId": df.get("BrcId", pd.Series(np.nan, index=df.index)),

    "true_diag": y_true,

    "P_AD": p1_all,

    "final_pred": pred_final

})

if idx_pred_nonad.size > 0:

    out_pred.loc[~is_ad, "P_Mixed_given_nonAD"] = p2_pred

    out_pred.to_csv(HIER_DIR / f"preds_hierarchical_thr{thr:.2f}.csv",
↳index=False)

# -

summary = pd.DataFrame(rows).sort_values("macro_f1", ascending=False)

summary.to_csv(HIER_DIR / "threshold_sweep_stage1_summary.csv", index=False)

print("\n[] Saved threshold sweep summary →", HIER_DIR /
↳"threshold_sweep_stage1_summary.csv")

print(summary.head(10))

```

[Hier] Assemble final 3-class predictions (threshold sweep) ...

```

[] Saved threshold sweep summary → hier_out/threshold_sweep_stage1_summary.csv
thr1 macro_f1 accuracy recall_AD

```


1	0.55	0.805205	0.864048	0.956827
2	0.60	0.798483	0.853776	0.921687
0	0.50	0.793629	0.859215	0.976908
3	0.65	0.785501	0.834441	0.872490
4	0.70	0.742765	0.783384	0.782631

```
[15]: # ----- 4)      Stage-1      sweep 0.50-0.60  10  +      -----

from pathlib import Path

import numpy as np, pandas as pd

import matplotlib.pyplot as plt

from sklearn.metrics import confusion_matrix, f1_score

from IPython.display import display

print("\n[Hier] Assemble final 3-class predictions (threshold sweep, 0.50-0.60,
      ↪random 10) ...")

#

try:

    HIER_DIR

except NameError:

    HIER_DIR = Path("hier_out")

HIER_DIR.mkdir(parents=True, exist_ok=True)

#

y_true_col = "target_diag" if "target_diag" in df.columns else ("diag3" if
      ↪"diag3" in df.columns else None)

assert y_true_col is not None, "      target_diag / diag3 "

y_true = df[y_true_col].values

labels3 = ["AD", "Mixed", "Non-AD"]

# Stage-1      P(AD)

pipe1.fit(X1, y1)
```

```

p1_all = pipe1.predict_proba(X1)[: , 1]

# Stage-2      "  AD  "

pipe2.fit(X2, y2)

# ----  0.50-0.60    10        3    ----

rng = np.random.default_rng(2025)

thr_list = np.round(np.sort(rng.uniform(0.50, 0.60, size=10)), 3)

print("Thresholds (Stage-1, P[AD]   thr):", thr_list.tolist())

rows = []

for thr in thr_list:

    # Stage-1

    is_ad = (p1_all >= thr)

    pred_final = np.array([""]*len(df), dtype=object)

    pred_final[is_ad] = "AD"

    # Stage-2  "  non-AD"  Mixed vs Non-AD

    idx_pred_nonad = np.where(~is_ad)[0]

    if idx_pred_nonad.size > 0:

        df_pred_nonad = df.iloc[idx_pred_nonad].reset_index(drop=True)

        #      build_stage2_matrix(df_slice, ref_cols=...)

        X2_pred = build_stage2_matrix(df_pred_nonad, ref_cols=X2.columns)

        p2_pred = pipe2.predict_proba(X2_pred)[: , 1]

        pred_final[idx_pred_nonad] = np.where(p2_pred >= 0.50, "Mixed",
↪ "Non-AD")

    else:

        p2_pred = np.array([], dtype=float)

```

```

#      +

cm3 = confusion_matrix(y_true, pred_final, labels=labels3)

cm3_df = pd.DataFrame(cm3, index=[f"true_{l}" for l in labels3],
                      columns=[f"pred_{l}" for l in labels3])

cm3_df.to_csv(HIER_DIR / f"final_confusion_matrix_thr{thr:.3f}.csv",
             index=True)

macro_f1 = f1_score(y_true, pred_final, average="macro", labels=labels3)

acc      = (pred_final == y_true).mean()

rec_AD   = (pred_final[y_true=="AD"]=="AD").mean() if (y_true=="AD").
             sum()>0 else np.nan

rows.append({"thr1": float(thr), "macro_f1": float(macro_f1),
            "accuracy": float(acc), "recall_AD": float(rec_AD)})

#      /

out_pred = pd.DataFrame({
    "BrcId": df.get("BrcId", pd.Series(np.nan, index=df.index)),
    "true_diag": y_true,
    "P_AD": p1_all,
    "final_pred": pred_final
})

if idx_pred_nonad.size > 0:
    out_pred.loc[~is_ad, "P_Mixed_given_nonAD"] = p2_pred

    out_pred.to_csv(HIER_DIR / f"preds_hierarchical_thr{thr:.3f}.csv",
                   index=False)

#      +

summary = pd.DataFrame(rows).sort_values("macro_f1", ascending=False)

```

```

summary.to_csv(HIER_DIR / "threshold_sweep_stage1_summary.csv", index=False)

print(f"\n[] Saved threshold sweep summary → {HIER_DIR/
↳ 'threshold_sweep_stage1_summary.csv'}")

display(summary)                                # <--

best = summary.iloc[0]

print(f"\nBest thr1 = {best['thr1']:.3f} | macro-F1={best['macro_f1']:.3f} | "
      f"acc={best['accuracy']:.3f} | recall_AD={best['recall_AD']:.3f}")

#

summary_sort = summary.sort_values("thr1")

fig, ax1 = plt.subplots(figsize=(6.6,4.6))

ax1.plot(summary_sort["thr1"], summary_sort["macro_f1"], "o-", label="Macro-F1")

ax1.plot(summary_sort["thr1"], summary_sort["accuracy"], "s--",
↳ label="Accuracy")

ax1.set_xlabel("Stage-1 threshold (P[AD]   thr)")

ax1.set_ylabel("Score")

ax1.grid(alpha=.3)

ax2 = ax1.twinx()

ax2.plot(summary_sort["thr1"], summary_sort["recall_AD"], "^-.", color="tab:
↳ red", label="Recall (AD)")

ax2.set_ylabel("Recall (AD)")

#

lines, labels = ax1.get_legend_handles_labels()

lines2, labels2 = ax2.get_legend_handles_labels()

ax1.legend(lines+lines2, labels+labels2, loc="best")

plt.tight_layout()

```

```
plt.show() # <--

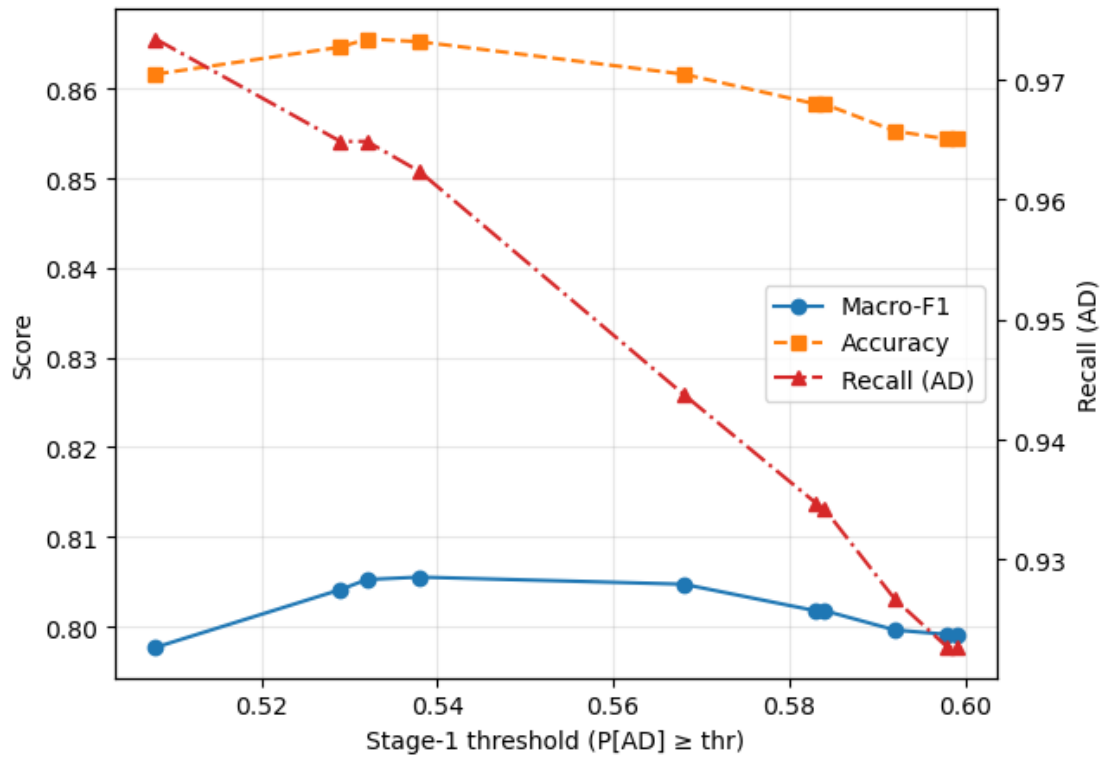
fig.savefig(HIER_DIR / "plot_threshold_sweep_stage1.png", dpi=300)
```

```
[Hier] Assemble final 3-class predictions (threshold sweep, 0.50-0.60 random 10)
...
Thresholds (Stage-1, P[AD] thr): [0.508, 0.529, 0.532, 0.538, 0.568, 0.583,
0.584, 0.592, 0.598, 0.599]
```

```
[ ] Saved threshold sweep summary → hier_out/threshold_sweep_stage1_summary.csv
```

	thr1	macro_f1	accuracy	recall_AD
3	0.538	0.805512	0.865257	0.962349
2	0.532	0.805236	0.865559	0.964859
4	0.568	0.804717	0.861631	0.943775
1	0.529	0.804108	0.864653	0.964859
6	0.584	0.801764	0.858308	0.934237
5	0.583	0.801754	0.858308	0.934739
7	0.592	0.799605	0.855287	0.926707
8	0.598	0.799113	0.854381	0.922691
9	0.599	0.799017	0.854381	0.922691
0	0.508	0.797667	0.861631	0.973394

```
Best thr1 = 0.538 | macro-F1=0.806 | acc=0.865 | recall_AD=0.962
```



```
[16]: # ===== 2.      Stage-1 / Stage-2 / Overall  1      =====

import numpy as np, pandas as pd, matplotlib.pyplot as plt

from pathlib import Path

from sklearn.metrics import (roc_auc_score, average_precision_score, roc_curve,
    precision_recall_curve,

                                confusion_matrix, classification_report,
    brier_score_loss, f1_score)

from sklearn.calibration import calibration_curve, CalibratedClassifierCV

from sklearn.preprocessing import label_binarize

OUTM = Path("surv_out")/"analysis_out"; OUTM.mkdir(parents=True, exist_ok=True)

# -----      ----- #

def metrics_binary_block(y_true, prob, thr=0.5):
```

```

pred = (prob >= thr).astype(int)

cm = confusion_matrix(y_true, pred, labels=[0,1])

rep = classification_report(y_true, pred, output_dict=True, zero_division=0)

fpr, tpr, _ = roc_curve(y_true, prob)

prec, rec, _ = precision_recall_curve(y_true, prob)

out = {

    "roc_auc": roc_auc_score(y_true, prob),

    "pr_auc": average_precision_score(y_true, prob),

    "accuracy": (pred==y_true).mean(),

    "precision_pos": rep["1"]["precision"],

    "recall_pos":    rep["1"]["recall"],

    "f1_pos":        rep["1"]["f1-score"],

    "brier": brier_score_loss(y_true, prob),

    "cm_TN":cm[0,0], "cm_FP":cm[0,1], "cm_FN":cm[1,0], "cm_TP":cm[1,1]

}

curves = {"roc":(fpr,tpr), "pr":(prec,rec)}

return out, curves

# ----- Stage-1 -----

# 1 fit pipe1 fit

try:

    P1_all = p1_all

except NameError:

    pipe1.fit(X1, y1)

    P1_all = pipe1.predict_proba(X1)[: ,1]

```

```

# 1 sweep best['thr1'] 0.60

try:

    thr1 = float(best["thr1"])

except Exception:

    thr1 = 0.60

# Stage-1 OOF/

m1, curves1 = metrics_binary_block(y1.values, P1_all, thr=thr1)

stage1_df = pd.DataFrame([m1])

# vs Isotonic

cal_x1, cal_y1 = calibration_curve(y1.values, P1_all, n_bins=10,
    ↪strategy="quantile")

iso1 = CalibratedClassifierCV(pipe1, method="isotonic", cv=3).fit(X1, y1)

prob1_iso = iso1.predict_proba(X1)[: ,1]

cal_x1b, cal_y1b = calibration_curve(y1.values, prob1_iso, n_bins=10,
    ↪strategy="quantile")

plt.figure(figsize=(4.8,4.2))

plt.plot([0,1],[0,1], "k--", alpha=.3)

plt.plot(cal_x1, cal_y1, "o-", label="uncalibrated")

plt.plot(cal_x1b, cal_y1b, "s-", label="isotonic")

plt.xlabel("Predicted probability"); plt.ylabel("Observed fraction")

plt.title("Stage-1 calibration"); plt.legend(); plt.grid(alpha=.3)

plt.tight_layout(); plt.savefig(OUTM/"calibration_stage1.png", dpi=300); plt.
    ↪close()

# ----- Stage-2 Non-AD + X2_all P2_all -----

# pipe2 fit fit

```



```

try:
    _ = pipe2 # noqa
except NameError:
    raise RuntimeError(" 1 pipe2 XGB for Stage-2 ")

# df2/X2/y2
pipe2.fit(X2, y2)
prob2 = pipe2.predict_proba(X2)[: ,1]
thr2 = 0.50
m2, curves2 = metrics_binary_block(y2.values, prob2, thr=thr2)
stage2_df = pd.DataFrame([m2])

#
cal_x2, cal_y2 = calibration_curve(y2.values, prob2, n_bins=10,
    ↪strategy="quantile")
iso2 = CalibratedClassifierCV(pipe2, method="isotonic", cv=3).fit(X2, y2)
prob2_iso = iso2.predict_proba(X2)[: ,1]
cal_x2b, cal_y2b = calibration_curve(y2.values, prob2_iso, n_bins=10,
    ↪strategy="quantile")
plt.figure(figsize=(4.8,4.2))
plt.plot([0,1],[0,1],"k--",alpha=.3)
plt.plot(cal_x2, cal_y2, "o-", label="uncalibrated")
plt.plot(cal_x2b, cal_y2b, "s-", label="isotonic")
plt.xlabel("Predicted probability"); plt.ylabel("Observed fraction")
plt.title("Stage-2 calibration"); plt.legend(); plt.grid(alpha=.3)
plt.tight_layout(); plt.savefig(OUTM/"calibration_stage2.png", dpi=300); plt.
    ↪close()

```

```

# " df" Stage-2 P2_all /

X2_all = build_stage2_matrix(df.reset_index(drop=True), ref_cols=X2.columns)

P2_all = pipe2.predict_proba(X2_all)[: ,1] # = len(df)

# ----- Overall thr1 + Stage-2 thr2=0.50 -----

#

is_ad = (P1_all >= thr1)

pred_final = np.array([""]*len(df), dtype=object)

pred_final[is_ad] = "AD"

pred_final[~is_ad] = np.where(P2_all[~is_ad] >= thr2, "Mixed", "Non-AD")

# n×3

P_AD = P1_all

P_Mixed = (1.0 - P1_all) * P2_all

P_NonAD = (1.0 - P1_all) * (1.0 - P2_all)

prob_mat = np.c_[P_AD, P_Mixed, P_NonAD] # shape: (n_samples, 3)

# + Macro-F1 + OVR AUC

y_true_col = "target_diag" if "target_diag" in df.columns else ("diag3" if
↳ "diag3" in df.columns else None)

y_true3 = df[y_true_col].values

labels3 = ["AD", "Mixed", "Non-AD"]

cm3 = confusion_matrix(y_true3, pred_final, labels=labels3)

cm3_df = pd.DataFrame(cm3, index=[f"true_{l}" for l in labels3],
↳ columns=[f"pred_{l}" for l in labels3])

rep3 = classification_report(y_true3, pred_final, labels=labels3,
↳ output_dict=True, zero_division=0)

overall_tab = pd.DataFrame([

```

```

    "macro_f1": np.mean([rep3[l]["f1-score"] for l in labels3]),

    "acc": rep3["accuracy"],

    "macro_roc_auc_ovr": roc_auc_score(label_binarize(y_true3, classes=labels3),

                                         prob_mat, average="macro",
↪multi_class="ovr")

}))

# Excel Stage-1/Stage-2/Overall
with pd.ExcelWriter(OUTM/"final.csv") as xw:

    stage1_df.to_excel(xw, sheet_name="Stage1_metrics", index=False)

    stage2_df.to_excel(xw, sheet_name="Stage2_metrics", index=False)

    overall_tab.to_excel(xw, sheet_name="Overall_summary", index=False)

    cm3_df.to_excel(xw, sheet_name="Overall_confusion", index=True)

print("[Metrics] Saved:", OUTM/"final.csv",

      "|", OUTM/"calibration_stage1.png", "|", OUTM/"calibration_stage2.png")

```

```

[Metrics] Saved: surv_out/analysis_out/final.csv |
surv_out/analysis_out/calibration_stage1.png |
surv_out/analysis_out/calibration_stage2.png

```

```

[17]: # ===== SHAP-style contributions for Stage-1 & Stage-2 XGBoost ↪
↪pred_contribs =====

import numpy as np, pandas as pd, matplotlib.pyplot as plt

from pathlib import Path

import xgboost as xgb

from sklearn.pipeline import Pipeline

OUTS = Path("surv_out")/"analysis_out"; OUTS.mkdir(parents=True, exist_ok=True)

def get_transformed_X_and_names(pipe: Pipeline, X):

```

```

""" sklearn Pipeline ColumnTransformer """

prep = pipe.named_steps["prep"]

Xmat = prep.transform(X)

try:

    names = prep.get_feature_names_out()

except Exception:

    #

    names = np.array([f"f{i}" for i in range(Xmat.shape[1])])

return Xmat, names

def shap_contrib_from_xgb(pipe: Pipeline, X, model_name:str, sample_n:int=2000):

""" pipeline XGBClassifier TreeSHAP pred_contribs top """

# fit

try:

    _ = pipe.named_steps["clf"].get_booster()

except Exception:

    pipe.fit(X, pipe.predict(X)) # fit

clf = pipe.named_steps["clf"]

#

Xmat, names = get_transformed_X_and_names(pipe, X)

n = Xmat.shape[0]

if sample_n and n>sample_n:

    rng = np.random.default_rng(2025)

    idx = rng.choice(n, size=sample_n, replace=False)

```

```

    Xmat_s = Xmat[idx]

else:

    idx = np.arange(n)

    Xmat_s = Xmat

    dmat = xgb.DMatrix(Xmat_s, feature_names=names.tolist())

    contrib = clf.get_booster().predict(dmat, pred_contribs=True) # shape: (n,
↪p+1) (n, p+1, C)

    # /

    if contrib.ndim == 2:

        phi = contrib[:, :-1] # bias

    elif contrib.ndim == 3:

        # (n, p+1, C) ↪

        phi = contrib[:, :-1, -1]

    else:

        raise RuntimeError(f"Unexpected contrib shape: {contrib.shape}")

    # /SHAP/

    imp = pd.DataFrame({

        "feature": names,

        "mean_abs_shap": np.abs(phi).mean(axis=0),

        "mean_shap": phi.mean(axis=0)

    }).sort_values("mean_abs_shap", ascending=False)

    # shap

    per_sample = pd.DataFrame(phi, columns=names)

    per_sample.insert(0, "sample_index_in_X", idx)

```

```

# Top-20

top20 = imp.head(20).iloc[::-1] #

plt.figure(figsize=(6.8, 6.0))

plt.barh(top20["feature"], top20["mean_abs_shap"])

plt.xlabel("mean(|SHAP|)"); plt.title(f"{model_name} contributions_
↳(Top-20)")

plt.grid(axis="x", alpha=.2)

fig_path = OUTS/f"{model_name.lower()}_shap_bar.png"

plt.tight_layout(); plt.savefig(fig_path, dpi=300); plt.close()

#

imp.to_csv(OUTS/f"{model_name.lower()}_shap_values.csv", index=False)

per_sample.to_csv(OUTS/f"{model_name.lower()}_shap_samples.csv",
↳index=False)

print(f"[SHAP] {model_name} saved:", fig_path, "|", OUTS/f"{model_name.
↳lower()}_shap_values.csv")

return imp, per_sample

# ----- Stage-1 AD vs Others pipe1 / X1, y1 -----

try:

    pipe1.named_steps["clf"].get_booster()

except Exception:

    pipe1.fit(X1, y1)

imp1, shap1_samples = shap_contrib_from_xgb(pipe1, X1, model_name="Stage1")

# ----- Stage-2 Mixed vs Non-AD pipe2 / X2, y2 -----

try:

    pipe2.named_steps["clf"].get_booster()

```

```

except Exception:

    pipe2.fit(X2, y2)

imp2, shap2_samples = shap_contrib_from_xgb(pipe2, X2, model_name="Stage2")

# Top-10

print("\n[Stage-1] Top-10 features by mean(|SHAP|):")

print(imp1.head(10).to_string(index=False))

print("\n[Stage-2] Top-10 features by mean(|SHAP|):")

print(imp2.head(10).to_string(index=False))

```

[SHAP] Stage1 saved: surv_out/analysis_out/stage1_shap_bar.png |
surv_out/analysis_out/stage1_shap_values.csv
[SHAP] Stage2 saved: surv_out/analysis_out/stage2_shap_bar.png |
surv_out/analysis_out/stage2_shap_values.csv

[Stage-1] Top-10 features by mean(|SHAP|):

	feature	mean_abs_shap	mean_shap
	num__MMSE	0.193538	0.009166
	num__Cerebrovascular_accident	0.144728	0.005919
	num__050_Right Inf Lat Vent__pct	0.140217	0.001491
	num__118_Left Ent entorhinal area__pct	0.134004	-0.007065
	num__051_Left Inf Lat Vent__pct	0.132246	0.001231
	num__Brain_Left_0_Frontal_99__pct	0.118064	0.004361
	num__Brain_Left_4_DeepNuclei_9__pct	0.092967	0.005101
	num__049_Left Hippocampus__pct	0.091276	0.007771
	num__172_Left PHG parahippocampal gyrus__pct	0.086784	-0.001344
	num__LI_PCu	0.086562	0.004348

[Stage-2] Top-10 features by mean(|SHAP|):

	feature	mean_abs_shap	mean_shap
	num__wmh_icv_pct	0.906033	0.009294
	num__wmh_log_icv	0.202818	0.000422
	num__Cerebrovascular_accident	0.196147	-0.005705
	num__Hypertensive_disorder	0.154991	-0.011313
	num__hypertension	0.131173	0.008345
	num__Heart_failure	0.093152	0.001439
	cat__Gender_ID_Female	0.075931	0.001702
	num__Diabetes_mellitus	0.069000	0.001603
	num__Atrial_fibrillation	0.068532	-0.000436
	num__Transient_ischemic_attack	0.059454	-0.000598

```
[18]: prep1 = pipe1.named_steps['prep']
      feat1_names = prep1.get_feature_names_out()
      print([c for c in feat1_names if 'age' in c.lower()])
```

```
['num__age']
```

```
[ ]:
```

```
[19]: ##AD P = 1827/(1827+46+20)=0.965 R=1827/1992=0.918 F1 0.941
      ##Mixed P = 473/(70+473+132)=0.701 R=473/645=0.733 F1 0.716
      ##Non-AD P = 521/(95+126+521)=0.702 R=521/673=0.774 F1 0.736
      ##Macro-F1 (0.941+0.716+0.736)/3 = 0.798
      #-----P=precision, R=recall F1=2PR/(P+R)-----#
```

```
[20]: import re, numpy as np, pandas as pd
      import matplotlib.pyplot as plt
      from math import ceil
      from pathlib import Path

      OUT_DIR = Path("surv_out"); OUT_DIR.mkdir(exist_ok=True, parents=True)

      # 1) / → 3 AD / Mixed / Non-AD
      if "target_diag" in df.columns:
          grp_raw = df["target_diag"].astype(str)
      elif "diag_3class" in df.columns:
          grp_raw = df["diag_3class"].astype(str)
      else:
          raise ValueError("df `target_diag` `diag_3class`")

      def collapse_to_3(x: str) -> str:
          s = str(x)
          if "Alzheimer" in s or s.strip().upper() in {"AD", "ALZHEIMER'S DISEASE", "ALZHEIMER'S DISEASE"}:
              return "AD"
          if "Mixed" in s or s.strip().upper()=="MIXED":
              return "Mixed"
          return "Non-AD"

      df = df.copy()
      df["diag_group3"] = grp_raw.map(collapse_to_3)

      # AD → Mixed → Non-AD
      GROUPS = ["AD", "Mixed", "Non-AD"]
```



```

[21]: ROI_LIST = [
    # MTL
    "048_Right Hippocampus__pct", "049_Left Hippocampus__pct",
    "117_Right Ent entorhinal area__pct", "118_Left Ent entorhinal area__pct",
    "171_Right PHG parahippocampal gyrus__pct", "172_Left PHG parahippocampal_
    ↪gyrus__pct",
    # Lateral/basal temporal
    "155_Right MTG middle temporal gyrus__pct", "156_Left MTG middle temporal_
    ↪gyrus__pct",
    "133_Right ITG inferior temporal gyrus__pct", "134_Left ITG inferior_
    ↪temporal gyrus__pct",
    "201_Right STG superior temporal gyrus__pct", "202_Left STG superior_
    ↪temporal gyrus__pct",
    "123_Right FuG fusiform gyrus__pct", "124_Left FuG fusiform gyrus__pct",
    "203_Right TMP temporal pole__pct", "204_Left TMP temporal pole__pct",
    # Medial parietal
    "167_Right PCgG posterior cingulate gyrus__pct", "168_Left PCgG posterior_
    ↪cingulate gyrus__pct",
    "169_Right PCu precuneus__pct", "170_Left PCu precuneus__pct",
    # Posterior parietal/occipital (PCA axis)
    "199_Right SPL superior parietal lobule__pct", "200_Left SPL superior_
    ↪parietal lobule__pct",
    "197_Right SOG superior occipital gyrus__pct", "198_Left SOG superior_
    ↪occipital gyrus__pct",
    "145_Right MOG middle occipital gyrus__pct", "146_Left MOG middle occipital_
    ↪gyrus__pct",
    "129_Right IOG inferior occipital gyrus__pct", "130_Left IOG inferior_
    ↪occipital gyrus__pct",
    # Ventricular markers
    "052_Right Lateral Ventricle__pct", "053_Left Lateral Ventricle__pct",
    "050_Right Inf Lat Vent__pct", "051_Left Inf Lat Vent__pct",
]

# + WMH/ICV
extras = []
if "age_at_scan_date" in df.columns:
    extras.append("age_at_scan_date")
elif "age_years" in df.columns:
    df["age_at_scan_date"] = pd.to_numeric(df["age_years"], errors="coerce")
    extras.append("age_at_scan_date")
if "wmh_icv_pct" in df.columns:
    extras.append("wmh_icv_pct")

FEATURES = [c for c in ROI_LIST + extras if c in df.columns]
assert len(FEATURES) > 0, " ROI/age/WMH df __pct "

```

```
#
for c in FEATURES:
    df[c] = pd.to_numeric(df[c], errors="coerce")
```

```
[22]: # 3) y
def global_ylim(cols, groups):
    lows, highs = [], []
    for c in cols:
        for g in groups:
            v = df.loc[df["diag_group3"]==g, c].dropna().values
            if v.size:
                lows.append(np.percentile(v, 1)); highs.append(np.percentile(v,
↪99))
    if len(lows)==0: return (0.0, 1.0)
    lo, hi = float(min(lows)), float(max(highs))
    rng = hi-lo
    return (lo-0.05*rng, hi+0.05*rng)

ymin, ymax = global_ylim(FEATURES, GROUPS)

def pretty(col):
    if col.endswith("__pct"):
        return col.replace("Brain_", "").replace("__pct", "").replace("_", " ") +
↪" (%)"
    return col.replace("_", " ")

# 4) violin + dot
n = len(FEATURES)
ncols = 3 if n>=3 else n
nrows = int(np.ceil(n/ncols))

fig, axes = plt.subplots(nrows, ncols, figsize=(ncols*4.6, nrows*4.3),
↪squeeze=False)
plt.subplots_adjust(wspace=0.35, hspace=0.45)
rng = np.random.default_rng(42)

for i, col in enumerate(FEATURES):
    r, c = divmod(i, ncols)
    ax = axes[r, c]
    data = [df.loc[df["diag_group3"]==g, col].dropna().values for g in GROUPS]
    #
    non_empty = [(j, v) for j, v in enumerate(data, start=1) if v.size]
    if not non_empty:
        ax.axis("off"); continue

    # violin
```

```

ax.violinplot([v for _,v in non_empty], showextrema=False, widths=0.9)
# dots
for j,v in non_empty:
    x = rng.normal(loc=j, scale=0.07, size=v.size)
    ax.plot(x, v, 'o', ms=3, alpha=0.6, color="#1f77b4")

ax.set_title(pretty(col), fontsize=10)
ax.set_xticks([j for j,_ in non_empty])
ax.set_xticklabels([GROUPS[j-1] for j,_ in non_empty], fontsize=9)
ax.set_ylim(ymin, ymax)
ax.grid(True, axis='y', alpha=0.3)

#
for k in range(n, nrows*ncols):
    rr,cc = divmod(k, ncols)
    axes[rr,cc].axis("off")

panel_path = OUT_DIR / f"fig_violin_selected_{n}.png"
plt.savefig(panel_path, dpi=300, bbox_inches="tight")
plt.close(fig)
print("Saved panel figure ->", panel_path)

# 5)
for col in FEATURES:
    fig = plt.figure(figsize=(4.2,4.0)); ax = plt.gca()
    data = [df.loc[df["diag_group3"]==g, col].dropna().values for g in GROUPS]
    non_empty = [(j,v) for j,v in enumerate(data, start=1) if v.size]
    if not non_empty:
        plt.close(fig); continue
    ax.violinplot([v for _,v in non_empty], showextrema=False, widths=0.9)
    for j,v in non_empty:
        x = np.random.normal(loc=j, scale=0.07, size=v.size)
        ax.plot(x, v, 'o', ms=3, alpha=0.6, color="#1f77b4")
    ax.set_title(pretty(col), fontsize=10)
    ax.set_xticks([j for j,_ in non_empty]); ax.set_xticklabels([GROUPS[j-1]
↪for j,_ in non_empty], fontsize=9)
    ax.set_ylim(ymin, ymax); ax.grid(True, axis='y', alpha=0.3)
    out_p = OUT_DIR / f"violin_{re.sub('[^0-9A-Za-z]+','_', col)}.png"
    plt.savefig(out_p, dpi=300, bbox_inches="tight"); plt.close(fig)

```

Saved panel figure -> surv_out/fig_violin_selected_2.png

```

[23]: # === Violin + jitter plots for AD-signature volumes (percent), grouped by AD /
↪Mixed / Non-AD ===
import numpy as np, pandas as pd, matplotlib.pyplot as plt, re
from pathlib import Path

```

```

from math import ceil

OUTP = Path("surv_out") / "violin_ad_signature"; OUTP.mkdir(parents=True,
↳exist_ok=True)

# 0)      df['diag3']      diag3
def to_diag3(x: str) -> str:
    s = str(x)
    if "Mixed" in s or s.strip().upper()=="MIXED": return "Mixed"
    if "Alzheimer" in s or s.strip().upper() in {"AD", "ALZHEIMER'S",
↳DISEASE", "ALZHEIMER'S DISEASE"}: return "AD"
    return "Non-AD"

if "diag3" in df.columns:
    df["diag3"] = df["diag3"].astype(str).map(to_diag3)
elif "target_diag" in df.columns:
    df["diag3"] = df["target_diag"].astype(str).map(to_diag3)
elif "diag_group3" in df.columns:
    df["diag3"] = df["diag_group3"].astype(str).map(to_diag3)
else:
    raise KeyError("    df    target_diag/diag3/diag_group3 ")

GROUPS = ["AD", "Mixed", "Non-AD"]
df = df[df["diag3"].isin(GROUPS)].copy().reset_index(drop=True)

# 1)      ( '048' )      __pct      Total
def get_roi_pct(code: str) -> pd.Series | None:
    #      " _"      '048_Right Hippocampus'
    base = next((c for c in df.columns if c.startswith(f"{code}_")), None)
    if base is None:
        return None
    pct_col = base + "__pct"
    if pct_col in df.columns:
        return pd.to_numeric(df[pct_col], errors="coerce")
    #      __pct      /Total*100
    if "Total" in df.columns:
        num = pd.to_numeric(df[base], errors="coerce")
        den = pd.to_numeric(df["Total"], errors="coerce").replace(0, np.nan)
        return (num/den) * 100.0
    #      %
    return pd.to_numeric(df[base], errors="coerce")

```

```

[24]: # 2)      bilateral
def bilateral(code_R: str, code_L: str, name: str) -> pd.Series:
    r = get_roi_pct(code_R); l = get_roi_pct(code_L)
    if r is None and l is None:

```

```

        return pd.Series(np.nan, index=df.index, name=name)
    if r is None: r = pd.Series(0.0, index=df.index)
    if l is None: l = pd.Series(0.0, index=df.index)
    s = (r.fillna(0) + l.fillna(0)).replace([np.inf, -np.inf], np.nan)
    s.name = name
    return s

# 3)
single_areas = {
    # Medial temporal lobe
    "Hippocampus_bi_pct"      : ("048", "049"),
    "Entorhinal_bi_pct"      : ("117", "118"),
    "Parahippocampal_bi_pct" : ("171", "172"),
    # Lateral/basal temporal
    "MTG_bi_pct"              : ("155", "156"),
    "ITG_bi_pct"              : ("133", "134"),
    "STG_bi_pct"              : ("201", "202"),
    "Fusiform_bi_pct"        : ("123", "124"),
    "TemporalPole_bi_pct"     : ("203", "204"),
    # Medial parietal
    "PosteriorCing_bi_pct"    : ("167", "168"),
    "Precuneus_bi_pct"        : ("169", "170"),
    # Posterior parietal/occipital (PCA axis)
    "SPL_bi_pct"              : ("199", "200"),
    "SOG_bi_pct"              : ("197", "198"),
    "MOG_bi_pct"              : ("145", "146"),
    "IOG_bi_pct"              : ("129", "130"),
    # Ventricles
    "LateralVentr_bi_pct"     : ("052", "053"),
    "InfLatVentr_bi_pct"      : ("050", "051"),
}

X_single = pd.DataFrame({name: bilateral(r, l, name) for name, (r,l) in
    ↪ single_areas.items()})

# 4)      Ash
def sum_cols(df_block: pd.DataFrame, names: list[str], out: str) -> pd.Series:
    s = pd.Series(0.0, index=df_block.index, dtype=float)
    for n in names:
        if n in df_block.columns:
            s = s.add(pd.to_numeric(df_block[n], errors="coerce").fillna(0),
    ↪ fill_value=0)
    s.replace([np.inf, -np.inf], np.nan, inplace=True)
    s.name = out
    return s

composites = {

```

```

    "MTL_total_pct"      :␣
    ↪["Hippocampus_bi_pct", "Entorhinal_bi_pct", "Parahippocampal_bi_pct"],
    "Temporal_lateral_pct" :␣
    ↪["MTG_bi_pct", "ITG_bi_pct", "STG_bi_pct", "Fusiform_bi_pct", "TemporalPole_bi_pct"],
    "Medial_parietal_pct" : ["PosteriorCing_bi_pct", "Precuneus_bi_pct"],
    "Posterior_total_pct" :␣
    ↪["SPL_bi_pct", "SOG_bi_pct", "MOG_bi_pct", "IOG_bi_pct"],
    "Ventricles_total_pct" : ["LateralVentr_bi_pct", "InfLatVentr_bi_pct"],
}
X_comp = pd.DataFrame({out: sum_cols(X_single, names, out) for out, names in␣
    ↪composites.items()})

```

```

[25]: # 5)
if "age_at_scan_date" in df.columns:
    age_series = pd.to_numeric(df["age_at_scan_date"], errors="coerce").
    ↪rename("Age_years")
elif "age_years" in df.columns:
    age_series = pd.to_numeric(df["age_years"], errors="coerce").
    ↪rename("Age_years")
else:
    age_series = None

```

```

[26]: # 6)      " "      +      +
to_plot = pd.concat([X_comp, X_single], axis=1)
if age_series is not None:
    to_plot = pd.concat([to_plot, age_series], axis=1)

```

```

[27]: # ===== " " +      y +      =====
import numpy as np, pandas as pd, matplotlib.pyplot as plt, re
from pathlib import Path
from math import ceil

OUTP = Path("surv_out") / "violin_ad_signature"; OUTP.mkdir(parents=True,␣
    ↪exist_ok=True)
GROUPS = ["AD", "Mixed", "Non-AD"]

#      to_plot +      + Age      df["diag3"]
assert "to_plot" in globals(), "to_plot      / /      to_plot"
assert "diag3" in df.columns, "df      diag3      "

#
def pretty(col: str) -> str:
    if col.endswith("__pct"):

```

```

        return col.replace("Brain_", "").replace("__pct", "").replace("_", " ") + "
↳" (%)
        return col.replace("_", " ")

# " " y 1-99 + 10%
def feature_yylim(series_all_groups: list[np.ndarray], pad=0.10):
    vals = np.concatenate([v for v in series_all_groups if v.size>0])
    if vals.size == 0:
        return (0.0, 1.0)
    lo, hi = np.percentile(vals, [1, 99])
    rng = hi - lo if hi > lo else max(1.0, hi) #
    return float(lo - pad*rng), float(hi + pad*rng)

#
def draw_quantiles(ax, data, xpositions):
    for x, v in zip(xpositions, data):
        if v.size == 0:
            continue
        q1, med, q3 = np.percentile(v, [25, 50, 75])
        ax.hlines([q1, q3], x-0.12, x+0.12, colors="#444", lw=1, alpha=0.9)
        ax.hlines(med, x-0.18, x+0.18, colors="#d62728", lw=1.8, alpha=0.9)

# -- A) y --
rng = np.random.default_rng(42)
saved = []
for col in to_plot.columns:
    fig = plt.figure(figsize=(4.2, 4.0)); ax = plt.gca()

    data = [pd.to_numeric(to_plot.loc[df["diag3"]==g, col], errors="coerce").
↳dropna().values
        for g in GROUPS]
    non_empty = [(j, v) for j, v in enumerate(data, start=1) if v.size]
    if not non_empty:
        plt.close(); continue

#
vals_list = [v for _, v in non_empty]
ax.violinplot(vals_list, showextrema=False, widths=0.8, bw_method=0.2)

# jitter
for j, v in non_empty:
    xj = rng.normal(loc=j, scale=0.06, size=v.size)
    ax.plot(xj, v, 'o', ms=2.8, alpha=0.55, color="#1f77b4")

# /
draw_quantiles(ax, vals_list, [j for j, _ in non_empty])

```

```

# y
ymin, ymax = feature_ylim(vals_list, pad=0.10)
ax.set_ylim(ymin, ymax)

ax.set_xticks([j for j, _ in non_empty])
ax.set_xticklabels([GROUPS[j-1] for j, _ in non_empty], fontsize=9)
ax.set_title(pretty(col), fontsize=10)
ax.grid(True, axis='y', alpha=0.3)

out_png = OUTP / f"violin_autoscale_{re.sub('[^0-9A-Za-z]+','_', col)}.png"
plt.savefig(out_png, dpi=300, bbox_inches="tight")
plt.close(fig); saved.append(out_png)

print(f"[A] Saved autoscale figs: {len(saved)} images to {OUTP}")

```

[A] Saved autoscale figs: 21 images to surv_out/violin_ad_signature

```

[28]: # =====
# Ash      ROI / / / + +
# surv_out/panels/panel_violin_dots.png / panel_violin_mmse.png / panel_stats.
# ↪ xlsx
# matplotlib/scipy/statsmodels  scikit-posthocs
# =====
import numpy as np, pandas as pd, re, sys
import matplotlib.pyplot as plt
from pathlib import Path
from math import ceil
from scipy import stats
from statsmodels.stats.multitest import multipletests

PAN_OUT = Path("surv_out") / "panels"; PAN_OUT.mkdir(parents=True,
# ↪ exist_ok=True)
DATA_FALLBACK = Path("surv_out") / "survival_dataset.csv"

def log(msg):
    print("[PAN]", msg)

# ----- 0) df -----
if "df" not in globals():
    if DATA_FALLBACK.exists():
        log(f"df {DATA_FALLBACK}");
        df = pd.read_csv(DATA_FALLBACK, low_memory=False)
    else:
        raise RuntimeError(" df surv_out/survival_dataset.csv ")

# ----- 1) -----

```



```

def map3(x:str)->str:
    s=str(x)
    if "Mixed" in s or s.strip().upper()=="MIXED": return "Mixed"
    if "Alzheimer" in s or s.strip().upper() in {"AD", "ALZHEIMER'S DISEASE", "ALZHEIMER'S DISEASE", "AD"}: return "AD"
    return "Non-AD"

if "diag3" in df.columns:
    grp_col = "diag3"; df["diag3"] = df[grp_col].astype(str).map(map3)
elif "target_diag" in df.columns:
    grp_col = "target_diag"; df["diag3"] = df[grp_col].astype(str).map(map3)
elif "diag_group3" in df.columns:
    grp_col = "diag_group3"; df["diag3"] = df[grp_col].astype(str).map(map3)
else:
    raise KeyError("    target_diag/diag3/diag_group3 ")

GROUPS = ["AD", "Mixed", "Non-AD"]
df = df[df["diag3"].isin(GROUPS)].reset_index(drop=True)
log(f"    \n{df['diag3'].value_counts().to_string()}")

# ----- 2)  '__pct'    +Total    -----
def get_pct_by_code(code:str):
    """    code    __pct    ROI/Total*100    """
    base = next((c for c in df.columns if c.startswith(f"{code}_")), None)
    if base is None:
        return None
    pct_col = base + "__pct"
    if pct_col in df.columns:
        return pd.to_numeric(df[pct_col], errors="coerce")
    if "Total" in df.columns:
        num = pd.to_numeric(df[base], errors="coerce")
        den = pd.to_numeric(df["Total"], errors="coerce").replace(0, np.nan)
        return (num/den) * 100.0
    return None

def bilateral(code_R, code_L, outname):
    r = get_pct_by_code(code_R); l = get_pct_by_code(code_L)
    if r is None and l is None:
        return pd.Series(np.nan, index=df.index, name=outname)
    if r is None: r = pd.Series(0.0, index=df.index)
    if l is None: l = pd.Series(0.0, index=df.index)
    s = (r.fillna(0)+l.fillna(0)).replace([np.inf, -np.inf], np.nan); s.name=outname; return s

#    Ash
single_defs = {
    # MTL

```

```

"Hippocampus_bi_pct"      : ("048","049"),
"Entorhinal_bi_pct"      : ("117","118"),
"Parahippocampal_bi_pct" : ("171","172"),
# Lateral/basal temporal
"MTG_bi_pct"             : ("155","156"),
"ITG_bi_pct"             : ("133","134"),
"STG_bi_pct"             : ("201","202"),
"Fusiform_bi_pct"        : ("123","124"),
"TemporalPole_bi_pct"    : ("203","204"),
# Medial parietal
"PosteriorCing_bi_pct"   : ("167","168"),
"Precuneus_bi_pct"       : ("169","170"),
# Posterior axis
"SPL_bi_pct"             : ("199","200"),
"SOG_bi_pct"             : ("197","198"),
"MOG_bi_pct"             : ("145","146"),
"IOG_bi_pct"             : ("129","130"),
# Ventricles
"LateralVentr_bi_pct"    : ("052","053"),
"InfLatVentr_bi_pct"     : ("050","051"),
}
singles = {}
for name,(r,l) in single_defs.items():
    s = bilateral(r,l,name); singles[name] = s

X_single = pd.DataFrame(singles)
present_single = [c for c in X_single.columns if X_single[c].notna().any()]
log(f"      {len(present_single)}")

#
def sum_cols(df_block, cols, out):
    s = pd.Series(0.0, index=df_block.index)
    for c in cols:
        if c in df_block.columns:
            s = s.add(pd.to_numeric(df_block[c], errors="coerce").fillna(0),
↪fill_value=0)
    s.replace([np.inf,-np.inf], np.nan, inplace=True); s.name = out; return s

composites = {
    "MTL_total_pct"      :
↪["Hippocampus_bi_pct","Entorhinal_bi_pct","Parahippocampal_bi_pct"],
    "Temporal_lateral_total_pct" :
↪["MTG_bi_pct","ITG_bi_pct","STG_bi_pct","Fusiform_bi_pct","TemporalPole_bi_pct"],
    "Medial_parietal_total_pct"  : ["PosteriorCing_bi_pct","Precuneus_bi_pct"],
    "Posterior_total_pct"       :
↪["SPL_bi_pct","SOG_bi_pct","MOG_bi_pct","IOG_bi_pct"],
    "Ventricles_total_pct"      : ["LateralVentr_bi_pct","InfLatVentr_bi_pct"],

```

```

}
X_comp = pd.DataFrame({out: sum_cols(X_single, cols, out) for out, cols in
    ↪composites.items()})
present_comp = [c for c in X_comp.columns if X_comp[c].notna().any()]
log(f"    {len(present_comp)}")

# ----- 2)          +          -----

def pct_or_raw_sum(patterns:list[str], outname:str):

    """

    patterns

    -   __pct

    -   Total   → (sum(raw)/Total)*100

    -   None

    """

    #   __pct

    pct_cols = []

    for p in patterns:

        pct_cols += [c for c in df.columns if re.match(p + r".*__pct$", str(c))]

    if pct_cols:

        s = df[pct_cols].apply(pd.to_numeric, errors="coerce").sum(axis=1)

        s.replace([np.inf,-np.inf], np.nan, inplace=True)

        s.name = outname

        return s

    #

    raw_cols = []

    for p in patterns:

```

```

        raw_cols += [c for c in df.columns if re.match(p + r".*$", str(c)) and
↳not str(c).endswith("__pct")]

    raw_cols = list(dict.fromkeys(raw_cols)) #

    if raw_cols and "Total" in df.columns:

        raw_sum = df[raw_cols].apply(pd.to_numeric, errors="coerce").sum(axis=1)

        total = pd.to_numeric(df["Total"], errors="coerce").replace(0, np.nan)

        s = (raw_sum / total) * 100.0

        s.replace([np.inf, -np.inf], np.nan, inplace=True)

        s.name = outname

    return s

return None

# Brain_Left_0_Frontal_0/1/2... Right

LOBE_PATTERNS = {

    "Frontal_lobe_pct": [r"^Brain_Left_0_Frontal_",
↳r"^Brain_Right_0_Frontal_"],

    "Parietal_lobe_pct": [r"^Brain_Left_2_Parietal_",
↳r"^Brain_Right_2_Parietal_"],

    "Occipital_lobe_pct":
↳[r"^Brain_Left_3_Occipital_", r"^Brain_Right_3_Occipital_"],

    "Temporal_lobe_pct": [r"^Brain_Left_1_Temporal_",
↳r"^Brain_Right_1_Temporal_"],

    # Cerebellum / Brainstem Brain_*

    "Cerebellum_pct": [

        r"^039_Right Cerebellum Exterior", r"^040_Left Cerebellum Exterior",

        r"^041_Right Cerebellum White Matter", r"^042_Left Cerebellum White
↳Matter",

```

```

        r"^072_Cerebellar Vermal Lobules I-V", r"^073_Cerebellar Vermal Lobules_
↪VI-VII",

        r"^074_Cerebellar Vermal Lobules VIII-X"

    ],

    "Brainstem_pct": [r"^035_Pons", r"^036_Brain Stem"]
}

lobes_new = {}

for name, pats in LOBE_PATTERNS.items():

    if name not in df.columns:

        s = pct_or_raw_sum(pats, name)

        if s is not None:

            lobes_new[name] = s

lobes_df = pd.DataFrame(lobes_new)

if not lobes_df.empty:

    df = pd.concat([df, lobes_df], axis=1)

#   to_plot / FEATURES   to_plot

to_plot = pd.concat([X_comp, X_single, lobes_df], axis=1)

FEATURES = [c for c in to_plot.columns if to_plot[c].notna().any()]

print("[PAN]      /      ", len(FEATURES))

# ----- 3)   desc + KW + posthoc   to_plot -----

desc_rows = []

stat_rows = []

try:

    import scikit_posthocs as sp

```

```

HAVE_DUNN = True

except Exception:

    HAVE_DUNN = False

    print("[PAN] scikit-posthocs      Mann-Whitney + Holm ")

for feat in FEATURES:

    #

    for g in GROUPS:

        mask = (df["diag3"]==g) & to_plot[feat].notna()

        v = pd.to_numeric(to_plot.loc[mask, feat], errors="coerce").dropna().
↪values

        if v.size:

            q25,q50,q75 = np.percentile(v,[25,50,75])

            desc_rows.append({"feature":feat,"group":g,"n":int(v.size),
                               "median":q50,"IQR":q75-q25,"mean":v.mean(),"sd":v.
↪std(ddof=1)})

        #

        arrays = [pd.to_numeric(to_plot.loc[(df["diag3"]==g) & to_plot[feat].
↪notna(), feat], errors="coerce").dropna().values

                    for g in GROUPS]

        if any(a.size==0 for a in arrays):

            stat_rows.append({"feature":feat,"kw_p":np.nan,"AD_vs_Mixed":np.
↪nan,"AD_vs_NonAD":np.nan,"Mixed_vs_NonAD":np.nan})

            continue

        kw = stats.kruskal(*arrays, nan_policy="omit")

        row = {"feature":feat,"kw_p":kw.pvalue}

        if HAVE_DUNN:

```

```

pmat = sp.posthoc_dunn(arrays, p_adjust="holm")

row.update({"AD_vs_Mixed": float(pmat.values[0,1]),
           "AD_vs_NonAD": float(pmat.values[0,2]),
           "Mixed_vs_NonAD": float(pmat.values[1,2])})

else:

    pairs=[(0,1),(0,2),(1,2)]

    p=[]

    for i,j in pairs:

        _, pv = stats.mannwhitneyu(arrays[i], arrays[j],
        ↪alternative="two-sided")

        p.append(pv)

    _, p_adj, _, _ = multipletests(p, method="holm")

    row.update({"AD_vs_Mixed": p_adj[0], "AD_vs_NonAD": p_adj[1],
    ↪"Mixed_vs_NonAD": p_adj[2]})

    stat_rows.append(row)

desc_df = pd.DataFrame(desc_rows)

stat_df = pd.DataFrame(stat_rows)

with pd.ExcelWriter(PAN_OUT/"panel_stats.xlsx", engine="xlsxwriter") as xw:

    desc_df.to_excel(xw, sheet_name="desc", index=False)

    stat_df.to_excel(xw, sheet_name="posthoc", index=False)

print("[PAN]      ", PAN_OUT/"panel_stats.xlsx")

```

```

[PAN]
diag3
AD      1992
Non-AD   673
Mixed    645

```

```
[PAN]      16
[PAN]      5
[PAN]      /      27
[PAN]  scikit-posthocs      Mann-Whitney + Holm
[PAN]      surv_out/panels/panel_stats.xlsx
```

```
[29]: # =====
#      +      y +      / MMSE
#      surv_out/panels_grouped/panel_all_groups_dots.png
#      surv_out/panels_grouped/panel_all_groups_mmse.png
# =====
import numpy as np, pandas as pd, matplotlib.pyplot as plt, re
from pathlib import Path
from math import ceil
from scipy import stats
from statsmodels.stats.multitest import multipletests

GOUT = Path("surv_out")/"panels_grouped"; GOUT.mkdir(parents=True,
↳exist_ok=True)

# ----- 0) -----
#
if "diag3" not in df.columns:
    if "target_diag" in df.columns:
        df["diag3"] = df["target_diag"]
    elif "diag_group3" in df.columns:
        df["diag3"] = df["diag_group3"]
    else:
        raise KeyError("      diag3/target_diag/diag_group3 ")

def norm3(x:str)->str:
    s=str(x)
    if "Mixed" in s or s.strip().upper()=="MIXED": return "Mixed"
    if "Alzheimer" in s or s.strip().upper() in {"AD","ALZHEIMER'S",
↳DISEASE","ALZHEIMER'S DISEASE","AD"}: return "AD"
    return "Non-AD"

df = df.copy()
df["diag3"] = df["diag3"].astype(str).map(norm3)
GROUPS = ["AD","Mixed","Non-AD"]

#
FEATURES = [c for c in to_plot.columns if to_plot[c].notna().any()]
assert len(FEATURES)>0, "to_plot      "

# MMSE
```



```

MMSE_CANDS = [
    ↪ ["Pre_Mini_Mental_Total", "Post_Mini_Mental_Total", "MMSE", "mmse", "MMSE_plot"]
mmse_col = next((c for c in MMSE_CANDS if c in df.columns), None)
df["MMSE_plot"] = pd.to_numeric(df[mmse_col], errors="coerce") if mmse_col else
    ↪ np.nan
mmse_avail = df["MMSE_plot"].notna().any()

# ----- 1)      p      Kruskal + -----
if "stat_df" not in globals():
    try:
        import scikit_posthocs as sp
        HAVE_DUNN = True
    except Exception:
        HAVE_DUNN = False
    print("[Panel-All] scikit-posthocs      Mann-Whitney + Holm      ")
    rows=[]
    for feat in FEATURES:
        arrays = [pd.to_numeric(to_plot.loc[(df["diag3"]==g)&to_plot[feat].
    ↪ notna(), feat],
                                errors="coerce").dropna().values for g in
    ↪ GROUPS]
        if any(a.size==0 for a in arrays):
            rows.append({"feature":feat, "kw_p":np.nan, "AD_vs_Mixed":np.
    ↪ nan, "AD_vs_NonAD":np.nan, "Mixed_vs_NonAD":np.nan})
            continue
        kw = stats.kruskal(*arrays, nan_policy="omit")
        rec = {"feature":feat, "kw_p":kw.pvalue}
        if HAVE_DUNN:
            pmat = sp.posthoc_dunn(arrays, p_adjust="holm")
            rec.update({"AD_vs_Mixed": float(pmat.values[0,1]),
                        "AD_vs_NonAD": float(pmat.values[0,2]),
                        "Mixed_vs_NonAD": float(pmat.values[1,2])})
        else:
            pairs=[(0,1),(0,2),(1,2)]; p=[]
            for i,j in pairs:
                _, pv = stats.
    ↪ mannwhitneyu(arrays[i],arrays[j],alternative="two-sided")
                p.append(pv)
            _, p_adj, _, _ = multipletests(p, method="holm")
            rec.update({"AD_vs_Mixed": p_adj[0], "AD_vs_NonAD": p_adj[1],
    ↪ "Mixed_vs_NonAD": p_adj[2]})
            rows.append(rec)
    stat_df = pd.DataFrame(rows)

def star(p):
    if pd.isna(p): return ""

```

```

if p<0.001: return "***"
if p<0.01 : return "**"
if p<0.05 : return "*"
return ""

```

```

[30]: # ----- 2) " " p95 y -----
#
MAG = {}
for f in FEATURES:
    v = pd.to_numeric(to_plot[f], errors="coerce").dropna().values
    MAG[f] = float(np.percentile(v,95)) if v.size else 0.0

ratio_factor = 2.0 # 2
s_feats = [k for k,_ in sorted(MAG.items(), key=lambda x:x[1])] #
groups = []
cur, cur_min, cur_max = [], None, None
for f in s_feats:
    p = MAG[f]
    if cur_min is None:
        cur=[f]; cur_min=cur_max=p
        continue
    new_min, new_max = min(cur_min,p), max(cur_max,p)
    if (new_max / max(new_min,1e-9)) <= ratio_factor:
        cur.append(f); cur_min,cur_max=new_min,new_max
    else:
        groups.append(cur); cur=[f]; cur_min=cur_max=p
if cur: groups.append(cur)

print(f"[Panel-All] {len(groups)} ", [len(g) for g in groups])
ordered_features = [f for g in groups for f in g] #

# y (1-99 + IQR )
def robust_yylim(values_list, pad=0.08, iqr_cap=8.0):
    vals = np.concatenate([v for v in values_list if v.size])
    p1,p99 = np.percentile(vals,[1,99]); med = np.median(vals)
    q1,q3 = np.percentile(vals,[25,75]); iqr=max(q3-q1, 1e-9)
    lo = max(p1, med - iqr_cap*iqr)
    hi = min(p99, med + iqr_cap*iqr)
    rng = hi - lo if hi>lo else max(1.0, hi)
    return float(lo - pad*rng), float(hi + pad*rng)

group_yylim = {}
for g in groups:
    vals_all=[]
    for f in g:
        for lab in GROUPS:

```

```

        v = pd.to_numeric(to_plot.loc[(df["diag3"]==lab)&to_plot[f].
↳notna(), f], errors="coerce").dropna().values
        if v.size: vals_all.append(v)
        group_ylim[tuple(g)] = robust_ylim(vals_all, pad=0.12, iqr_cap=12.0)

```

[Panel-All] 5 [6, 9, 4, 7, 1]

```

[31]: from matplotlib.colors import Normalize

import numpy as np, pandas as pd, matplotlib.pyplot as plt

from math import ceil

from pathlib import Path

def draw_one_big_panel(mmse: bool=False,

                        out_name: str="panel_all_groups_dots.png",

                        ncols: int=6,

                        pad_top_ax: float=0.02,          #          padding

                        ylayer_step_ax: float=0.045,      #

                        show_connect_line: bool=True,     #

                        show_overflow_tag: bool=True,     #      "+N"

                        dpi: int=300) -> Path:

    """

        ordered_features      y      group_ylim

        MMSE

        "      "      +N

    """

    assert ("ordered_features" in globals()
            and "group_of" in globals()
            and "group_ylim" in globals()), \
        " ordered_features / group_of / group_ylim      y      "

```

```

GOUT = Path("surv_out")/"panels_grouped"; GOUT.mkdir(parents=True,
↪exist_ok=True)

n = len(ordered_features)

nrows = int(np.ceil(n / ncols))

fig, axes = plt.subplots(nrows, ncols, figsize=(ncols*5.2, nrows*4.6),
↪squeeze=False)

plt.subplots_adjust(wspace=0.48, hspace=0.62)

# colormap for MMSE

if mmse and mmse_avail:

    cmap = plt.get_cmap("RdYlGn"); norm = Normalize(vmin=0, vmax=30)

# x=1/2/3

def data_to_axes_x(ax, x_data):

    inv = ax.transAxes.inverted()

    x_pix = ax.transData.transform((x_data, 0))[0]

    x_ax = inv.transform((x_pix, 0))[0]

    return x_ax

#

y_layers_ax = [1.00 + pad_top_ax + i*ylayer_step_ax for i in range(3)]

pairs = {"AD_vs_Mixed": (1,2), "AD_vs_NonAD": (1,3), "Mixed_vs_NonAD":
↪(2,3)}

order = ["AD_vs_Mixed", "AD_vs_NonAD", "Mixed_vs_NonAD"]

#

def title_of(col):

    try:

        return LONG_NAME.get(col, col.replace("__pct", " (%)").replace("_", "
↪"))

```

```

except NameError:

    return col.replace("__pct", " (%)").replace("_", " ")

for i, feat in enumerate(ordered_features):

    r, c = divmod(i, ncols)

    ax = axes[r, c]

    data = [pd.to_numeric(to_plot.loc[(df["diag3"]==lab)&to_plot[feat].
↪notna(), feat],

                                errors="coerce").dropna().values for lab in_
↪GROUPS]

    non_empty = [(j, v) for j, v in enumerate(data, start=1) if v.size]

    if not non_empty:

        ax.axis("off"); continue

    vals = [v for _, v in non_empty]

    # violin

    ax.violinplot(vals, showextrema=False, widths=0.82, bw_method=0.25)

    # dots

    for j, _ in non_empty:

        mask = (df["diag3"]==GROUPS[j-1]) & to_plot[feat].notna()

        yv = pd.to_numeric(to_plot.loc[mask, feat], errors="coerce").
↪values

        x = np.random.default_rng(42).normal(loc=j, scale=0.06, size=yv.
↪size)

        if mmse and mmse_avail:

            mm = df.loc[mask, "MMSE_plot"].values

            colors = plt.get_cmap("RdYlGn")(np.clip(mm/30.0, 0, 1))

```

```

        ax.scatter(x, yv, c=colors, s=10, alpha=0.65, edgecolors='none')

    else:

        ax.plot(x, yv, 'o', ms=3, alpha=0.55, color="#1f77b4")

# / IQR

for j, arr in non_empty:

    q1, med, q3 = np.percentile(arr,[25,50,75])

    ax.hlines([q1,q3], j-0.12, j+0.12, colors="#444", lw=1, alpha=0.9)

    ax.hlines(med, j-0.18, j+0.18, colors="#d62728", lw=1.8,
↪alpha=0.9)

# y

ylo, yhi = group_ylim[group_of[feat]]

ax.set_ylim(ylo, yhi)

# [ylo, yhi]

if show_overflow_tag:

    all_vals = pd.to_numeric(to_plot[feat], errors="coerce").dropna().
↪values

    n_upper = int((all_vals > yhi).sum())

    if n_upper > 0:

        ax.text(0.98, 1.00 + pad_top_ax*0.6, f"+{n_upper}",
↪transform=ax.transAxes,

                ha="right", va="bottom", fontsize=9, color="#555",
↪clip_on=False)

#

srow = stat_df.loc[stat_df["feature"]==feat]

for idx, key in enumerate(order):

    a, b = pairs[key]

```

```

        p = float(srow[key].iloc[0]) if (len(srow) and key in srow.columns)
    else np.nan

    mark = "***" if p<0.001 else ("**" if p<0.01 else ("*" if p<0.05
    else ""))

    if not mark:

        continue

    y_ax = y_layers_ax[idx]

    ax_a = data_to_axes_x(ax, a)

    ax_b = data_to_axes_x(ax, b)

    x_mid = (ax_a + ax_b) / 2.0

    if show_connect_line:

        ax.plot([ax_a, ax_a, ax_b, ax_b], [y_ax-0.01, y_ax, y_ax,
    y_ax-0.01],

                transform=ax.transAxes, lw=1.0, c="#444", clip_on=False)

        ax.text(x_mid, y_ax + 0.005, mark, transform=ax.transAxes,

                ha="center", va="bottom", fontsize=10, color="#d62728",
    clip_on=False)

    #

    ax.set_title(title_of(feats), fontsize=10)

    ax.set_xticks([j for j, _ in non_empty]); ax.set_xticklabels(GROUPS,
    fontsize=9)

    ax.set_ylabel("%", fontsize=9)

    ax.grid(True, axis='y', alpha=0.3)

    #

    for k in range(len(ordered_features), n_rows*n_cols):

        rr, cc = divmod(k, n_cols)

```

```

        axes[rr, cc].axis("off")

    out_path = GOUT / out_name

    plt.savefig(out_path, dpi=dpi, bbox_inches="tight")

    plt.close(fig)

    print("[Panel-All] Saved:", out_path)

    return out_path

```

```

[32]: # ===== 3.          & MMSE          =====
#      one-way ANOVA + Tukey HSD

# -   AD / Mixed / Non-AD

# -   Tukey HSD      p      *, **, ***

#use_parametric = True # True: ANOVA+Tukey False: Kruskal+ Dunn

import numpy as np, pandas as pd, matplotlib.pyplot as plt

from matplotlib.colors import Normalize

from math import ceil

from pathlib import Path

PANF = Path("surv_out")/"panels_final"; PANF.mkdir(parents=True, exist_ok=True)

# 3.1      to_plot      wmh_icv_pct

FEATURES_PANEL = [c for c in to_plot.columns if to_plot[c].notna().any()]

assert len(FEATURES_PANEL)>0, "to_plot      "

# 3.2      MMSE

GROUPS = ["AD", "Mixed", "Non-AD"]

df = df[df["diag3"].isin(GROUPS)].reset_index(drop=True)

```



```

mmse_col = next((c for c in
    ↪["Pre_Mini_Mental_Total", "Post_Mini_Mental_Total", "MMSE", "mmse", "MMSE_plot"]
    ↪if c in df.columns), None)

df["MMSE_plot"] = pd.to_numeric(df[mmse_col], errors="coerce") if mmse_col else
    ↪np.nan

mmse_avail = df["MMSE_plot"].notna().any()

def title_of(col):

    try:

        return LONG_NAME.get(col, col.replace("__pct", " (%)").replace("_", " "))

    except NameError:

        return col.replace("__pct", " (%)").replace("_", " ")

# 3.3      / MMSE

def draw_two_page_panel(mmse=False, out_name="panel_roi_dist.png", ncols=6,
    ↪max_n_per_group=400):

    feats = FEATURES_PANEL

    n = len(feats); nrows = ceil(n/ncols)

    fig, axes = plt.subplots(nrows, ncols, figsize=(ncols*5.2, nrows*4.6),
    ↪squeeze=False)

    plt.subplots_adjust(wspace=0.45, hspace=0.60)

    if mmse and mmse_avail:

        cmap = plt.get_cmap("RdYlGn"); norm = Normalize(vmin=0, vmax=30)

        cbar_added = False

        for i, feat in enumerate(feats):

            r,c = divmod(i, ncols); ax = axes[r,c]

            data = [pd.to_numeric(to_plot.loc[(df["diag3"]==lab)&to_plot[feat].
    ↪notna(), feat],

```

```

errors="coerce").dropna().values for lab in
GROUPS]

non_empty = [(j,v) for j,v in enumerate(data, start=1) if v.size]

if not non_empty:
    ax.axis("off"); continue

# y 1-99 +IQR

allv = np.concatenate([v for _,v in non_empty])

p1,p99 = np.percentile(allv,[1,99]); med=np.median(allv)

q1,q3 = np.percentile(allv,[25,75]); iqr=max(q3-q1,1e-9)

ylo = max(p1, med - 8.0*iqr); yhi = min(p99, med + 8.0*iqr)

pad = 0.10*(yhi-ylo); ylo -= pad; yhi += pad

# violin

ax.violinplot([v for _,v in non_empty], showextrema=False, widths=0.82,
bw_method=0.25)

# dots

for j,_ in non_empty:
    mask = (df["diag3"]==GROUPS[j-1]) & to_plot[feat].notna()

    yv = pd.to_numeric(to_plot.loc[mask, feat], errors="coerce").values

    x = np.random.default_rng(42).normal(loc=j, scale=0.06,
size=len(yv))

# max_n_per_group

if max_n_per_group and len(yv)>max_n_per_group:
    sel = np.random.default_rng(0).choice(len(yv),
size=max_n_per_group, replace=False)

    yv = yv[sel]; x = x[sel]

```

```

        mm = df.loc[mask, "MMSE_plot"].values[sel] if mmse and
↪mmse_avail else None

        else:

            mm = df.loc[mask, "MMSE_plot"].values if mmse and mmse_avail
↪else None

            if mmse and mmse_avail:

                sc = ax.scatter(x, yv, c=np.clip((mm/30.0),0,1), cmap="RdYlGn",
↪s=10, alpha=0.65, edgecolors='none', vmin=0,
↪vmax=1)

                if not cbar_added:

                    cbar = fig.colorbar(sc, ax=axes.ravel().tolist(),
↪fraction=0.02, pad=0.01)

                    cbar.set_label("MMSE (0-30)")

                    cbar_added = True

                else:

                    ax.plot(x, yv, 'o', ms=3, alpha=0.55, color="#1f77b4")

# /IQR

for j, arr in non_empty:

    q1,med,q3 = np.percentile(arr,[25,50,75])

    ax.hlines([q1,q3], j-0.12, j+0.12, colors="#444", lw=1, alpha=0.9)

    ax.hlines(med,      j-0.18, j+0.18, colors="#d62728", lw=1.8,
↪alpha=0.9)

    ax.set_title(title_of(feats), fontsize=10)

    ax.set_xticks([j for j, _ in non_empty]); ax.set_xticklabels(GROUPS,
↪fontsize=9)

    ax.set_ylabel("%", fontsize=9); ax.set_ylim(ylo, yhi); ax.grid(True,
↪axis='y', alpha=0.3)

```

```

for k in range(len(feats), nrows*ncols):

    rr,cc = divmod(k, ncols); axes[rr,cc].axis("off")

    out = PANF / out_name

    plt.savefig(out, dpi=300, bbox_inches="tight"); plt.close(fig)

    print("[Panel-Final] Saved:", out)

# 1

draw_two_page_panel(mmse=False, out_name="panel_roi_dist.png", ncols=6,
    ↪max_n_per_group=400)

# 2 MMSE

draw_two_page_panel(mmse=True, out_name="panel_roi_mmse_colored.png", ncols=6,
    ↪max_n_per_group=400)

```

[Panel-Final] Saved: surv_out/panels_final/panel_roi_dist.png

[Panel-Final] Saved: surv_out/panels_final/panel_roi_mmse_colored.png

```

[33]: # ===== 4. ANOVA/Tukey Kruskal/Dunn + =====

from scipy import stats

from statsmodels.stats.multitest import multipletests

import statsmodels.api as sm

import statsmodels.formula.api as smf

from pathlib import Path

OUTS = Path("surv_out")/"analysis_out"; OUTS.mkdir(parents=True, exist_ok=True)

# Cliff's delta 2 ANOVA

def cliffs_delta(x, y):

    x=np.asarray(x); y=np.asarray(y)

    nx,ny=len(x),len(y)

```

```

gt = sum((xi > y).sum() for xi in x)

lt = sum((xi < y).sum() for xi in x)

return float((gt - lt)/(nx*ny))

def eta_squared_anova(model):

    anov = sm.stats.anova_lm(model, typ=2)

    ss_between = anov.loc["C(diag3)", "sum_sq"]

    ss_total = anov["sum_sq"].sum()

    return float(ss_between/ss_total) if ss_total>0 else np.nan

rows=[]

for feat in FEATURES_PANEL:

    #

    arr = [pd.to_numeric(to_plot.loc[(df["diag3"]==g)&to_plot[feat].notna()],
↳feat], errors="coerce").dropna().values

        for g in GROUPS]

    # / Shapiro/Levene Shapiro

    try_norm = all([stats.shapiro(a[:500]).pvalue>0.05 for a in arr if a.
↳size>=5]) # 500

    try_lev = stats.levene(*arr, center="median").pvalue>0.05 if all([a.size>5,
↳for a in arr]) else False

    method = "ANOVA+Tukey" if (try_norm and try_lev) else "Kruskal+Dunn"

    if method=="ANOVA+Tukey":

        tmp = pd.DataFrame({"y": to_plot[feat], "diag3": df["diag3"]}).dropna()

        mod = smf.ols("y ~ C(diag3)", data=tmp).fit()

        eta2 = eta_squared_anova(mod)

        # Tukey HSD

```

```

from statsmodels.stats.multicomp import pairwise_tukeyhsd

tk = pairwise_tukeyhsd(endog=tmp["y"].values, groups=tmp["diag3"].
↪values, alpha=0.05)

#      p

def tukey_p(a,b):

    #      a-b

    mask =(
        ((tk.groupsunique[tk._multicomp.pairindices[0]]==a) & (tk.
↪groupsunique[tk._multicomp.pairindices[1]]==b))
        |
        ((tk.groupsunique[tk._multicomp.pairindices[0]]==b) & (tk.
↪groupsunique[tk._multicomp.pairindices[1]]==a))
    )

    # statsmodels      summary

    for row in tk.summary().data[1:]:

        if set(row[:2])==set([a,b]):

            return float(row[-1])

    return np.nan

p_am = tukey_p("AD", "Mixed"); p_an = tukey_p("AD", "Non-AD"); p_mn =
↪tukey_p("Mixed", "Non-AD")

rows.append({"feature":feat, "method":method, "effect":
↪"eta2", "effect_value":eta2,

            "AD_vs_Mixed":p_am, "AD_vs_NonAD":p_an, "Mixed_vs_NonAD":
↪p_mn})

else:

    # Kruskal + Dunn

    kw = stats.kruskal(*arr, nan_policy="omit").pvalue

    try:

```

```

import scikit_posthocs as sp

pmat = sp.posthoc_dunn(arr, p_adjust="fdr_bh") # BH

p_am = float(pmat.values[0,1])

p_an = float(pmat.values[0,2])

p_mn = float(pmat.values[1,2])

except Exception:

    # MW + BH

    pairs=[(0,1),(0,2),(1,2)]

    p_raw=[]

    for i,j in pairs:

        _, pv = stats.
↪mannwhitneyu(arr[i],arr[j],alternative="two-sided")

        p_raw.append(pv)

        _, p_bh, _, _ = multipletests(p_raw, method="fdr_bh")

        p_am, p_an, p_mn = p_bh

    # Cliff's

    d_am = cliffs_delta(arr[0],arr[1]); d_an = cliffs_delta(arr[0],arr[2]); ↪
↪d_mn = cliffs_delta(arr[1],arr[2])

    rows.append({"feature":feat,"method":method,"effect":
↪"cliffs_delta","effect_value":np.nan,

                "AD_vs_Mixed":p_am, "AD_vs_NonAD":p_an, "Mixed_vs_NonAD":
↪p_mn,

                "delta_AD_Mixed":d_am, "delta_AD_NonAD":d_an, ↪
↪"delta_Mixed_NonAD":d_mn})

posthoc_effects = pd.DataFrame(rows)

posthoc_effects.to_csv(OUTS/"posthoc_effects.csv", index=False)

```

```
#

md = """# Statistical methods for group comparisons

- If normality and homoskedasticity hold → **ANOVA + Tukey's HSD**; effect size
  ↳ reported as ** $\eta^2$ **.

- Otherwise → **Kruskal-Wallis + Dunn**_. Pairwise p-values are **FDR (Benjamini-
  Hochberg)** corrected.

- Plots display **violin + jitter**_, with **median (red)** and **IQR (black)**_.
  ↳ Significance is shown as stars (* p<.05, ** p<.01, *** p<.001).

- For non-parametric comparisons, we report **Cliff's  $\eta$  as the effect size.

"""

(Path("surv_out")/"analysis_out"/"stats_methods.md").write_text(md,
  ↳ encoding="utf-8")

print("[Stats] Saved:", OUTS/"posthoc_effects.csv", "|", OUTS/"stats_methods.
  ↳ md")
```

```
[Stats] Saved: surv_out/analysis_out/posthoc_effects.csv |
surv_out/analysis_out/stats_methods.md
```

```
[34]: # ===== 5.      SOP      =====

from pathlib import Path

SOP = Path("surv_out")/"analysis_out"/"preprocessing_SOP.md"

mmse_rule = "- **MMSE**_: take the score closest to the **scan date** per
  ↳ subject; if `Pre` and `Post` both available, prefer `Pre`."

dedup_rule= "- **De-duplication**_: per-scan one row (prefer `ScanID`; otherwise
  ↳ `BrcId+Scan_Date`); keep the earliest or most complete record."

pct_rule = "- **Percent volume**_: `ROI_pct = ROI_volume / Total × 100`_.
  ↳ Bilateral ROIs are summed before percent; lobe-level composites sum all ROI%
  ↳ in the lobe."

wmh_rule = "- **WMH**_: merge by `BrcId` from `WMH.xlsx` (columns `WMH(77)`,
  ↳ `Intracranial-volume`). If multiple rows per `BrcId`, take **max** WMH and
  ↳ **median** ICV; derive `wmh_icv_pct`, `wmh_log_icv`, `wmh_missing`."
```



```

vars_stage1 = list(X1.columns)

vars_stage2 = list(X2.columns)

md = f"""# Preprocessing SOP

## Labels

- Three-class diagnosis: **AD / Mixed / Non-AD** from `Final_Diagnosis` mapping.

- Stage-1 label (binary): **AD vs Others**; Stage-2 label: **Mixed vs Non-AD**
  ↳(within non-AD subset).

## De-duplication

{dedup_rule}

## MMSE policy

{mmse_rule}

## Variables

- **Stage-1 features**: {vars_stage1}

- **Stage-2 features**: {vars_stage2}

## Percent volumes

{pct_rule}

## WMH

{wmh_rule}

## Missing data

- Numerical: **median imputation** (fit inside fold); Categorical:
  ↳**most-frequent** + One-Hot (`handle_unknown='ignore'`).

## Cross-validation & thresholding

- 5-fold **grouped** CV by `BrcId`; **Stage-1 threshold** selected by inner CV
  ↳(objective: macro-F1); external evaluation on outer folds.

"""

```

```
SOP.write_text(md, encoding="utf-8")
```

```
print("[SOP] Saved:", SOP)
```

[SOP] Saved: surv_out/analysis_out/preprocessing_SOP.md

```
[35]: # ===== 7. Two-stage overall evaluation + interpretation =====

import numpy as np, pandas as pd, matplotlib.pyplot as plt

from pathlib import Path

from sklearn.metrics import (roc_auc_score, average_precision_score, roc_curve,
                             precision_recall_curve, confusion_matrix,
                             ↪classification_report)

from sklearn.preprocessing import label_binarize

from sklearn.inspection import permutation_importance

OUT7 = Path("surv_out")/"analysis_out"; OUT7.mkdir(parents=True, exist_ok=True)

# 7.1          P1_all/P2_all

try:

    P1_all

except NameError:

    pipe1.fit(X1, y1); P1_all = pipe1.predict_proba(X1)[: ,1]

try:

    X2_all = build_stage2_matrix(df.reset_index(drop=True), ref_cols=X2.columns)

    pipe2.fit(X2, y2); P2_all = pipe2.predict_proba(X2_all)[: ,1]

except Exception as e:

    raise RuntimeError(" Stage-2      build_stage2_matrix ") from e

P_AD      = P1_all
```

```

P_Mixed = (1.0 - P1_all)*P2_all

P_NonAD = (1.0 - P1_all)*(1.0 - P2_all)

prob_mat = np.c_[P_AD, P_Mixed, P_NonAD]

labels3 = ["AD", "Mixed", "Non-AD"]

y_true_col = "target_diag" if "target_diag" in df.columns else ("diag3" if
↳ "diag3" in df.columns else None)

y_true3 = df[y_true_col].values

# 7.2 ROC/PR OVR

Y_bin = label_binarize(y_true3, classes=labels3)

macro_roc = roc_auc_score(Y_bin, prob_mat, average="macro", multi_class="ovr")

macro_pr = average_precision_score(Y_bin, prob_mat, average="macro")

# 7.3 overall OVR ROC + OVR PR + AD vs Others

fig, axes = plt.subplots(1,3, figsize=(14,4.2))

# ROC

for i,lab in enumerate(labels3):

    fpr,tpr,_ = roc_curve(Y_bin[:,i], prob_mat[:,i])

    axes[0].plot(fpr,tpr,label=lab)

axes[0].plot([0,1],[0,1], 'k--', alpha=.3)

axes[0].set_title(f"OVR ROC (macro AUC={macro_roc:.3f})"); axes[0].
↳ set_xlabel("FPR"); axes[0].set_ylabel("TPR"); axes[0].legend(); axes[0].
↳ grid(alpha=.2)

# PR

for i,lab in enumerate(labels3):

    prec,rec,_ = precision_recall_curve(Y_bin[:,i], prob_mat[:,i])

    axes[1].plot(rec,prec,label=lab)

```

```

axes[1].set_title(f"OVR PR (macro AP={macro_pr:.3f})"); axes[1].
    ↪set_xlabel("Recall"); axes[1].set_ylabel("Precision"); axes[1].legend();
    ↪axes[1].grid(alpha=.2)

# AD vs Others

from sklearn.calibration import calibration_curve

y_ad = (y_true3=="AD").astype(int)

cal_x, cal_y = calibration_curve(y_ad, P_AD, n_bins=10, strategy="quantile")

axes[2].plot([0,1],[0,1], 'k--', alpha=.3)

axes[2].plot(cal_x, cal_y, 'o-')

axes[2].set_title("Calibration: AD vs Others"); axes[2].set_xlabel("Predicted_
    ↪P(AD)"); axes[2].set_ylabel("Observed AD"); axes[2].grid(alpha=.2)

plt.tight_layout()

plt.savefig(OUT7/"pipeline_overall_metrics.png", dpi=300); plt.close(fig)

# 7.4 Stage-1 & Stage-2          Stage-1 Stage-2

# Stage-1

prep1 = pipe1.named_steps["prep"]; clf1 = pipe1.named_steps["clf"]

X1_mat = prep1.transform(X1)

feat1_names = prep1.get_feature_names_out()

pi1 = permutation_importance(clf1, X1_mat, y1, n_repeats=10, random_state=0,
    ↪scoring="f1")

imp1 = pd.DataFrame({"stage": "stage1", "feature": feat1_names, "perm_importance":
    ↪pi1.importances_mean})

# Stage-2

prep2 = pipe2.named_steps["prep"]; clf2 = pipe2.named_steps["clf"]

X2_mat = prep2.transform(X2)

feat2_names = prep2.get_feature_names_out()

```

```

pi2 = permutation_importance(clf2, X2_mat, y2, n_repeats=10, random_state=0,
    ↪scoring="f1")

imp2 = pd.DataFrame({"stage": "stage2", "feature": feat2_names, "perm_importance":
    ↪pi2.importances_mean})

imp_all = pd.concat([imp1, imp2], ignore_index=True)

imp_all.to_csv(OUT7/"importances_stage1_stage2.csv", index=False)

# 7.5 shap_global.png Stage-1 Top-20 + "age vs imaging"

# "age" "imaging"

age_cols = [c for c in feat1_names if "age" in c.lower()]

imaging_cols = [c for c in feat1_names if ("_pct" in c.lower()) or ("Hipp" in
    ↪c or "Ventr" in c or "Temporal" in c)]

# /

imaging_cols = list(set(list(imaging_cols) + [c for c in feat1_names if any(k
    ↪in c for k in
    ↪["MTL", "Posterior", "Parietal", "Occipital", "Temporal_lobe", "Frontal_lobe"]]))))

age_gain = float(imp1.loc[imp1["feature"].isin(age_cols), "perm_importance"].
    ↪sum())

img_gain = float(imp1.loc[imp1["feature"].isin(imaging_cols),
    ↪"perm_importance"].sum())

other_gain = float(imp1["perm_importance"].sum() - age_gain - img_gain)

top20 = imp1.sort_values("perm_importance", ascending=False).head(20)

fig, axes = plt.subplots(1, 2, figsize=(12, 4))

axes[0].barh(top20["feature"][:, -1], top20["perm_importance"][:, -1])

axes[0].set_title("Stage-1 permutation importance (Top-20)")

axes[0].set_xlabel("ΔF1 (mean, permutation)")

axes[0].grid(axis='x', alpha=.2)

```

```

axes[1].pie([img_gain, age_gain, max(other_gain,0.0)],
    labels=["Imaging", "Age", "Other"], autopct="%.1f%%", startangle=90)

axes[1].set_title("Relative contributions (Stage-1)")

plt.tight_layout()

plt.savefig(OUT7/"shap_global.png", dpi=300); plt.close(fig)

print("[Overall] Saved:", OUT7/"pipeline_overall_metrics.png", "|", OUT7/
    "shap_global.png", "|", OUT7/"importances_stage1_stage2.csv")

```

[Overall] Saved: surv_out/analysis_out/pipeline_overall_metrics.png |
 surv_out/analysis_out/shap_global.png |
 surv_out/analysis_out/importances_stage1_stage2.csv

```

[36]: # ===== 8. Interaction / Multidimensional visual probes =====

import numpy as np, pandas as pd, matplotlib.pyplot as plt

from pathlib import Path

from sklearn.linear_model import LogisticRegression

from mpl_toolkits.mplot3d import Axes3D # noqa

OUT8 = Path("surv_out")/"analysis_out"/"interaction_plots"; OUT8.
    mkdir(parents=True, exist_ok=True)

pairs_2d = [

    ("Ventricles_total_pct", "MTL_total_pct"),

    ("LateralVentr_bi_pct", "Posterior_total_pct"),

    ("wmh_icv_pct", "Temporal_lateral_total_pct") #   wmh_icv_pct

]

triples_3d = [

    ("Ventricles_total_pct", "MTL_total_pct", "wmh_icv_pct")

]

for a,b in pairs_2d:

```

```

if a not in to_plot.columns or b not in to_plot.columns:

    continue

    tmp = pd.DataFrame({"a":to_plot[a], "b":to_plot[b], "y":
↪(df["diag3"]=="Mixed").astype(int)}).dropna()

    if tmp.empty: continue

    X = tmp[["a","b"]].values; y = tmp["y"].values

    lr = LogisticRegression().fit(X,y)

    xa = np.linspace(np.quantile(X[:,0], .01), np.quantile(X[:,0], .99), 200)

    xb = np.linspace(np.quantile(X[:,1], .01), np.quantile(X[:,1], .99), 200)

    XX,YY = np.meshgrid(xa,xb); ZZ = lr.predict_proba(np.c_[XX.ravel(),YY.
↪ravel()])[:,1].reshape(XX.shape)

    plt.figure(figsize=(5.0,4.2))

    cs = plt.contour(XX,YY,ZZ, levels=[0.25,0.5,0.75],
↪colors=["#aaa", "#444", "#aaa"])

    plt.clabel(cs, fmt={0.5:"0.5"}, inline=True, fontsize=8)

    for g,c in zip(["AD","Mixed","Non-AD"], ["#1f77b4", "#d62728", "#2ca02c"]):

        m = df["diag3"]==g

        plt.scatter(to_plot.loc[m,a], to_plot.loc[m,b], s=6, alpha=0.4,
↪label=g, c=c)

    plt.xlabel(a); plt.ylabel(b); plt.legend(markerscale=2, frameon=False)

    plt.title("Mixed boundary in 2D feature space")

    plt.tight_layout(); plt.savefig(OUT8/f"interact_2D_{a}_{b}.png", dpi=300);
↪plt.close()

for a,b,c in triples_3d:

    if any(col not in to_plot.columns for col in [a,b,c]):

        continue

```

```

    tmp = pd.DataFrame({"a":to_plot[a], "b":to_plot[b], "c":to_plot[c], "group":
↳df["diag3"]}).dropna()

    if tmp.empty: continue

    fig = plt.figure(figsize=(5.0,4.2)); ax = fig.add_subplot(111,
↳projection='3d')

    colors = {"AD": "#1f77b4", "Mixed": "#d62728", "Non-AD": "#2ca02c"}

    for g in ["AD", "Mixed", "Non-AD"]:

        d = tmp[tmp["group"]==g]

        ax.scatter(d["a"], d["b"], d["c"], s=8, alpha=0.4, c=colors[g], label=g)

        ax.set_xlabel(a); ax.set_ylabel(b); ax.set_zlabel(c); ax.legend(loc="upper
↳left")

    plt.tight_layout(); plt.savefig(OUT8/f"interact_3D_{a}_{b}_{c}.png",
↳dpi=300); plt.close()

print("[Interact] Saved to:", OUT8)

print("One-liner: Mixed separates best in (Ventricles_total_pct vs
↳MTL_total_pct) plane; "

      "WMH/ICV adds orthogonal separation when combined with posterior/temporal
↳composites.")

```

[Interact] Saved to: surv_out/analysis_out/interaction_plots
One-liner: Mixed separates best in (Ventricles_total_pct vs MTL_total_pct)
plane; WMH/ICV adds orthogonal separation when combined with posterior/temporal
composites.

```

[37]: import os

import json

import time

import datetime as dt

import pandas as pd

```



```

import matplotlib.pyplot as plt

import matplotlib.image as mpimg

# ----- " " -----

HOURS_BACK = 24

now_ts = time.time()

cutoff_ts = now_ts - HOURS_BACK * 3600

print("\n" + "="*80)

print(f"    out_dir / hier_dir / surv_out    {HOURS_BACK}    /    ")

print("CSV/XLSX >100    10    PNG/JPG    ")

print("="*80 + "\n")

def format_ts(ts):

    return dt.datetime.fromtimestamp(ts).strftime("%Y-%m-%d %H:%M:%S")

# ----- / -----

def print_csv_limited(path, max_full_rows=100, head_rows=10):

    df = pd.read_csv(path)

    n_rows, n_cols = df.shape

    if n_rows <= max_full_rows:

        print(f"[CSV]    {n_rows}    × {n_cols}    \n")

        print(df.to_string(index=False))

    else:

        print(f"[CSV]    {n_rows}    × {n_cols}    {head_rows}    \n")

        print(df.head(head_rows).to_string(index=False))

def print_excel_limited(path, max_full_rows=100, head_rows=10):

    df = pd.read_excel(path)

```

```

n_rows, n_cols = df.shape

if n_rows <= max_full_rows:

    print(f"[XLSX] {n_rows} × {n_cols} \n")

    print(df.to_string(index=False))

else:

    print(f"[XLSX] {n_rows} × {n_cols} {head_rows} \n")

    print(df.head(head_rows).to_string(index=False))

def print_txt(path):

    with open(path, "r", encoding="utf-8", errors="ignore") as f:

        print(f.read())

def print_json(path):

    with open(path, "r", encoding="utf-8", errors="ignore") as f:

        obj = json.load(f)

        print(json.dumps(obj, indent=2, ensure_ascii=False))

def show_image(path):

    img = mpimg.imread(path)

    plt.figure(figsize=(6, 4))

    plt.imshow(img)

    plt.axis("off")

    plt.title(os.path.basename(path))

    plt.tight_layout()

    plt.show()

# ----- 24 -----

```

```

def dump_root(root_path, title):

    if root_path is None or not os.path.isdir(root_path):

        print(f"[WARN]          {title} ({root_path})")

        return

    print("\n" + "#"*80)

    print(f"#      : {title} ({root_path})")

    print("#"*80 + "\n")

    for dirpath, dirnames, filenames in os.walk(root_path):

        rel_dir = os.path.relpath(dirpath, root_path)

        if rel_dir == ".":

            rel_dir = "(root)"

        print("\n" + "-"*80)

        print(f"      : {rel_dir}")

        print("-"*80 + "\n")

        printed_any = False

        for fname in sorted(filenames):

            fpath = os.path.join(dirpath, fname)

            ext = os.path.splitext(fname)[1].lower()

            #

            if ext not in [".csv", ".txt", ".json", ".xlsx", ".xls",

                           ".png", ".jpg", ".jpeg"]:

                continue

            try:

                mtime = os.path.getmtime(fpath)

```

```

except OSError:

    continue

if mtime < cutoff_ts:

    # 24

    continue

if not printed_any:

    printed_any = True

print("\n" + "-"*40)

print(f" : {fname}")

print(f" : {format_ts(mtime)}")

print("-"*40 + "\n")

try:

    if ext == ".csv":

        print_csv_limited(fpath)

    elif ext in [".xlsx", ".xls"]:

        print_excel_limited(fpath)

    elif ext == ".txt":

        print_txt(fpath)

    elif ext == ".json":

        print_json(fpath)

    elif ext in [".png", ".jpg", ".jpeg"]:

        show_image(fpath)

except Exception as e:

```

```

        print(f"[ERROR]    {fname}    : {e}")

    if not printed_any:

        print("[INFO]        24        ")

# -----      surv_out -----

OUT_DIR_ = globals().get("OUT_DIR") or globals().get("out_dir")

HIER_DIR_ = globals().get("HIER_DIR") or globals().get("hier_dir")

SURV_OUT_ = globals().get("surv_out")

dump_root(OUT_DIR_, "out_dir /      24h ")

dump_root(HIER_DIR_, "hier_dir        24h ")

dump_root(SURV_OUT_, "surv_out /      24h ")

print("\n" + "="*80)

print(f"    {HOURS_BACK}        ")

print("="*80 + "\n")

```

```

=====
    out_dir / hier_dir / surv_out    24    /
CSV/XLSX >100    10  PNG/JPG
=====

```

```

#####
#   : out_dir /      24h (surv_out)
#####

```

```

-----
: (root)
-----

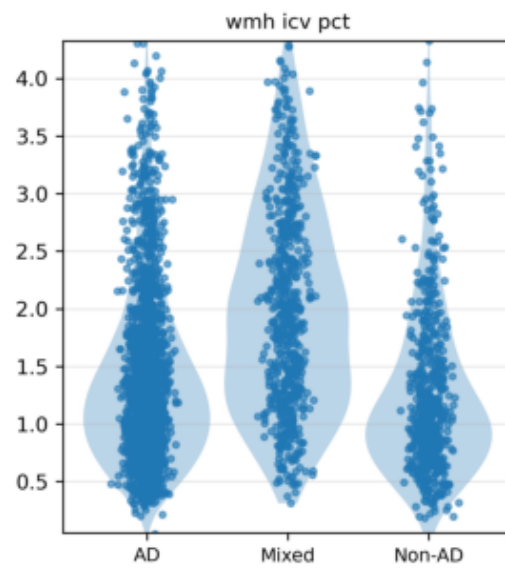
```

```

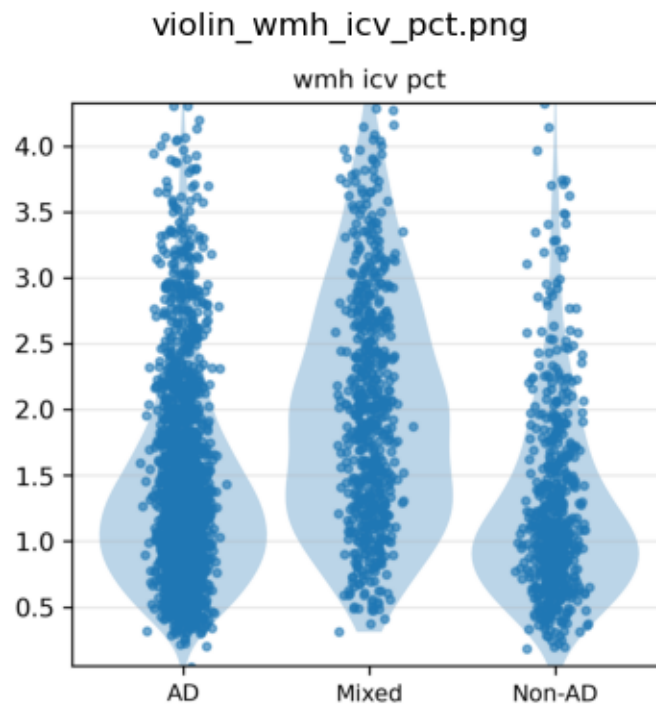
-----
: fig_violin_selected_2.png
: 2025-12-11 06:39:49

```

fig_violin_selected_2.png



: violin_wmh_icv_pct.png
: 2025-12-11 06:39:49

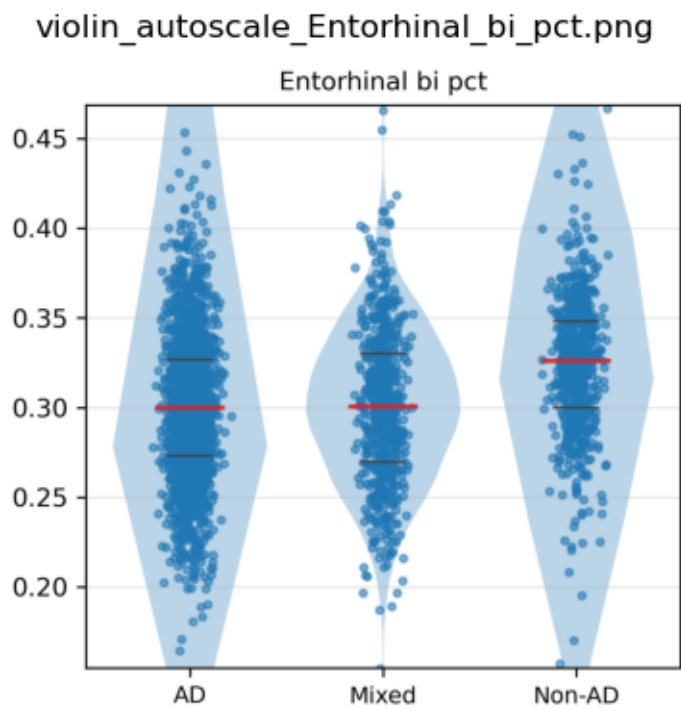


```
: .ipynb_checkpoints
```

[INFO] 24

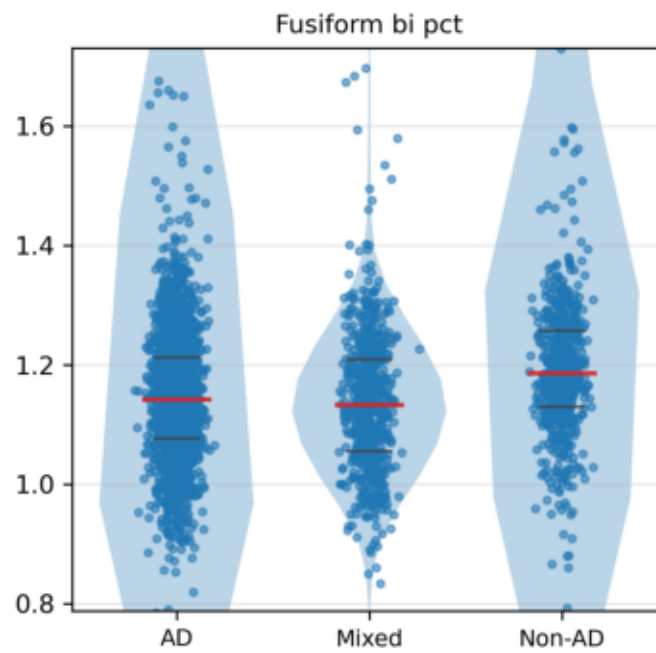
```
: violin_ad_signature
```

```
: violin_autoscale_Entorhinal_bi_pct.png  
: 2025-12-11 06:39:51
```



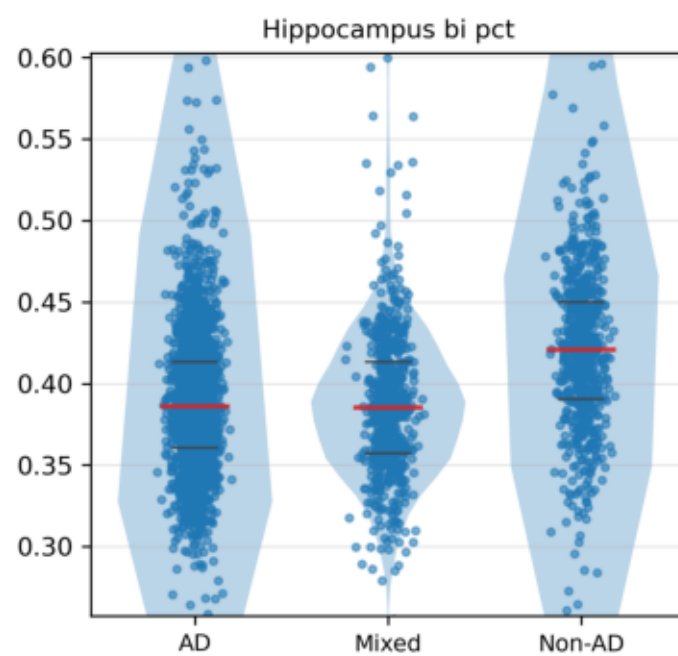
: violin_autoscale_Fusiform_bi_pct.png
: 2025-12-11 06:39:52

violin_autoscale_Fusiform_bi_pct.png



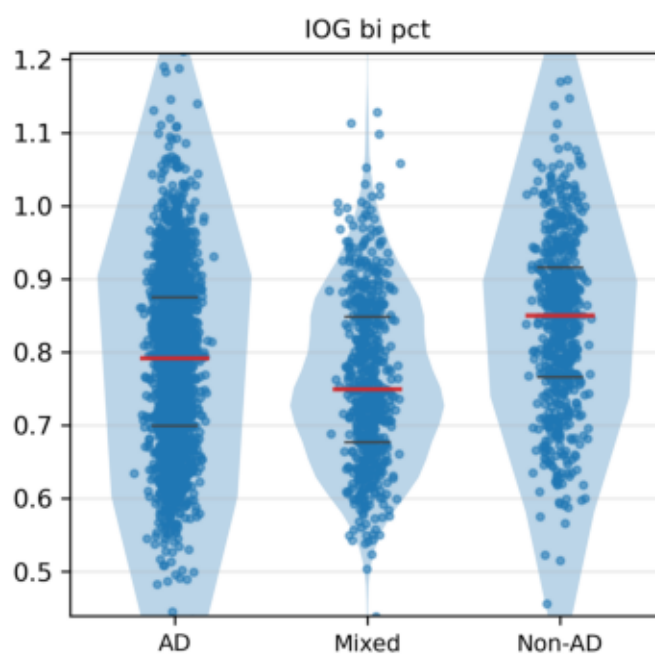
: violin_autoscale_Hippocampus_bi_pct.png
: 2025-12-11 06:39:51

violin_autoscale_Hippocampus_bi_pct.png



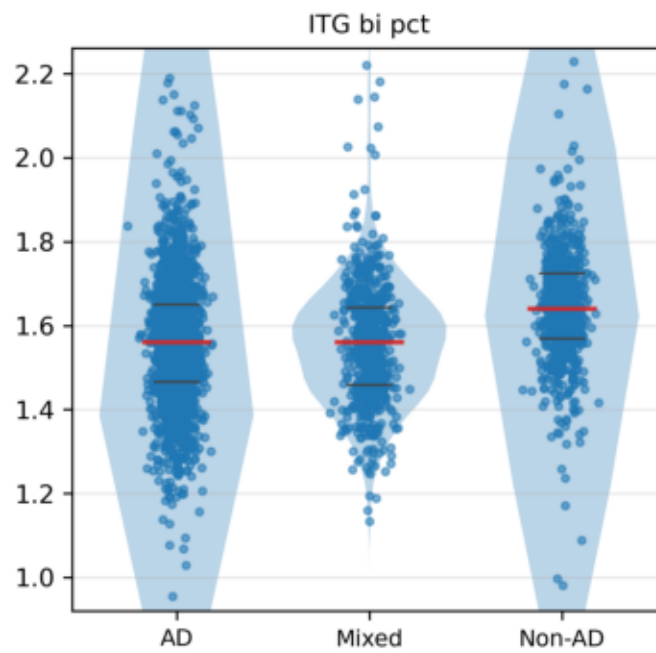
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: 2025-12-11 06:39:53

violin_autoscale_IQG_bi_pct.png



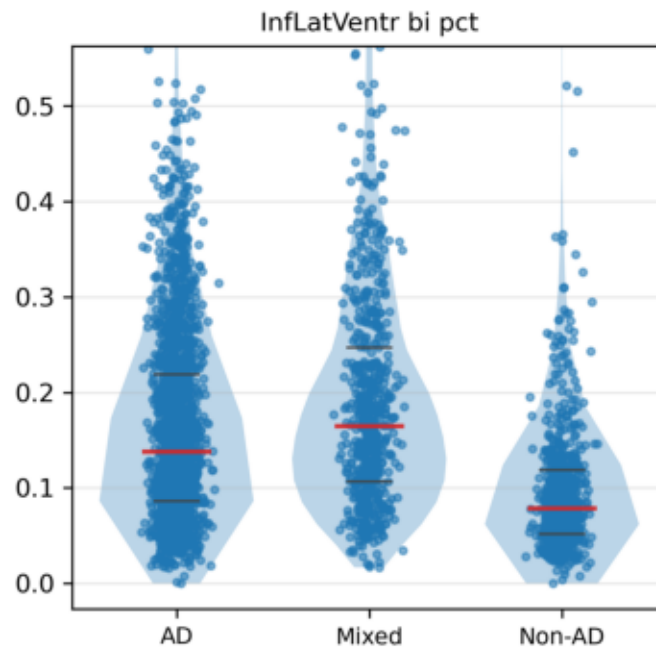
: violin_autoscale_ITG_bi_pct.png
: 2025-12-11 06:39:51

violin_autoscale_ITG_bi_pct.png



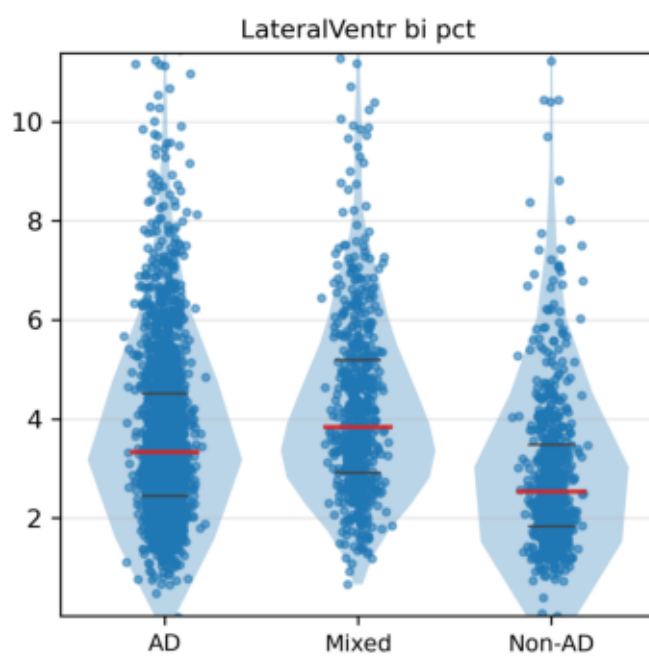
: violin_autoscale_InfLatVentr_bi_pct.png
: 2025-12-11 06:39:54

violin_autoscale_InfLatVentr_bi_pct.png



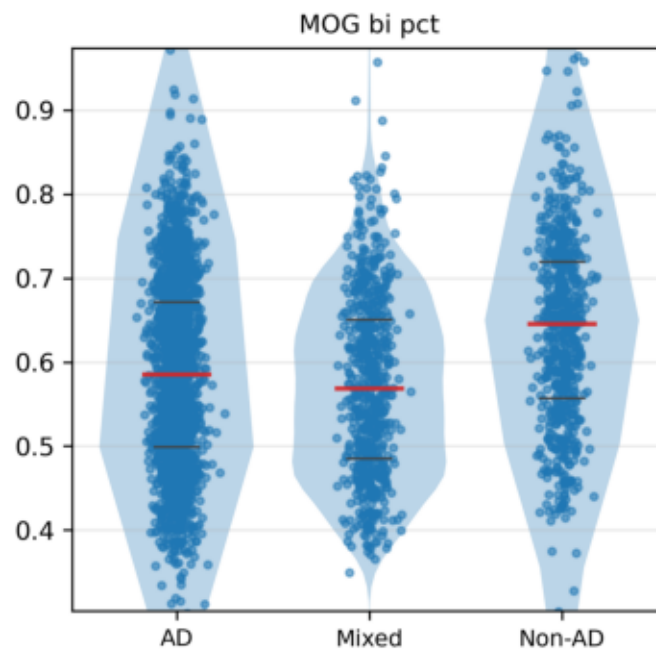
: violin_autoscale_LateralVentr_bi_pct.png
: 2025-12-11 06:39:54

violin_autoscale_LateralVentr_bi_pct.png



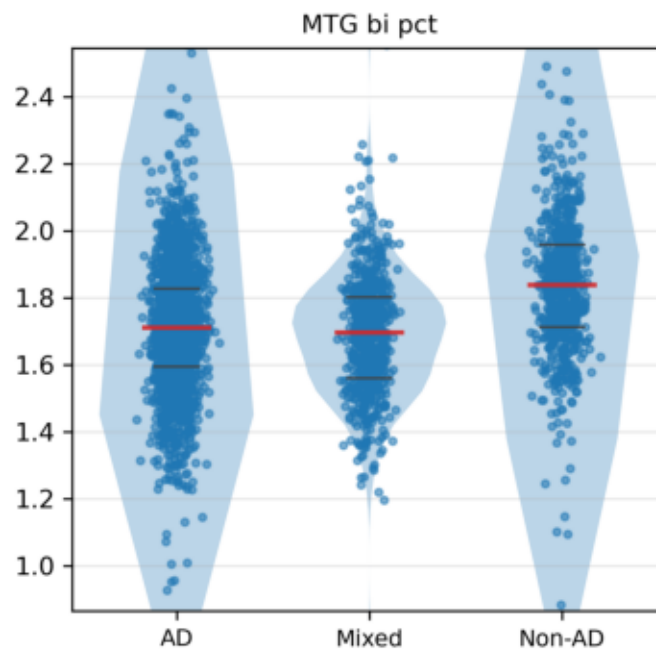
: violin_autoscale_MOG_bi_pct.png
: 2025-12-11 06:39:53

violin_autoscale_MOG_bi_pct.png



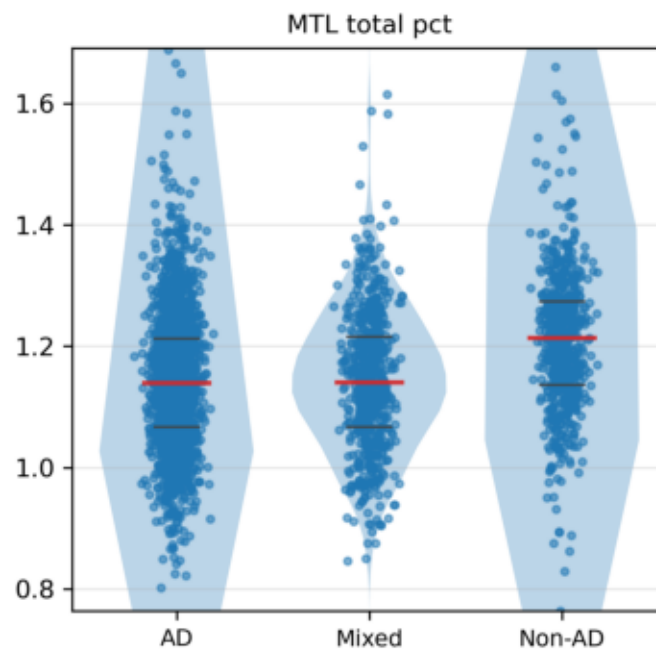
: violin_autoscale_MTG_bi_pct.png
: 2025-12-11 06:39:51

violin_autoscale_MTG_bi_pct.png



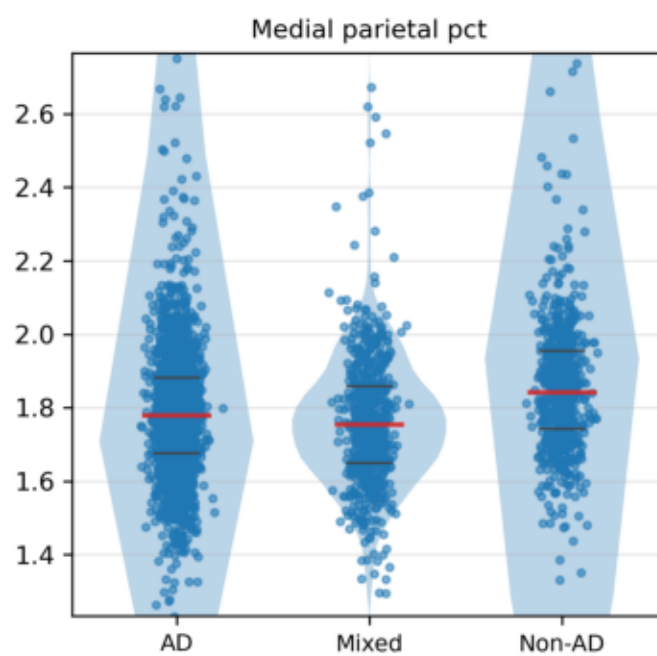
: violin_autoscale_MTL_total_pct.png
: 2025-12-11 06:39:50

violin_autoscale_MTL_total_pct.png



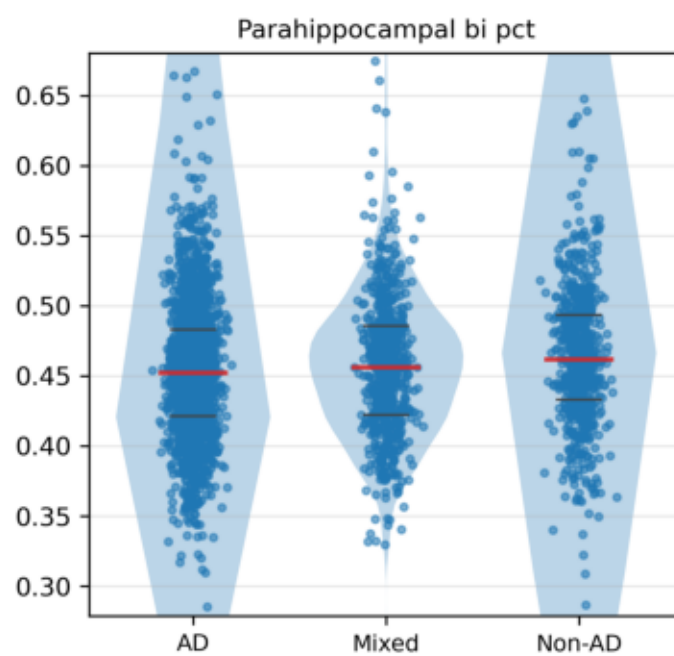
: violin_autoscale_Medial_parietal_pct.png
: 2025-12-11 06:39:50

violin_autoscale_Medial_parietal_pct.png



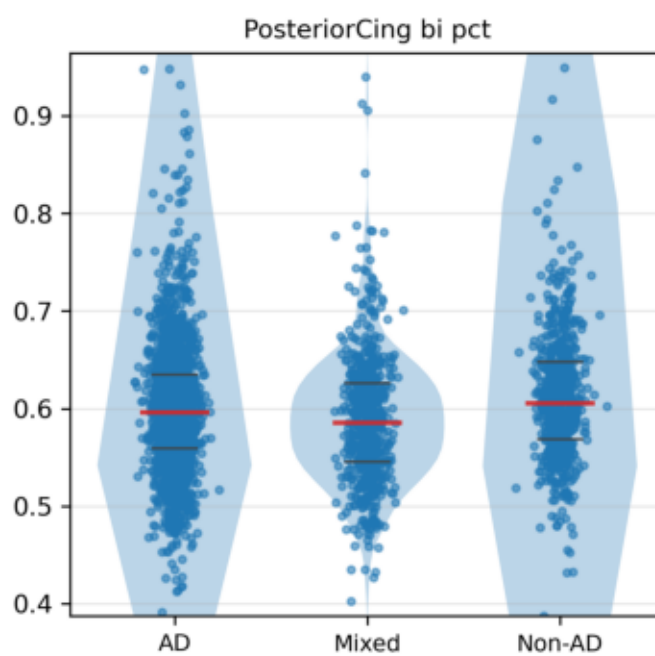
: violin_autoscale_Parahippocampal_bi_pct.png
: 2025-12-11 06:39:51

violin_autoscale_Parahippocampal_bi_pct.png



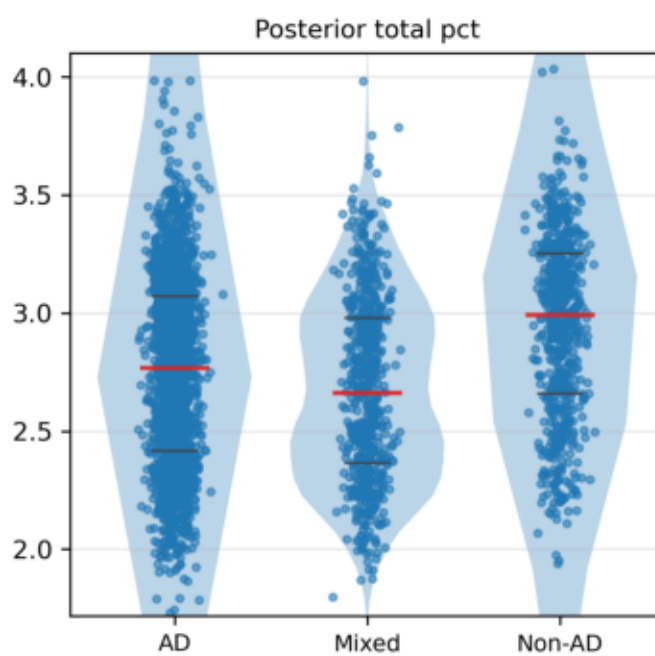
: violin_autoscale_PosteriorCing_bi_pct.png
: 2025-12-11 06:39:52

violin_autoscale_PosteriorCing_bi_pct.png



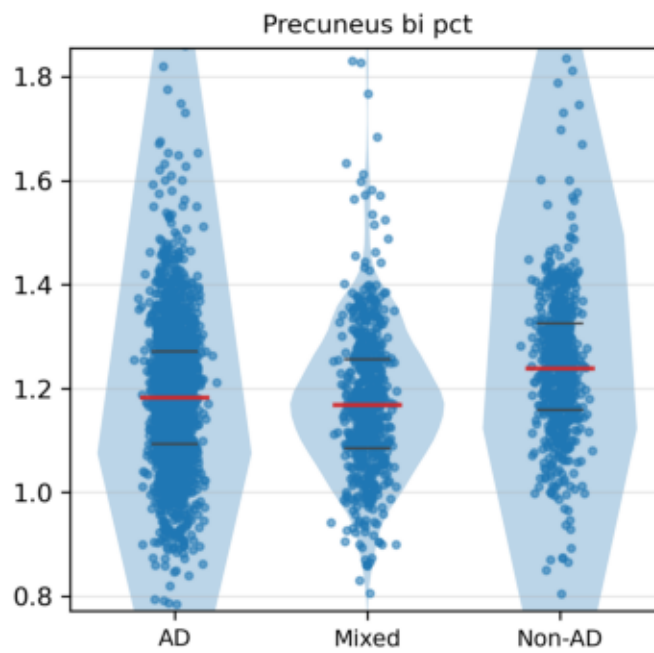
: violin_autoscale_Posterior_total_pct.png
: 2025-12-11 06:39:50

violin_autoscale_Posterior_total_pct.png

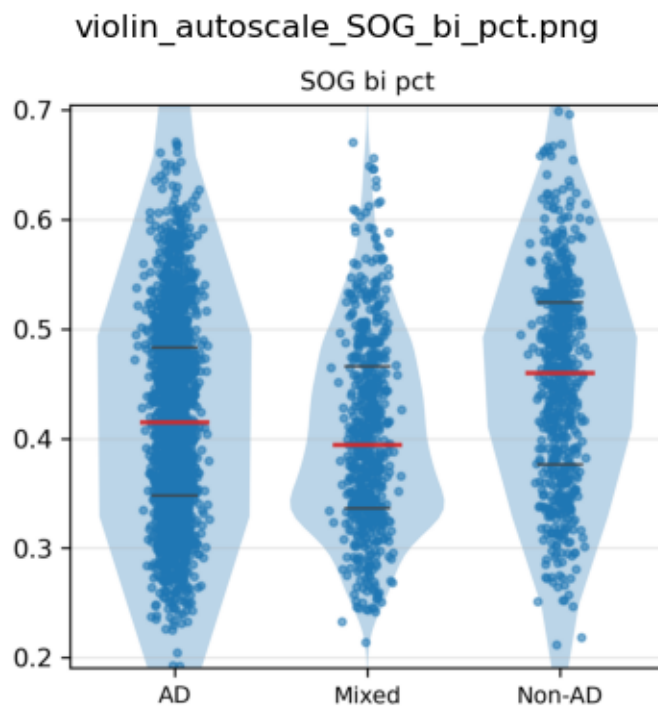


: violin_autoscale_Precuneus_bi_pct.png
: 2025-12-11 06:39:52

violin_autoscale_Precuneus_bi_pct.png

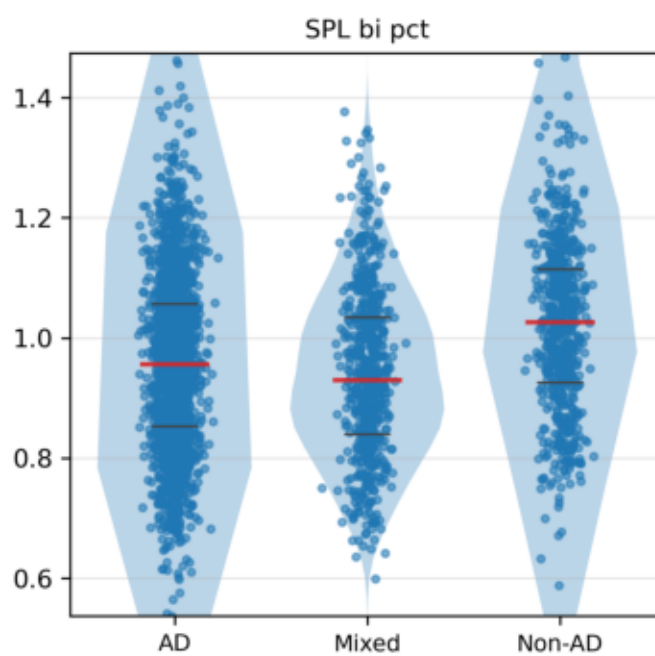


: violin_autoscale_SOG_bi_pct.png
: 2025-12-11 06:39:53



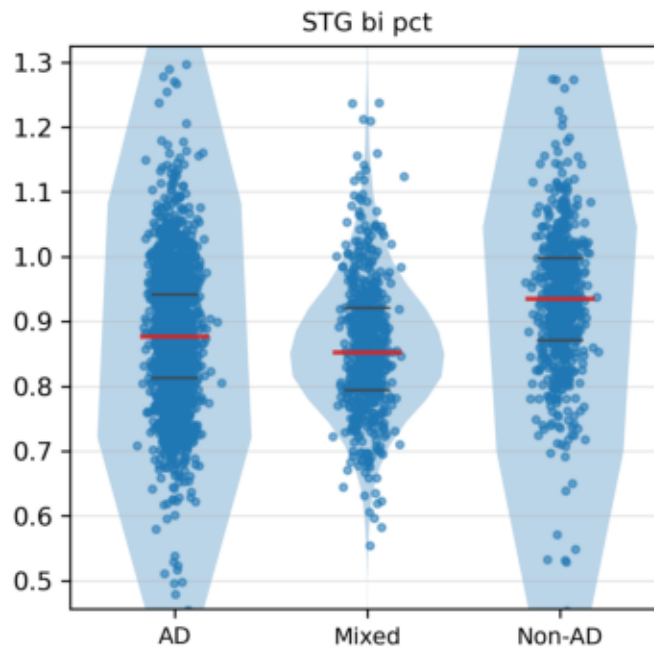
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: 2025-12-11 06:39:53

violin_autoscale_SPL_bi_pct.png



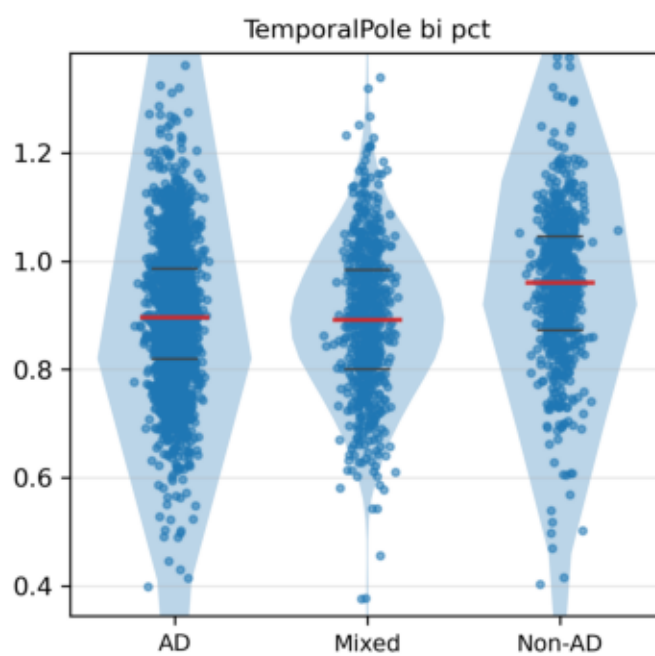
: violin_autoscale_STG_bi_pct.png
: 2025-12-11 06:39:52

violin_autoscale_STG_bi_pct.png



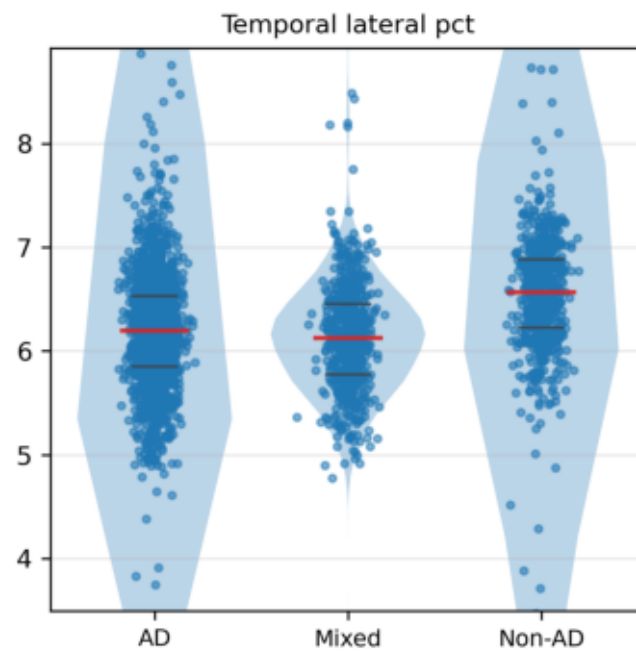
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: 2025-12-11 06:39:52

violin_autoscale_TemporalPole_bi_pct.png



: violin_autoscale_Temporal_lateral_pct.png
: 2025-12-11 06:39:50

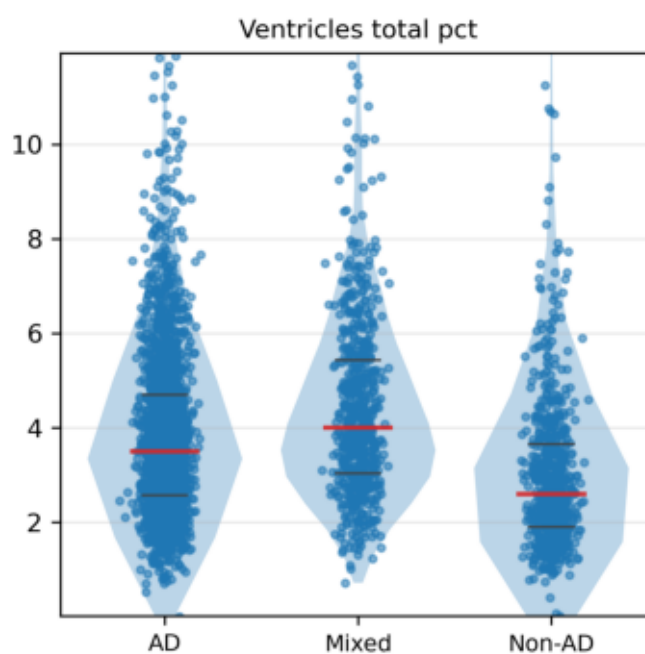
violin_autoscale_Temporal_lateral_pct.png



: violin_autoscale_Ventricles_total_pct.png

: 2025-12-11 06:39:50

violin_autoscale_Ventricles_total_pct.png



```
: violin_ad_signature/.ipynb_checkpoints
```

```
[INFO]      24
```

```
: panels
```

```
: panel_stats.xlsx
: 2025-12-11 06:39:56
```

```
[XLSX]  81   × 7
```

	feature	group	n	median	IQR	mean	sd
	MTL_total_pct	AD	1992	1.139259	0.145325	1.201319	1.526429
	MTL_total_pct	Mixed	645	1.140277	0.148799	1.146916	0.162011
	MTL_total_pct	Non-AD	673	1.213147	0.137617	1.297637	1.686869
	Temporal_lateral_total_pct	AD	1992	6.195232	0.679280	6.496795	8.282181

Temporal_lateral_total_pct	Mixed	645	6.127592	0.680406	6.159007	0.841304
Temporal_lateral_total_pct	Non-AD	673	6.563137	0.657565	7.032315	8.829971
Medial_parietal_total_pct	AD	1992	1.778761	0.206711	1.879352	2.363406
Medial_parietal_total_pct	Mixed	645	1.753844	0.207630	1.761621	0.210522
Medial_parietal_total_pct	Non-AD	673	1.842116	0.212410	2.018839	2.939572
Posterior_total_pct	AD	1992	2.767136	0.655610	2.886503	3.387500
Posterior_total_pct	Mixed	645	2.661584	0.614563	2.688156	0.466122
Posterior_total_pct	Non-AD	673	2.991168	0.592856	3.111213	2.889485
Ventricles_total_pct	AD	1992	3.496242	2.131109	4.075819	5.872265
Ventricles_total_pct	Mixed	645	3.999202	2.402378	4.491929	2.830929
Ventricles_total_pct	Non-AD	673	2.601969	1.756048	3.345217	6.512640
Hippocampus_bi_pct	AD	1992	0.386053	0.052670	0.408027	0.487231
Hippocampus_bi_pct	Mixed	645	0.385176	0.055795	0.388308	0.060990
Hippocampus_bi_pct	Non-AD	673	0.420978	0.059567	0.451873	0.587622
Entorhinal_bi_pct	AD	1992	0.299949	0.053490	0.314979	0.409513
Entorhinal_bi_pct	Mixed	645	0.300647	0.060372	0.302333	0.056461
Entorhinal_bi_pct	Non-AD	673	0.326027	0.048150	0.345304	0.404680
Parahippocampal_bi_pct	AD	1992	0.452037	0.061883	0.478312	0.631902
Parahippocampal_bi_pct	Mixed	645	0.456191	0.063376	0.456275	0.064864
Parahippocampal_bi_pct	Non-AD	673	0.461998	0.060181	0.500460	0.702435
MTG_bi_pct	AD	1992	1.709775	0.232766	1.792930	2.293839
MTG_bi_pct	Mixed	645	1.694928	0.240371	1.695152	0.259014
MTG_bi_pct	Non-AD	673	1.837126	0.243986	1.973532	2.574655
ITG_bi_pct	AD	1992	1.559783	0.184607	1.636937	2.111106
ITG_bi_pct	Mixed	645	1.560474	0.184199	1.557358	0.196197
ITG_bi_pct	Non-AD	673	1.640526	0.154963	1.756533	2.107073
STG_bi_pct	AD	1992	0.877108	0.128775	0.919214	1.152872
STG_bi_pct	Mixed	645	0.852414	0.126692	0.864929	0.159662
STG_bi_pct	Non-AD	673	0.935208	0.126567	1.017377	1.565971
Fusiform_bi_pct	AD	1992	1.142027	0.135805	1.204811	1.519547
Fusiform_bi_pct	Mixed	645	1.132731	0.154208	1.145425	0.228966
Fusiform_bi_pct	Non-AD	673	1.185327	0.127597	1.276276	1.586081
TemporalPole_bi_pct	AD	1992	0.896349	0.165823	0.942903	1.226623
TemporalPole_bi_pct	Mixed	645	0.891029	0.183899	0.896143	0.194418
TemporalPole_bi_pct	Non-AD	673	0.959916	0.173394	1.008596	1.058556
PosteriorCing_bi_pct	AD	1992	0.595853	0.075455	0.631169	0.801846
PosteriorCing_bi_pct	Mixed	645	0.585491	0.080165	0.588833	0.078716
PosteriorCing_bi_pct	Non-AD	673	0.605608	0.079314	0.676934	1.180881
Precuneus_bi_pct	AD	1992	1.182792	0.177988	1.248183	1.565205
Precuneus_bi_pct	Mixed	645	1.168490	0.171812	1.172788	0.156704
Precuneus_bi_pct	Non-AD	673	1.238770	0.166474	1.341905	1.767684
SPL_bi_pct	AD	1992	0.956389	0.203734	1.009458	1.202711
SPL_bi_pct	Mixed	645	0.929858	0.194135	0.943586	0.177207
SPL_bi_pct	Non-AD	673	1.025704	0.188953	1.076687	1.093838
SOG_bi_pct	AD	1992	0.414528	0.135477	0.438237	0.536496
SOG_bi_pct	Mixed	645	0.393944	0.129822	0.404220	0.092303
SOG_bi_pct	Non-AD	673	0.459814	0.148263	0.469221	0.342198
MOG_bi_pct	AD	1992	0.584834	0.172287	0.614614	0.725199

MOG_bi_pct	Mixed	645	0.568642	0.165141	0.574576	0.120535
MOG_bi_pct	Non-AD	673	0.645096	0.162763	0.675756	0.678962
IOG_bi_pct	AD	1992	0.791794	0.175393	0.824195	0.952378
IOG_bi_pct	Mixed	645	0.749055	0.171568	0.765774	0.149784
IOG_bi_pct	Non-AD	673	0.849820	0.149495	0.889550	0.810581
LateralVentr_bi_pct	AD	1992	3.333679	2.065164	3.904633	5.652007
LateralVentr_bi_pct	Mixed	645	3.838285	2.270582	4.299402	2.719391
LateralVentr_bi_pct	Non-AD	673	2.530190	1.648096	3.240460	6.283593
InfLatVentr_bi_pct	AD	1992	0.137946	0.132496	0.171186	0.244964
InfLatVentr_bi_pct	Mixed	645	0.164789	0.139927	0.192527	0.140319
InfLatVentr_bi_pct	Non-AD	673	0.078636	0.066628	0.104757	0.242632
Frontal_lobe_pct	AD	1992	11.315142	1.092529	11.908631	14.717127
Frontal_lobe_pct	Mixed	645	10.960051	1.224867	11.024246	1.234187
Frontal_lobe_pct	Non-AD	673	11.825900	1.226606	12.609408	14.704461
Parietal_lobe_pct	AD	1992	6.016160	0.815502	6.330346	7.933474
Parietal_lobe_pct	Mixed	645	5.853023	0.827523	5.893747	0.679606
Parietal_lobe_pct	Non-AD	673	6.312448	0.728582	6.785547	8.716028
Occipital_lobe_pct	AD	1992	4.476996	0.770140	4.704412	5.625371
Occipital_lobe_pct	Mixed	645	4.346759	0.712566	4.370506	0.618659
Occipital_lobe_pct	Non-AD	673	4.733904	0.793287	4.986190	4.853155
Temporal_lobe_pct	AD	1992	8.194915	0.790056	8.617716	10.933475
Temporal_lobe_pct	Mixed	645	8.112879	0.790070	8.174911	1.089876
Temporal_lobe_pct	Non-AD	673	8.711527	0.787113	9.318653	11.621893
Cerebellum_pct	AD	1992	8.966131	1.201418	9.210946	10.645446
Cerebellum_pct	Mixed	645	8.859635	1.306490	8.668051	1.850111
Cerebellum_pct	Non-AD	673	9.028581	1.179957	9.633571	13.621774
Brainstem_pct	AD	1992	1.223703	0.169722	1.287620	1.513234
Brainstem_pct	Mixed	645	1.211597	0.180940	1.222858	0.302610
Brainstem_pct	Non-AD	673	1.238692	0.172759	1.341089	1.732629

```
: panels/.ipynb_checkpoints
```

```
[INFO]      24
```

```
: panels_grouped
```

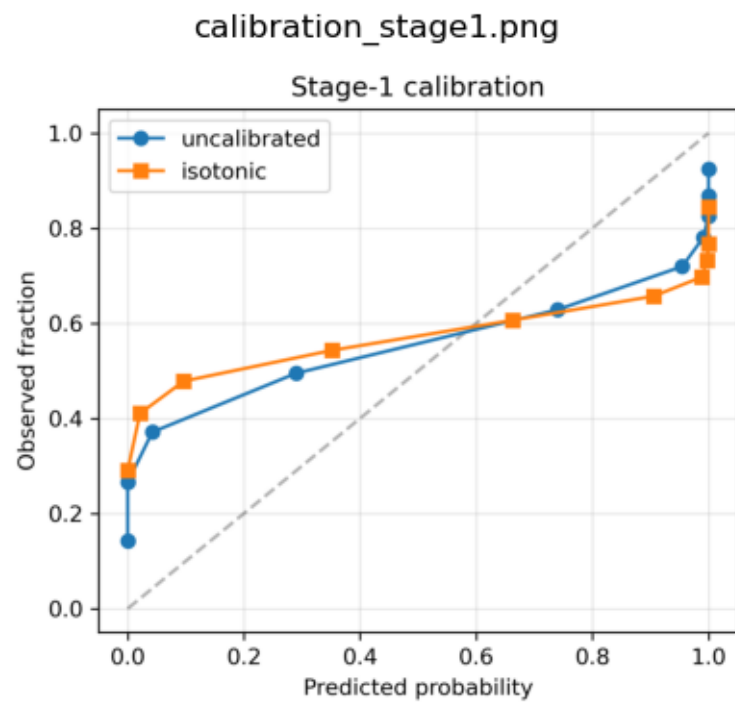
```
[INFO]      24
```

```
: panels_grouped/.ipynb_checkpoints
```

```
[INFO]      24
```

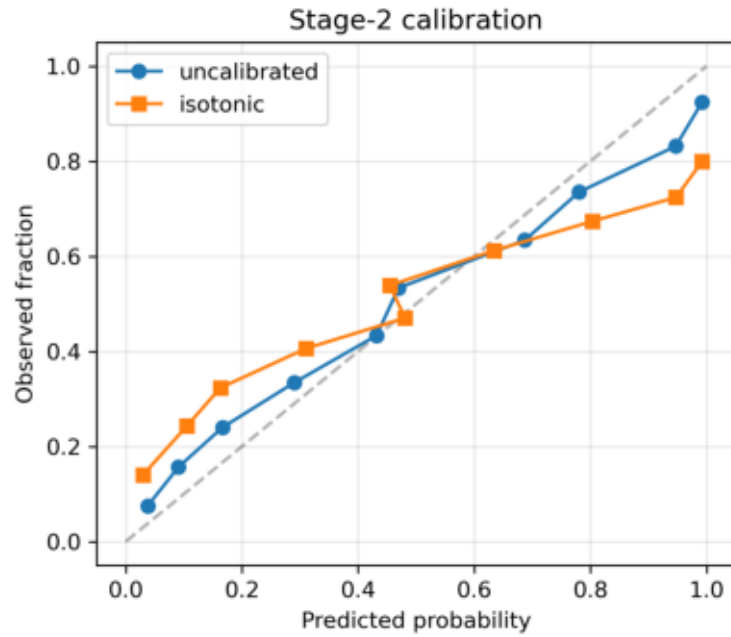
: analysis_out

: calibration_stage1.png
: 2025-12-11 06:39:47



: calibration_stage2.png
: 2025-12-11 06:39:47

calibration_stage2.png



```
-----
: final.csv
: 2025-12-11 06:39:48
-----
```

```
[ERROR] final.csv : 'utf-8' codec can't decode byte 0xfd in position 15:
invalid start byte
```

```
-----
: importances_stage1_stage2.csv
: 2025-12-11 06:40:20
-----
```

```
[CSV] 118 × 3 10
```

stage	feature	perm_importance
stage1	num__048_Right Hippocampus__pct	0.001998
stage1	num__049_Left Hippocampus__pct	0.002747
stage1	num__117_Right Ent entorhinal area__pct	0.004600
stage1	num__118_Left Ent entorhinal area__pct	0.011946
stage1	num__171_Right PHG parahippocampal gyrus__pct	0.007468
stage1	num__172_Left PHG parahippocampal gyrus__pct	0.005891
stage1	num__155_Right MTG middle temporal gyrus__pct	0.004018

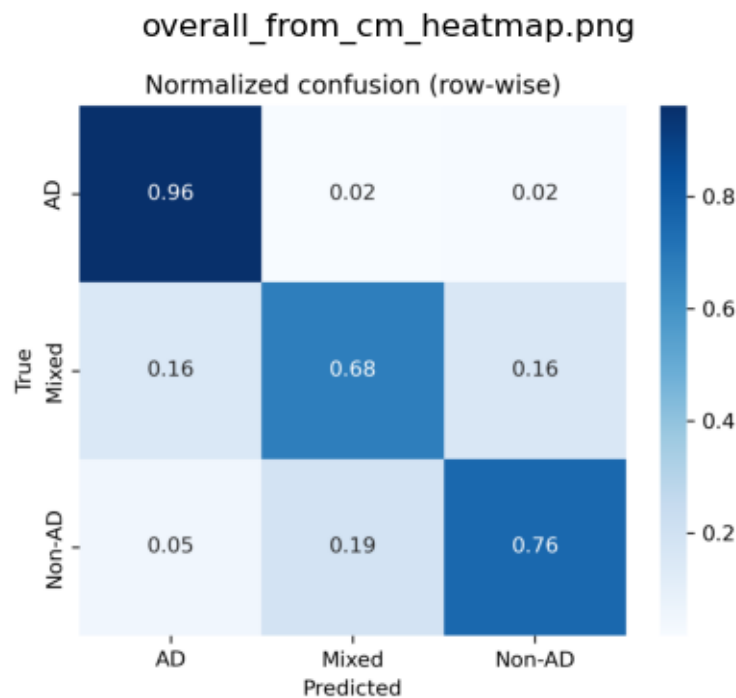

```
stage1    num__156_Left MTG middle temporal gyrus__pct      0.003773
stage1 num__133_Right ITG inferior temporal gyrus__pct      0.004667
stage1  num__134_Left ITG inferior temporal gyrus__pct      0.004838
```

: overall_from_cm_confusion.csv
: 2025-12-10 12:36:56

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1917	33	42
true_Mixed	101	438	106
true_Non-AD	34	130	509

: overall_from_cm_heatmap.png
: 2025-12-10 12:36:56



```
: overall_from_cm_per_class.csv
: 2025-12-10 12:36:56
```

[CSV] 3 × 5

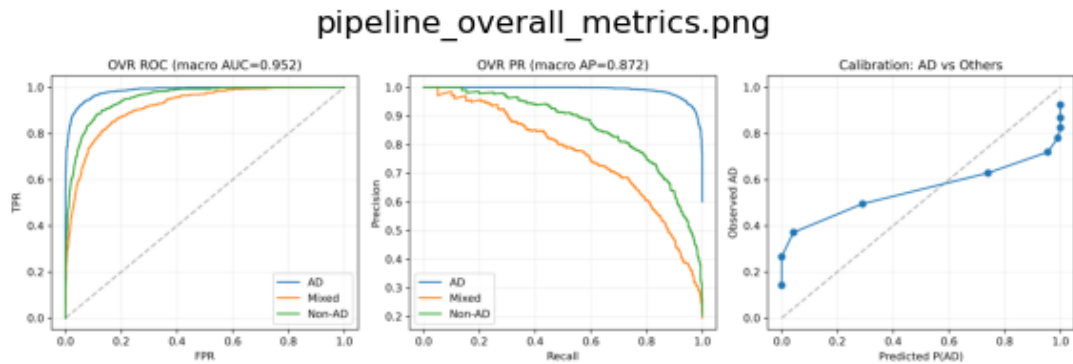
class	support	precision	recall	f1
AD	1992	0.934211	0.962349	0.948071
Mixed	645	0.728785	0.679070	0.703050
Non-AD	673	0.774734	0.756315	0.765414

```
: overall_from_cm_summary.csv
: 2025-12-10 12:36:56
```

[CSV] 1 × 7

accuracy	balanced_accuracy	macro_precision	macro_recall	macro_f1
weighted_f1	N			
0.865257	0.799245	0.812577	0.799245	0.805512
0.863187	3310			

```
: pipeline_overall_metrics.png
: 2025-12-11 06:40:07
```



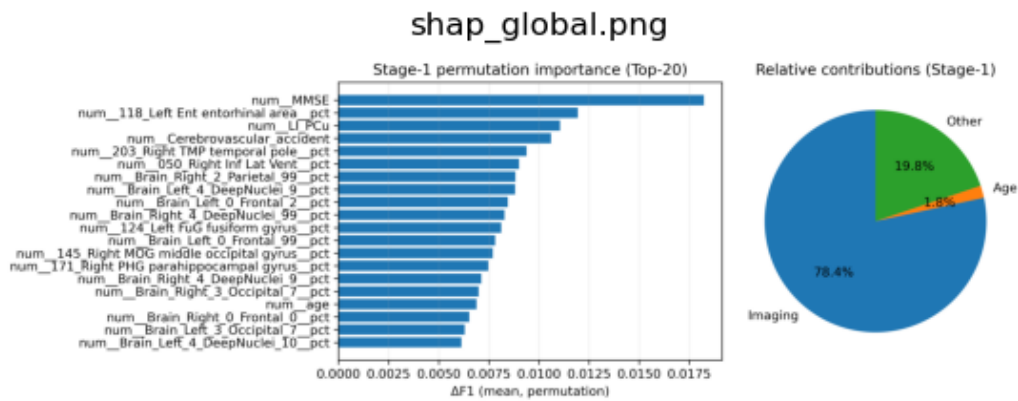
```
: posthoc_effects.csv
: 2025-12-11 06:40:07
```

[CSV] 27 × 10

	feature	method	effect	effect_value	AD_vs_Mixed
AD_vs_NonAD	Mixed_vs_NonAD	delta_AD_Mixed	delta_AD_NonAD	delta_Mixed_NonAD	
	MTL_total_pct	Kruskal+Dunn	cliffs_delta	NaN	7.356779e-01
1.308308e-38	6.863209e-24	-0.008832	-0.336730		-0.322027
	Temporal_lateral_total_pct	Kruskal+Dunn	cliffs_delta	NaN	3.162468e-03
3.453051e-53	1.822434e-47	0.077214	-0.397082		-0.461465
	Medial_parietal_total_pct	Kruskal+Dunn	cliffs_delta	NaN	7.761681e-04
3.403544e-19	1.176805e-22	0.087932	-0.231704		-0.315263
	Posterior_total_pct	Kruskal+Dunn	cliffs_delta	NaN	2.437196e-05
1.365917e-25	3.149690e-31	0.110414	-0.270210		-0.372883
	Ventricles_total_pct	Kruskal+Dunn	cliffs_delta	NaN	9.806081e-13
2.418328e-37	1.375395e-52	-0.186612	0.329589		0.487990
	Hippocampus_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	3.566490e-01
1.545801e-52	5.696907e-39	0.024115	-0.394582		-0.416570
	Entorhinal_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	5.148307e-01
4.093116e-41	6.644804e-23	-0.017040	-0.347836		-0.314876
	Parahippocampal_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	2.699707e-01
8.054004e-06	7.315053e-03	-0.028860	-0.120844		-0.089593
	MTG_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	7.093597e-03
9.478035e-46	3.132182e-42	0.070437	-0.367516		-0.434341
	ITG_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	2.952550e-01
1.824305e-40	5.876624e-32	0.027382	-0.344992		-0.375532
	STG_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	6.249700e-05
1.109280e-35	7.288792e-38	0.104727	-0.321851		-0.412060
	Fusiform_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	4.722359e-02
5.081099e-22	1.742540e-19	0.051912	-0.251252		-0.288743
	TemporalPole_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	1.918256e-01
8.897487e-22	2.913916e-17	0.034146	-0.249784		-0.270429
	PosteriorCing_bi_pct	Kruskal+Dunn	cliffs_delta	NaN	8.293257e-05
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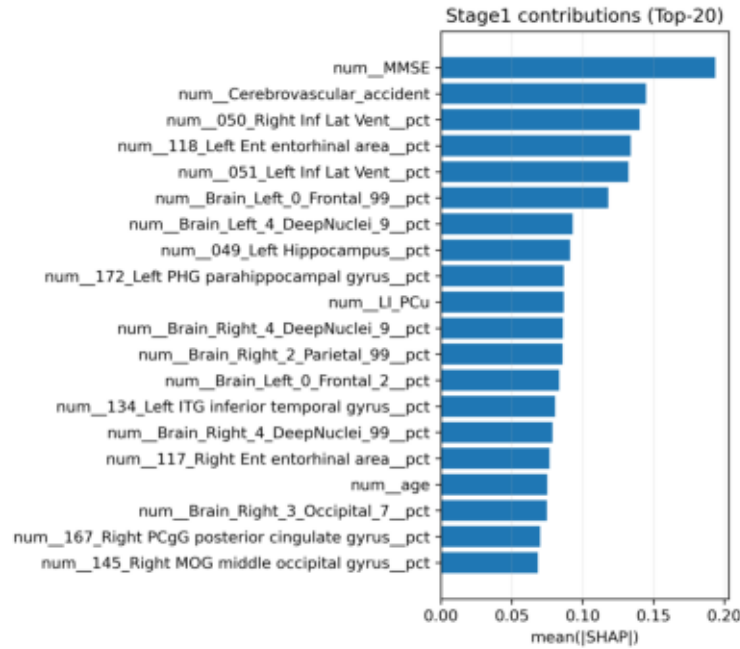
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	Brainstem_pct	Kruskal+Dunn	cliffs_delta	NaN 2.995326e-02
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stage1_shap_bar.png



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inferior temporal gyrus__pct num_134_Left ITG inferior temporal gyrus__pct
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-0.035911		0.0	-0.000311		0.005153
-0.419104	0.002868			0.015937	0.011542
-0.018184	0.000255		0.021415		
0.000241		0.001040		-0.000121	
0.0		0.001482	0.015178	-0.097325	0.024521
0.007182					
	2197		0.034944		
0.083222			0.060063		
-0.116525			0.016164		
0.107576			-0.024387		
0.001446			0.003256		
-0.050202			-0.019796		
-0.001366			0.041300		
0.011476			-0.001806		
-0.003060				-0.083605	
-0.025260			-0.025824		-0.027072
0.095004				-0.070873	
0.044068				0.026996	
-0.020919				0.008966	
-0.047444				-0.011139	
0.122259			0.042817		-0.064387
0.042349			-0.011475		0.047553
0.014952			0.120074		0.048191
0.003417			-0.002175		0.010953
-0.026762			-0.010429		
0.059882			-0.025657		
0.041734			-0.003657		
-0.077244		0.028377		0.096056	
0.036277			-0.037512		0.152744
0.014964			0.013123		-0.014347
0.015821			0.003974		0.047494
0.048975			0.033550		
0.007351			0.017934		
0.063196			-0.039389		0.062957
0.044780			0.004784		0.011402
-0.003528		-0.012218	0.167282	-0.167123	0.002220
-0.061773	-0.006455	0.000181	-0.020467		0.0
0.090515	-0.000254		0.038641		0.001613
0.0	-0.012063			0.0	-0.007332
0.001785		0.040587		0.002682	
0.020673	-0.005472		-0.017140	0.000440	
0.013637		0.000241			0.002053
-0.000108		0.0		0.001482	0.058006
-0.021367		-0.013837		-0.004504	
	2125		0.032695		
-0.001140			0.048574		

-0.122320				0.010575	
-0.157058				-0.055247	
0.006476				-0.024198	
-0.004033				-0.054361	
-0.001798			-0.090650		
0.005080			-0.040494		
-0.038136				0.045453	
-0.012863		-0.037800			-0.062272
0.001405				-0.050698	
0.026467				0.039803	
-0.086003				0.018722	
-0.053730				-0.004403	
0.034906		-0.011419			-0.087098
0.044551		-0.019152			-0.009545
-0.048225		0.195843			-0.019403
0.005563		-0.135147			-0.038808
-0.030394		-0.013683			
-0.049435		0.008123			
0.200610		-0.023889			
-0.091178		0.055712		-0.013228	
-0.058888		0.029637			-0.005206
0.060710		0.013123			-0.021744
-0.061050		0.000642			
-0.020826		-0.241329			
-0.124323		-0.001091			
0.159584		0.075970			
-0.005820		0.049777		0.019224	
-0.024588		0.010413		0.033451	
-0.014392	-0.040721	-0.015724	-0.043229	-0.070458	-0.012003
-0.002050	0.007733		0.0		-0.330416
-0.000503		-0.026581		0.001572	0.0
-0.018617		0.0	0.041194		0.001785
0.035909	0.002664			-0.190455	0.011779
0.109529	-0.003117		0.015190		
-0.038217		0.003987		-0.000108	
0.0		-0.008058	-0.043345	-0.000267	-0.053101
-0.010146					

```

-----
: stage1_shap_values.csv
: 2025-12-11 06:39:48
-----

```

[CSV] 103 × 3 10

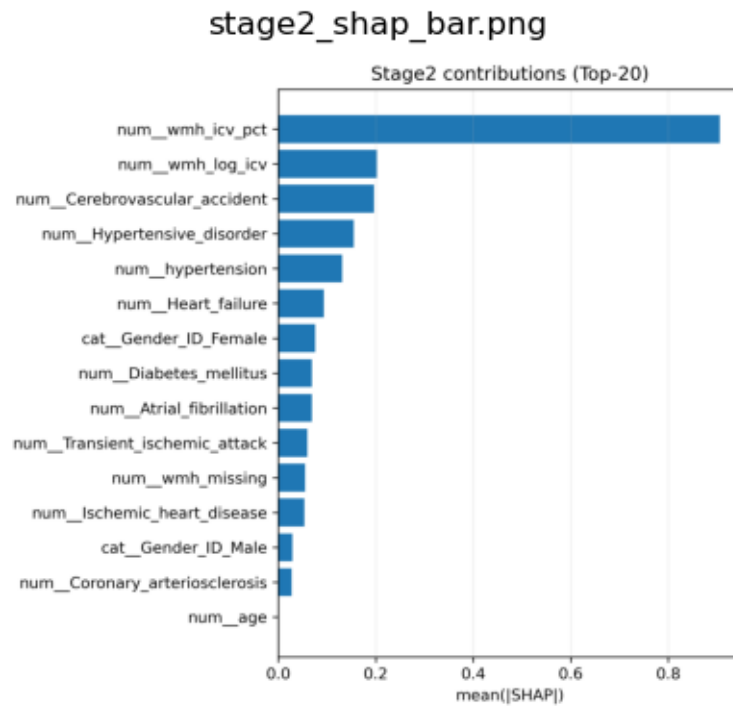
	feature	mean_abs_shap	mean_shap
	num_MMSE	0.193538	0.009166
	num_Cerebrovascular_accident	0.144728	0.005919

num__050_Right Inf Lat Vent__pct	0.140217	0.001491
num__118_Left Ent entorhinal area__pct	0.134004	-0.007065
num__051_Left Inf Lat Vent__pct	0.132246	0.001231
num__Brain_Left_0_Frontal_99__pct	0.118064	0.004361
num__Brain_Left_4_DeepNuclei_9__pct	0.092967	0.005101
num__049_Left Hippocampus__pct	0.091276	0.007771
num__172_Left PHG parahippocampal gyrus__pct	0.086784	-0.001344
num__LI_PCu	0.086562	0.004348

```

-----
: stage2_shap_bar.png
: 2025-12-11 06:39:49
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```



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-----
: stage2_shap_samples.csv
: 2025-12-11 06:39:49
-----

```

[CSV] 1318 × 16 10

sample_index_in_X num__wmh_icv_pct num__wmh_log_icv num__wmh_missing

num__hypertension	num__Hypertensive_disorder	num__Diabetes_mellitus	num__Atrial_fibrillation	num__Ischemic_heart_disease	num__Cerebrovascular_accident	num__Transient_ischemic_attack	num__Heart_failure	num__Coronary_arteriosclerosis	num__age	cat__Gender_ID_Female	cat__Gender_ID_Male
	0	3.069893	0.545670	-0.012984							
0.152238		-0.136515	0.121644								
-0.023649		0.274102		0.178153							
-0.047183	0.575292		0.408033	0.0							
0.115974	0.052113										
	1	-0.847595	-0.146412	-0.030086							
-0.136204		0.143394	0.036404								
-0.053536		-0.056558		0.560236							
0.349379	-0.091912		-0.006170	0.0							
-0.009646	-0.020595										
	2	0.575597	0.197067	-0.029766							
-0.131319		0.122634	-0.030062								
-0.053286		0.007277		-0.113992							
-0.020116	-0.046308		-0.028739	0.0							
-0.019336	0.025279										
	3	-0.679651	-0.016201	-0.019955							
0.049933		-0.215406	-0.021051								
-0.074895		-0.028163		-0.162663							
-0.046192	-0.046924		-0.003415	0.0							
0.032710	-0.025772										
	4	1.742481	0.485864	-0.032974							
-0.115195		0.128924	0.080774								
-0.053546		0.024693		-0.106026							
-0.011531	-0.040631		-0.027795	0.0							
0.030277	-0.020672										
	5	0.747782	0.096549	-0.033888							
-0.155981		0.108594	0.053536								
-0.040164		0.042554		-0.141239							
-0.009484	-0.042210		-0.013010	0.0							
-0.136344	-0.037059										
	6	0.697491	0.089717	-0.027392							
-0.146860		0.075124	-0.178983								
-0.019314		-0.049149		0.239779							
-0.032204	-0.043487		-0.023511	0.0							
0.115507	0.039513										
	7	1.601996	0.436930	-0.023142							
0.106064		0.149975	-0.047425								
-0.047623		0.037244		-0.079909							
-0.020372	-0.042874		-0.026448	0.0							
0.036821	0.038206										
	8	-1.249282	-0.190036	-0.099193							
0.130209		-0.137611	0.129513								
-0.030535		-0.003799		-0.185395							

-0.047456	-0.038352	-0.008922	0.0
0.099186	0.027035		
	9	-1.479216	-0.288324
0.054810		-0.094243	0.076457
-0.029136		-0.030028	-0.150663
-0.066874	-0.037551	-0.009346	0.0
-0.010489	-0.020419		

```

-----
: stage2_shap_values.csv
: 2025-12-11 06:39:49
-----

```

[CSV] 15 × 3

	feature	mean_abs_shap	mean_shap
	num_wmh_icv_pct	0.906033	0.009294
	num_wmh_log_icv	0.202818	0.000422
num_Cerebrovascular_accident		0.196147	-0.005705
num_Hypertensive_disorder		0.154991	-0.011313
num_hypertension		0.131173	0.008345
num_Heart_failure		0.093152	0.001439
cat_Gender_ID_Female		0.075931	0.001702
num_Diabetes_mellitus		0.069000	0.001603
num_Atrial_fibrillation		0.068532	-0.000436
num_Transient_ischemic_attack		0.059454	-0.000598
num_wmh_missing		0.055096	-0.002030
num_Ischemic_heart_disease		0.053870	0.000513
cat_Gender_ID_Male		0.028831	-0.000288
num_Coronary_arteriosclerosis		0.027562	0.001512
num_age		0.000000	0.000000

```

-----
: analysis_out/.ipynb_checkpoints
-----

```

[INFO] 24

```

-----
: analysis_out/interaction_plots
-----

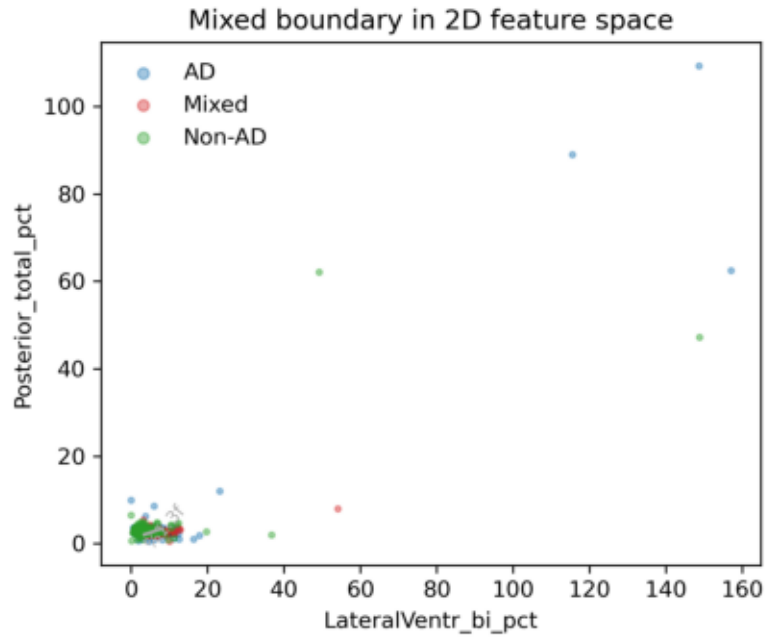
```

```

-----
: interact_2D_LateralVentr_bi_pct_Posterior_total_pct.png
: 2025-12-11 06:40:21
-----

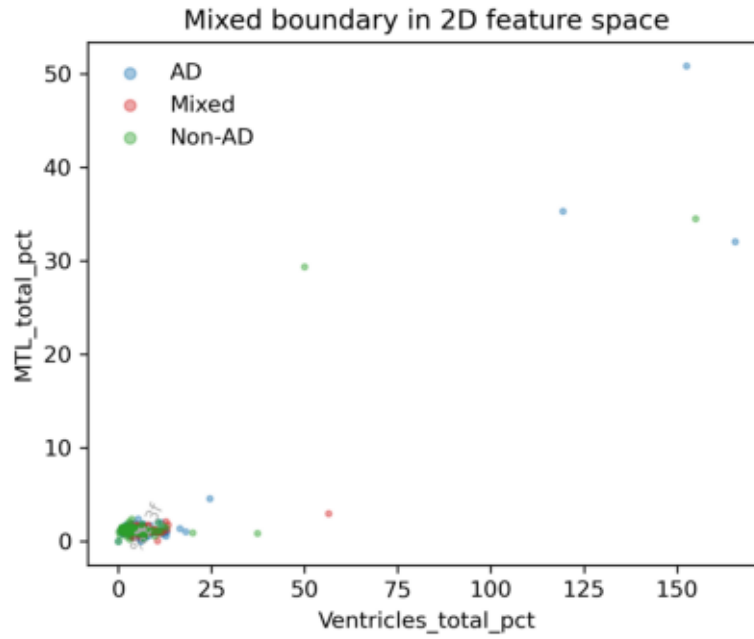
```

interact_2D_LateralVentr_bi_pct_Posterior_total_pct.png



: interact_2D_Ventricles_total_pct_MTL_total_pct.png
: 2025-12-11 06:40:21

interact_2D_Ventricles_total_pct_MTL_total_pct.png



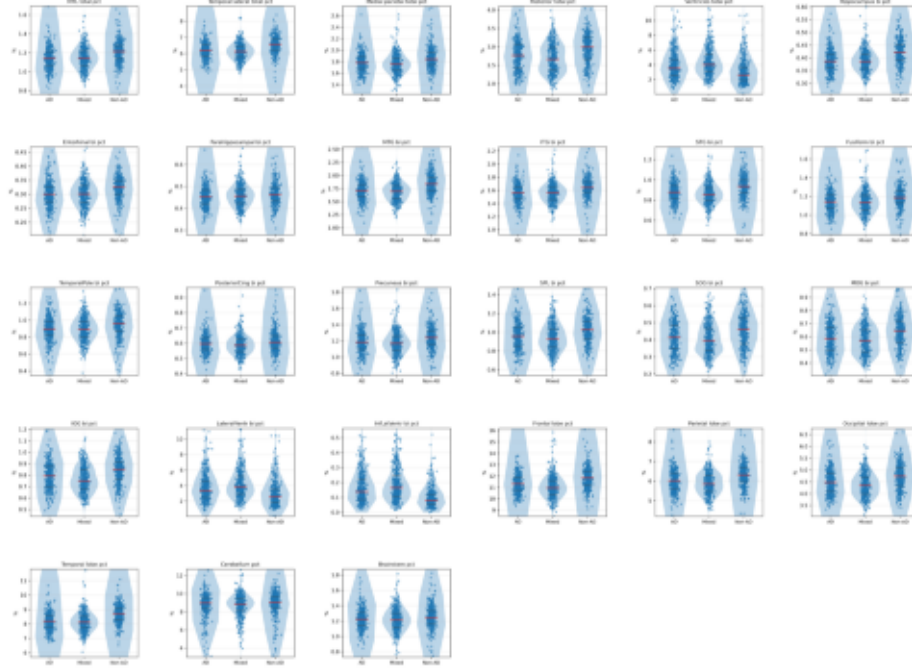
: analysis_out/interaction_plots/.ipynb_checkpoints

[INFO] 24

: panels_final

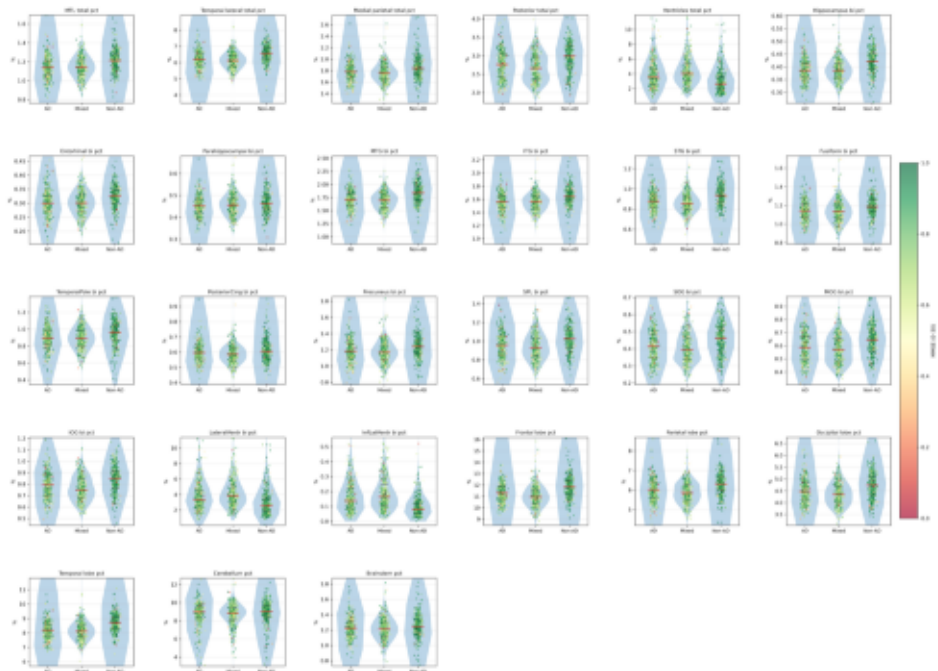
: panel_roi_dist.png
: 2025-12-11 06:40:00

panel_roi_dist.png



: panel_roi_mmse_colored.png
: 2025-12-11 06:40:05

panel_roi_mmse_colored.png



: panels_final/.ipynb_checkpoints

[INFO] 24

: fairness

: final3_ethnicity_perclass.csv
: 2025-12-11 06:30:28

[CSV] 57 × 7

ethnicity_code	ethnicity_label	class	precision	recall	f1	support
0	0	AD	0.921275	0.975198	0.947470	1008
0	0	Mixed	0.771812	0.660920	0.712074	348
0	0	Non-AD	0.760714	0.737024	0.748682	289
1	1	AD	0.958333	0.932432	0.945205	74

1	1	Mixed	0.571429	0.827586	0.676056	29
1	1	Non-AD	0.838710	0.619048	0.712329	42
2	2	AD	0.921053	0.933333	0.927152	75
2	2	Mixed	0.862069	0.757576	0.806452	33
2	2	Non-AD	0.652174	0.750000	0.697674	20
3	3	AD	0.930233	0.952381	0.941176	42
3	3	Mixed	0.692308	0.818182	0.750000	11
3	3	Non-AD	0.875000	0.636364	0.736842	11
4	4	AD	0.940741	0.954887	0.947761	133
4	4	Mixed	0.787879	0.702703	0.742857	37
4	4	Non-AD	0.807692	0.840000	0.823529	50
5	5	AD	0.959044	0.959044	0.959044	293
5	5	Mixed	0.773810	0.691489	0.730337	94
5	5	Non-AD	0.758929	0.833333	0.794393	102
6	6	AD	0.917647	0.951220	0.934132	82
6	6	Mixed	0.814815	0.687500	0.745763	32
6	6	Non-AD	0.666667	0.736842	0.700000	19
7	7	AD	1.000000	1.000000	1.000000	6
7	7	Mixed	0.000000	0.000000	0.000000	0
7	7	Non-AD	1.000000	0.500000	0.666667	4
8	8	AD	0.966102	0.966102	0.966102	59
8	8	Mixed	0.769231	0.526316	0.625000	19
8	8	Non-AD	0.640000	0.842105	0.727273	19
9	9	AD	0.939024	0.974684	0.956522	79
9	9	Mixed	0.523810	0.611111	0.564103	18
9	9	Non-AD	0.880000	0.709677	0.785714	31
10	10	AD	1.000000	1.000000	1.000000	17
10	10	Mixed	0.000000	0.000000	0.000000	0
10	10	Non-AD	1.000000	0.666667	0.800000	6
11	11	AD	0.960000	0.888889	0.923077	27
11	11	Mixed	0.714286	0.625000	0.666667	8
11	11	Non-AD	0.642857	0.818182	0.720000	11
12	12	AD	1.000000	0.800000	0.888889	10
12	12	Mixed	0.000000	0.000000	0.000000	0
12	12	Non-AD	1.000000	0.750000	0.857143	4
13	13	AD	0.900000	0.818182	0.857143	11
13	13	Mixed	0.333333	0.500000	0.400000	2
13	13	Non-AD	0.888889	0.888889	0.888889	9
14	14	AD	0.800000	1.000000	0.888889	4
14	14	Mixed	0.333333	0.333333	0.333333	3
14	14	Non-AD	0.857143	0.750000	0.800000	8
15	15	AD	1.000000	0.833333	0.909091	6
15	15	Mixed	0.000000	0.000000	0.000000	0
15	15	Non-AD	1.000000	0.500000	0.666667	2
16	16	AD	1.000000	1.000000	1.000000	4
16	16	Mixed	0.000000	0.000000	0.000000	0
16	16	Non-AD	0.000000	0.000000	0.000000	0
17	17	AD	0.000000	0.000000	0.000000	0

17	17	Mixed	0.000000	0.000000	0.000000	0
17	17	Non-AD	1.000000	0.500000	0.666667	2
18	18	AD	1.000000	1.000000	1.000000	1
18	18	Mixed	0.000000	0.000000	0.000000	0
18	18	Non-AD	0.000000	0.000000	0.000000	0

```

-----
: final3_ethnicity_summary.csv
: 2025-12-11 06:30:28
-----

```

[CSV] 19 × 6

ethnicity_code	ethnicity_label	n	accuracy	macro_recall	macro_f1
0		0 1645	0.866869	0.791047	0.802742
1		1 145	0.820690	0.793022	0.777864
2		2 128	0.859375	0.813636	0.810426
3		3 64	0.875000	0.802309	0.809340
4		4 220	0.886364	0.832530	0.838049
5		5 489	0.881391	0.827956	0.827925
6		6 133	0.857143	0.791854	0.793298
7		7 10	0.800000	0.500000	0.555556
8		8 97	0.855670	0.778174	0.772791
9		9 128	0.859375	0.765157	0.768780
10		10 23	0.913043	0.555556	0.600000
11		11 46	0.826087	0.777357	0.769915
12		12 14	0.785714	0.516667	0.582011
13		13 22	0.818182	0.735690	0.715344
14		14 15	0.733333	0.694444	0.674074
15		15 8	0.750000	0.444444	0.525253
16		16 4	1.000000	0.333333	0.333333
17		17 2	0.500000	0.166667	0.222222
18		18 1	1.000000	0.333333	0.333333

```

-----
: final3_gender_perclass.csv
: 2025-12-11 06:30:28
-----

```

[CSV] 6 × 7

gender_code	gender_label	class	precision	recall	f1	support
0	Gender_0	AD	0.947771	0.961240	0.954458	774
0	Gender_0	Mixed	0.752768	0.733813	0.743169	278
0	Gender_0	Non-AD	0.768683	0.757895	0.763251	285
1	Gender_1	AD	0.925809	0.963054	0.944064	1218
1	Gender_1	Mixed	0.709091	0.637602	0.671449	367
1	Gender_1	Non-AD	0.779255	0.755155	0.767016	388

```
-----
: final3_gender_summary.csv
: 2025-12-11 06:30:28
-----
```

[CSV] 2 × 6

gender_code	gender_label	n	accuracy	macro_recall	macro_f1
0	Gender_0	1337	0.870606	0.817649	0.820293
1	Gender_1	1973	0.861632	0.785270	0.794176

```
-----
: stage1_stage1_ethnicity.csv
: 2025-12-11 06:30:28
-----
```

[CSV] 19 × 9

ethnicity_code	ethnicity_label	n	accuracy	precision_pos	recall_pos
0		1645	0.933739	0.921275	0.975198
0.947470	0.921665	0.986122			
1		145	0.944828	0.958333	0.932432
0.945205	0.945089	0.983251			
2		128	0.914062	0.921053	0.933333
0.927152	0.910063	0.986415			
3		64	0.921875	0.930233	0.952381
0.941176	0.908009	0.965368			
4		220	0.936364	0.940741	0.954887
0.947761	0.931467	0.983666			
5		489	0.950920	0.959044	0.959044
0.959044	0.948910	0.987915			
6		133	0.917293	0.917647	0.951220
0.934132	0.906982	0.981110			
7		10	1.000000	1.000000	1.000000
1.000000	1.000000	1.000000			
8		97	0.958763	0.966102	0.966102
0.966102	0.956735	0.989741			
9		128	0.945312	0.939024	0.974684
0.956522	0.936321	0.994833			
10		23	1.000000	1.000000	1.000000
1.000000	1.000000	1.000000			
11		46	0.913043	0.960000	0.888889
0.923077	0.918129	0.986355			
12		14	0.857143	1.000000	0.800000
0.888889	0.900000	1.000000			
13		22	0.863636	0.900000	0.818182

0.857143	0.863636	0.950413					
	14		14	15	0.933333	0.800000	1.000000
0.888889	0.954545	1.000000					
	15		15	8	0.875000	1.000000	0.833333
0.909091	0.916667	1.000000					
	16		16	4	1.000000	1.000000	1.000000
1.000000	1.000000	NaN					
	17		17	2	1.000000	NaN	NaN
NaN	1.000000	NaN					
	18		18	1	1.000000	1.000000	1.000000
1.000000	1.000000	NaN					

```
-----
: stage1_stage1_gender.csv
: 2025-12-11 06:30:28
-----
```

[CSV] 2 × 9

gender_code	gender_label	n	accuracy	precision_pos	recall_pos	f1_pos
balanced_acc	roc_auc					
0	Gender_0	1337	0.946896	0.947771	0.961240	0.954458
0.944208	0.988893					
1	Gender_1	1973	0.929549	0.925809	0.963054	0.944064
0.919275	0.983137					

```
-----
: stage2_stage2_ethnicity.csv
: 2025-12-11 06:30:28
-----
```

[CSV] 17 × 9

ethnicity_code	ethnicity_label	n	accuracy	precision_pos	recall_pos
f1_pos	balanced_acc	roc_auc			
0		0	637	0.511774	0.566308
0.503987	0.517669	0.518504			
1		1	71	0.577465	0.487179
0.558824	0.589491	0.627258			
2		2	53	0.509434	0.666667
0.518519	0.537121	0.587121			
3		3	22	0.500000	0.500000
0.476190	0.500000	0.471074			
4		4	87	0.540230	0.461538
0.473684	0.533243	0.484595			
5		5	196	0.540816	0.520833
0.526316	0.540467	0.503598			
6		6	51	0.470588	0.608696
					0.437500

0.509091	0.481908	0.490132					
	7		7	4	0.250000	0.000000	NaN
NaN	0.250000	NaN					
	8		8	38	0.421053	0.421053	0.421053
0.421053	0.421053	0.396122					
	9		9	49	0.551020	0.409091	0.500000
0.450000	0.540323	0.603943					
	10		10	6	0.833333	0.000000	NaN
NaN	0.833333	NaN					
	11		11	19	0.368421	0.250000	0.250000
0.250000	0.352273	0.250000					
	12		12	4	0.750000	0.000000	NaN
NaN	0.750000	NaN					
	13		13	11	0.636364	0.000000	0.000000
NaN	0.388889	0.444444					
	14		14	11	0.454545	0.000000	0.000000
NaN	0.312500	0.375000					
	15		15	2	0.500000	0.000000	NaN
NaN	0.500000	NaN					
	17		17	2	0.000000	0.000000	NaN
NaN	0.000000	NaN					

```
-----
: stage2_stage2_gender.csv
: 2025-12-11 06:30:28
-----
```

[CSV] 2 × 9

gender_code	gender_label	n	accuracy	precision_pos	recall_pos	f1_pos
0	Gender_0	563	0.507993	0.502075	0.435252	0.466281
0.507100	0.519658					
1	Gender_1	755	0.521854	0.508523	0.487738	0.497914
0.520931	0.503810					

```
-----
: fairness_WMH
-----
```

[INFO] 24

```
#####
# : hier_dir      24h (hier_out)
#####
```

: (root)

: feature_list_stage2.csv
: 2025-12-11 06:39:37

[CSV] 14 × 1

feature_name
wmh_icv_pct
wmh_log_icv
wmh_missing
hypertension
Hypertensive_disorder
Diabetes_mellitus
Atrial_fibrillation
Ischemic_heart_disease
Cerebrovascular_accident
Transient_ischemic_attack
Heart_failure
Coronary_arteriosclerosis
age
Gender_ID

: final_confusion_matrix.csv
: 2025-12-11 06:39:26

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1946	17	29
true_Mixed	137	302	206
true_Non-AD	52	226	395

: final_confusion_matrix_thr0.50.csv
: 2025-12-11 06:39:40

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1946	19	27

true_Mixed	137	407	101
true_Non-AD	52	130	491

: final_confusion_matrix_thr0.508.csv
: 2025-12-11 06:39:42

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1939	23	30
true_Mixed	125	416	104
true_Non-AD	46	130	497

: final_confusion_matrix_thr0.529.csv
: 2025-12-11 06:39:42

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1922	32	38
true_Mixed	107	433	105
true_Non-AD	36	130	507

: final_confusion_matrix_thr0.532.csv
: 2025-12-11 06:39:42

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1922	32	38
true_Mixed	104	435	106
true_Non-AD	35	130	508

: final_confusion_matrix_thr0.538.csv
: 2025-12-11 06:39:42

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1917	33	42

true_Mixed	101	438	106
true_Non-AD	34	130	509

: final_confusion_matrix_thr0.55.csv
: 2025-12-11 06:39:40

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1906	36	50
true_Mixed	92	445	108
true_Non-AD	34	130	509

: final_confusion_matrix_thr0.568.csv
: 2025-12-11 06:39:42

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1880	47	65
true_Mixed	75	456	114
true_Non-AD	27	130	516

: final_confusion_matrix_thr0.583.csv
: 2025-12-11 06:39:43

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1862	60	70
true_Mixed	65	459	121
true_Non-AD	22	131	520

: final_confusion_matrix_thr0.584.csv
: 2025-12-11 06:39:43

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1861	60	71

true_Mixed	64	459	122
true_Non-AD	21	131	521

: final_confusion_matrix_thr0.592.csv
: 2025-12-11 06:39:43

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1846	67	79
true_Mixed	60	462	123
true_Non-AD	19	131	523

: final_confusion_matrix_thr0.598.csv
: 2025-12-11 06:39:43

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1838	70	84
true_Mixed	56	464	125
true_Non-AD	15	132	526

: final_confusion_matrix_thr0.599.csv
: 2025-12-11 06:39:43

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1838	70	84
true_Mixed	55	464	126
true_Non-AD	15	132	526

: final_confusion_matrix_thr0.60.csv
: 2025-12-11 06:39:41

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1836	70	86

true_Mixed	55	464	126
true_Non-AD	15	132	526

```
-----
: final_confusion_matrix_thr0.65.csv
: 2025-12-11 06:39:41
-----
```

[CSV] 3 × 4

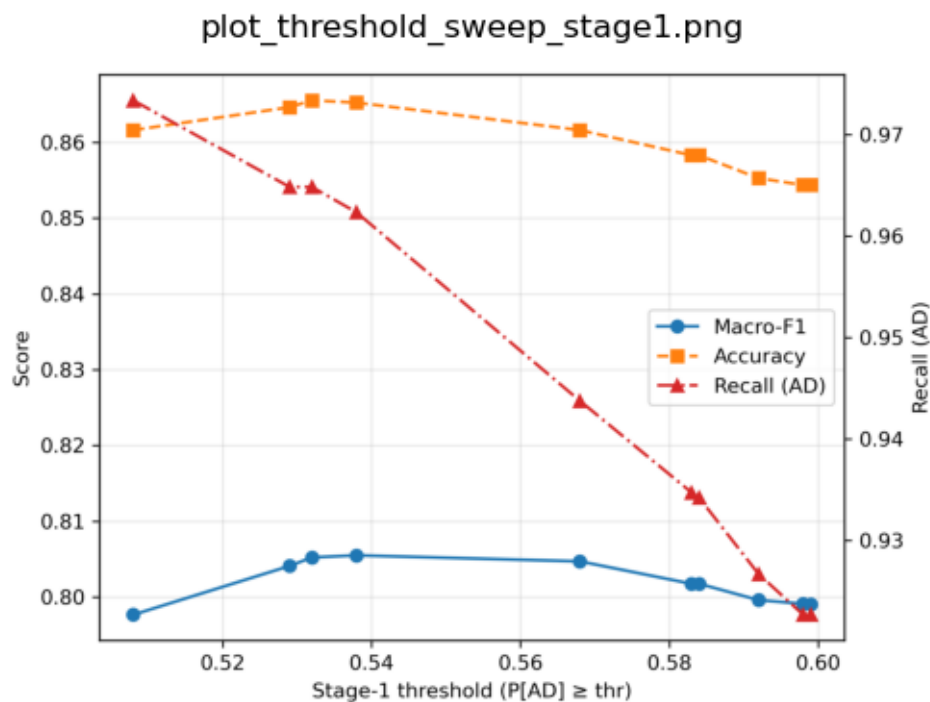
Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1738	111	143
true_Mixed	25	490	130
true_Non-AD	6	133	534

```
-----
: final_confusion_matrix_thr0.70.csv
: 2025-12-11 06:39:41
-----
```

[CSV] 3 × 4

Unnamed: 0	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1559	192	241
true_Mixed	11	497	137
true_Non-AD	2	134	537

```
-----
: plot_threshold_sweep_stage1.png
: 2025-12-11 06:39:43
-----
```



```
-----
: preds_hierarchical.csv
: 2025-12-11 06:39:26
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.718330
10331478	Non-AD	0.178709	Mixed	0.661639
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Non-AD	0.445762
10425175	Non-AD	0.243892	Non-AD	0.419252
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	AD	NaN
10426958	Mixed	0.135618	Non-AD	0.448564
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.50.csv
: 2025-12-11 06:39:40
-----
```


[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	AD	NaN
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

: preds_hierarchical_thr0.508.csv
: 2025-12-11 06:39:42

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	AD	NaN
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

: preds_hierarchical_thr0.529.csv
: 2025-12-11 06:39:42

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541

10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.532.csv
: 2025-12-11 06:39:42
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.538.csv
: 2025-12-11 06:39:42
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.55.csv
: 2025-12-11 06:39:40
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

: preds_hierarchical_thr0.568.csv
: 2025-12-11 06:39:43

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

: preds_hierarchical_thr0.583.csv
: 2025-12-11 06:39:43

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN

10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.584.csv
: 2025-12-11 06:39:43
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.592.csv
: 2025-12-11 06:39:43
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.598.csv
: 2025-12-11 06:39:43
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.599.csv
: 2025-12-11 06:39:43
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.60.csv
: 2025-12-11 06:39:41
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153

10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.65.csv
: 2025-12-11 06:39:41
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	AD	NaN
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: preds_hierarchical_thr0.70.csv
: 2025-12-11 06:39:41
-----
```

[CSV] 3310 × 5 10

BrcId	true_diag	P_AD	final_pred	P_Mixed_given_nonAD
10309796	AD	0.828000	AD	NaN
10415039	Mixed	0.149271	Mixed	0.994657
10331478	Non-AD	0.178709	Non-AD	0.412081
10426976	AD	0.751963	AD	NaN
10019261	Mixed	0.244223	Mixed	0.600816
10425175	Non-AD	0.243892	Non-AD	0.213541
10432969	AD	0.699629	Non-AD	0.474046
10433396	AD	0.516806	Non-AD	0.040153
10426958	Mixed	0.135618	Mixed	0.884788
10418222	AD	0.705930	AD	NaN

```
-----
: stage1_metrics.json
: 2025-12-11 06:39:37
-----
```

```
{
  "f1": 0.6453259104616098,
```

```

    "balanced_acc": 0.49709802792353025,
    "auc": 0.4974893885100341
}

```

```

-----
: stage1_report.txt
: 2025-12-11 06:39:37
-----

```

	precision	recall	f1-score	support
0	0.394	0.296	0.338	1318
1	0.600	0.698	0.645	1992
accuracy			0.538	3310
macro avg	0.497	0.497	0.492	3310
weighted avg	0.518	0.538	0.523	3310

```

-----
: stage2_best_params.json
: 2025-12-11 06:39:24
-----

```

```

{
  "clf__subsample": 0.7,
  "clf__scale_pos_weight": 1.0434108527131782,
  "clf__reg_lambda": 1.0,
  "clf__reg_alpha": 0.5,
  "clf__n_estimators": 900,
  "clf__min_child_weight": 7,
  "clf__max_depth": 2,
  "clf__learning_rate": 0.03,
  "clf__gamma": 1.0,
  "clf__colsample_bytree": 0.6
}

```

```

-----
: stage2_metrics.json
: 2025-12-11 06:39:38
-----

```

```

{
  "f1": 0.5093167701863354,
  "balanced_acc": 0.5202368199776541,
  "auc": 0.516654572261193
}

```

: stage2_report.txt
: 2025-12-11 06:39:38

	precision	recall	f1-score	support
0	0.530	0.532	0.531	673
1	0.510	0.509	0.509	645
accuracy			0.520	1318
macro avg	0.520	0.520	0.520	1318
weighted avg	0.520	0.520	0.520	1318

: threshold_sweep_stage1_summary.csv
: 2025-12-11 06:39:43

[CSV] 10 × 4

thr1	macro_f1	accuracy	recall_AD
0.538	0.805512	0.865257	0.962349
0.532	0.805236	0.865559	0.964859
0.568	0.804717	0.861631	0.943775
0.529	0.804108	0.864653	0.964859
0.584	0.801764	0.858308	0.934237
0.583	0.801754	0.858308	0.934739
0.592	0.799605	0.855287	0.926707
0.598	0.799113	0.854381	0.922691
0.599	0.799017	0.854381	0.922691
0.508	0.797667	0.861631	0.973394

: .ipynb_checkpoints

[INFO] 24
[WARN] surv_out / 24h (None)

=====
24
=====


```

[38]: # =====

import numpy as np

import pandas as pd

import matplotlib.pyplot as plt

import seaborn as sns

from pathlib import Path

OUT = Path("surv_out") / "analysis_out"

OUT.mkdir(parents=True, exist_ok=True)

#      =true =pred

labels = ["AD", "Mixed", "Non-AD"]

cm = np.array([

    [1917, 33, 42],    # true_AD

    [ 101, 438, 106],  # true_Mixed

    [ 34, 130, 509]    # true_Non-AD

], dtype=int)

cm_df = pd.DataFrame(cm, index=[f"true_{l}" for l in labels],

                      columns=[f"pred_{l}" for l in labels])

print("Confusion matrix:\n", cm_df)

#

support = cm.sum(axis=1)

total   = cm.sum()

# Precision / Recall / F1

per_cls_rows = []

for i, lab in enumerate(labels):

```

```

TP = cm[i, i]

FN = cm[i, :].sum() - TP

FP = cm[:, i].sum() - TP

TN = total - TP - FN - FP

prec = TP / (TP + FP) if (TP + FP) > 0 else 0.0

rec = TP / (TP + FN) if (TP + FN) > 0 else 0.0

f1 = 2 * prec * rec / (prec + rec) if (prec + rec) > 0 else 0.0

per_cls_rows.append({

    "class": lab,

    "support": int(support[i]),

    "precision": float(prec),

    "recall": float(rec),

    "f1": float(f1)

})

per_cls = pd.DataFrame(per_cls_rows)

per_cls.set_index("class", inplace=True)

print("\nPer-class metrics:\n", per_cls)

#

macro_precision = per_cls["precision"].mean()

macro_recall = per_cls["recall"].mean()

macro_f1 = per_cls["f1"].mean()

weighted_f1 = (per_cls["f1"] * per_cls["support"]).sum() / support.sum()

balanced_acc = macro_recall

```

```

overall_acc      = np.trace(cm) / total

overall_rows = [{

    "accuracy": float(overall_acc),

    "balanced_accuracy": float(balanced_acc),

    "macro_precision": float(macro_precision),

    "macro_recall": float(macro_recall),

    "macro_f1": float(macro_f1),

    "weighted_f1": float(weighted_f1),

    "N": int(total)

}]

overall = pd.DataFrame(overall_rows)

print("\nOverall:\n", overall)

#

cm_df.to_csv(OUT / "overall_from_cm_confusion.csv", index=True)

per_cls.to_csv(OUT / "overall_from_cm_per_class.csv", index=True)

overall.to_csv(OUT / "overall_from_cm_summary.csv", index=False)

print("\n[Saved]")

print(OUT / "overall_from_cm_confusion.csv")

print(OUT / "overall_from_cm_per_class.csv")

print(OUT / "overall_from_cm_summary.csv")

#

norm_cm = cm / support.reshape(-1, 1)

plt.figure(figsize=(5.2, 4.4))

sns.heatmap(norm_cm, annot=True, fmt=".2f", cmap="Blues",

```

```

xticklabels=labels, yticklabels=labels)

plt.xlabel("Predicted"); plt.ylabel("True"); plt.title("Normalized confusion_
↪(row-wise)")

plt.tight_layout()

plt.savefig(OUT / "overall_from_cm_heatmap.png", dpi=300)

plt.close()

print(OUT / "overall_from_cm_heatmap.png")

```

Confusion matrix:

	pred_AD	pred_Mixed	pred_Non-AD
true_AD	1917	33	42
true_Mixed	101	438	106
true_Non-AD	34	130	509

Per-class metrics:

	support	precision	recall	f1
class				
AD	1992	0.934211	0.962349	0.948071
Mixed	645	0.728785	0.679070	0.703050
Non-AD	673	0.774734	0.756315	0.765414

Overall:

	accuracy	balanced_accuracy	macro_precision	macro_recall	macro_f1	\
0	0.865257	0.799245	0.812577	0.799245	0.805512	

	weighted_f1	N
0	0.863187	3310

[Saved]

```

surv_out/analysis_out/overall_from_cm_confusion.csv
surv_out/analysis_out/overall_from_cm_per_class.csv
surv_out/analysis_out/overall_from_cm_summary.csv
surv_out/analysis_out/overall_from_cm_heatmap.png

```

[]:

[39]: !jupyter nbconvert --to pdf baseline_dignosis.ipynb

```

[NbConvertApp] Converting notebook baseline_dignosis.ipynb to pdf
[NbConvertApp] Support files will be in baseline_dignosis_files/

```

```
[NbConvertApp] Making directory ./baseline_dignosis_files
[NbConvertApp] Writing 804643 bytes to notebook.tex
[NbConvertApp] Building PDF
[NbConvertApp] Running xelatex 3 times: ['xelatex', 'notebook.tex', '-quiet']
[NbConvertApp] Running bibtex 1 time: ['bibtex', 'notebook']
[NbConvertApp] WARNING | bibtex had problems, most likely because there were no
citations
[NbConvertApp] PDF successfully created
[NbConvertApp] Writing 2511765 bytes to baseline_dignosis.pdf
```

```
[ ]:
```

```
[40]: !pip install python-pptx
```

```
Requirement already satisfied: python-pptx in /opt/conda/lib/python3.11/site-
packages (1.0.2)
Requirement already satisfied: Pillow>=3.3.2 in /opt/conda/lib/python3.11/site-
packages (from python-pptx) (10.1.0)
Requirement already satisfied: XlsxWriter>=0.5.7 in
/opt/conda/lib/python3.11/site-packages (from python-pptx) (3.2.9)
Requirement already satisfied: lxml>=3.1.0 in /opt/conda/lib/python3.11/site-
packages (from python-pptx) (6.0.2)
Requirement already satisfied: typing-extensions>=4.9.0 in
/opt/conda/lib/python3.11/site-packages (from python-pptx) (4.15.0)
```

```
[41]: import matplotlib.pyplot as plt

import seaborn as sns

from sklearn.model_selection import cross_val_predict, StratifiedGroupKFold

from sklearn.calibration import calibration_curve, CalibratedClassifierCV

from sklearn.metrics import brier_score_loss

import numpy as np

import pandas as pd

# =====

# Daniel's diagnostic module:

# 1. Probability histogram (bimodality check)

# 2. Calibration curve comparison (Raw vs Isotonic)
```

```

# 3. Misclassification analysis (missing data & threshold proximity)

# =====

def daniel_diagnostic_suite(model, X, y, groups, df_raw, stage_name,
    ↪best_thresh=0.5):

    print(f"\n{'='*50}")

    print(f"Running diagnostics for: {stage_name} (Threshold = {best_thresh})")

    print(f"{'='*50}")

    # -----

    # A. Obtain out-of-fold prediction probabilities

    # -----

    print("[*] Computing cross-validated prediction probabilities...")

    cv = StratifiedGroupKFold(n_splits=5, shuffle=True, random_state=42)

    try:

        y_prob = cross_val_predict(

            model, X, y, groups=groups, cv=cv,

            method="predict_proba", n_jobs=-1

       )[: , 1]

    except Exception as e:

        print(f"[!] CV prediction failed. Falling back to in-sample predictions,
    ↪(may overfit). Error: {e}")

        y_prob = model.predict_proba(X)[: , 1]

    # -----

    # B. Probability histogram (bimodality inspection)

    # -----

    plt.figure(figsize=(10, 5))

```

```

sns.histplot(y_prob[y == 0], color="tab:blue", label="True Negative",
             kde=True, stat="density", alpha=0.5)

sns.histplot(y_prob[y == 1], color="tab:orange", label="True Positive",
             kde=True, stat="density", alpha=0.5)

plt.axvline(best_thresh, color="red", linestyle="--",
            label=f"Threshold = {best_thresh}")

plt.title(f"{stage_name}: Prediction Probability Histogram\n(Ideal shape = bimodal)")

plt.xlabel("Predicted Probability")

plt.legend()

plt.show()

hesitation_ratio = ((y_prob > 0.4) & (y_prob < 0.6)).mean()

print(f"> Histogram summary: {hesitation_ratio:.1%} of samples fall in the ambiguous range (0.4-0.6).")

if hesitation_ratio > 0.2:

    print(" -> Warning: Model displays notable uncertainty around the mid-probability range.")

else:

    print(" -> Good: Model displays clear bimodality (confident predictions).")

# -----

# C. Calibration curve analysis

# -----

print("\n[*] Comparing calibration: Raw vs Isotonic...")

prob_true_raw, prob_pred_raw = calibration_curve(y, y_prob, n_bins=10)

```

```

brier_raw = brier_score_loss(y, y_prob)

iso_model = CalibratedClassifierCV(model, method="isotonic", cv=cv)

try:

    y_prob_iso = cross_val_predict(

        iso_model, X, y, groups=groups, cv=cv,

        method="predict_proba", n_jobs=-1

    )[:, 1]

    prob_true_iso, prob_pred_iso = calibration_curve(y, y_prob_iso,
↪ n_bins=10)

    brier_iso = brier_score_loss(y, y_prob_iso)

    plt.figure(figsize=(6, 6))

    plt.plot([0, 1], [0, 1], "k:", label="Perfect calibration")

    plt.plot(prob_pred_raw, prob_true_raw, "s-", label=f"Raw
↪ (Brier={brier_raw:.4f})")

    plt.plot(prob_pred_iso, prob_true_iso, "o--", label=f"Isotonic
↪ (Brier={brier_iso:.4f})")

    plt.title(f"{stage_name}: Calibration Comparison")

    plt.xlabel("Mean Predicted Probability")

    plt.ylabel("Observed Fraction of Positives")

    plt.legend()

    plt.show()

    print(f"> Brier Score - Raw: {brier_raw:.4f}, Isotonic: {brier_iso:.
↪ 4f}")

    if brier_iso >= brier_raw:

        print(" -> Isotonic calibration does NOT improve model reliability.
↪ Raw probabilities preferred.")

```



```

else:

    print(" -> Isotonic calibration improves calibration.")

except Exception as e:

    print(f"[!] Isotonic calibration skipped due to error: {e}")

# -----

# D. Misclassification analysis

# -----

print(f"\n[*] Misclassification analysis...")

y_pred_class = (y_prob >= best_thresh).astype(int)

mis_mask = (y != y_pred_class)

correct_mask = ~mis_mask

print(f" - Total misclassified samples: {mis_mask.sum()}")

# ---- D1. Missing data check ----

try:

    raw_subset = df_raw.loc[X.index]    # index alignment

    missing_counts = raw_subset.isnull().sum(axis=1)

    avg_miss_wrong = missing_counts[mis_mask].mean()

    avg_miss_right = missing_counts[correct_mask].mean()

    print(f" - Avg missing values (misclassified): {avg_miss_wrong:.2f}")

    print(f" - Avg missing values (correct): {avg_miss_right:.2f}")

    plt.figure(figsize=(5, 4))

    sns.boxplot(

        x=[0]*correct_mask.sum() + [1]*mis_mask.sum(),

```

```

        y=pd.concat([missing_counts[correct_mask],
↪missing_counts[mis_mask]]),

        palette=["green", "red"]

    )

    plt.xticks([0, 1], ["Correct", "Misclassified"])

    plt.ylabel("Count of Missing Values (NaNs)")

    plt.title(f"{stage_name}: Missing Data Distribution")

    plt.show()

    if avg_miss_wrong > avg_miss_right * 1.1:

        print(" -> Misclassified cases tend to have substantially more
↪missing data.")

    else:

        print(" -> Missingness is similar between correct and incorrect
↪cases.")

    except Exception as e:

        print(f"[!] Missing-data analysis skipped (index alignment issue): {e}")

# ---- D2. Threshold-distance analysis ----

distance = np.abs(y_prob - best_thresh)

close_call_mask = distance < 0.05

wrong_and_close = (mis_mask & close_call_mask).sum()

proportion_close = wrong_and_close / mis_mask.sum()

    print(f" - Fraction of misclassifications near threshold (<0.05):
↪{proportion_close:.1%}")

plt.figure(figsize=(6, 4))

sns.kdeplot(distance[mis_mask], color="red", label="Misclassified",

            fill=True, alpha=0.3)

```

```

sns.kdeplot(distance[correct_mask], color="green", label="Correct",
             fill=True, alpha=0.3)

plt.xlabel(f"Distance to threshold {best_thresh}")

plt.title(f"{stage_name}: Threshold-Proximity Error Analysis")

plt.legend()

plt.show()

if proportion_close > 0.2:

    print(" -> A notable portion of errors occur near the decision_
↳threshold (model uncertainty).")

else:

    print(" -> Errors are widely distributed; not primarily due to the_
↳threshold cut-off.")

# =====

# Execute diagnostics

# =====

# Stage-1

if "pipe1" in locals() and "X1" in locals():

    daniel_diagnostic_suite(

        model=pipe1,

        X=X1,

        y=y1,

        groups=groups,

        df_raw=df,

        stage_name="Stage-1 (AD vs Others)",

```

```

        best_thresh=0.538

    )

# Stage-2
if "pipe2" in locals() and "X2" in locals():

    daniel_diagnostic_suite(

        model=pipe2,

        X=X2,

        y=y2,

        groups=groups2,

        df_raw=df2,

        stage_name="Stage-2 (Mixed vs Non-AD)",

        best_thresh=0.5

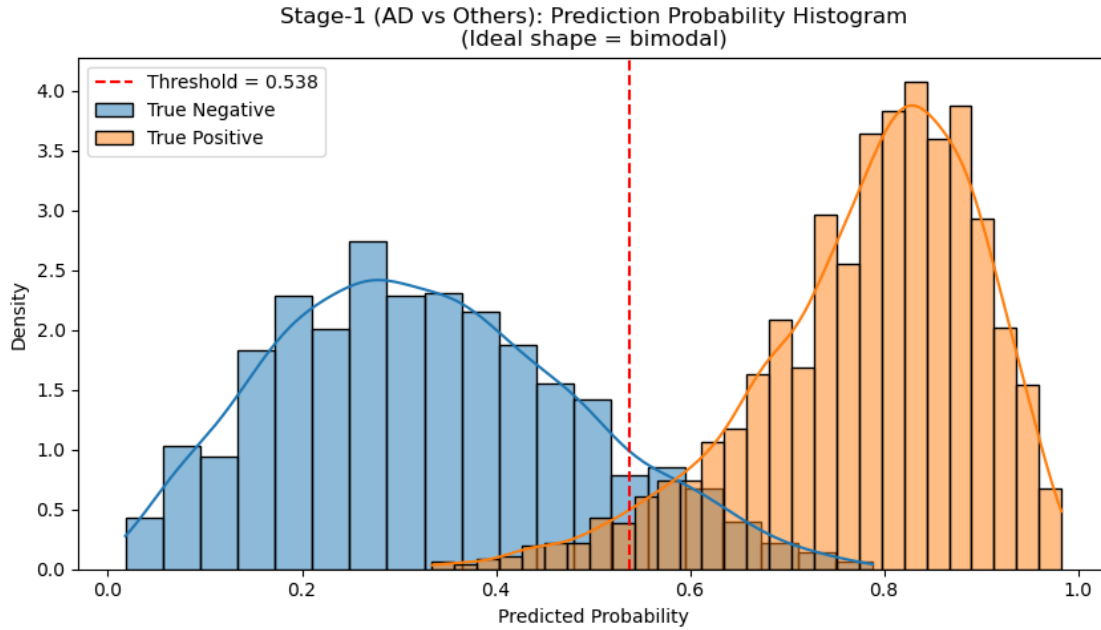
    )

```

```

=====
Running diagnostics for: Stage-1 (AD vs Others) (Threshold = 0.538)
=====
[*] Computing cross-validated prediction probabilities...
[!] CV prediction failed. Falling back to in-sample predictions (may overfit).
Error: Found input variables with inconsistent numbers of samples: [3310, 3310,
5]

```



```
> Histogram summary: 14.7% of samples fall in the ambiguous range (0.4-0.6).
  -> Good: Model displays clear bimodality (confident predictions).
```

```
[*] Comparing calibration: Raw vs Isotonic...
```

```
[!] Isotonic calibration skipped due to error: Found input variables with
inconsistent numbers of samples: [3310, 3310, 5]
```

```
[*] Misclassification analysis...
```

```
  - Total misclassified samples: 210
```

```
[!] Missing-data analysis skipped (index alignment issue): '[3318, 3326, 3333,
3340, 3341, 3349, 3350, 3351, 3362, 3368, 3379, 3389, 3402, 3409, 3416, 3433,
3441, 3444, 3451, 3454, 3462, 3464, 3473, 3479, 3482, 3506, 3522, 3524, 3527,
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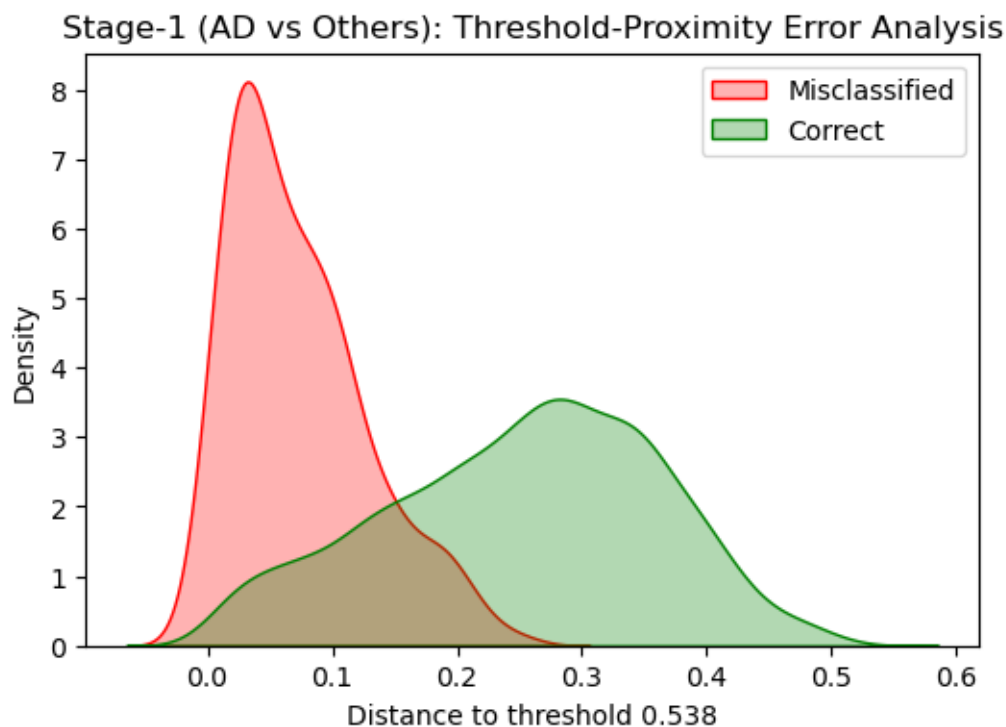
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10763, 10768, 10771, 10774, 10777, 10779, 10783, 10789, 10798, 10802, 10806,
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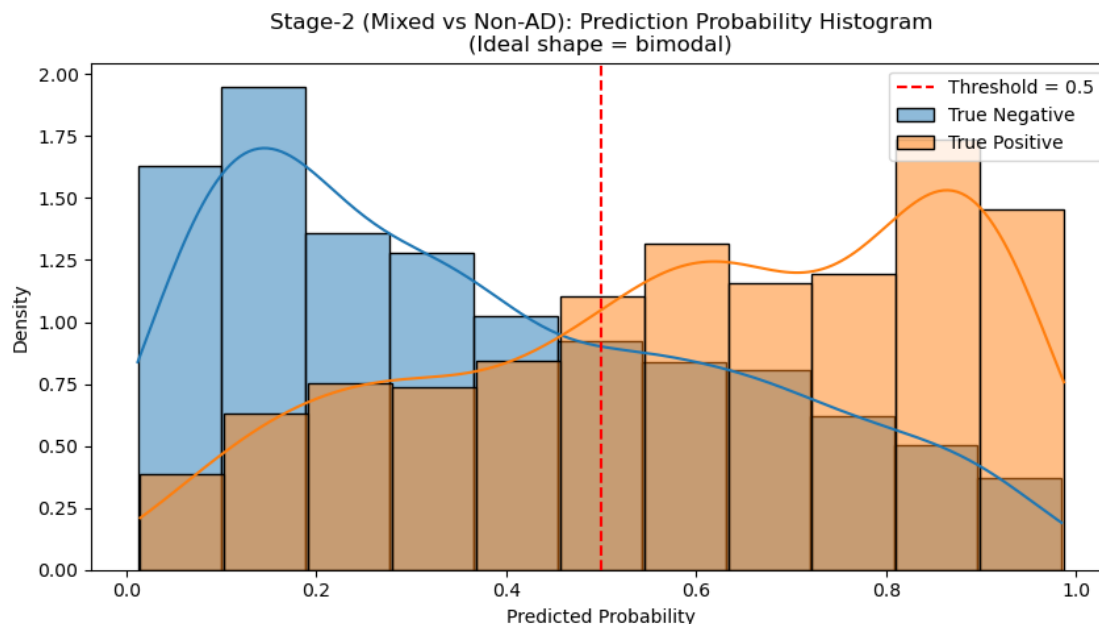
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- Fraction of misclassifications near threshold (<0.05): 44.3%



-> A notable portion of errors occur near the decision threshold (model uncertainty).

```
=====
Running diagnostics for: Stage-2 (Mixed vs Non-AD) (Threshold = 0.5)
=====
[*] Computing cross-validated prediction probabilities...
```



```
> Histogram summary: 19.0% of samples fall in the ambiguous range (0.4-0.6).
-> Good: Model displays clear bimodality (confident predictions).
```

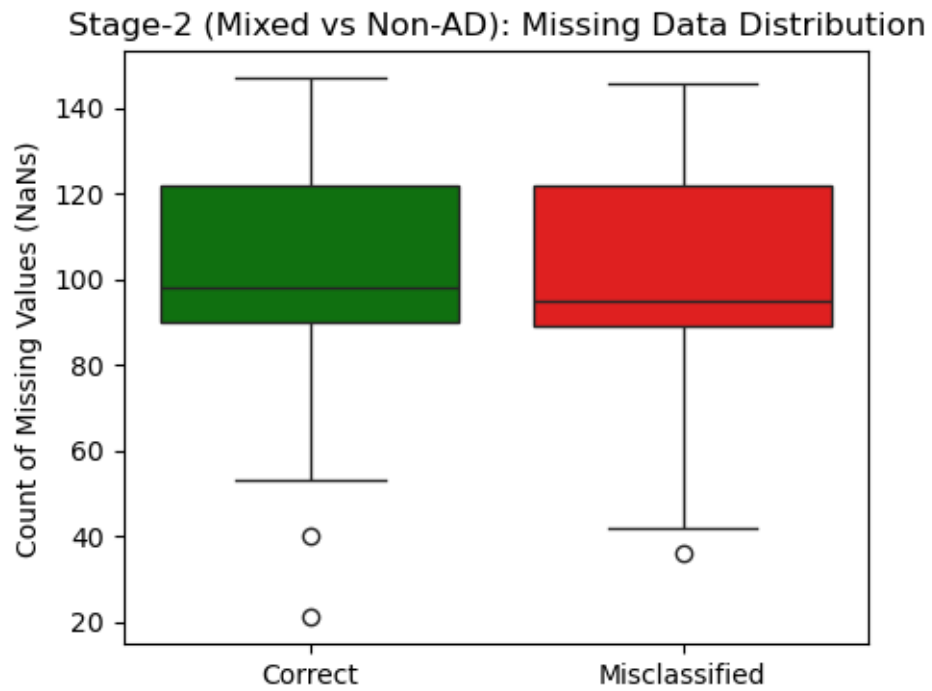
```
[*] Comparing calibration: Raw vs Isotonic...
[!] Isotonic calibration skipped due to error: Found array with 0 sample(s)
(shape=(0, 13)) while a minimum of 1 is required by SimpleImputer.
```

```
[*] Misclassification analysis...
- Total misclassified samples: 430
- Avg missing values (misclassified): 99.59
- Avg missing values (correct): 100.66
```

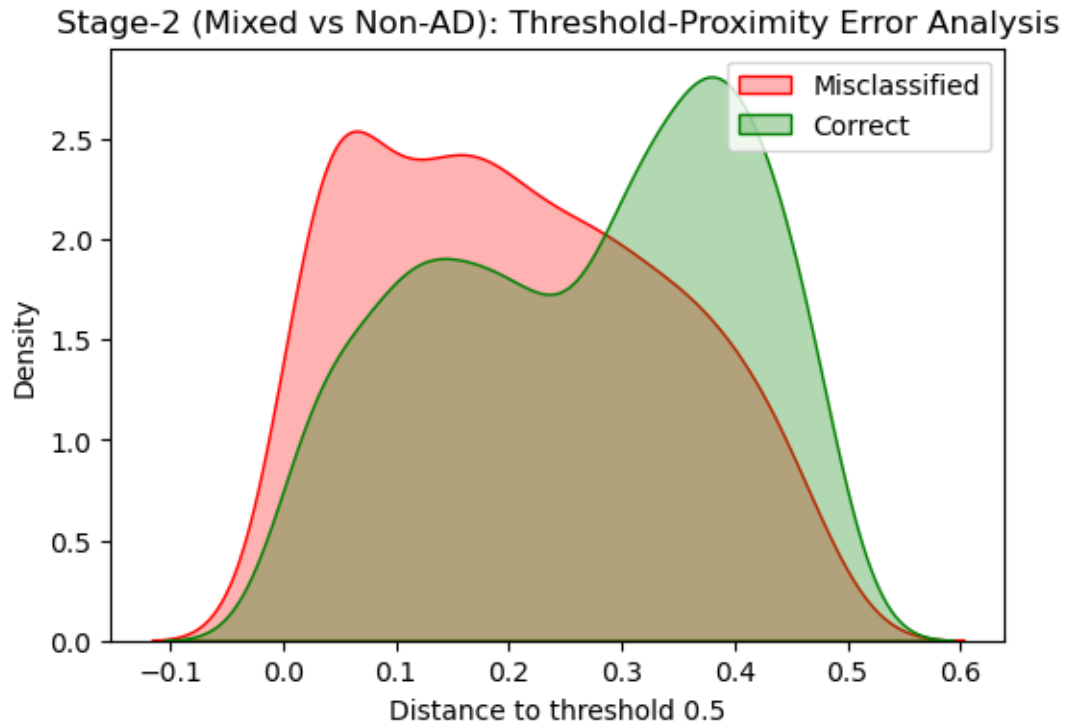
```
/tmp/ipykernel_1118297/625832797.py:195: FutureWarning:
```

```
Passing `palette` without assigning `hue` is deprecated and will be removed in
v0.14.0. Assign the `x` variable to `hue` and set `legend=False` for the same
effect.
```

```
sns.boxplot(
```



- > Missingness is similar between correct and incorrect cases.
- Fraction of misclassifications near threshold (<0.05): 15.1%



-> Errors are widely distributed; not primarily due to the threshold cut-off.

```
[42]: import pandas as pd

from sklearn.metrics import confusion_matrix

# =====
#      (Final Confusion Matrix)
# =====

print("\n" + "="*50)

print("      (Final Hierarchical Confusion Matrix)")

print("="*50)

# 1.

labels3 = ["AD", "Mixed", "Non-AD"]

# 2.      (      )
```

```

if 'y_true3' in locals():
    y_target = y_true3
elif 'y_true' in locals():
    y_target = y_true
else:
    #         df

    target_col = "target_diag" if "target_diag" in df.columns else "diag3"

    y_target = df[target_col].values

# 3.

# pred_final

cm_final_array = confusion_matrix(y_target, pred_final, labels=labels3)

# 4.      DataFrame

cm_final_df = pd.DataFrame(

    cm_final_array,

    index=[f"True {label}" for label in labels3],      #

    columns=[f"Pred {label}" for label in labels3]      #

)

# 5.

print(cm_final_df)

print("-" * 50)

#

acc = (y_target == pred_final).mean()

print(f"Overall Accuracy: {acc:.4f} ({cm_final_array.trace()}/{len(y_target)})")

```

```
print("="*50)
```

```
=====
      (Final Hierarchical Confusion Matrix)
=====
      Pred AD   Pred Mixed   Pred Non-AD
True AD      1917          33          42
True Mixed    101         438         106
True Non-AD    34         130         509
-----
Overall Accuracy: 0.8653 (2864/3310)
=====
```

```
[46]: import numpy as np
import pandas as pd
from pathlib import Path
from sklearn.metrics import (
    classification_report,
    balanced_accuracy_score,
    roc_auc_score,
    accuracy_score,
)

# -----
FAIR_DIR = Path("surv_out") / "fairness"
FAIR_DIR.mkdir(parents=True, exist_ok=True)

# -----
required_vars = ["df", "y_true3", "pred_final"]
for v in required_vars:
    if v not in globals():
        raise RuntimeError(f" {v} ")

if "Gender_ID_num" not in df.columns or "ethnicitycleaned_code" not in df.
    columns:
    raise RuntimeError("df   Gender_ID_num   ethnicitycleaned_code   ")

# -----
labels_3    = ["AD", "Mixed", "Non-AD"]    #
y_true_all  = np.array(y_target)
y_pred_all  = np.array(pred_final)

gender_codes = df["Gender_ID_num"].to_numpy()
eth_codes    = df["ethnicitycleaned_code"].to_numpy()
```

```

#
gender_label_map = {
    -1: "Missing_gender",
    0: "Gender_0", # "Female"
    1: "Gender_1", # "Male"
}
#
eth_label_map = {
    -1: "Missing_eth",
}

# =====
# 1
# =====
def fairness_multiclass_by_group(y_true, y_pred, group_codes,
                                group_label_map, labels_3, group_name,
                                skip_code=-1, min_n=30,
                                save_prefix=""):
    """
    group /
    - accuracy
    - macro recall (balanced accuracy for multiclass)
    - macro F1
      (AD/Mixed/Non-AD) precision/recall/F1
    """
    summary_rows = []
    perclass_rows = []

    unique_codes = np.unique(group_codes)
    for g in unique_codes:
        if g == skip_code:
            #
            continue

        mask = (group_codes == g)
        n = int(mask.sum())
        if n < min_n:
            print(f"[{group_name}] {g} {group_label_map.get(g, g)} {n} <
↪{min_n} / ")
            if n == 0:
                continue

        yt = y_true[mask]
        yp = y_pred[mask]

        rep = classification_report(
            yt, yp, labels=labels_3,

```



```

        output_dict=True, zero_division=0
    )

    #    accuracy / macro F1 / macro recall
    acc      = accuracy_score(yt, yp)
    macro_f1 = np.mean([rep[c]["f1-score"] for c in labels_3])
    macro_rec = np.mean([rep[c]["recall"]   for c in labels_3])

    summary_rows.append({
        f"{group_name}_code": int(g),
        f"{group_name}_label": group_label_map.get(g, str(g)),
        "n": n,
        "accuracy": acc,
        "macro_recall": macro_rec,
        "macro_f1": macro_f1,
    })

    for c in labels_3:
        perclass_rows.append({
            f"{group_name}_code": int(g),
            f"{group_name}_label": group_label_map.get(g, str(g)),
            "class": c,
            "precision": rep[c]["precision"],
            "recall": rep[c]["recall"],
            "f1": rep[c]["f1-score"],
            "support": int(rep[c]["support"]),
        })

    summary_df = pd.DataFrame(summary_rows).sort_values(f"{group_name}_code")
    perclass_df = pd.DataFrame(perclass_rows).
    ↪sort_values([f"{group_name}_code", "class"])

    if save_prefix:
        summary_path = FAIR_DIR / f"{save_prefix}_{group_name}_summary.csv"
        perclass_path = FAIR_DIR / f"{save_prefix}_{group_name}_perclass.csv"
        summary_df.to_csv(summary_path, index=False)
        perclass_df.to_csv(perclass_path, index=False)
        print(f"[{group_name}]      ")
        print("  -", summary_path)
        print("  -", perclass_path)

    return summary_df, perclass_df

# =====
#    2      Stage-1 / Stage-2
# =====

```

```

from sklearn.metrics import (

    accuracy_score,

    balanced_accuracy_score,

    roc_auc_score,

    confusion_matrix,

)

def fairness_binary_by_group(y_true, y_prob, group_codes, group_label_map,

                            group_name, task_name, thr=0.5,

                            skip_code=-1, min_n=30,

                            save_prefix=""):

    """

    group

    - accuracy

    - precision / recall / F1

    - balanced accuracy

    - ROC AUC          NaN

    """

    rows = []

    unique_codes = np.unique(group_codes)

    for g in unique_codes:

        if g == skip_code:

            continue

        mask = (group_codes == g)

        n = int(mask.sum())

```

```

if n < min_n:

    print(f"[{task_name}]-{group_name}]   {g} {group_label_map.get(g, 'u
↪g)} {n} < {min_n} / ")

    if n == 0:

        continue

yt = y_true[mask]

prob = y_prob[mask]

yp = (prob >= thr).astype(int)

# ---- labels=[0,1] 2x2 ----

cm = confusion_matrix(yt, yp, labels=[0, 1])

# cm 2x2 0

tn, fp, fn, tp = cm.ravel()

# accuracy

acc = (tp + tn) / max(tp + tn + fp + fn, 1)

# precision / recall / F1

if tp + fp > 0:

    precision_pos = tp / (tp + fp)

else:

    precision_pos = np.nan

if tp + fn > 0:

    recall_pos = tp / (tp + fn)

else:

    recall_pos = np.nan

```

```

        if precision_pos is not np.nan and recall_pos is not np.nan and
↪(precision_pos + recall_pos) > 0:

            f1_pos = 2 * precision_pos * recall_pos / (precision_pos +
↪recall_pos)

        else:

            f1_pos = np.nan

        # balanced accuracy

        bal_acc = balanced_accuracy_score(yt, yp)

        # AUC          NaN

        try:

            auc = roc_auc_score(yt, prob)

        except Exception:

            auc = np.nan

        rows.append({

            f"{group_name}_code": int(g),

            f"{group_name}_label": group_label_map.get(g, str(g)),

            "n": n,

            "accuracy": acc,

            "precision_pos": precision_pos,

            "recall_pos": recall_pos,

            "f1_pos": f1_pos,

            "balanced_acc": bal_acc,

            "roc_auc": auc,

        })

df_out = pd.DataFrame(rows).sort_values(f"{group_name}_code")

```

```

if save_prefix:

    path = FAIR_DIR / f"{save_prefix}_{task_name}_{group_name}.csv"

    df_out.to_csv(path, index=False)

    print(f"[{task_name}-{group_name}]          {path}")

return df_out

# =====
# 1.      /
# =====
print("\n=====")
gender_mask_valid = (gender_codes != -1)      #      np.
↳ ones_like(gender_codes, dtype=bool)
gender_summary, gender_perclass = fairness_multiclass_by_group(
    y_true=y_true_all[gender_mask_valid],
    y_pred=y_pred_all[gender_mask_valid],
    group_codes=gender_codes[gender_mask_valid],
    group_label_map=gender_label_map,
    labels_3=labels_3,
    group_name="gender",
    skip_code=-1,
    min_n=30,
    save_prefix="final3"
)
print(gender_summary)

print("\n=====")
eth_mask_valid = (eth_codes != -1)
eth_summary, eth_perclass = fairness_multiclass_by_group(
    y_true=y_true_all[eth_mask_valid],
    y_pred=y_pred_all[eth_mask_valid],
    group_codes=eth_codes[eth_mask_valid],
    group_label_map=eth_label_map,
    labels_3=labels_3,
    group_name="ethnicity",
    skip_code=-1,
    min_n=30,
    save_prefix="final3"
)
print(eth_summary)

```

```

=====
[gender]
- surv_out/fairness/final3_gender_summary.csv
- surv_out/fairness/final3_gender_perclass.csv
  gender_code gender_label      n accuracy macro_recall macro_f1
0           0   Gender_0  1337  0.870606      0.817649  0.820293
1           1   Gender_1  1973  0.861632      0.785270  0.794176

```

```

=====
[ethnicity]  7 7   10 < 30 /
[ethnicity] 10 10  23 < 30 /
[ethnicity] 12 12  14 < 30 /
[ethnicity] 13 13  22 < 30 /
[ethnicity] 14 14  15 < 30 /
[ethnicity] 15 15   8 < 30 /
[ethnicity] 16 16   4 < 30 /
[ethnicity] 17 17   2 < 30 /
[ethnicity] 18 18   1 < 30 /
[ethnicity]
- surv_out/fairness/final3_ethnicity_summary.csv
- surv_out/fairness/final3_ethnicity_perclass.csv
  ethnicity_code ethnicity_label      n accuracy macro_recall macro_f1
0              0              0  1645  0.866869      0.791047  0.802742
1              1              1   145  0.820690      0.793022  0.777864
2              2              2   128  0.859375      0.813636  0.810426
3              3              3    64  0.875000      0.802309  0.809340
4              4              4   220  0.886364      0.832530  0.838049
5              5              5   489  0.881391      0.827956  0.827925
6              6              6   133  0.857143      0.791854  0.793298
7              7              7    10  0.800000      0.500000  0.555556
8              8              8    97  0.855670      0.778174  0.772791
9              9              9   128  0.859375      0.765157  0.768780
10             10             10    23  0.913043      0.555556  0.600000
11             11             11    46  0.826087      0.777357  0.769915
12             12             12    14  0.785714      0.516667  0.582011
13             13             13    22  0.818182      0.735690  0.715344
14             14             14    15  0.733333      0.694444  0.674074
15             15             15     8  0.750000      0.444444  0.525253
16             16             16     4  1.000000      0.333333  0.333333
17             17             17     2  0.500000      0.166667  0.222222
18             18             18     1  1.000000      0.333333  0.333333

```

[]: