Hi, we wish to start pertaining homework #2.

Theoretical section

```
from IPython.display import IFrame, display
filepath = "Theoretical_part_hw2.pdf"
IFrame(filepath, width=700, height=1000)
```

Out[1]:

Jupyter Notebook requires JavaScript.

Please enable it to proceed.

404: Not Found

You are requesting a page that does not exist!

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Practical Section

Q1:

Importing the data

```
In [2]:
         import pandas as pd
         import numpy as np
         from pathlib import Path
         import matplotlib.pyplot as plt
         import matplotlib as mpl
         import seaborn as sns
         from sklearn import metrics
         from sklearn import svm
         from sklearn.model selection import train test split as tts
         from sklearn.linear model import LogisticRegression
         from sklearn.model_selection import StratifiedKFold as SKFold
         from sklearn.model selection import GridSearchCV
         from sklearn.metrics import confusion matrix
         from matplotlib.ticker import (AutoMinorLocator, MultipleLocator)
         from sklearn.preprocessing import OneHotEncoder
         from sklearn.preprocessing import StandardScaler as stsc
         from sklearn.ensemble import RandomForestClassifier as RFC
         from sklearn.decomposition import PCA
         filepath = Path.cwd().joinpath('HW2 data.csv')
         T1D = pd.read csv(filepath)
```

hw2

Now we want to exclude patients that did not answer all the questions. and then assign the Diagnosis to a distinct dataframe

```
In [3]:
    T1D_raw_exnan = T1D.dropna()
    diagnosis = T1D_raw_exnan['Diagnosis']
    diagnosis = diagnosis.replace('Positive', True).replace('Negative', False)
    age = T1D_raw_exnan['Age']
    feat_col = set(T1D_raw_exnan.columns)
    feat_col.remove('Diagnosis')
    T1D_exnan_age = T1D_raw_exnan[feat_col]
    T1D_exnan_age = T1D_raw_exnan[feat_col]
    T1D_exnan_age = T1D_exnan_age.replace('Yes', 1)
    T1D_exnan_age = T1D_exnan_age.replace('No', 0)
    T1D_exnan_age = T1D_exnan_age.replace('Female', 1)
    T1D_exnan_age = T1D_exnan_age.replace('Male', 0)
```

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```
feat_col.remove('Age')
T1D_exnan = T1D_exnan_age[feat_col]
```

explanation for preprocessing:

we have ommitted every patient that has a NaN feature. we should check that this doesn't affect the distribution and force a bias. furthermore, for simplicity we turned the data into numeric values, yes to 1 no to 0, and Gender as well, 1 to female 0 to male (Done that before reading Q4). and diagnosis Vector turned into True and False

Q2:

Perfroming test-train split

```
In [4]: xtr, xte, ytr, yte = tts(T1D_exnan, np.ravel(diagnosis), test_size=0.2, random_state=7,
```

Q3:

Visualization and exploration of the data:

3. a.

```
vis_table = pd.DataFrame(index=feat_col, columns=['Train [%]', 'Test [%]', 'Delta [%]']
for ii, col in enumerate(feat_col):
    vis_table.loc[col, 'Train [%]'] = 100 * xtr[col].sum() / xtr[col].count()
    vis_table.loc[col, 'Test [%]'] = 100 * xte[col].sum() / xte[col].count()
    vis_table.loc[col, 'Delta [%]'] = vis_table.loc[col, 'Train [%]'] - vis_table.loc[c
    vis_table=vis_table.rename_axis('Positive Feature', axis=1)
    display(vis_table)
```

Hair Loss 33.9713 37.1429 -3.17157 Sudden Weight Loss 41.3876 41.9048 -0.517202 Partial Paresis 42.823 42.8571 -0.0341763 Increased Thirst 44.2584 45.7143 -1.45591 Family History 50.4785 52.381 -1.90248 Gender 37.799 33.3333 4.46571 Genital Thrush 21.2919 25.7143 -4.42242 Irritability 23.445 27.619 -4.17407
Partial Paresis 42.823 42.823 -0.0341763 Increased Thirst 44.2584 45.7143 -1.45591 Family History 50.4785 52.381 -1.90248 Gender 37.799 33.3333 4.46571 Genital Thrush 21.2919 25.7143 -4.42242
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Gender 37.799 33.3333 4.46571 Genital Thrush 21.2919 25.7143 -4.42242
Genital Thrush 21.2919 25.7143 -4.42242
Irritability 23.445 27.619 -4.17407
Muscle Stiffness 37.5598 36.1905 1.36933
Increased Urination 48.5646 53.3333 -4.76874
Increased Hunger 46.1722 42.8571 3.31511
Visual Blurring 44.7368 43.8095 0.927318
Obesity 16.2679 19.0476 -2.77968
Itching 48.3254 49.5238 -1.19845

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Positive Feature	Train [%]	Test [%]	Delta [%]
Weakness	59.5694	54.2857	5.28366
Delayed Healing	46.4115	43.8095	2.60196

We can see that the the representation of the various features are very similar in the train and test sets.

i. An imbalance of features between train and test might force a bias to the results because of under-representation or over representation to a particular phenomena that might build a model that doesn't fit the test data or correctly predict the futural data.

ii. If the medical condition is not complex, meaning that only exclusive set of sympotms are indicative/predictive for the condition, then stratifying the test-train data based on the diagnosis guarantees balanced representation of these features, meaning we can ignore other features that are imbalanced. For more complex datasets, we should try to stratify the data based on the features, i.e. looking at the n-dimesional distribution and try to split the data in a balanced way along with trying to balance the diagnosis.

3. b. in order to observe the relationship between features and labels we've constructed:

```
fig = plt.figure(figsize=(30,30))
sns.set_theme(style="ticks", color_codes=True)
for ii, col in enumerate(feat_col):
    ax = fig.add_subplot(4,4,ii+1)
    plot = sns.countplot(y=col, hue="Diagnosis", data=T1D_raw_exnan, palette="rocket",
```

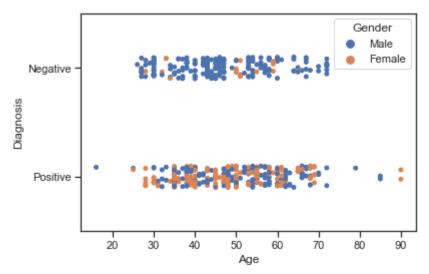
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Further, in the next plot we wanted to find any correlation between age and diagnosis, since the Age data is not binary, and further discriminated between females and males in order to observe any biases.

```
fig = plt.figure()
plot = sns.stripplot(x="Age", y="Diagnosis", hue="Gender", data=T1D_raw_exnan)
```

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- 1. d. i. first, we noticed that nearly all the females in this dataset have T1D, which might bias the model/predictions. afterwards we further investigated the countplots with two different perspectives:
 - 1- Picking the symptoms (features) that are present mainly in the T1D positive patients (https://www.mayoclinic.org/diseases-conditions/type-1-diabetes/symptoms-causes/syc-20353011), we see that if the symptom is present, most probably the patient is T1D-positive.
 - 2- looking at the positives patients and noticing that not all of them are showing the same symptoms, meaning that there is a T1D-positive patient that doesn't suffer from increased-thirst for instance. you may look at visual blurring symptom, we can see that there is nearly 50-50 partition in T1D-positive patients. Meaning, a model based mainly on this feature is not sensitive enough and will miss-classify a significant amount of T1D-positive patients.
- d. ii. Yes, we scanned the categorical plots above and searched for large gaps between Negative and Positive diagnosis correlated with specific features. For instance, look at Increased Urination, where you can see that most of the participants that have this symptom are positive for T1D so we can infer that this symptom is indicative/predictive for T1D. other than Increased Urination, there are: Increased Hunger, Increased Thirst, Irritibaility, Prtial Paresis and Visual Paresis.

Q4:

We have already done the transformation for hot vectors manually, as we binarized the yes/no dataset. We decided to neglect the Age feature at first, due to a broad distribution that might be detrimental for the performance, and we further corroborated the decision by showing above that there is no clear observed correlation between Age and Diagnosis. to observe our manual work excluding age feature:

```
In [8]: display(T1D_exnan)
```

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	Hair Loss	Sudden Weight Loss	Partial Paresis	Increased Thirst	Family History	Gender	Genital Thrush	Irritability	Muscle Stiffness	Increased Urination	Increa Hui
0	1	0	1	0	0	0	0	0	0	0	
1	1	0	0	0	0	0	0	0	0	0	
2	1	0	0	1	0	0	1	0	0	1	
3	0	0	0	0	1	1	0	0	0	0	
4	0	1	1	1	0	1	0	0	1	1	
•••											
518	1	0	1	0	0	0	1	1	1	1	
519	1	0	0	0	0	0	1	1	0	1	
562	1	0	0	0	1	1	0	1	0	1	
563	1	0	0	0	0	0	0	0	0	0	
564	0	0	0	0	0	0	0	0	0	0	

523 rows × 16 columns

←

Notwithstanding, we decided to test whether the age criteria might influence the model, we will include it in a parallel analysis, here we encode the whole data with one-hot vectors as offered by sklearn (pseudo-binarized the age feature, by trnasforming it for multiple 'pseudo-features'- every bit in the age sub-field indicates one number).

```
onehot = OneHotEncoder(handle_unknown='ignore')
onehot.fit(T1D_exnan_age)
T1D_exnan_age_oh = onehot.transform(T1D_exnan_age)
T1D_exnan_age_oh = pd.DataFrame(T1D_exnan_age_oh.toarray())
display(T1D_exnan_age_oh)
```

	0	1	2	3	4	5	6	7	8	9	•••	73	74	75	76	77	78	79	80	81	82
0	0.0	1.0	1.0	0.0	0.0	1.0	1.0	0.0	0.0	0.0		1.0	0.0	1.0	0.0	0.0	1.0	0.0	1.0	1.0	0.0
1	0.0	1.0	1.0	0.0	1.0	0.0	1.0	0.0	0.0	0.0		1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0
2	0.0	1.0	1.0	0.0	1.0	0.0	0.0	1.0	0.0	0.0		1.0	0.0	1.0	0.0	1.0	0.0	0.0	1.0	0.0	1.0
3	1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0	0.0	0.0		1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0
4	1.0	0.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0	0.0		0.0	1.0	1.0	0.0	0.0	1.0	0.0	1.0	1.0	0.0
•••																					
518	0.0	1.0	1.0	0.0	0.0	1.0	1.0	0.0	0.0	0.0		0.0	1.0	0.0	1.0	0.0	1.0	0.0	1.0	1.0	0.0
519	0.0	1.0	1.0	0.0	1.0	0.0	1.0	0.0	0.0	0.0		1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0	0.0	1.0
520	0.0	1.0	1.0	0.0	1.0	0.0	1.0	0.0	0.0	0.0		1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0	1.0	0.0
521	0.0	1.0	1.0	0.0	1.0	0.0	1.0	0.0	0.0	0.0		1.0	0.0	1.0	0.0	0.0	1.0	0.0	1.0	0.0	1.0

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```
    0
    1
    2
    3
    4
    5
    6
    7
    8
    9
    ...
    73
    74
    75
    76
    77
    78
    79
    80
    81
    82

    522
    1.0
    0.0
    1.0
    0.0
    1.0
    0.0
    0.0
    0.0
    0.0
    1.0
    0.0
    1.0
    0.0
    1.0
    0.0
```

523 rows × 83 columns

In order to input the one-hot vectors to the model construction, we would like to stratify it for train and test data accordingly to the same random_state we have already done

```
In [10]: xtr_age, xte_age, ytr_age, yte_age = tts(T1D_exnan_age_oh, np.ravel(diagnosis), test_si
```

Q5:

1. a. We will apply two different models starting with a linear model: Logistic regression.

```
In [11]:
          L = 120
          K = 5
          C = np.linspace(3/L, 3, L)
          C[0] = 0.000001
          indnum = L*2
          penalty = ['l1','l2']
          se, sp = np.zeros((L,1)), np.zeros((L,1))
          fig = plt.figure(figsize=(30,30))
          kf = SKFold(n_splits=K)
          validation dict lr = []
          validation dict age lr = []
          valdf_lr = pd.DataFrame(columns=('C', 'penalty', 'AUC', 'ACC', 'LogLoss', 'F1'))
          valdf_age_lr = pd.DataFrame(columns=('C', 'penalty', 'AUC', 'ACC', 'LogLoss', 'F1'))
          ind = 0
          for ii, cc in enumerate(C):
                  for p in penalty:
                      logloss, logloss_age = np.zeros(K), np.zeros(K)
                      acc, acc_age = np.zeros(K), np.zeros(K)
                      auc, auc_age = np.zeros(K), np.zeros(K)
                      f1, f1_age = np.zeros(K), np.zeros(K)
                      k = 0
                      for train idx, val idx in kf.split(xtr, ytr):
                          x_train, x_val, x_age_train, x_age_val = xtr.iloc[train_idx], xtr.iloc[
                           logreg = LogisticRegression(solver='saga', penalty=p, max_iter=10000, C
                          logreg_age = LogisticRegression(solver='saga', penalty=p, max_iter=1000
                           logreg.fit(x train, ytr[train idx])
                          logreg age.fit(x age train, ytr age[train idx])
                          y_pred_log, y_age_pred_log = logreg.predict(x_val), logreg_age.predict(
                          y_predproba_log, y_age_predproba_log = logreg.predict_proba(x_val), log
                          y_val_log, y_age_val_log = ytr[val_idx], ytr_age[val_idx]
                          acc[k], acc_age[k] = metrics.accuracy_score(y_val_log, y_pred_log), met
                          f1[k], f1_age[k] = metrics.f1_score(y_val_log, y_pred_log), metrics.f1_
                          logloss[k], logloss age[k] = metrics.log loss(y val log, y predproba lo
                          auc[k], auc_age[k] = metrics.roc_auc_score(y_val_log, y_predproba_log[:
                      validation_dict_lr.append({'C': cc, 'penalty': p, 'logloss' : logloss.mean(
                                              'AUC' : auc.mean()})
                      validation_dict_lr.append({'C': cc, 'penalty': p, 'logloss' : logloss_age.m
                                              'AUC' : auc_age.mean()})
                      valdf_lr.loc[ind, :] = cc, p, auc.mean(), acc.mean(), logloss.mean(), f1.me
                      valdf_age_lr.loc[ind, :] = cc, p, auc_age.mean(), acc_age.mean(), logloss_a
```

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ind+=1

<Figure size 2160x2160 with 0 Axes>

In [12]:

```
display(valdf_lr)
```

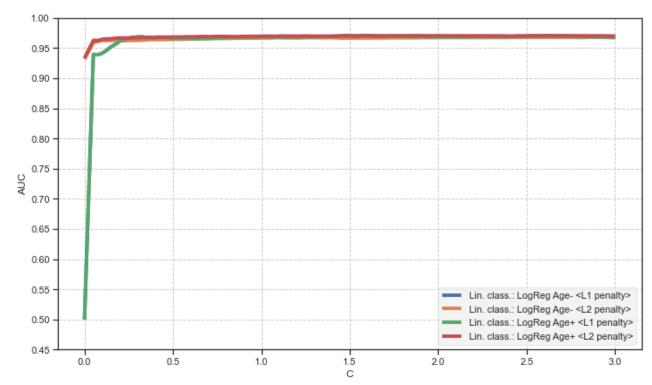
	С	penalty	AUC	ACC	LogLoss	F1
0	1e-06	l1	0.5	0.614831	0.670326	0.761471
1	1e-06	12	0.931582	0.614831	0.66654	0.761471
2	0.05	I1	0.939332	0.851434	0.405501	0.873607
3	0.05	12	0.958847	0.88973	0.351774	0.909689
4	0.075	I1	0.93884	0.875301	0.350019	0.900933
•••						
235	2.95	12	0.967815	0.925674	0.216197	0.938708
236	2.975	I1	0.968331	0.911388	0.21843	0.926859
237	2.975	12	0.967815	0.925674	0.216177	0.938708
238	3	I1	0.968331	0.911388	0.218457	0.926859
239	3	12	0.967815	0.925674	0.216157	0.938708

240 rows × 6 columns

```
fig, ax = plt.subplots(figsize=(12, 7))
    mpl.style.use('fivethirtyeight')
    plt1=plt.plot(valdf_lr['C'].loc[valdf_lr['penalty']=='l1'], valdf_lr['AUC'].loc[valdf_l
    plt2=plt.plot(valdf_lr['C'].loc[valdf_lr['penalty']=='l2'], valdf_lr['AUC'].loc[valdf_l
    plt3=plt.plot(valdf_age_lr['C'].loc[valdf_age_lr['penalty']=='l1'], valdf_age_lr['AUC']
    plt4=plt.plot(valdf_age_lr['C'].loc[valdf_age_lr['penalty']=='l2'], valdf_age_lr['AUC']
    plt.xlabel('C')
    plt.ylabel('AUC')
    plt.legend()
    ax.yaxis.set_major_locator(MultipleLocator(0.05))
    ax.grid(which='major', color='#CCCCCC', linestyle='--')
    ax.grid(which='minor', color='#CCCCCC', linestyle=':-')
    ax.set_ylim(0.45, 1)
```

Out[13]: (0.45, 1.0)

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untill now, we conducted manual parameter tuning and observed the above graph and noticed that the best model would be without taking into consideration the Age feature and using L2 penalty (for the logistic regression model). Now we want to apply an objective and automatic model selection based on GridSearchCV, also two parallel computations with and without age feature.

Excluding the Age feature:

Best parameters: C= 1.58, penalty=11 with score=0.96929

Including the Age feature:

```
estimator_lr_age = LogisticRegression(solver='saga', max_iter=10000)
params_dict = {
    'C' : np.linspace(0.01, 3, 100),
    'penalty' : ['l1', 'l2']
}
clf_lr_age = GridSearchCV(estimator_lr_age, params_dict, scoring='roc_auc')
clf_lr_age.fit(xtr_age, ytr_age)
```

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```
clf_lr_age.best_estimator_
best_lr_age = clf_lr_age.best_params_
```

Best parameters: C= 1.58, penalty=l1 with score=0.97098

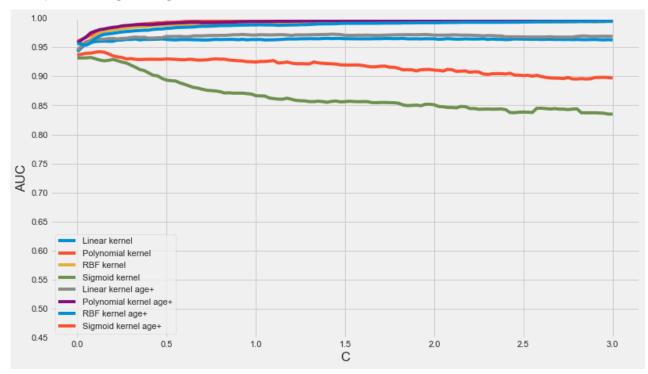
Next, we will apply a non-linear model to the dataset -> SVM:

```
In [18]:
          L = 120
          K = 5
          C = np.linspace(3/L, 3, L)
          C[0] = 0.000001
          kernel=['linear', 'poly', 'rbf', 'sigmoid']
          indnum = L*np.shape(kernel)
          se, sp = np.zeros((L,1)), np.zeros((L,1))
          fig = plt.figure(figsize=(30,30))
          kf = SKFold(n splits=K)
          validation dict svm = []
          validation_age_dict_svm = []
          valdf_svm = pd.DataFrame(columns=('C', 'kernel', 'AUC', 'ACC', 'Hinge Loss', 'F1'))
          valdf_age_svm = pd.DataFrame(columns=('C', 'kernel', 'AUC', 'ACC', 'Hinge Loss', 'F1'))
          ind = 0
          for ii, cc in enumerate(C):
                  for kk in kernel:
                      hingeloss, hingeloss_age = np.zeros(K), np.zeros(K)
                      acc, acc_age = np.zeros(K), np.zeros(K)
                      auc, auc_age = np.zeros(K), np.zeros(K)
                      f1, f1 age = np.zeros(K), np.zeros(K)
                      k = 0
                      for train_idx, val_idx in kf.split(xtr, ytr):
                          x_train, x_val, x_age_train, x_age_val = xtr.iloc[train_idx], xtr.iloc[
                          clf = svm.SVC(C=cc, kernel=kk, probability=True)
                          clf age = svm.SVC(C=cc, kernel=kk, probability=True)
                          clf.fit(x_train, ytr[train_idx])
                          clf age.fit(x age train, ytr age[train idx])
                          y_pred_svm = clf.predict(x_val)
                          y_predproba_svm = clf.predict_proba(x_val)
                          y age pred svm = clf age.predict(x age val)
                          y age predproba svm = clf age.predict proba(x age val)
                          y_val_svm, y_age_val_svm = ytr[val_idx], ytr_age[val_idx]
                          acc[k], acc_age[k] = metrics.accuracy_score(y_val_svm, y_pred_svm), met
                          f1[k], f1_age[k] = metrics.f1_score(y_val_svm, y_pred_svm), metrics.f1_
                          hingeloss[k], hingeloss age[k] = metrics.hinge loss(y val svm, y pred s
                          auc[k], auc age[k] = metrics.roc auc score(y val svm, y predproba svm[
                      validation dict svm.append({'C': cc, 'kernel': kk, 'Hinge Loss': logloss.m
                                              'AUC' : auc.mean()})
                      validation_age_dict_svm.append({'C': cc, 'kernel': kk, 'Hinge Loss' : loglo
                                              'AUC' : auc age.mean()})
                      valdf svm.loc[ind, :] = cc, kk, auc.mean(), acc.mean(), hingeloss.mean(), f
                      valdf_age_svm.loc[ind, :] = cc, kk, auc_age.mean(), acc_age.mean(), hingelo
                      ind+=1
```

<Figure size 2160x2160 with 0 Axes>

```
fig2, ax2 = plt.subplots(figsize=(12, 7))
plt5 = plt.plot(valdf svm['C'].loc[valdf svm['kernel']=='linear'], valdf svm['AUC'].loc
plt6 = plt.plot(valdf_svm['C'].loc[valdf_svm['kernel']=='poly'], valdf_svm['AUC'].loc[v
plt7 = plt.plot(valdf_svm['C'].loc[valdf_svm['kernel']=='rbf'], valdf_svm['AUC'].loc[valdf_svm['AUC'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[valdf_svm['Nuc'].loc[val
plt8 = plt.plot(valdf_svm['C'].loc[valdf_svm['kernel']=='sigmoid'], valdf_svm['AUC'].lo
plt5 = plt.plot(valdf_age_svm['C'].loc[valdf_age_svm['kernel']=='linear'], valdf_age_sv
plt6 = plt.plot(valdf_age_svm['C'].loc[valdf_age_svm['kernel']=='poly'], valdf_age_svm[
plt7 = plt.plot(valdf age svm['C'].loc[valdf age svm['kernel']=='rbf'], valdf age svm['
plt8 = plt.plot(valdf_age_svm['C'].loc[valdf_age_svm['kernel']=='sigmoid'], valdf_age_s
mpl.style.use('fivethirtyeight')
plt.xlabel('C')
plt.ylabel('AUC')
ax2.yaxis.set major locator(MultipleLocator(0.05))
ax.grid(which='major', color='#CCCCCC', linestyle='--')
ax.grid(which='minor', color='#CCCCCC', linestyle=':')
ax2.set_ylim(0.45, 1)
plt.legend()
```

Out[19]: <matplotlib.legend.Legend at 0x2dbf8aaf808>



Now, after we've done the preliminary comparison of the SVM models (based on their various kernels and varying C), we want to further test the complex kernels (poly, rbf, sigmoid) which leverage another hyperparameter- Gamma.

excluding age feature:

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```
In [21]:
          print("Best parameters: C = " + str("{0:.2f}".format(best svm['C'])) + " gamma= " + str
                ", kernel=" + best svm['kernel'] + " with score=" + str("{0:.5f}".format(clf svm
         Best parameters: C = 2.64 gamma= 0.31, kernel=rbf with score=0.99623
        including age feature:
In [23]:
          estimator svm age = svm.SVC()
          params dict = {
               'kernel' : ['poly', 'sigmoid', 'rbf'],
                       : np.linspace(0.01, 3, 100),
                       : np.linspace(0.01, 3, 100),
          clf svm age = GridSearchCV(estimator svm age, params dict)
          clf svm age.fit(xtr age, ytr age)
          clf svm age.best estimator
          best_svm_age = clf_svm_age.best_params_
In [25]:
          print("Best parameters: C = " + str("{0:.2f}".format(best_svm_age['C'])) + ", gamma = "
                ", kernel=" + str(best svm age['kernel']) + " with score=" + str("{0:.5f}".forma
         Best parameters: C = 1.25, gamma = 0.28, kernel=rbf with score=0.98087
 In [ ]:
          #In order to observe the contrast we shall overlay the AUC plots
 In [ ]:
          #from collections import OrderedDict
          #fig, ax = plt.subplots(figsize=(12, 7))
          #mpl.style.use('fivethirtyeight')
          #ax.set title('Receiver-operator AUC of all Classifiers'.format('seaborn'), color='C3')
          #plt1 = plt.plot(valdf_lr['C'].loc[valdf_lr['penalty']=='l1'], valdf_lr['AUC'].loc[vald]
          #plt2 = plt.plot(valdf_lr['C'].loc[valdf_lr['penalty']=='l2'], valdf_lr['AUC'].loc[vald]
          #plt3 = plt.plot(valdf_age_lr['C'].loc[valdf_age_lr['penalty']=='l1'], valdf_age_lr['AU
          #plt4 = plt.plot(valdf age lr['C'].loc[valdf age lr['penalty']=='l2'], valdf age lr['AU
          #plt5 = plt.plot(valdf_svm['C'].loc[valdf_svm['kernel']=='linear'], valdf_svm['AUC'].lo
          #plt6 = plt.plot(valdf svm['C'].loc[valdf svm['kernel']=='poly'], valdf svm['AUC'].loc[
          #plt7 = plt.plot(valdf_svm['C'].loc[valdf_svm['kernel']=='rbf'], valdf_svm['AUC'].loc[v
          #plt8 = plt.plot(valdf_svm['C'].loc[valdf_svm['kernel']=='sigmoid'], valdf_svm['AUC'].l
          #plt9 = plt.plot(valdf_age_svm['C'].loc[valdf_age_svm['kernel']=='linear'], valdf_age_s
          #plt10 = plt.plot(valdf age svm['C'].loc[valdf age svm['kernel']=='poly'], valdf age sv
          #plt11 = plt.plot(valdf age svm['C'].loc[valdf age svm['kernel']=='rbf'], valdf age svm
          #plt12 = plt.plot(valdf_age_svm['C'].loc[valdf_age_svm['kernel']=='sigmoid'], valdf_age
          #plt.xlabel('C')
          #plt.ylabel('AUC')
          #ax.yaxis.set major locator(MultipleLocator(0.05))
          #ax.grid(which='major', color='#CCCCCC', linestyle='--')
          #ax.grid(which='minor', color='#CCCCCC', linestyle=':')
          #ax.set_ylim(0.45, 1)
          #ax.Legend()
```

1. b. the interim results based on the training data are listed: note that including or excluding the Age feature gave much similar results and due to it's computational labor, we decided to neglect the age.

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```
In [26]: print("i- Lin. Class.: Logistic regression with " + best_lr['penalty'] +" penalty and
    print("ii- Nonlin. Class.: SVM classifier with " + best_svm['kernel'] +" kernel, regula
```

i- Lin. Class.: Logistic regression with l1 penalty and regularization parameter C = 1.5 $\,^{8}$ ii- Nonlin. Class.: SVM classifier with rbf kernel, regularization parameter C = 2.64 an d gamma = $\,^{0.31}$

Next, we will report the evaluation metrics for the models mentioned above. First, we will train the models on the training set then report the metrics for classifying the train set itself. Second, we will report the metrics for the test set classification.

Now, let us report the metrics for classifying the two best models:

• for the linear classifier LR:

```
In [27]:
          C = best lr['C']
          penalty=best lr['penalty']
          testdf_lr = pd.DataFrame(index=('Train', 'Test'), columns=('C', 'penalty', 'AUC', 'ACC'
          LR = LogisticRegression(solver='saga', penalty=penalty, max_iter=10000, C=C)
          LR.fit(xtr, ytr)
          y_pred_lr = LR.predict(xtr)
          y predproba lr = LR.predict proba(xtr)
          acc = metrics.accuracy_score(ytr, y_pred_lr)
          f1 = metrics.f1 score(ytr, y pred lr)
          logloss = metrics.log_loss(ytr, y_predproba_lr)
          auc = metrics.roc_auc_score(ytr, y_predproba_lr[:,1])
          testdf_lr.loc['Train', :] = C, penalty, auc, acc, logloss, f1
          y_pred_lr = LR.predict(xte)
          y predproba lr = LR.predict proba(xte)
          acc = metrics.accuracy score(yte, y pred lr)
          f1 = metrics.f1_score(yte, y_pred_lr)
          logloss = metrics.log_loss(yte, y_predproba_lr)
          auc = metrics.roc_auc_score(yte, y_predproba_lr[:,1])
          testdf_lr.loc['Test', :] = C, penalty, auc, acc, logloss, f1
          display(testdf_lr)
```

	C	penalty	AUC	ACC	LogLoss	F1
Train	1.58051	I1	0.979336	0.933014	0.180184	0.944882
Test	1.58051	I1	0.98247	0.942857	0.165017	0.952381

note: the test set got better results from the train, which can be surprising. But, let us recall that the training and searching for the best model used a K-fold procedure and not the raw train set.

• for the Nonlinear classifier SVM:

```
In [28]:
    C = best_svm['C']
    kernel=best_svm['kernel']
    gamma=best_svm['gamma']
    testdf_svm = pd.DataFrame(index=('Train', 'Test'), columns=('C', 'kernel', 'gamma', 'AU
```

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```
clf_svm = svm.SVC(C=C, kernel=kernel, gamma=gamma, probability=True)
clf_svm.fit(xtr, ytr)
y pred svm = clf svm.predict(xtr)
y_predproba_svm = clf_svm.predict_proba(xtr)
acc = metrics.accuracy_score(ytr, y_pred_svm)
f1 = metrics.f1_score(ytr, y_pred_svm)
hingeloss = metrics.hinge loss(ytr, y pred svm)
auc = metrics.roc_auc_score(ytr, y_predproba_svm[:,1])
testdf_svm.loc['Train', :] = C, kernel, gamma, auc, acc, hingeloss, f1
y pred svm = clf svm.predict(xte)
y predproba svm = clf svm.predict proba(xte)
acc = metrics.accuracy_score(yte, y_pred_svm)
f1 = metrics.f1 score(yte, y pred svm)
hingeloss = metrics.hinge_loss(yte, y_pred_svm)
auc = metrics.roc_auc_score(yte, y_predproba_svm[:,1])
testdf svm.loc['Test', :] = C, kernel, gamma, auc, acc, hingeloss, f1
display(testdf svm)
```

	C	kernel	gamma	AUC	ACC	HingeLoss	F1
Train	2.63758	rbf	0.31202	0.997728	0.995215	0.389952	0.996124
Test	2.63758	rbf	0.31202	0.984375	0.980952	0.409524	0.984615

5. c. Not surprisinigly, the Nonlinear Classifier performs a bit better than the Linear Classifier. Generally, Linear classifiers are sub-type of nonlinear classifier, so usually nonlinear classifiers perform better (but more computationally expensive). Pertaining this data of binary features, it may result in a multi-dimensional saw-like hyper-groups that may only be seperated wisely by nonlinear classifier.

Q6: Feature Selection

```
rfc = RFC(n_estimators=1000, criterion='gini')
rfc.fit(xtr, ytr)
y_rfc_pred = rfc.predict(xte)
rfc_acc = metrics.accuracy_score(yte, y_rfc_pred)
print("The accuracy of the random forest classifier is: " + str("{0:.5f}".format(rfc_ac)
```

The accuracy of the random forest classifier is: 0.98095

We see that the accuracy of this model is the highest we observed so far (equal to the accuracy of the svm on test data). Next, we will try to understand which features are of great importance and then check for correlation with the features we explored manually:

```
rfc_res = pd.DataFrame(columns = (T1D_exnan.columns))
rfc_res.loc[0,:] = rfc.feature_importances_

display(rfc_res)
```

Hair Loss	Sudden Weight Loss	Partial Paresis	Increased Thirst	Family History	Gender	Genital Thrush	Irritability	Muscle Stiffness	
-----------	--------------------------	--------------------	---------------------	-------------------	--------	-------------------	--------------	---------------------	--

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	Hair Loss	Sudden Weight Loss	Partial Paresis	Increased Thirst	Family History	Gender	Genital Thrush	Irritability	Muscle Stiffness	
0	0.0452326	0.0599982	0.0492593	0.194483	0.0110446	0.127512	0.0241703	0.0375492	0.0300864	0
4										•

- 1. a. i. The 2 most important features according to the random forest are **Increased Thirst** and **Increased Urination** (based on feature importance score provided by random forest model). If we inspect further, we can observe that the third important feature is the Gender feature, which can be explained by the bias seen beforehand- most of the Females in this dataset are T1D positive.
 - a. ii. This matches up exactly with the feature exploration we did above.

Q7: Data Separability Visualization

Note: using PCA to reduce dimensioms with binary data could be problematic and is not optimal. we show below the results for a scaled PCA on the binary data and a more usable method for dimensionality reduction on binary data -> Logistic PCA.

```
In [31]:
    scaler = stsc() #standard scaling
    xtr_sc = scaler.fit_transform(xtr)
    xte_sc = scaler.transform(xte)

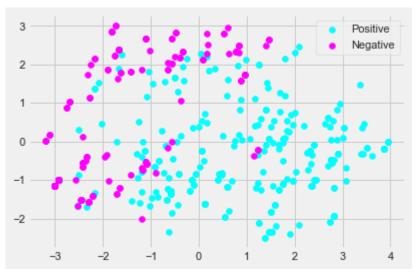
In [32]:
    pca = PCA(n_components=2)
    pca.fit(xtr_sc)
    pca.explained_variance_ratio_
    #display(pca.components_)

    pca_res = pca.transform(xtr_sc)

    plt.scatter(pca_res[ytr==1,0], pca_res[ytr==1,1], color='cyan', label='Positive')
    plt.scatter(pca_res[ytr==0,0], pca_res[ytr==0,1], color='magenta', label='Negative')
    plt.legend()
```

Out[32]: <matplotlib.legend.Legend at 0x2dbf88ad988>

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1. b. As observed in the scatter plot above, the two groups are generally distinct but are volatile and diffusive, in other words, it is apparent that there is 'geographical' seperability, but the border is blurred. Either way, a perfect separation between this two groups is not feasible using these features only.

1. c. + d.

```
In [33]:
    d = {'PC1': pca_res[:,0], 'PC2': pca_res[:,1], 'Diagnosis': ytr}
    pca_df = pd.DataFrame(data=d)
    #pca_df['PC1','PC2']= pca_res
    pca_df
```

Out[33]:		PC1	PC2	Diagnosis
	0	3.380113	-0.305302	True
	1	-3.008438	-1.147374	False
	2	2.751914	0.033196	True
	3	1.837806	-0.977971	True
	4	0.595998	2.948969	False
	•••			
	413	2.805743	0.528134	True
	414	-0.008371	0.391407	True
	415	-2.347778	-0.388015	False
	416	-2.432786	-0.656536	False
	417	-0.407392	2.175156	False

418 rows × 3 columns

Firstly, we want to split our data into train and test subsets (stratified by the diagnosis).

```
In [34]: #xtr_pca, xte_pca, ytr_pca, yte_pca = tts(pca_df.iloc[:,0:2], np.ravel(diagnosis), test
```

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```
xtr_rf, xte_rf, ytr_rf, yte_rf = tts(T1D_exnan.loc[:,['Increased Thirst','Increased Uri
```

c. Testing the models on the PCA-extracted features (PC1, PC2)

• Logistic Regression (Linear Classifier):

```
In [35]:
    C = best_lr['C']
    penalty=best_lr['penalty']
    testdf_pca_lr = pd.DataFrame(columns=('C', 'penalty', 'AUC', 'ACC', 'LogLoss', 'F1'))
    LR = LogisticRegression(solver='saga', penalty=penalty, max_iter=10000, C=C)
    LR.fit(xtr_sc, ytr)
    y_pred_lr = LR.predict(xte_sc)
    y_predproba_lr = LR.predict_proba(xte_sc)

acc = metrics.accuracy_score(yte, y_pred_lr)
    f1 = metrics.f1_score(yte, y_pred_lr)
    logloss = metrics.log_loss(yte, y_predproba_lr)
    auc = metrics.roc_auc_score(yte, y_predproba_lr[:,1])
    testdf_pca_lr.loc[0, :] = C, penalty, auc, acc, logloss, f1
    display(testdf_pca_lr)
```

```
        C
        penalty
        AUC
        ACC
        LogLoss
        F1

        0
        1.58051
        I1
        0.976372
        0.942857
        0.170027
        0.952381
```

• SVM (Nonlinear Classifier):

```
In [36]:

C = best_svm['C']
kernel = best_svm['gamma']
testdf_pca_svm = pd.DataFrame(columns=('C', 'kernel','gamma', 'AUC', 'ACC', 'Hinge Loss
clf = svm.SVC(C=C, kernel=kernel, probability=True)
clf.fit(xtr_sc, ytr)
y_pred_svm = clf.predict(xte_sc)
y_predproba_svm = clf.predict_proba(xte_sc)
acc = metrics.accuracy_score(yte, y_pred_svm)
f1 = metrics.f1_score(yte, y_pred_svm)
hingeloss = metrics.hinge_loss(yte, y_pred_svm)
auc = metrics.roc_auc_score(yte, y_pred_svm[:,1])
testdf_pca_svm.loc[0, :] = C, kernel, gamma, auc, acc, hingeloss, f1
display(testdf_pca_svm)
```

```
        C
        kernel
        gamma
        AUC
        ACC
        Hinge Loss
        F1

        0
        2.63758
        rbf
        0.31202
        0.985899
        0.971429
        0.419048
        0.976744
```

d. Testing the models on the Random Forest extracted features (Increased Thirst and Urination)

Logistic Regression (Linear Classifier):

```
In [37]:
    C = best_lr['C']
    penalty = best_lr['penalty']
```

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```
testdf_rf_lr = pd.DataFrame(columns=('C', 'penalty', 'AUC', 'ACC', 'LogLoss', 'F1'))
LR = LogisticRegression(solver='saga', penalty=penalty, max_iter=10000, C=C)
LR.fit(xtr_rf, ytr_rf)
y_pred_lr = LR.predict(xte_rf)
y_predproba_lr = LR.predict_proba(xte_rf)
acc = metrics.accuracy_score(yte_rf, y_pred_lr)
f1 = metrics.f1_score(yte_rf, y_pred_lr)
logloss = metrics.log_loss(yte_rf, y_predproba_lr)
auc = metrics.roc_auc_score(yte_rf, y_predproba_lr[:,1])
testdf_rf_lr.loc[0, :] = C, penalty, auc, acc, logloss, f1
display(testdf_rf_lr)
```

```
        C
        penalty
        AUC
        ACC
        LogLoss
        F1

        0
        1.58051
        I1
        0.962843
        0.942857
        0.231786
        0.953125
```

• SVM (Nonlinear Classifier):

```
In [38]:

C = best_svm['C']
kernel = best_svm['gamma']
testdf_rf_svm = pd.DataFrame(columns=('C', 'kernel', 'gamma', 'AUC', 'ACC', 'Hinge Loss
clf = svm.SVC(C=C, kernel=kernel, probability = True)
clf.fit(xtr_rf, ytr_rf)
y_pred_svm = clf.predict(xte_rf)
y_predproba_svm = clf.predict_proba(xte_rf)
acc = metrics.accuracy_score(yte_rf, y_pred_svm)
f1 = metrics.f1_score(yte_rf, y_pred_svm)
hingeloss = metrics.hinge_loss(yte_rf, y_pred_svm)
auc = metrics.roc_auc_score(yte_rf, y_predproba_svm[:,1])
testdf_rf_svm.loc[0, :] = C, kernel, gamma, auc, acc, hingeloss, f1
display(testdf_rf_svm)
```

```
        C
        kernel
        gamma
        AUC
        ACC
        Hinge Loss
        F1

        0
        2.63758
        rbf
        0.31202
        0.947599
        0.942857
        0.447619
        0.953125
```

e. As we have seen, the model trained on dimensionality reduced training set performed better. This result was predictable, as in the random forest procedure we have picked the two most "important" features, when in the dimensionality reduced procedure we used the two "most" principal components which include information from the whole set of features therefore is more significant for the model and helps the model to predict better. Being sufficed by 2 features may be lucrative, because for every person we most obtain only a small number of features which is less time consuming and cheaper, and the computing process will be fast. On the other hand, obtaining a large number of features will add information which most probably will assist the model to predict better, although it is more time consuming and more expensive.

Bonus: We tried to think on a good way to perform a dimensionallity reduction on the binary dataset. we saw online some possible solutions but tried to think on another one on are own. We will give every patient two scores, one relying on all of the features (weighted by the importance vector output by the random forest classifier) and the other on the MayoClinic

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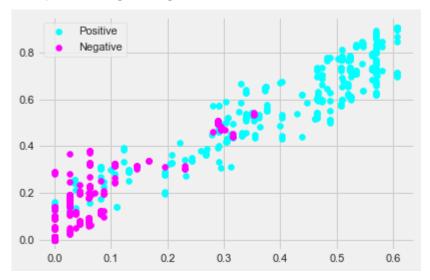
(https://www.mayoclinic.org/diseases-conditions/type-1-diabetes/symptoms-causes/syc-20353011) ones.

```
In [39]:
    rfc_scale = rfc.feature_importances_
    T1D_scaled = T1D_exnan * rfc_scale
    all_feat_score = T1D_scaled.sum(axis=1)
#all_feat_score = T1D_scaled.sum(axis=1) / len(feat_col)

T1D_mayo = T1D_scaled.loc[:,{'Increased Thirst', 'Increased Urination', 'Increased Hung
#T1D_exnan
mayo_feat_score = T1D_mayo.sum(axis=1)
#mayo_feat_score = T1D_mayo.sum(axis=1) / len(T1D_mayo.columns)

plt.scatter(mayo_feat_score[diagnosis==1], all_feat_score[diagnosis==1], color='cyan',
    plt.scatter(mayo_feat_score[diagnosis==0], all_feat_score[diagnosis==0], color='magenta
    plt.legend()
```

Out[39]: <matplotlib.legend.Legend at 0x2dbe6d69748>



```
In [40]:
          xtr, xte, ytr, yte = tts(T1D scaled, np.ravel(diagnosis), test size=0.2, random state=7
          T1D_mayo_tr = xtr.loc[:,{'Increased Thirst', 'Increased Urination', 'Increased Hunger',
          T1D_mayo_te = xte.loc[:,{'Increased Thirst', 'Increased Urination', 'Increased Hunger',
          all_feat_score_tr = xtr.sum(axis=1)
          all feat score te = xte.sum(axis=1)
          mayo_feat_score_tr = T1D_mayo_tr.sum(axis=1)
          mayo feat score te = T1D mayo te.sum(axis=1)
          xtrain = pd.DataFrame(columns=('all', 'mayo'))
          xtrain['all'] = all feat score tr
          xtrain['mayo'] = mayo_feat_score_tr
          xtest = pd.DataFrame(columns=('all', 'mayo'))
          xtest['all'] = all_feat_score_te
          xtest['mayo'] = mayo_feat_score_te
          #scaler = stsc() #standard scaling
          #xtr sc = scaler.fit transform(xtrain)
          #xte_sc = scaler.transform(xtest)
```

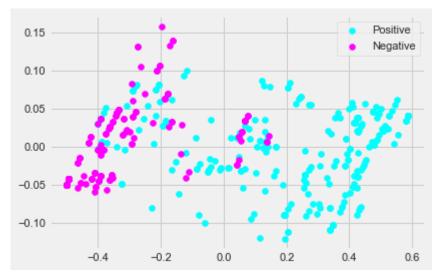
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```
In [41]:
    xtr_sc = xtrain
    pca = PCA(n_components=2)
    pca.fit(xtr_sc)
    pca.explained_variance_ratio_
    #dispLay(pca.components_)

pca_res = pca.transform(xtr_sc)

plt.scatter(pca_res[ytr==1,0], pca_res[ytr==1,1], color='cyan', label='Positive')
    plt.scatter(pca_res[ytr==0,0], pca_res[ytr==0,1], color='magenta', label='Negative')
    plt.legend()
```

Out[41]: <matplotlib.legend.Legend at 0x2dbf86f3548>



To summarize, we managed to project tha binary data on a single axis which represents them all. we can see that although the sepereation is not perfect, we got pretty nice visual results by making binary data of multiple features spreaded on a single score meter.



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