# diagnostic-predictor

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## 1 Predicting Patient Diagnosis from Natural Language Symptoms

#### 1.1 AAI-501 Team 3 Final Project

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GitHub Repository: https://github.com/t4ai/AAI-501-Team3

```
[1]: import re
     import numpy as np
     import pandas as pd
     import matplotlib.pyplot as plt
     import scipy.stats as stats
     import random
     import statistics
     import spacy
     import nltk
     import pickle
     # SciKit
     from sklearn.feature_extraction.text import CountVectorizer
     from sklearn.feature_extraction.text import TfidfTransformer
     from sklearn.naive_bayes import MultinomialNB
     from sklearn.linear_model import LogisticRegression
     from sklearn.linear_model import Ridge
     from sklearn.model_selection import GridSearchCV, RandomizedSearchCV, __
      →train_test_split, cross_val_score, cross_val_predict
     from sklearn.metrics import accuracy_score, confusion_matrix, precision_score,_
      Grecall_score, ConfusionMatrixDisplay, make_scorer, f1_score, □

¬classification_report
     from sklearn import preprocessing
     from nltk.corpus import stopwords
     import tensorflow_hub as hub
     import tensorflow.compat.v1 as tf
     tf.disable_eager_execution()
```

```
import keras
from keras.layers import Input, Lambda, Dense
from keras.models import Model
import keras.backend as K
```

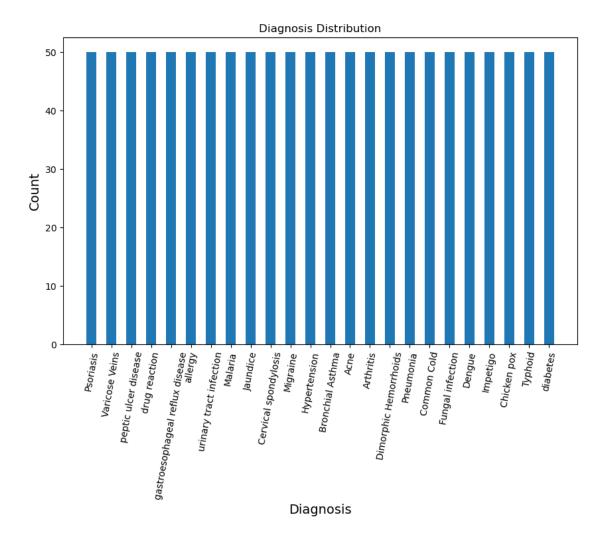
2023-08-07 16:42:45.111790: I tensorflow/core/platform/cpu\_feature\_guard.cc:193] This TensorFlow binary is optimized with oneAPI Deep Neural Network Library (oneDNN) to use the following CPU instructions in performance-critical operations: SSE4.1 SSE4.2 AVX AVX2 FMA To enable them in other operations, rebuild TensorFlow with the appropriate compiler flags.

#### 1.2 Exploratory Data Analysis

- Load data into dataframe
- Generate and review descriptive statistics of the dataset/variables
- Plot visualization of data spread for each variable

```
[120]: # Load dataset into dataframe and peek at head
symptoms_disease_df = pd.read_csv('./Symptom2Disease.csv')
symptoms_disease_df.head()
```

```
[120]:
                     label
          index
                                                                         text
             O Psoriasis I have been experiencing a skin rash on my arm...
             1 Psoriasis My skin has been peeling, especially on my kne...
             2 Psoriasis I have been experiencing joint pain in my fing...
             3 Psoriasis There is a silver like dusting on my skin, esp...
             4 Psoriasis My nails have small dents or pits in them, and...
[121]: # plot distribution of diagnoses
       fig, ax = plt.subplots(figsize=(10, 6))
       categories = symptoms_disease_df['label'].value_counts().index
       counts = symptoms_disease_df['label'].value_counts().values
       plt.bar(categories, counts, width=0.5)
       # Add labels
       plt.ylabel('Count',fontsize=14)
       plt.xlabel('Diagnosis',fontsize=14)
       plt.xticks(fontsize=10, rotation = 80)
       plt.yticks(fontsize=10)
       # Add Title
       plt.title('Diagnosis Distribution',fontsize=12);
       plt.show()
```



### 1.3 Data Cleanup

- Perform routine cleanup on data:
  - remove punctuation marks
  - convert to lowercase
  - remove numbers
  - remove whitespace
- Remove stop words
- Lemmatize the text
  - Normalize to base words

```
[122]: # helper function to remove special characters/numbers and normalzie to lower_

case
def string_clean(df, column_name):
    # remove punctuation marks
punctuation = '!"#$%&()*+-/:;<=>?@[\\]^_`{|}~'
```

```
df[column_name] = df[column_name].apply(lambda x: ''.join(ch for ch in x if_
        ⇔ch not in set(punctuation)))
           # convert text to lowercase
           df[column_name] = df[column_name].str.lower()
           # remove numbers
           df[column_name] = df[column_name].str.replace("[0-9]", " ", regex=True)
           # remove whitespaces
           df[column_name] = df[column_name].apply(lambda x:' '.join(x.split()))
           return df
       # helper function to remove common stop words from the text
       def remove_stop_words(df, column_name):
           nltk.download('stopwords', quiet=True)
           stop = stopwords.words('english')
           df[column_name] = df[column_name].apply(lambda x: " ".join(x for x in x.
        ⇔split() if x not in stop))
           return df
       # helper function to combine both preprocessing steps above into one function
       def preprocess_text(text_arr):
           temp_df = pd.DataFrame({'text': text_arr})
           temp_df = string_clean(temp_df, 'text')
           temp df = remove stop words(temp df, 'text')
           return temp_df['text'].to_numpy()
 [5]: # test the preprocessing function
       txt = preprocess_text(["WHY heLLo Mr. $Soul I WALKED BY? To Pick UP a %reason!!!
       print(txt)
      ['hello mr. soul walked pick reason']
[123]: # run text cleanup on dataset
       symptoms_disease_df = string_clean(symptoms_disease_df, 'text')
       symptoms_disease_df['text'].head()
[123]: 0
            i have been experiencing a skin rash on my arm...
           my skin has been peeling, especially on my kne...
       1
            i have been experiencing joint pain in {\tt my} fing...
       3
           there is a silver like dusting on my skin, esp...
           my nails have small dents or pits in them, and...
      Name: text, dtype: object
```

**Lemmatization helper functions** Lemmatization will normalize the text into base dictionary form or root words

```
[125]: # Lemmatize to normalize words - use only for Embeddings below
nlp = spacy.load('en_core_web_sm', disable=['parser', 'ner'])

# function to lemmatize symptoms text
def lemmatization(symptoms):
    output = []
    for i in symptoms:
        s = [token.lemma_ for token in nlp(i)]
        output.append(' '.join(s))
    return output
```

#### 1.4 Data Preparation

- Split data into test/train/validate datasets (80/10/10)
- Create 3 datasets for experimentation:
  - 1. Vectorize natrual language text using TFIDF
  - 2. Setup embeddings using ELMo (Embeddings from Language Models)
- Encode labels with one-hot encoding
- For each of the above, ensure no data leakage by separating train/test

```
[126]: # extract symptom description text to df X (features)
X = symptoms_disease_df['text'].copy()

# extract diagnosis into df for y (labels)
y_raw = symptoms_disease_df['label'].copy()
```

```
[10]: # Helper functions for one-hot encode/decode for labels (y)
def encode(en, labels):
    enc = en.transform(labels)
    return keras.utils.to_categorical(enc)

def decode(en, one_hot):
    dec = np.argmax(one_hot, axis=1)
    return en.inverse_transform(dec)
```

```
# one hot encode y
encoder = preprocessing.LabelEncoder()
encoder.fit(y_raw)
```

#### [10]: LabelEncoder()

```
[112]: # Create one-hot ecoded vectors for y (for NN model)
y_train_enc = encode(encoder, y_train)
y_val_enc = encode(encoder, y_val)
y_test_enc = encode(encoder, y_test)
```

#### 1.4.1 Prepare Lemmatized copies of datasets

Lemmatize train, validation and test datsets. This will be used for experiments with NBC.

```
[]: # Copy datasets for lemmatization
X_train_lemm = X_train.copy()
X_val_lemm = X_val.copy()
X_test_lemm = X_test.copy()

# Lemmatize the datasets
X_train_lemm = pd.Series(lemmatization(X_train_lemm))
X_val_lemm = pd.Series(lemmatization(X_val_lemm))
X_test_lemm = pd.Series(lemmatization(X_test_lemm))
```

#### 1.4.2 Prepare TF-IDF Vectorized datasets

Vectorize train, validation and test datsets. First, use CountVectorizer to vectorize the text based on word frequency. Next, use TF-IDF embedding to create dense vectors based on term frequency and in-document frequency. The vectorizer and transformer will be trained only on the training dataset to avoid data leakage.

```
[145]: # Start with count vecotrizer to build vocabulary - fit on train data first count_vectorizer = CountVectorizer()

# Vectorize training data to create bag of words - fit the vectorizor on the_u training set only to avoid data leakage

X_train_count = count_vectorizer.fit_transform(X_train_lemm)

X_train_count.shape

# Vectorize test and validation data
```

```
X_val_count = count_vectorizer.transform(X_val_lemm)
X_test_count = count_vectorizer.transform(X_test_lemm)

# Fit tfidf vectorizer on training count only to avoid data leakage
tf_transformer = TfidfTransformer(use_idf=True).fit(X_train_count)

# Vectorize training, val, test data to TFIDF
X_train_tfidf = tf_transformer.transform(X_train_count)
X_val_tfidf = tf_transformer.transform(X_val_count)
X_test_tfidf = tf_transformer.transform(X_test_count)
```

#### 1.4.3 Setup ELMo Embeddings Helper Functions

Helper functions for use with ELMo classifier. These functions intialize the embedder for use in the DNN layer. Also supporting helper functions to experiement with ELMo separate from DNN.

```
[15]: # Load pre trained ELMo model
elmo = hub.Module("https://tfhub.dev/google/elmo/3", trainable=True)
```

```
[16]: # ELMo Embedding - use as embedding layer in NN
      def elmo_embedding(x):
          return elmo(tf.squeeze(tf.cast(x, tf.string)), signature="default", ___
       →as_dict=True) ["default"]
      def get_elmo_batches(dataset, batch_size):
        return [dataset[i:i+batch size] for i in range(0,len(dataset),batch size)]
      # save generated embedding to pickle file
      def save_embedding(embedding, file_name):
        # save elmo_train_new
       pickle_out = open(file_name,"wb")
       pickle.dump(embedding, pickle_out)
       pickle_out.close()
      # load previously generated embedding from pickle file
      def load_saved_embedding(file_name):
       pickle_in = open(file_name, "rb")
       return pickle.load(pickle_in)
      # use for generating stand alone embeddings (ie: to train shallow classifiers)
      def elmo preprocess embeddings(x):
        embeddings = elmo(x, signature="default", as_dict=True)["elmo"]
       with tf.Session() as sess:
          sess.run(tf.global_variables_initializer())
          sess.run(tf.tables_initializer())
          # return average of ELMo features
```

```
return sess.run(tf.reduce_mean(embeddings,1))
```

#### 1.4.4 Lemmatize copies of the datasets for use with ELMo experiements

These copies will be used as part of experiments to train the ELMo classifer model on different preprocessing techniques

```
[111]: # Copy datasets for elmo
X_train_elmo = X_train.copy()
X_val_elmo = X_val.copy()
X_test_elmo = X_test.copy()

# Lemmatize the datasets
X_train_elmo = pd.Series(lemmatization(X_train_elmo))
X_val_elmo = pd.Series(lemmatization(X_val_elmo))
X_test_elmo = pd.Series(lemmatization(X_test_elmo))
```

```
[18]: # peek at the lemmatized data
print(X_train.values[2])
print(X_train_elmo.values[2])
```

standing walking long periods time causing lot pain legs. feels like cramp becomes worse longer feet.

stand walk long period time cause lot pain  $\log$  . feel like cramp become worse long foot .

#### 1.5 Model Training and Tuning

- Run two experiments with different model architectures and data preparation:
  - Experiment 1: Train a Naive Bayes Classifier usin TF-IDF vectorized features
  - Experiment 2: Train a neural network classifier with an ELMo text embedding
- For each experiment:
  - Train the model on each experimental dataset
  - Validate against validation dataset
  - Tune hyperparameters as necessary to optimize performance
  - Repeat until optimized
  - Measure model performance

#### 1.5.1 Experiment 1: Train Naive Bayes Classifer on TF-IDF embedded data

Fit the Naive Bayes MultiNomial classifer using the TF-IDF transformed dataset.

Evaluate accuracy performance and performance across classes.

```
[97]: # Initialize Naive Bayes Classifier
clf1 = MultinomialNB()

# Train classifier
clf1.fit(X_train_tfidf, y_train)
```

#### [97]: MultinomialNB()

[98]: # get accuracy score
score\_val = clf1.score(X\_val\_tfidf, y\_val)
print("Accuracy score: {}".format(round(score\_val, 4)))

Accuracy score: 0.9167

[99]: # run predictions and display classification report to see performance across

classes

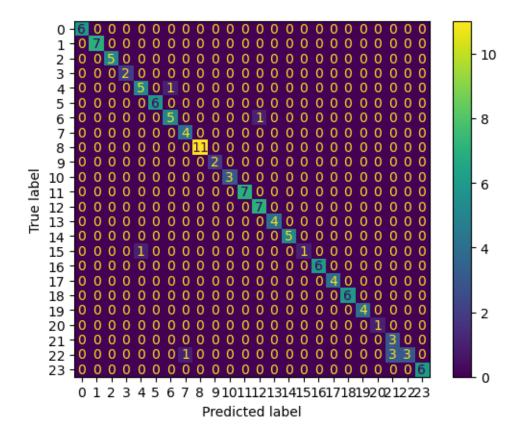
predictions = clf1.predict(X\_val\_tfidf)

print(classification\_report(y\_val, predictions))

	precision	recall	f1-score	support
Acne	1.00	1.00	1.00	6
Arthritis	1.00	1.00	1.00	7
Bronchial Asthma	1.00	1.00	1.00	5
Cervical spondylosis	1.00	1.00	1.00	2
Chicken pox	1.00	0.83	0.91	6
Common Cold	1.00	1.00	1.00	6
Dengue	0.67	0.67	0.67	6
Dimorphic Hemorrhoids	0.80	1.00	0.89	4
Fungal infection	1.00	1.00	1.00	11
Hypertension	1.00	1.00	1.00	2
Impetigo	1.00	1.00	1.00	3
Jaundice	1.00	1.00	1.00	7
Malaria	0.88	1.00	0.93	7
Migraine	1.00	1.00	1.00	4
Pneumonia	1.00	1.00	1.00	5
Psoriasis	1.00	0.50	0.67	2
Typhoid	0.86	1.00	0.92	6
Varicose Veins	1.00	1.00	1.00	4
allergy	1.00	0.83	0.91	6
diabetes	1.00	1.00	1.00	4
drug reaction	1.00	1.00	1.00	1
gastroesophageal reflux disease	0.38	1.00	0.55	3
peptic ulcer disease	1.00	0.29	0.44	7
urinary tract infection	1.00	1.00	1.00	6
accuracy			0.92	120
macro avg	0.94	0.92	0.91	120
weighted avg	0.95	0.92	0.91	120

[24]: # plot the confusion matrix
ConfusionMatrixDisplay(confusion\_matrix=confusion\_matrix(y\_val, predictions)).

plot();



Confusion Matrix Classes: [0:Acne], [1:Arthritis], [2:Bronchial Asthma], [3:Cervical spondylosis], [4:Chicken pox], [5:Common Cold], [6:Dengue], [7:Dimorphic Hemorrhoids], [8:Fungal infection], [9:Hypertension], [10:Impetigo], [11:Jaundice], [12:Malaria], [13:Migraine], [14:Pneumonia], [15:Psoriasis], [16:Typhoid], [17:Varicose Veins], [18:allergy], [19:diabetes], [20:drug reaction], [21:gastroesophageal reflux disease], [22:peptic ulcer disease]

#### 1.5.2 Experiment 2: Train Neural Network Classifier with ELMo Embeddings

Build a DNN with the following architecture:

- Input Layer with raw text input
- ELMo embedding layer with 1024 vector output
- Hidden layer with 256 units and ReLu activation
- Output layer with 24 units and Softwmax activation

**Define Model Architecture and Train** Define a model with a text input layer feeding into ELMo embedding layer, one additional hidden layer and an output layer predicting the 24 classes (diagnoses). Once trained, weights are saved to a file so training is not necessary every run.

Note: this step is only necessary for model re-training

```
[31]: # Build Model
      def build_nn_model():
          input_text = Input(shape=(1,), dtype=tf.string, name="Text_Input")
          embedding = Lambda(elmo_embedding, output_shape=(1024, ),__

¬name="ELMo_Embedding")(input_text)

          dense = Dense(256, activation='relu', name="Dense_Hidden")(embedding)
          pred = Dense(24, activation='softmax', name="Dense_Predict")(dense)
          model = Model(inputs=[input_text], outputs=pred)
          model.compile(loss='categorical_crossentropy', optimizer='adam', __
       →metrics=['accuracy'])
          return model
[113]: # Train the model and save to weights file
      model = build_nn_model()
      with tf.Session() as session:
          K.set session(session)
          session.run(tf.global_variables_initializer())
          session.run(tf.tables_initializer())
          history = model.fit(X_train_elmo, y_train_enc, epochs=20, batch_size=16)
          model.save_weights('./weights/symptoms-elmo-model-no-proc.h5')
     INFO:tensorflow:Saver not created because there are no variables in the graph to
     restore
     INFO:tensorflow:Saver not created because there are no variables in the graph to
     Train on 960 samples
     Epoch 1/20
     960/960 [=========== ] - 199s 207ms/sample - loss: 2.4544 -
     accuracy: 0.3562
     Epoch 2/20
     960/960 [============ ] - 199s 208ms/sample - loss: 1.2369 -
     accuracy: 0.7333
     Epoch 3/20
     960/960 [============ ] - 173s 180ms/sample - loss: 0.7196 -
     accuracy: 0.8448
     Epoch 4/20
     960/960 [=========== ] - 175s 182ms/sample - loss: 0.4789 -
     accuracy: 0.9083
     Epoch 5/20
     960/960 [============= ] - 161s 168ms/sample - loss: 0.3279 -
     accuracy: 0.9500
     Epoch 6/20
     960/960 [========== ] - 159s 166ms/sample - loss: 0.2481 -
     accuracy: 0.9698
     Epoch 7/20
     960/960 [============ ] - 172s 179ms/sample - loss: 0.1836 -
     accuracy: 0.9812
```

```
960/960 [=========== ] - 186s 194ms/sample - loss: 0.1435 -
    accuracy: 0.9844
    Epoch 9/20
    960/960 [=========== ] - 181s 188ms/sample - loss: 0.1127 -
    accuracy: 0.9958
    Epoch 10/20
    960/960 [============= ] - 186s 194ms/sample - loss: 0.0903 -
    accuracy: 0.9979
    Epoch 11/20
    960/960 [=========== ] - 202s 211ms/sample - loss: 0.0729 -
    accuracy: 0.9990
    Epoch 12/20
    960/960 [============ ] - 182s 190ms/sample - loss: 0.0606 -
    accuracy: 0.9979
    Epoch 13/20
    960/960 [============ ] - 187s 194ms/sample - loss: 0.0480 -
    accuracy: 1.0000
    Epoch 14/20
    960/960 [=========== ] - 182s 190ms/sample - loss: 0.0425 -
    accuracy: 0.9990
    Epoch 15/20
    960/960 [============= ] - 172s 180ms/sample - loss: 0.0341 -
    accuracy: 1.0000
    Epoch 16/20
    960/960 [=========== ] - 165s 172ms/sample - loss: 0.0303 -
    accuracy: 1.0000
    Epoch 17/20
    960/960 [============ ] - 185s 192ms/sample - loss: 0.0259 -
    accuracy: 1.0000
    Epoch 18/20
    960/960 [============ ] - 173s 181ms/sample - loss: 0.0219 -
    accuracy: 1.0000
    Epoch 19/20
    960/960 [=========== ] - 166s 173ms/sample - loss: 0.0197 -
    accuracy: 1.0000
    Epoch 20/20
    960/960 [============= ] - 166s 173ms/sample - loss: 0.0177 -
    accuracy: 1.0000
[32]: # graphically display model architecture
     model1 = build_nn_model()
     keras.utils.plot_model(
        model1,
        show_shapes=True,
        show_dtype=True,
        show_layer_names=True,
```

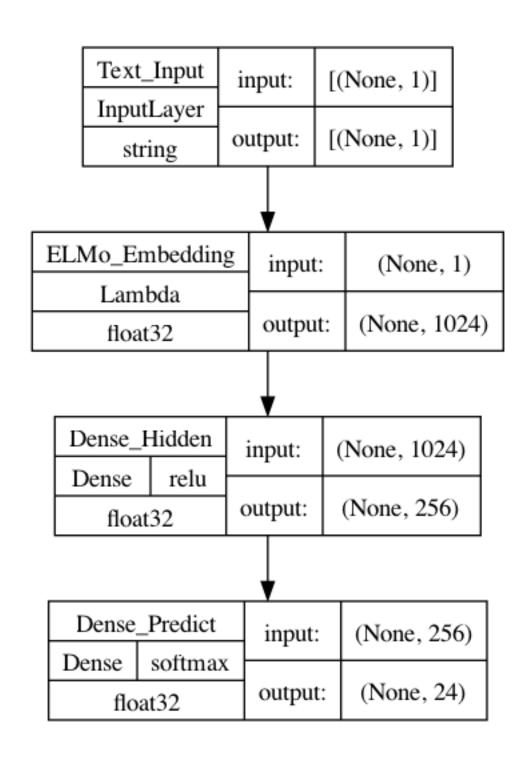
Epoch 8/20

```
rankdir="TB",
  expand_nested=True,
  dpi=96,
  layer_range=None,
  show_layer_activations=True
)
```

INFO:tensorflow:Saver not created because there are no variables in the graph to restore

INFO:tensorflow:Saver not created because there are no variables in the graph to restore

[32]:



```
[61]: # visualize training progression

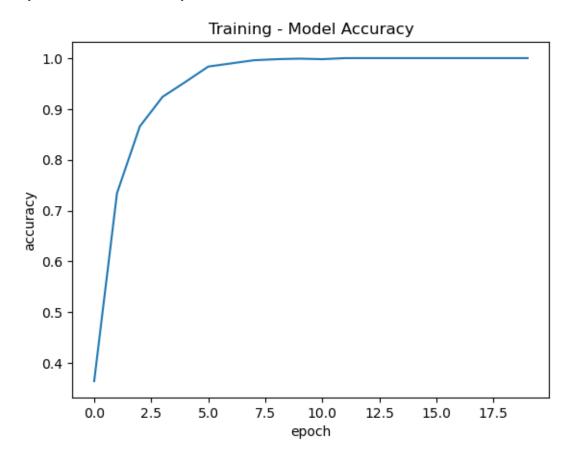
print(history.history.keys())

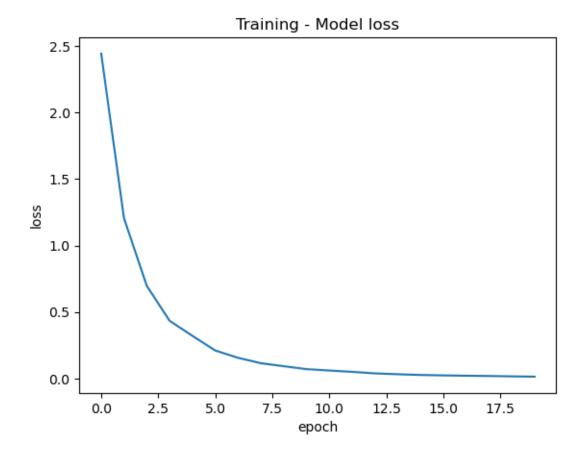
# summarize history for accuracy
plt.plot(history.history['accuracy'])
```

```
plt.title('Training - Model Accuracy')
plt.ylabel('accuracy')
plt.xlabel('epoch')
plt.show()

# summarize history for loss
plt.plot(history.history['loss'])
plt.title('Training - Model loss')
plt.ylabel('loss')
plt.xlabel('epoch')
plt.show()
```

dict\_keys(['loss', 'accuracy'])





Helper fuctions for running experiment result analysis Re-usable functions to run predictions and report on the results for the ELMo classifier

```
[115]: def run_elmo_predictions(X):
    model = build_nn_model()
    with tf.Session() as session:
        K.set_session(session)
        session.run(tf.global_variables_initializer())
        session.run(tf.tables_initializer())
        model.load_weights('./weights/symptoms-elmo-model-no-proc.h5')
        return model.predict(X, batch_size=16)

def run_elmo_report(y_in, predictions):
    # decode validation labels
    y_in_dec = decode(encoder, y_in)

# decode predicted labels
    y_preds_dec = decode(encoder, predictions)

print(classification_report(y_in_dec, y_preds_dec))
```

```
def run_confusion_matrix_elmo(y_in, predictions):
    ConfusionMatrixDisplay(confusion_matrix=confusion_matrix(y_in,
→predictions)).plot();
```

Assess model with validation data With the trained model, run against validation dataset and measure performance. Note, we have done two training runs - one with non-Lemmatized data and one with Lemmatized data. The weight files are as follows, load the approrpate one:

- Lemmatized Input Data: symptoms-elmo-model-lemm.h5
- Non-Lemmatized Input Data: symptoms-elmo-model.h5

```
[116]: # Run model on validation dataset
val_predictions = run_elmo_predictions(X_val_elmo)
```

INFO:tensorflow:Saver not created because there are no variables in the graph to restore

INFO:tensorflow:Saver not created because there are no variables in the graph to restore

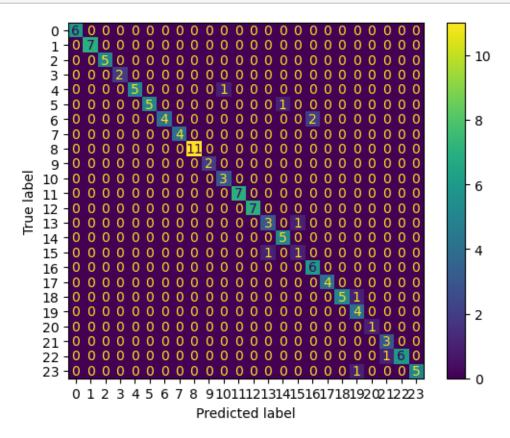
/Users/tylerforeman/anaconda3/envs/base-ml-39/lib/python3.9/site-packages/keras/engine/training\_v1.py:2356: UserWarning: `Model.state\_updates` will be removed in a future version. This property should not be used in TensorFlow 2.0, as `updates` are applied automatically. updates=self.state\_updates,

# [117]: # Display classification report run\_elmo\_report(y\_val\_enc, val\_predictions)

	precision	recall	f1-score	support
Acne	1.00	1.00	1.00	6
Arthritis	1.00	1.00	1.00	7
Bronchial Asthma	0.83	1.00	0.91	5
Cervical spondylosis	1.00	1.00	1.00	2
Chicken pox	0.80	0.67	0.73	6
Common Cold	1.00	1.00	1.00	6
Dengue	0.75	0.50	0.60	6
Dimorphic Hemorrhoids	1.00	1.00	1.00	4
Fungal infection	1.00	1.00	1.00	11
Hypertension	1.00	1.00	1.00	2
Impetigo	0.75	1.00	0.86	3
Jaundice	1.00	1.00	1.00	7
Malaria	1.00	1.00	1.00	7
Migraine	1.00	1.00	1.00	4
Pneumonia	1.00	0.80	0.89	5
Psoriasis	1.00	0.50	0.67	2
Typhoid	0.75	1.00	0.86	6
Varicose Veins	1.00	1.00	1.00	4

alle	ergy	1.00	0.83	0.91	6
diabe	etes	0.80	1.00	0.89	4
drug react	tion	0.50	1.00	0.67	1
gastroesophageal reflux dise	ease	0.50	1.00	0.67	3
peptic ulcer dise	ease	0.75	0.43	0.55	7
urinary tract infect	tion	1.00	1.00	1.00	6
accur	racy			0.90	120
macro	avg	0.89	0.91	0.88	120
weighted	avg	0.91	0.90	0.90	120

[31]: # plot the confusion matrix run\_confusion\_matrix\_elmo(y\_val, decode(encoder, val\_predictions))



Confusion Matrix Classes: [0:Acne], [1:Arthritis], [2:Bronchial Asthma], [3:Cervical spondylosis], [4:Chicken pox], [5:Common Cold], [6:Dengue], [7:Dimorphic Hemorrhoids], [8:Fungal infection], [9:Hypertension], [10:Impetigo], [11:Jaundice], [12:Malaria], [13:Migraine], [14:Pneumonia], [15:Psoriasis], [16:Typhoid], [17:Varicose Veins], [18:allergy], [19:diabetes], [20:drug reaction], [21:gastroesophageal reflux disease], [22:peptic ulcer disease]

# 1.6 Model Comparison and Selection

- Test both models against test dataset
- Compare model performance

Assess NBC model with test data With the trained model, run against test dataset and measure performance.

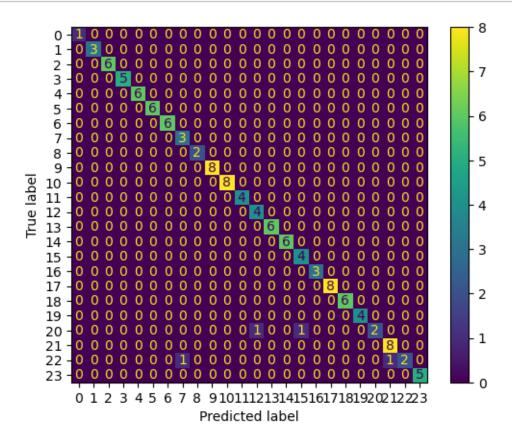
```
[32]: # get score
score_test = clf1.score(X_test_tfidf, y_test)
print("[Test] Accuracy score: {}".format(round(score_test, 4)))
```

[Test] Accuracy score: 0.9667

[33]: # run predictions and display report
 nbc\_test\_predictions = clf1.predict(X\_test\_tfidf)
 print(classification\_report(y\_test, nbc\_test\_predictions))

	precision	recall	f1-score	support
Acne	1.00	1.00	1.00	1
Arthritis	1.00	1.00	1.00	3
Bronchial Asthma	1.00	1.00	1.00	6
Cervical spondylosis	1.00	1.00	1.00	5
Chicken pox	1.00	1.00	1.00	6
Common Cold	1.00	1.00	1.00	6
Dengue	1.00	1.00	1.00	6
Dimorphic Hemorrhoids	0.75	1.00	0.86	3
Fungal infection	1.00	1.00	1.00	2
Hypertension	1.00	1.00	1.00	8
Impetigo	1.00	1.00	1.00	8
Jaundice	1.00	1.00	1.00	4
Malaria	0.80	1.00	0.89	4
Migraine	1.00	1.00	1.00	6
Pneumonia	1.00	1.00	1.00	6
Psoriasis	0.80	1.00	0.89	4
Typhoid	1.00	1.00	1.00	3
Varicose Veins	1.00	1.00	1.00	8
allergy	1.00	1.00	1.00	6
diabetes	1.00	1.00	1.00	4
drug reaction	1.00	0.50	0.67	4
gastroesophageal reflux disease	0.89	1.00	0.94	8
peptic ulcer disease	1.00	0.50	0.67	4
urinary tract infection	1.00	1.00	1.00	5
accuracy			0.97	120
macro avg	0.97	0.96	0.95	120
weighted avg	0.97	0.97	0.96	120

```
[34]: # plot the confusion matrix
ConfusionMatrixDisplay(confusion_matrix=confusion_matrix(y_test,__
_nbc_test_predictions)).plot();
```



Assess ELMo model with test data With the trained model, run against test dataset and measure performance. Note, we have done two training runs - one with non-Lemmatized data and one with Lemmatized data. The weight files are as follows, load the appropriate one:

```
[35]: # Run model on validation dataset test_predictions = run_elmo_predictions(X_test_elmo)
```

INFO:tensorflow:Saver not created because there are no variables in the graph to restore

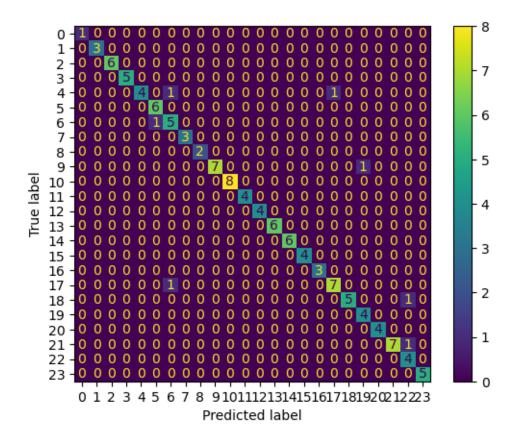
INFO:tensorflow:Saver not created because there are no variables in the graph to restore

/Users/tylerforeman/anaconda3/envs/base-ml-39/lib/python3.9/site-packages/keras/engine/training\_v1.py:2356: UserWarning: `Model.state\_updates` will be removed in a future version. This property should not be used in TensorFlow 2.0, as `updates` are applied automatically. updates=self.state\_updates,

[36]: # Display classification report run\_elmo\_report(y\_test\_enc, test\_predictions)

	precision	recall	f1-score	support
Acne	1.00	1.00	1.00	1
Arthritis	1.00	1.00	1.00	3
Bronchial Asthma	1.00	1.00	1.00	6
Cervical spondylosis	1.00	1.00	1.00	5
Chicken pox	1.00	0.67	0.80	6
Common Cold	0.86	1.00	0.92	6
Dengue	0.71	0.83	0.77	6
Dimorphic Hemorrhoids	1.00	1.00	1.00	3
Fungal infection	1.00	1.00	1.00	2
Hypertension	1.00	0.88	0.93	8
Impetigo	1.00	1.00	1.00	8
Jaundice	1.00	1.00	1.00	4
Malaria	1.00	1.00	1.00	4
Migraine	1.00	1.00	1.00	6
Pneumonia	1.00	1.00	1.00	6
Psoriasis	1.00	1.00	1.00	4
Typhoid	1.00	1.00	1.00	3
Varicose Veins	0.88	0.88	0.88	8
allergy	1.00	0.83	0.91	6
diabetes	0.80	1.00	0.89	4
drug reaction	1.00	1.00	1.00	4
gastroesophageal reflux disease	1.00	0.88	0.93	8
peptic ulcer disease	0.67	1.00	0.80	4
urinary tract infection	1.00	1.00	1.00	5
accuracy			0.94	120
macro avg	0.95	0.96	0.95	120
weighted avg	0.95	0.94	0.94	120

[37]: # plot the confusion matrix run\_confusion\_matrix\_elmo(y\_test, decode(encoder, test\_predictions))



#### 1.7 Create inference pipeline with NBC

- Data prep pipeline
- Trained model pipeline

[131]: # train the pipeline
nbc\_model = pipeline.fit(X\_train, y\_train)

[132]: # test performance on test data to ensure it is the same
 nbc\_pipeline\_predictions = nbc\_model.predict(X\_test)
 print(classification\_report(y\_test, nbc\_pipeline\_predictions))

	precision	recall	f1-score	support
Acne	1.00	1.00	1.00	1
Arthritis	1.00	1.00	1.00	3
Bronchial Asthma	1.00	1.00	1.00	6
Cervical spondylosis	1.00	1.00	1.00	5
Chicken pox	1.00	1.00	1.00	6
Common Cold	1.00	1.00	1.00	6
Dengue	1.00	1.00	1.00	6
Dimorphic Hemorrhoids	0.75	1.00	0.86	3
Fungal infection	1.00	1.00	1.00	2
Hypertension	1.00	1.00	1.00	8
Impetigo	1.00	1.00	1.00	8
Jaundice	1.00	1.00	1.00	4
Malaria	0.80	1.00	0.89	4
Migraine	1.00	1.00	1.00	6
Pneumonia	1.00	1.00	1.00	6
Psoriasis	0.80	1.00	0.89	4
Typhoid	1.00	1.00	1.00	3
Varicose Veins	1.00	1.00	1.00	8
allergy	1.00	1.00	1.00	6
diabetes	1.00	1.00	1.00	4
drug reaction	1.00	0.50	0.67	4
gastroesophageal reflux disease	0.89	1.00	0.94	8
peptic ulcer disease	1.00	0.50	0.67	4
urinary tract infection	1.00	1.00	1.00	5
accuracy			0.97	120
macro avg	0.97	0.96	0.95	120
weighted avg	0.97	0.97	0.96	120

Run some interactive tests with the NBC pipeline Test some predictions made by the pipeline with ad-hoc input text

```
[135]: # Run some random symptom descriptions and get diagnosis
run_diagnosis("I have red itchy spots all over my skin. I have a fever and a

→headache.")
run_diagnosis("My nose is really stuffy. I am sneezing alot. My throat is

→sore and I am starting to cough.")
run_diagnosis("My legs are sore and bruised. I can't stand for too long.")
```

Based on your sypmtom descriptions, you have Chicken pox Based on your sypmtom descriptions, you have Common Cold Based on your sypmtom descriptions, you have Varicose Veins

Export the model pipeline for use in API

```
[140]: import joblib
joblib.dump(nbc_model, './models/nbc_pipeline.pkl')
```

[140]: ['./models/nbc\_pipeline.pkl']

#### 1.8 Test NBC accuracy across different mixes to train/test/val

Since class samples were small in each train/test/val split, test overall accuracy with different random mixes and get average accuracy score

```
[159]: # Run with 100 random samples of train/val/test splits
scores = []
for x in range(0, 100):

# Split dataset into train, validate, test
X_train, X_test, y_train, y_test = train_test_split(X, np.asarray(y_raw),u)
test_size=0.2)

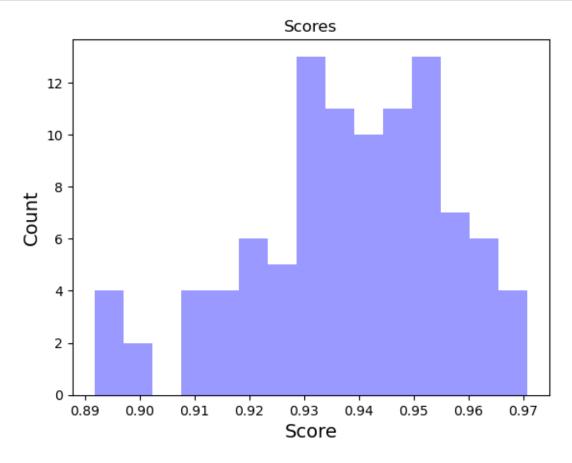
# train the pipeline
nbc_model = pipeline.fit(X_train, y_train)

# test performance on test data to ensure it is the same
score = pipeline.score(X_test, y_test)
scores.append(score)
```

```
[160]: #plot histogram
plt.hist(scores, alpha=0.4, color='blue', bins=15)
```

```
plt.ylabel('Count',fontsize=14)
plt.xlabel('Score',fontsize=14)
plt.xticks(fontsize=10)
plt.yticks(fontsize=10)

# Add Title
plt.title('Scores',fontsize=12);
plt.show()
```



```
[162]: # get mean accuracy score
print("Mean accuracy score of NBC classifier: {}".format(round(np.mean(scores),
$\u2\)))
```

Mean accuracy score of NBC classifier: 0.94