```
In [1]: from scgenome import tantalus
        import pandas as pd
        from IPython.display import display
        from scgenome import utils, cncluster, simulation, cnplot
        import scipy.cluster.hierarchy as sch
        import matplotlib.pyplot as plt
        import numpy as np
        import seaborn as sns
        import time
In [2]: all cn data fp = "/Users/massoudmaher/data/sc 1935 1936 1937 cn data qc.
        csv"
        all cn data = pd.read csv(all cn data fp)
        all cn data = all_cn_data.iloc[:,1:]
In [3]: hmmcopy tickets = ['SC-1935', 'SC-1936', 'SC-1937']
        sample ids = [["SA922"], ['SA921'], ['SA1090']]
        # spike in params
        total ncells = 100
        proportions = [0.3, 0.3, 0.4]
        # bhc params
        n states = 12
        alpha = 0.3
        prob cn change = 0.8
        bhc incon = 2 # inconsistent score used for making clusters from bhc
        bhc depth = 2
        # naive clusering params
        naive method = "complete"
        naive metric = "cityblock"
        naive incon = 1.1
        naive depth = 2
        # Params for testing threshold values
        params = simulation.expand grid({"transform":["log", "none"], "criterion"
        : ["inconsistent"], "threshold": np.arange(0.025, 2, step=0.05)})
        params = pd.concat([params, simulation.expand grid({"transform":["log",
        "none"], "criterion": ["distance"], "threshold": np.arange(3, 20, step=1
        ) } ) ] )
```

```
In [4]: subsample = utils.get cn_data_submixture(all_cn_data, total_ncells, hmmc
         opy tickets, sample ids, proportions=proportions)
         mixed_cn_data = subsample["mixed_cn_data"]
         mixed cn data["origin id int"] = mixed cn data["origin id"].factorize()[
         0 1
         cell counts = subsample["cell counts"]
         /Users/massoudmaher/Documents/Code/scgenome/scgenome/utils.py:169: Sett
         ingWithCopyWarning:
         A value is trying to be set on a copy of a slice from a DataFrame.
         Try using .loc[row_indexer,col_indexer] = value instead
         See the caveats in the documentation: http://pandas.pydata.org/pandas-d
         ocs/stable/indexing.html#indexing-view-versus-copy
           jira_cn_data[origin_field_name] = hmmcopy_tickets[i]
 In [5]: start = time.time()
         bhc linkage, bhc root, bhc cell ids, matrix data, measurement, variances
             cncluster.bayesian_cluster(mixed_cn_data, n_states=n_states, alpha=a
         lpha, prob cn change=prob cn change)
         print(f"{time.time()-start}s for BHC on {total_ncells} cells")
         155.69063305854797s for BHC on 100 cells
In [17]: bhc linkage, bhc plot data = simulation.get plot data(bhc linkage)
```

lbhc plot data[:,2] = np.log(lbhc plot data[:,2]) # Log because the high

naive linkage = sch.linkage(measurement, method=naive method, metric=nai

lbhc plot data = bhc plot data.copy()

lnaive linkage = naive linkage.copy()

lnaive linkage[:,2] = np.log(lnaive linkage[:,2])

est link is way higher

ve metric)

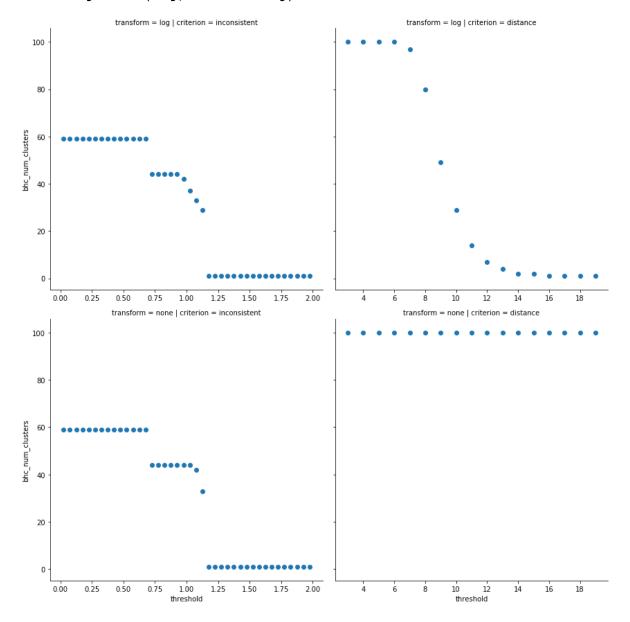
```
In [18]: def apply_fn(row):
             if row["transform"] == "log":
                 df = lbhc_plot_data
             else:
                 df = bhc_plot_data
             return sch.fcluster(df, row["threshold"], criterion=row["criterion"
         ])
         params["bhc_fcluster"] = params.apply(apply_fn, axis=1)
         params["bhc_num_clusters"] = params["bhc_fcluster"].apply(lambda x: len(
         set(x)))
         def apply_fn(row):
             if row["transform"] == "log":
                 df = lnaive linkage
             else:
                 df = naive_linkage
             return sch.fcluster(df, row["threshold"], criterion=row["criterion"
         ])
         params["naive_fcluster"] = params.apply(apply_fn, axis=1)
         params["naive_num_clusters"] = params["naive_fcluster"].apply(lambda x:
         len(set(x)))
         params.head()
```

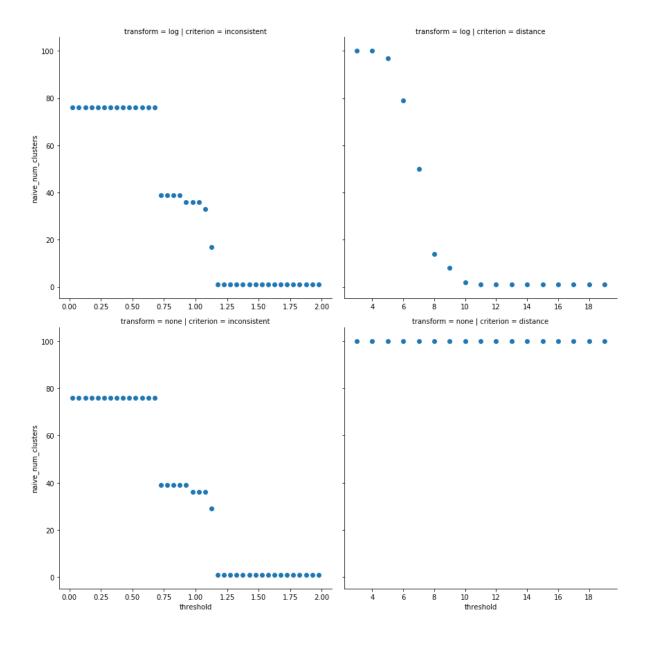
Out[18]:

	transform	criterion	threshold	fcluster	num_clusters	bhc_fcluster	bhc_num_clusters	naiv
0	log	inconsistent	0.025	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[13, 5,
1	log	inconsistent	0.075	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[13, 5,
2	log	inconsistent	0.125	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[13, 5,
3	log	inconsistent	0.175	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[13, 5,
4	log	inconsistent	0.225	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[1, 9, 3, 14, 15, 23, 7, 26, 11, 3, 26, 11, 10	59	[13, 5,

/Users/massoudmaher/Documents/Code/scgenome/scg/lib/python3.7/site-pack ages/seaborn/axisgrid.py:230: UserWarning: The `size` paramter has been renamed to `height`; please update your code.

warnings.warn(msg, UserWarning)



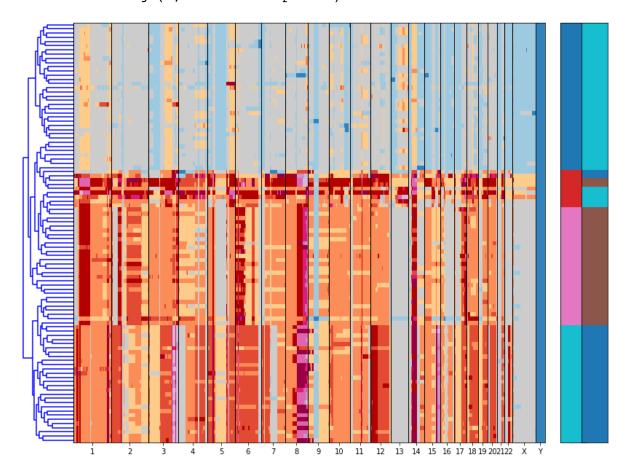


Rightmost bar represents where sample originally came frome

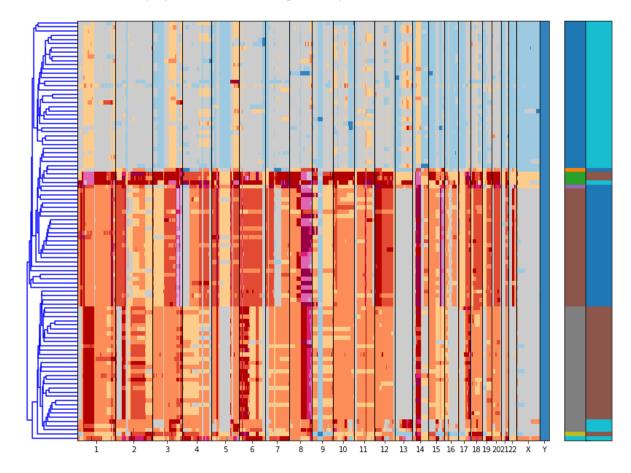
```
In [20]: cmixed_cn_data = mixed_cn_data.copy()
    clustering = sch.fcluster(lbhc_plot_data, 13, criterion="distance")
    cmixed_cn_data = cncluster.prune_cluster(clustering, bhc_cell_ids, mixed
    _cn_data)

fig = plt.figure(figsize=(10, 8))
    bimatrix_data = cnplot.plot_clustered_cell_cn_matrix_figure(
        fig, cmixed_cn_data, "state", cluster_field_name="bhc_cluster_id",
        linkage=lbhc_plot_data, origin_field_name="origin_id_int")
```

/Users/massoudmaher/Documents/Code/scgenome/scgenome/cnplot.py:50: Clus terWarning: scipy.cluster: The symmetric non-negative hollow observation matrix looks suspiciously like an uncondensed distance matrix Y = sch.linkage(D, method='complete')



/Users/massoudmaher/Documents/Code/scgenome/scgenome/cnplot.py:50: Clus terWarning: scipy.cluster: The symmetric non-negative hollow observation matrix looks suspiciously like an uncondensed distance matrix Y = sch.linkage(D, method='complete')



```
umap_params = utils.expand_grid({"n_neighbors": np.arange(3,18,step=1)})
In [34]:
         def apply fn(row):
             return cncluster.umap hdbscan_cluster(matrix_data["state"], n_neighb
         ors=row["n_neighbors"])
         umap params["umap clusters"] = umap params.apply(apply fn, axis=1)
         umap params["umap num clusters"] = umap params["umap clusters"].apply(la
         mbda x: len(set(x["cluster id"])))
         sns.scatterplot(data=umap params, x="n_neighbors", y="umap num clusters"
         )
         umap df = cncluster.umap hdbscan cluster(matrix data["state"], n_neighbo
         rs=10)
         umixed_cn_data = mixed_cn_data.merge(umap_df, how="inner")
         fig = plt.figure(figsize=(10, 8))
         bimatrix data = cnplot.plot clustered_cell_cn matrix figure(
             fig, umixed_cn_data, "state", cluster_field_name="cluster_id",
             linkage=None, origin field name="origin id int")
         #def umap hdbscan cluster(
         #
                  cn,
         #
                  n components=2,
         #
                  n neighbors=15,
                  min dist=0.1,
         #):
```

/Users/massoudmaher/Documents/Code/scgenome/scg/lib/python3.7/site-pack ages/umap/spectral.py:229: UserWarning: Embedding a total of 8 separate connected components using meta-embedding (experimental)

n components

/Users/massoudmaher/Documents/Code/scgenome/scg/lib/python3.7/site-pack ages/sklearn/manifold/spectral_embedding_.py:237: UserWarning: Graph is not fully connected, spectral embedding may not work as expected.

warnings.warn("Graph is not fully connected, spectral embedding" /Users/massoudmaher/Documents/Code/scgenome/scg/lib/python3.7/site-pack ages/umap/spectral.py:229: UserWarning: Embedding a total of 8 separate connected components using meta-embedding (experimental)

n_components

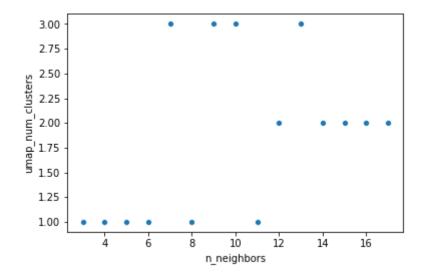
/Users/massoudmaher/Documents/Code/scgenome/scg/lib/python3.7/site-pack ages/sklearn/manifold/spectral_embedding_.py:237: UserWarning: Graph is not fully connected, spectral embedding may not work as expected.

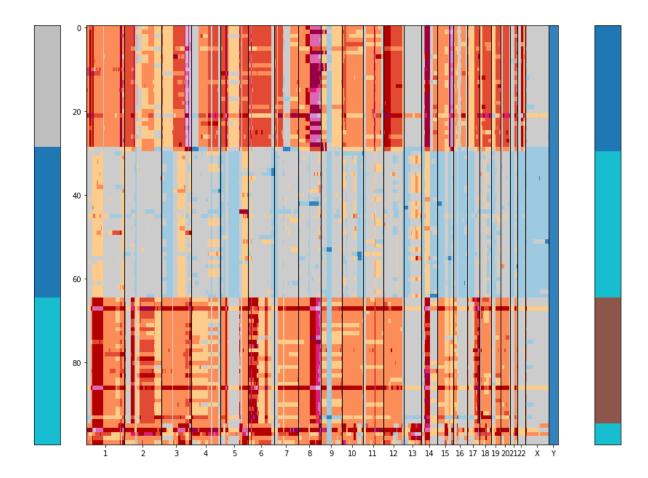
warnings.warn("Graph is not fully connected, spectral embedding" /Users/massoudmaher/Documents/Code/scgenome/scg/lib/python3.7/site-pack ages/umap/spectral.py:229: UserWarning: Embedding a total of 4 separate connected components using meta-embedding (experimental)

n_components

/Users/massoudmaher/Documents/Code/scgenome/scgenome/cnplot.py:50: Clus terWarning: scipy.cluster: The symmetric non-negative hollow observation matrix looks suspiciously like an uncondensed distance matrix

Y = sch.linkage(D, method='complete')





Umap does better-- especially considering that it runs way faster

```
In [ ]:
```