Breast cancer classification challenge

Part 0: All the packages

Step 1: The essentials

```
!pip install pandas
!pip install numpy
!pip install matplotlib
```

Step 2: The "some things that not everyone has installed"

```
!pip install json
!pip install sklearn
!pip install eli5
!pip install tensorflow
!pip install keras
```

Step 3: Some news to read while waiting

Best overall result:

Area under ROC curve: 0.9408866995073892

Recall: 0.7931034482758621

Precision: 0.8518518518519

Best recall and AUC:

Area under ROC curve: 0.9584017515051998

Recall: 0.8275862068965517

Precision: 0.7741935483870968

Part 1: Data import and preparation

Step 1: Read in JSON measurements

```
# Sanity check
print(recs['Left_Temps'][0]) # first left breast
```

[32.356388888889, 29.65194444444446, 30.4308333333325, 31.21722222222226, 30.3730555555555, 31.196944444444437, 31.413333333333, 31

Step 2: Put measurements in a dataframe and omit post-surgery cases

```
import pandas as pd

dataset = pd.DataFrame(recs)
dataset.head()
```

	ID	Surgery	Body_Temp	Cancer	Left_Temps	Right_Temps
0	115	0	35.8	0	[32.356388888888889, 29.651944444444446, 30.430	[30.34972222222226, 28.3255555555555, 28.115
1	154	0	35.6	0	[31.0913888888888886, 29.1699999999999, 28.63	[28.538888888888889, 27.670833333333334, 28.504
2	20	0	35.0	0	[32.24555555555557, 30.2777777777778, 30.0636	[33.01444444444445, 30.63055555555555, 29.5402
3	98	1	36.1	0	[31.711944444444444, 28.13416666666667, 27.1297	[29.64055555555556, 28.33027777777777, 28.401
4	210	0	36.7	1	[34.37111111111105, 32.69611111111111, 31.474	[34.271944444444444, 31.85416666666668, 30.913

```
# Taking rows where the column value is no surgery
dataset = dataset.loc[dataset['Surgery'] == 0]
dataset = dataset.reset_index(drop=True) # reset indices
dataset.head()
```

	ID	Surgery	Body_Temp	Cancer	Left_Temps	Right_Temps
0	115	0	35.8	0	[32.356388888888889, 29.651944444444446, 30.430	[30.34972222222226, 28.3255555555555, 28.115
1	154	0	35.6	0	[31.0913888888888886, 29.1699999999999, 28.63	[28.538888888888889, 27.670833333333334, 28.504
2	20	0	35.0	0	[32.24555555555557, 30.2777777777778, 30.0636	[33.01444444444445, 30.6305555555555, 29.5402
3	210	0	36.7	1	[34.37111111111105, 32.69611111111111, 31.474	[34.271944444444444, 31.85416666666668, 30.913
4	61	0	35.0	0	[31.337500000000002, 29.98722222222222, 29.69	[31.66611111111111, 29.7949999999998, 29.053

1. Rewrite left and right individual sensors' temperatures into separate columns

```
import numpy as np

temps_to_cols = {
    'Left_Temps': None,
    'Right_Temps': None
}

col_names = {
    'Left_Temps': [],
    'Right_Temps': []
}

for side in ['Left_Temps', 'Right_Temps']:
    num_meas, num_sensors = len(dataset[side]), len(dataset[side][0])
    temps_to_cols[side] = np.array(
        [dataset[side][x] for x in range(num_meas)])
    col_names[side] = list(map(lambda idx: side[:side.find('T')].lower() + str(idx), range(num_sensors)))
```

```
# Sanity check
pd.DataFrame(temps_to_cols['Left_Temps'], columns=col_names['Left_Temps']).head()
```

	left_0	left_1	left_2	left_3	left_4	left_5	left_6	left_7	left_8	left_9	 left_86	lef
0	32.356389	29.651944	30.430833	31.217222	30.373056	31.196944	31.413333	31.148611	31.811667	31.304167	 28.477778	28.26
1	31.091389	29.170000	28.633611	28.805278	29.184444	30.118056	30.163889	29.694722	29.980556	29.998889	 30.279167	28.74
2	32.245556	30.277778	30.063611	29.356667	29.539722	29.766944	29.575278	29.540000	30.222222	30.460556	 27.842222	28.418
3	34.371111	32.696111	31.474722	31.261389	32.046111	31.988889	31.983056	31.178333	31.789444	32.158611	 31.933889	31.724
4	31.337500	29.987222	29.697500	29.725278	30.421389	30.700278	30.270278	30.508611	31.275278	30.761111	 30.780833	28.80

5 rows × 96 columns

1. Put back to the main dataframe

```
for side in ['Left_Temps', 'Right_Temps']:
    temps_df = pd.DataFrame(temps_to_cols[side], columns=col_names[side])
    dataset = pd.concat([dataset, temps_df], axis=1) # bind dataframes column-wise
dataset.head()
```

	ID	Surgery	Body_Temp	Cancer	Left_Temps	Right_Temps	left_0	left_1	left_2	left_3	 righ
0	115	0	35.8	0	[32.35638888888889, 29.651944444444446, 30.430	[30.34972222222226, 28.32555555555555, 28.115	32.356389	29.651944	30.430833	31.217222	 29.15
1	154	0	35.6	0	[31.091388888888886, 29.169999999999998, 28.63	[28.5388888888889, 27.670833333333334, 28.504	31.091389	29.170000	28.633611	28.805278	 30.08
2	20	0	35.0	0	[32.2455555555557, 30.27777777777778, 30.0636	[33.0144444444445, 30.63055555555555, 29.5402	32.245556	30.277778	30.063611	29.356667	 27.77
3	210	0	36.7	1	[34.37111111111105, 32.69611111111111, 31.474	[34.27194444444444, 31.8541666666666668, 30.913	34.371111	32.696111	31.474722	31.261389	 30.75
4	61	0	35.0	0	[31.337500000000002, 29.98722222222222, 29.69	[31.66611111111111, 29.794999999999998, 29.053	31.337500	29.987222	29.697500	29.725278	 29.51

5 rows × 198 columns

```
print("Size of dataset before dropping:", len(dataset))
tmp = dataset.dropna(how='any', axis=0, inplace=False)
print("Size of dataset after dropping:", len(tmp))
```

Size of dataset before dropping: 156 Size of dataset after dropping: 155

Just one wrong entry, let's neglect it since we cannot restore information from NaN.

```
dataset = tmp
dataset.reset_index(drop=True)
dataset.head()
```

	ID	Surgery	Body_Temp	Cancer	Left_Temps	Right_Temps	left_0	left_1	left_2	left_3	 righ
0	115	0	35.8	0	[32.35638888888889, 29.651944444444446, 30.430	[30.34972222222226, 28.32555555555555, 28.115	28.3255555555555555 32.356389		30.430833	31.217222	 29.15
1	154	0	35.6	0	[31.091388888888886, 29.169999999999998, 28.63	[28.5388888888889, 27.6708333333333334, 28.504	31.091389	29.170000	28.633611	28.805278	 30.08
2	20	0	35.0	0	[32.2455555555557, 30.27777777777778, 30.0636	[33.0144444444445, 30.63055555555555, 29.5402	32.245556	30.277778	30.063611	29.356667	 27.77
3	210	0	36.7	1	[34.37111111111105, 32.69611111111111, 31.474	[34.27194444444444, 31.8541666666666668, 30.913	34.371111	32.696111	31.474722	31.261389	 30.750
4	61	0	35.0	0	[31.337500000000002, 29.98722222222222, 29.69	[31.66611111111111, 29.794999999999998, 29.053	31.337500	29.987222	29.697500	29.725278	 29.514

5 rows × 198 columns

Step 3: Form data and targets

```
targets = dataset['Cancer']
targets.head()

0     0
1     0
2     0
3     1
4     0
Name: Cancer, dtype: int64

data_cols = np.hstack([['Body_Temp'], dataset.columns.get_values()[6:]])
data = dataset[data_cols]
```

	Body_Temp	left_0	left_1	left_2	left_3	left_4	left_5	left_6	left_7	left_8	 right_86	riç
0	35.8	32.356389	29.651944	30.430833	31.217222	30.373056	31.196944	31.413333	31.148611	31.811667	 29.157500	29.3
1	35.6	31.091389	29.170000	28.633611	28.805278	29.184444	30.118056	30.163889	29.694722	29.980556	 30.085000	28.6
2	35.0	32.245556	30.277778	30.063611	29.356667	29.539722	29.766944	29.575278	29.540000	30.222222	 27.778889	27.8
3	36.7	34.371111	32.696111	31.474722	31.261389	32.046111	31.988889	31.983056	31.178333	31.789444	 30.750000	30.5
4	35.0	31.337500	29.987222	29.697500	29.725278	30.421389	30.700278	30.270278	30.508611	31.275278	 29.514722	28.9

5 rows × 193 columns

data.head()

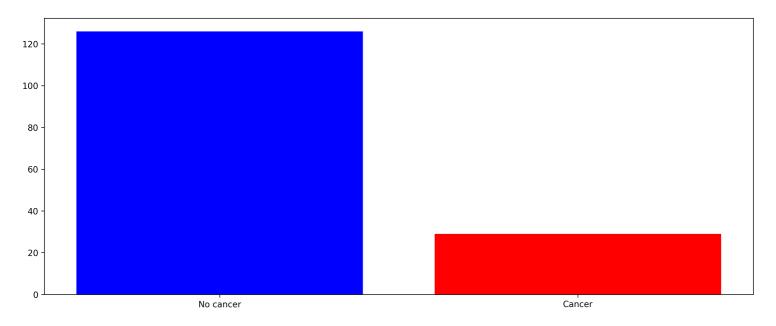
Part 2: Exploratory data analysis

Step 1: Some basic stats

1. Cancer dataset balance

```
from matplotlib import pyplot as plt

plt.figure(figsize=(15, 6), dpi=300)
plt.bar(x=['No cancer', 'Cancer'], height=[sum(targets == 0), sum(targets == 1)], color=["blue", "red"])
plt.show()
```

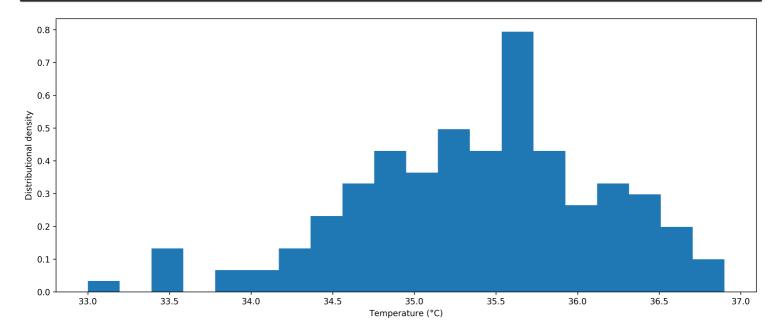


Quite a large imbalance of data, so we have to be looking into recall, precision, and AUC-ROC.

1. Distribution of body temperatures

```
bd_temps = list(data['Body_Temp'])

plt.figure(figsize=(15, 6), dpi=300)
plt.hist(bd_temps, density=True, bins=20)
plt.xlabel('Temperature (°C)')
plt.ylabel('Distributional density')
plt.show()
```



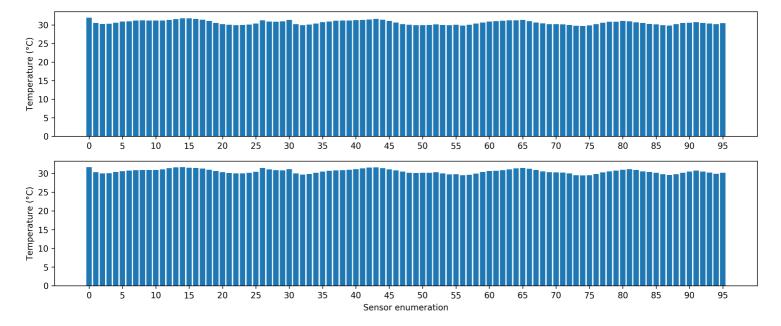
1. Average measurements on each sensor

```
left_avgs = np.mean([data.iloc[idx][1:97] for idx in range(len(data))], axis=0) # column-wise mean
right_avgs = np.mean([data.iloc[idx][97:] for idx in range(len(data))], axis=0) # column-wise mean

plt.figure(figsize=(15, 6), dpi=300)
plt.subplot(2, 1, 1)
plt.bar(range(96), height=left_avgs, width=0.8)
plt.xticks(np.arange(0, 100, 5))
plt.ylabel('Temperature (°C)')

plt.subplot(2, 1, 2)
plt.bar(range(96), height=right_avgs, width=0.8)
plt.xticks(np.arange(0, 100, 5))
plt.xlabel('Sensor enumeration')
plt.xlabel('Temperature (°C)')

plt.show()
```



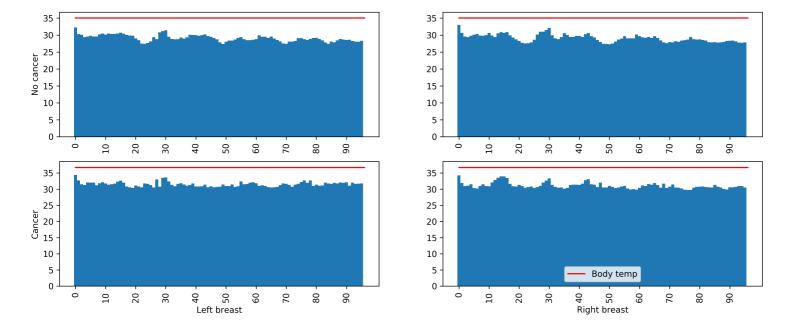
Should be pretty similar, since the sensors are located symmetrically on both cups!

Step 2: Point temperatures of no cancer and cancer breasts

1. Let's first compare how the temperatures distributions look like in sample normal breasts and sample breasts with cancer.

```
Custom enumerator to make indices for graph positions
def my_enum(xs, start=0, step=1):
        yield (start, x)
        start += step
\label{thm:condition} \mbox{Temperature distribution grapher for each cup}
\tt def \ graph\_rl\_temps(row\_idx, \ data=data, \ targets=targets, \ show\_bd\_temp=True):
    plt.figure(figsize=(15, 6), dpi=300)
    num_rows = len(row_idx)
    for plot_id, row_id in my_enum(row_idx, start=1, step=2):
        rec = data.iloc[row_id]
        cancer_status = 'Cancer' if list(targets)[row_id] == 1 else 'No cancer'
        last_row = (plot_id == num_rows * 2 - 1)
        plt.subplot(num_rows, 2, plot_id)
plt.bar(range(96), height=rec[1:1+num_sensors_per_cup], width=1)
        if show_bd_temp:
            bd_temp_ln, = plt.plot([0, 96], [rec[0], rec[0]], color="red", label='Body temp')
        plt.yticks(np.arange(0, 40, 5))
plt.xticks(np.arange(0, 97, 10), rotation='vertical')
        plt.ylabel(cancer_status)
        if last_row:
        plt.subplot(num_rows, 2, plot_id + 1)
        plt.bar(range(96), height=rec[1+num_sensors_per_cup:], width=1)
        if show_bd_temp:
            plt.plot([0, 96], [rec[0], rec[0]], color="red")
        plt.yticks(np.arange(0, 40, 5))
        plt.xticks(np.arange(0, 97, 10), rotation='vertical')
        if last_row:
            plt.xlabel('Right breast')
    if show_bd_temp:
        plt.legend(handles=[bd_temp_ln], loc='lower center')
    plt.show()
```

graph_rl_temps([2, 3])



We can observe that in normal breasts, the sensors show a "wavy" fashion as different parts of the breast have different temperatures depending on the distance from body and exposure to body heat. Observational studies show that only 2-5% of breast cancers affect both breasts, so in most cases, we should see weird things happening in one of the breasts.

We can see that in the person with breast cancer, the left breast temperature measurements behave a bit "too flat" in some places. This can indicate that because of tumor obtaining blood and heat, the temperature near the tumor is pretty much even and similar to the tumor temperature. Of course, this line of reasoning can also be subject to confirmation bias, so I will go more in depth.

1. It would be nice to know in which breast the cancer tumor is, but even if we do not know, when we average out the measurements in cancer left or right breasts, if our hypothesis is correct, we should get a generally flatter profile of the temperature distribution.

```
data_nocancer = data.loc[np.nonzero(targets == 0)]
data_cancer = data.loc[np.nonzero(targets == 1)]

avg_nocancer = np.mean(data_nocancer, axis=0)
avg_cancer = np.mean(data_cancer, axis=0)

avg_comp_df = pd.DataFrame(np.vstack([avg_nocancer, avg_cancer]))
avg_comp_df

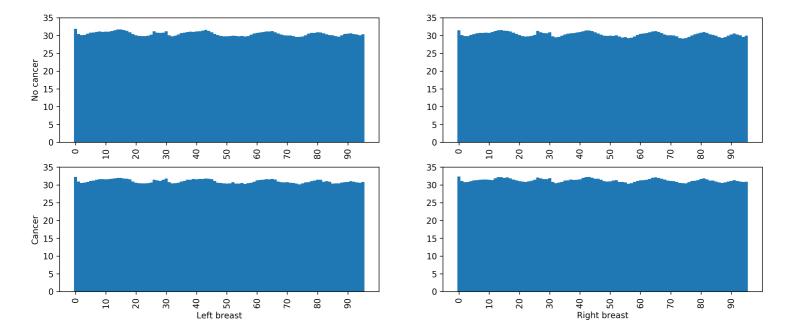
/usr/local/lib/python3.6/site-packages/ipykernel_launcher.py:1: FutureWarning:
Passing list-likes to .loc or [] with any missing label will raise
KeyError in the future, you can use .reindex() as an alternative.

See the documentation here:
https://pandas.pydata.org/pandas-docs/stable/indexing.html#deprecate-loc-reindex-listlike
"""Entry point for launching an IPython kernel.
```

	0	1	2	3	4	5	6	7	8	9	 183	
0	35.407200	31.883387	30.420836	30.146149	30.202138	30.502342	30.810491	30.856298	31.058171	31.134871	 29.547209	29.33
1	35.396552	32.235833	30.947088	30.557232	30.641513	30.815623	31.098477	31.225000	31.461772	31.617807	 30.696571	30.56

2 rows × 193 columns

graph_rl_temps([0, 1], data=avg_comp_df, targets=[0, 1], show_bd_temp=False)



Unclear what is going on here. The challenge is probably that different people have tumors at diffent places, so considering that our dataset is pretty small, the "hot" parts pretty much evenly distribute out without showing obvious patterns. Sad.

Part 3: Validation scheme

Step 1: LOOCV for lighter models

```
from sklearn.model_selection import LeaveOneOut
from sklearn.metrics import recall_score, precision_score, roc_auc_score, roc_curve

def loocv_preds(model, X, y, threshold=0):
    loocv = LeaveOneOut()
    loocv.get_n_splits(X)

# Since we're doing LOOCV, calculating scores on the spot is not possible
dcsn_fn = []
preds = []

for train_idx, test_idx in loocv.split(X):
    trgt = np.array(y) # target value
    model.fit(X.iloc[train_idx], trgt[train_idx])

    dist_from_bound = model.decision_function(X.iloc[test_idx])
    dcsn_fn.append(dist_from_bound)
    preds.append(int(dist_from_bound >= threshold))

return dcsn_fn, preds
```

Step 2: K-fold for neural networks

For larger neural networks, LOOCV will take forever to validate, so I'm creating this. I make sure to reset model weights each time I change training and validation sets, so that we don't have indirect information flow and so that the network would not suspect of anything in learned in the fold before.

Step 3: Metrics

Part 4: Separability evaluation

Step 1: Normalize the data

```
means = data.mean(axis=0)
stds = data.std(axis=0)

norm_data = (data - means) / (stds + 1e-16)
norm_data.head()
```

	Body_Temp	left_0	left_1	left_2	left_3	left_4	left_5	left_6	left_7	left_8		right_86	right_8	
0	0.514501	0.249376	-0.587489	0.130107	0.694777	-0.155391	0.232957	0.353118	0.001181	0.442206		-0.366109	-0.14401	
1	0.253888	-0.537865	-0.907441	-1.125057	-1.143731	-1.047553	-0.562022	-0.579386	-1.129996	-0.986758		0.185001	-0.56062	
2	-0.527952	0.180402	-0.172013	-0.126357	-0.723434	-0.780884	-0.820738	-1.018686	-1.250376	-0.798166		-1.185264	-1.01457	
3	1.687261	1.503189	1.433461	0.859150	0.728443	1.100390	0.816500	0.778321	0.024306	0.424864		0.580137	0.558786	
4	-0.527952	-0.384704	-0.364906	-0.382046	-0.442459	-0.119113	-0.133012	-0.499984	-0.496762	0.023618		-0.153851	-0.34355	

Step 2: PCA with two eigenvectors for visualization

```
from sklearn import decomposition

n_components = 2

pca = decomposition.PCA(n_components=n_components, svd_solver='randomized', whiten=True)

pca.fit(norm_data)

print("Explained variance of components:", pca.explained_variance_ratio_) # how much information is retained

Explained variance of components: [0.69780441 0.04143069]
```

Seems like 2 is a reasonable amount since eigenvectors that come after these will only be able to explain a tiny bit of variance (less than 4%), so we are retaining a good amount of information. This makes sense because when there is no tumor, the temperatures tend to behave correlatedly (e.g., if the body gets hotter, point A and point B on the cup will both capture slightly higher temperatures, just of varying degree of temperature increase). On the other hand, when there is tumor, areas around the tumor heat up just with slightly varying speeds.

1. Calculating back the single-value decomposition of original vectors to plot

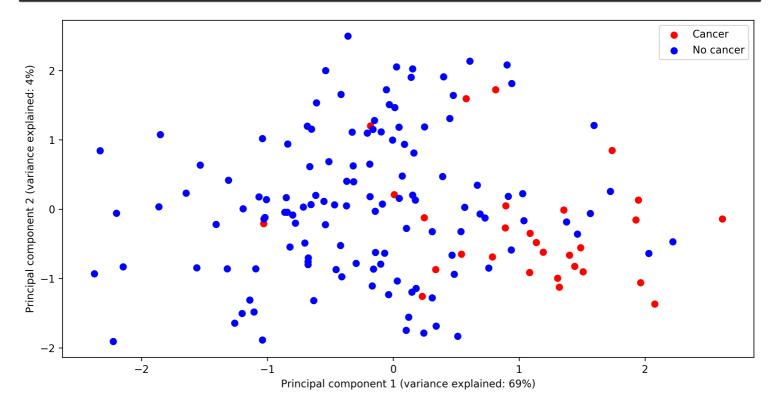
```
data_svd = pca.transform(norm_data)
data_svd[0] # check
```

array([-0.0316851 , 1.50810304])

1. Plotting to see have a decision boundary overview

```
nocancer_idx = np.where(targets == 0)
cancer_idx = np.where(targets == 1)

plt.figure(figsize=(12, 6), dpi=300)
plt.scatter(*zip(*data_svd[cancer_idx]), color="red", label='Cancer')
plt.scatter(*zip(*data_svd[nocancer_idx]), color="blue", label='No cancer')
plt.xlabel('Principal component 1 (variance explained: {0}%)'.format(int(pca.explained_variance_ratio_[0] * 100)))
plt.ylabel('Principal component 2 (variance explained: {0}%)'.format(int(pca.explained_variance_ratio_[1] * 100)))
plt.legend(loc="upper right")
plt.show()
```



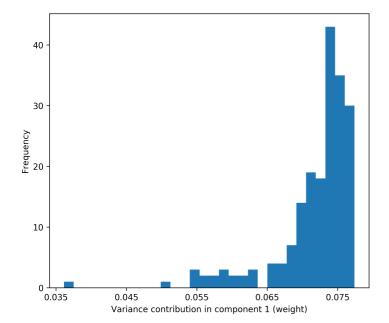
We can see that there are some "deep" overlaps of red dots inside the cluster of blue dots, and some blue dots in the cluster of red dots, thus, linear separation will create many false positives and negatives. Most likely, we care more about flagging the red dots inside the blue cluster at a cost of the blue dots in the red cluster. It's better to have a good

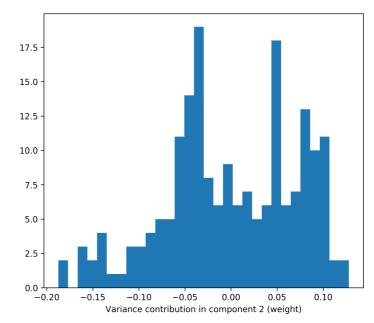
recall score because the cost of in-hospital check-up is lower than the cost of life.

1. Let's check out what the principal components found out about the "interesting" spots

```
plt.figure(figsize=(15, 6), dpi=300)
plt.subplot(1, 2, 1)
plt.hist(pca.components_[0], bins=30)
plt.xticks(np.arange(0.035, 0.085, 0.01))
plt.xlabel('Variance contribution in component 1 (weight)')
plt.ylabel('Frequency')

plt.subplot(1, 2, 2)
plt.hist(pca.components_[1], bins=30)
plt.xlabel('Variance contribution in component 2 (weight)')
plt.show()
```





We can see that for PC1 with 69.78% variance explained, most of the weights are in the maximum range of 0.075, meaning that we have no particularly "outstanding" feature. It mostly makes sense because tumors can appear anywhere, so it is hard to observe a pin-point of "Hey, sensor 5 has higher temperature, and most of the time it is the one that detects cancer." Maybe we have to do some manipulations with combination of features to come up with separation.

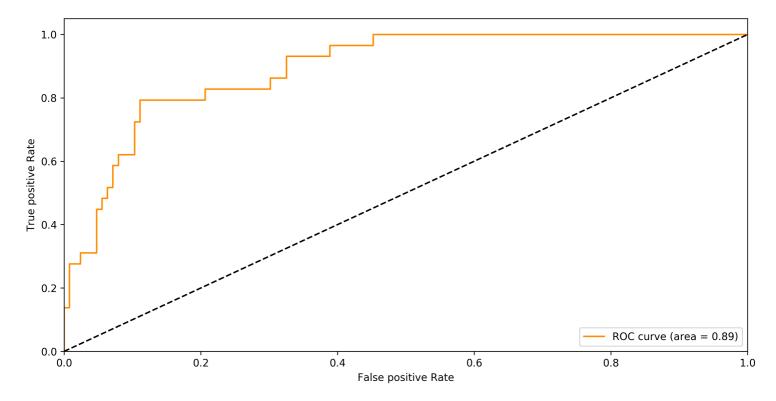
Part 5: Learning models

Step 1: Naive linear separation

```
from sklearn import svm

clf = svm.SVC(kernel='linear', gamma='scale', probability=True)

dcsn_fn, preds = loocv_preds(clf, norm_data, targets)
print("Area under ROC curve:", auc_roc(targets, dcsn_fn))
```



```
Area under ROC curve: 0.8924466338259442

rec, prec = rec_prec(list(targets), np.array(preds).flatten())
print("Recall:", rec)
print("Precision:", prec)

Recall: 0.5517241379310345
Precision: 0.64
```

Pretty low recall because we were not able to flag cases when it is actually cancer.

1. Let's try playing around with the threshold

```
dcsn_fn, preds = loocv_preds(clf, norm_data, targets, threshold=-0.9)
rec, prec = rec_prec(list(targets), np.array(preds).flatten())
print("Recall:", rec)
print("Precision:", prec)
Recall: 0.8275862068965517
Precision: 0.47058823529411764
```

27% gain in recall and 17% drop in precision. Not bad, considering that we would like to prioritize recall in this case.

2. We can play around a bit more to find the optimal threshold

```
# better to use genetic algorithm to find convergence
# and having more data would be super helpful!

dcsn_fn, preds = loocv_preds(clf, norm_data, targets, threshold=-0.4)
rec, prec = rec_prec(list(targets), np.array(preds).flatten())
print("Recall:", rec)
print("Precision:", prec)
```

Recall: 0.7931034482758621 Precision: 0.6216216216216216

Relative to last measurement: 3% drop in recall and 15% gain in precision. Compared to no threshold: 24% gain in recall and 2% drop in precision.

1. Let's look at which features had a more impact

```
from eli5.sklearn import PermutationImportance
from sklearn.model_selection import train_test_split
X_train, X_test, y_train, y_test = train_test_split(
  norm_data, targets, test_size=0.2, random_state=42)
clf.fit(X_train, y_train)
perm = PermutationImportance(clf, random_state=1).fit(X_test, y_test)
eli5.show_weights(clf, feature_names=list(norm_data.columns), top=50)
table.eli5-weights tr:hover {
  filter: brightness(85%);
  <h>
  y=1
top features
<table class="eli5-weights"
    style="border-collapse: collapse; border: none; margin-top: 0em; table-layout: auto; margin-bottom: 2em;">
  <thead>
  <th style="padding: 0 1em 0 0.5em; text-align: right; border: none;" title="Feature weights. Note that weights do not account for
        Weight<sup>?</sup>
    Feature
  </thead>
    +0.445
 left_71
right_74
+0.261
```

left_9

```
+0.233
left_29
+0.231
right_80
+0.223
right_41
+0.222
left_46
+0.208
right_28
+0.202
left_80
+0.197
left 51
+0.195
left_78
```

```
+0.193
right_40
+0.192
right_25
+0.187
right_64
+0.187
left_59
+0.176
left_87
+0.176
right_87
+0.171
right_34
+0.169
right_81
```

```
+0.165
left_74
+0.164
left_54
+0.162
left_70
+0.162
left_21
+0.156
right_14
+0.154
left_61
+0.153
right_33
```

```
<i>&hellip; 82 more positive &hellip;</i>
 <i>%hellip; 62 more negative %hellip;</i>
 -0.154
right_6
left_76
-0.155
right_59
-0.159
right_70
-0.163
right_86
-0.163
left_42
```

```
-0.165
right_62
-0.171
left 2
-0.173
left_68
-0.184
left_13
-0.194
right_7
-0.195
left_36
-0.195
right_53
-0.195
right_71
```

```
-0.195
right_27
-0.208
left_32
-0.209
right_36
-0.227
right_79
-0.233
right_2
-0.239
left_1
-0.302
Body_Temp
-0.327
left_82
```

```
    -0.372

    style="padding: 0 0.5em 0 0.5em; text-align: left; border: none;">
        left_40
```

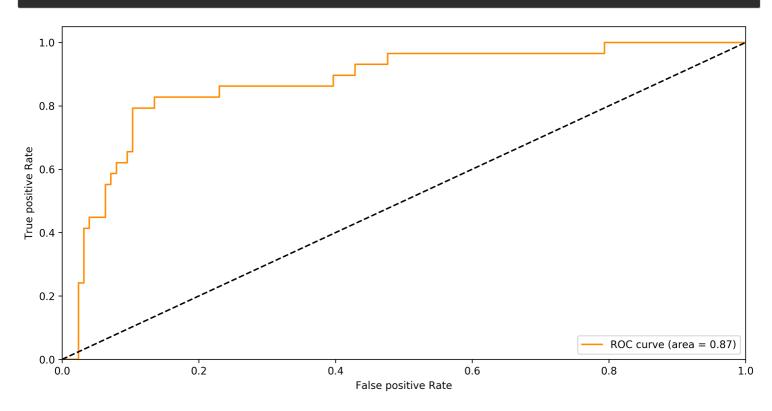
```
    -1.356

    style="padding: 0 0.5em 0 0.5em; text-align: left; border: none;">
        <BIAS&gt;
```

Step 2: Bring in the kernels!

1. Radial-basis function (RBF) kernel

```
clf2 = svm.SVC(kernel='rbf', gamma='auto')
dcsn_fn, preds = loocv_preds(clf2, norm_data, targets, threshold=0)
print("Area under ROC curve:", auc_roc(targets, dcsn_fn))
```

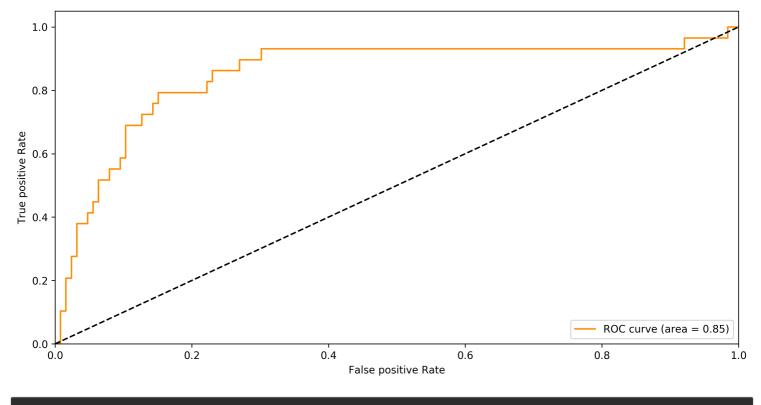


```
Area under ROC curve: 0.8732895457033388
```

```
dcsn_fn, preds = loocv_preds(clf2, norm_data, targets, threshold=-0.55)
rec, prec = rec_prec(list(targets), np.array(preds).flatten())
print("Recall:", rec)
print("Precision:", prec)
```

Recall: 0.7931034482758621 Precision: 0.6216216216216216

```
clf2 = svm.SVC(kernel='poly', degree=3, coef0=1, gamma='auto')
dcsn_fn, preds = loocv_preds(clf2, norm_data, targets, threshold=0)
print("Area under ROC curve:", auc_roc(targets, dcsn_fn))
```



```
Area under ROC curve: 0.8524904214559387

dcsn_fn, preds = loocv_preds(clf2, norm_data, targets, threshold=-0.75)
rec, prec = rec_prec(list(targets), np.array(preds).flatten())
print("Recall:", rec)
print("Precision:", prec)

Recall: 0.8275862068965517
Precision: 0.46153846153846156
```

I have tried to vary the degree of polynomial and 3 seemed to work best for degrees 2-10. I tried to vary the threshold to find a good balance for recall and precision, but the recall drops significantly when I move the threshold closer to zero (less negative). Seems like it is hard to project our data to a dimension that would make sense without overfitting (e.g., number of dimensions = number of data points).

Step 3: Time for neural nets

1. Experiment #1 with a toy network

```
import keras
from keras.layers import Dense
np.random.seed(1337) # for reproducible results
from keras.models import Sequential

# Building a multi-layer perceptron model
from keras.optimizers import Adam

adam = Adam(0.01)

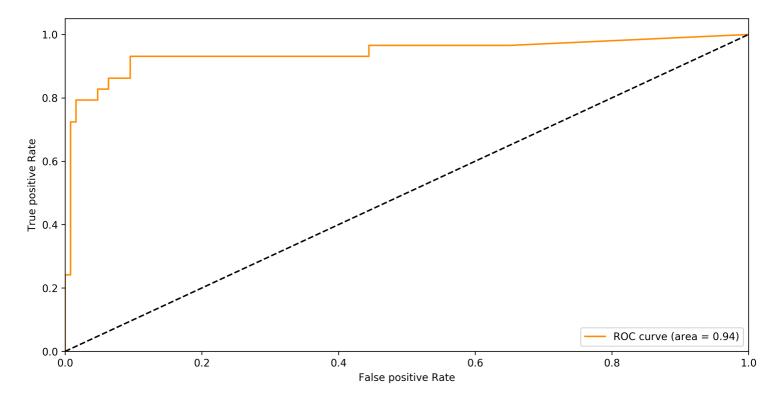
nn_model = Sequential()
nn_model.add(Dense(8, input_shape=(193,), activation='relu'))
nn_model.add(Dense(8, input_shape=(8,), activation='relu'))
nn_model.add(Dense(1, activation='sigmoid'))
```

```
dcsn_fn, preds = kfold_preds(nn_model, norm_data, targets, threshold=0.4, epochs=15)
    1.0
    0.8
 True positive Rate
    0.6
    0.4
    0.2
                                                                                                                        — ROC curve (area = 0.96)
    0.0
                                     0.2
                                                                 0.4
                                                                                               0.6
                                                                                                                            0.8
       0.0
                                                                                                                                                         1.0
                                                                        False positive Rate
  Area under ROC curve: 0.9584017515051998
  rec, prec = rec_prec(list(targets), np.array(preds).flatten())
  Recall: 0.8275862068965517
  Precision: 0.7741935483870968
Not too shaby, the best result so far.
 1. Experiment #2 with a bit more serious network. Gotta look out for overfitting because out dataset is small.
```

```
adam = Adam(0.01)

nn_model_2 = Sequential()
nn_model_2.add(Dense(16, input_shape=(193,), activation='relu'))
nn_model_2.add(Dense(16, input_shape=(16,), activation='relu'))
nn_model_2.add(Dense(16, input_shape=(16,), activation='relu'))
nn_model_2.add(Dense(1, activation='sigmoid'))

dcsn_fn, preds = kfold_preds(nn_model_2, norm_data, targets, threshold=0.5, epochs=15)
print("Area under ROC curve:", auc_roc(targets, dcsn_fn))
```



Area under ROC curve: 0.9408866995073892

rec, prec = rec_prec(list(targets), np.array(preds).flatten())
print("Recall:", rec)
print("Precision:", prec)

Recall: 0.7931034482758621 Precision: 0.8518518518518519

Best cumulative result (although a bit worse on recall than the previous model due to elevated threshold).

Part 6: Alternative approach to classification

So far, we have been utilizing all the features to come up with a decision boundary. However, as I mentioned before, it is very rare (2-5%) that someone has breast cancer on both sides. Thus, using sensor information from both sides might add more noise to the decision function. In the training dataset, the cancer in left or right breasts can be and should be evenly distributed, the right temperatures will also have some weights.

Example: there's a tumor in the left breast, but since we use both breasts to determine cancer, the right one would be "cooler" and more normally "temperaturized." Then, when we put together in a dot product, we can get "overwhelmed" by the right breast, for example, some abnormal patterns in the left breast might be averaged out by the weights in the right breast.

Step 1: Classify which breast is more likely to have cancer

We'll see which breast temperature distribution is furthest from the averaged temperature distribution of corresponding healthy breasts. Whichever breast that is furthest from its respective mean, it will be chosen as the "likely." The other one we will put to the mean healthy breast value to not affect during classification.

Step 2: Classify the breast with higher likelihood of cancer

Part 7: Works-to-be-done

In **Part 5**, I used the undergrad student descent to figure our the optimal threshold (e.g. at 0.5 the recall was too low, at 0.3 the precision was too low, go to the middle). The hyperparameter selection could be done automated with grid or random selection in a pre-determined range, and then the range would be modified to be more precise. Techniques like genetic algorithm could also be used to avoid local minima.

For the neural network, if I had more time, I would play around with network depth to see at which point it will start overfitting and settle with that architecture. To counterplay

overfitting, I would do data augmentation (oversampling) (via small wiggle around existing datapoints) especially on the cancer dataset to get more defined boundaries for this class. Dropout would also help the problem of overfitting on training dataset.

In this part, I have not looked into representing the sensor measurements as an image array. I would imagine that representing that way, we would avoid the correlated features (nearby sensors) better and find an assessment of temperature landscape as a group of sensors and not only individual sensors multiplied by some weights and added up together (losing information about locality).

Also, I think Part 6 might help boost a couple points in recall since we're "exposing" the irregularities more by not letting the non-affected breast confound the dot-product of measurements.

Last but not least, let me know if you have any questions!