

#### Background

This tutorial covers the analysis of medical notes using Spacy, SciSpacy and ClinicalBERT.

The objective of the analysis is to generate extracted entities using Spacy and SciSpacy, word2vec, and tSNE plots.

Additionally this tutorial will use ClinicalBert to generate the same analysis to determine whether there is increased efficacy and information relative to the other two processes.

The first step is to download and extract the MIMIC-III files from https://physionet.org/. The NOTEEVENTS file is then uploaded to Google drive for loading into the Jupyter notebook while other files are stored locally and uploaded as needed.

#### Data Loading and Prep

These steps are designed to create a reusable file for analysis so as to minimize the processing load and increase analysis speed.

```
import pandas as pd
import numpy as np
#connect to google drive to read in NOTEEVENTS file
from google.colab import drive
drive.mount('/content/drive', force remount=True)
%cd drive/MyDrive
noteevents df = pd.read csv('NOTEEVENTS.csv', low memory=False) #.set index('ROW ID')
# keep discharge summary text and create subject admission id key for merging
dischargetext df = noteevents df.loc[noteevents df['CATEGORY'] == 'Discharge summary',
                                     ['SUBJECT ID', 'HADM ID', 'TEXT']]
dischargetext df['subj hadm'] = list(zip(dischargetext df['SUBJECT ID'].astype(int),
                                         dischargetext df['HADM ID'].astype(int)))
#upload the DIAGNOSES ICD in a dataframe
from google.colab import files
uploaded = files.upload()
diagnoses icd df = pd.read csv('DIAGNOSES ICD.csv').set index('ROW ID')
#keeping with my SQL assignment, pulling data for Wolf-Parkinson White (WPW) Syndrome
disease list = ['4267']
disease df = diagnoses icd df[diagnoses icd df['ICD9 CODE'].isin(disease list)].copy()
disease df['subj hadm'] = list(zip(disease df['SUBJECT ID'].astype(int),
                                   disease df['HADM ID'].astype(int)))
# Join discharge summary text WITH disease subset ON (subj id, hadm id)
patients df = dischargetext df[['TEXT', 'subj hadm']]\
    .join(disease df.set index('subj hadm')['ICD9 CODE'], on='subj hadm', how='left')\
    .dropna()\
    .drop(columns=['subj hadm', 'ICD9 CODE'])
# save the WPW subset so no need to regenerate; saving as a tsv file given the large number of commas in medical notes
patients df.to csv("wpw notes.tsv", sep = "\t")
```

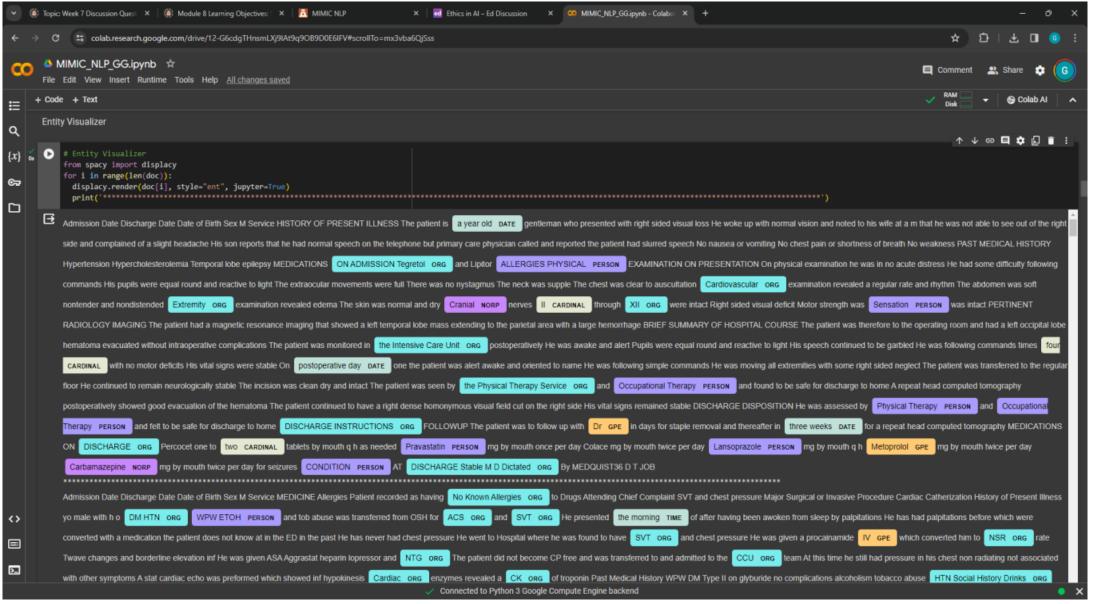
#### Spacy Tutorial – loading and building notes for processing

```
#install spacy
! pip install -U pip setuptools wheel
! pip install -U spacy
! python -m spacy download en core web sm
#load patient notes
import pandas as pd
import numpy as np
#connect to google drive
from google.colab import drive
drive.mount("/content/drive", force remount=True)
%cd drive/MyDrive
notes cardiac df = pd.read csv('wpw notes.tsv', low memory=False, sep='\t')
notes cardiac df.shape
#build notes
import spacy
nlp = spacy.load('en core web sm')
#clean up the notes by removing excess punctuation, characters, and numbers
temp = []
def clean up(text series):
    return (text series
            .str.replace('\[\*\*[^\]]*\*\*\]','')
            .str.replace('<[^>]*>', '')
            .str.replace('[\W]+', ' ')
            .str.replace(' \d+', ' '))
temp = clean up(notes cardiac df['TEXT'])
temp.shape
doc = []
for i in range(len(temp)):
 doc.append(nlp(temp[i]))
 print(doc[-1])
```

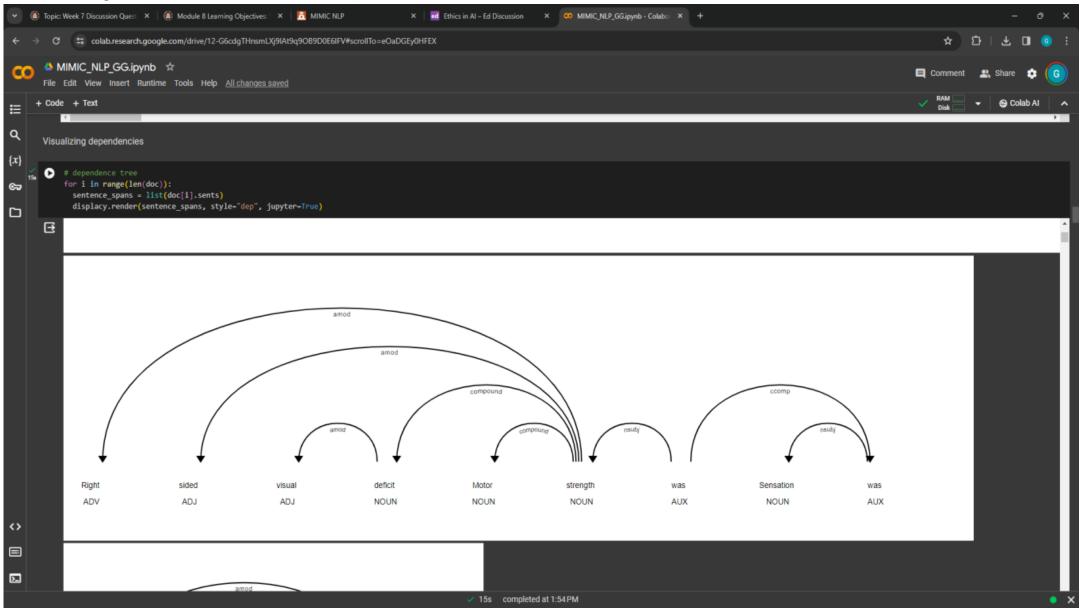
#### Spacy Tutorial – analyzing notes and visualization

```
#Get tokens without punctuations or white space for all patient notes
token without punct = []
for i in range(len(doc)):
 token without punct.append([token.orth for token in nlp(doc[i]) if not token.is punct or token.is space])
 print(token without punct[-1])
 for i in range(len(doc)):
 for token in doc[i]:
   print(token.text, token.pos )
 #Name Entity Recognition
entity doc = []
for i in range(len(doc)):
 entity doc.append(nlp(doc[i]))
 for ent in entity doc[-1].ents:
   print(ent.text, ent.start char, ent.end char, ent.label )
 # Entity Visualizer
from spacy import displacy
for i in range(len(doc)):
 displacy.render(doc[i], style="ent", jupyter=True)
#sentence identifier
for i in range(len(doc)):
 for ix, sent in enumerate(doc[i].sents, 1):
   print("Sentence number {}:{}".format(ix, sent))
# Visualizing dependencies: dependence tree
for i in range(len(doc)):
 sentence spans = list(doc[i].sents)
 displacy.render(sentence spans, style="dep", jupyter=True)
```

# Spacy Tutorial – Entity Visualizer snapshot



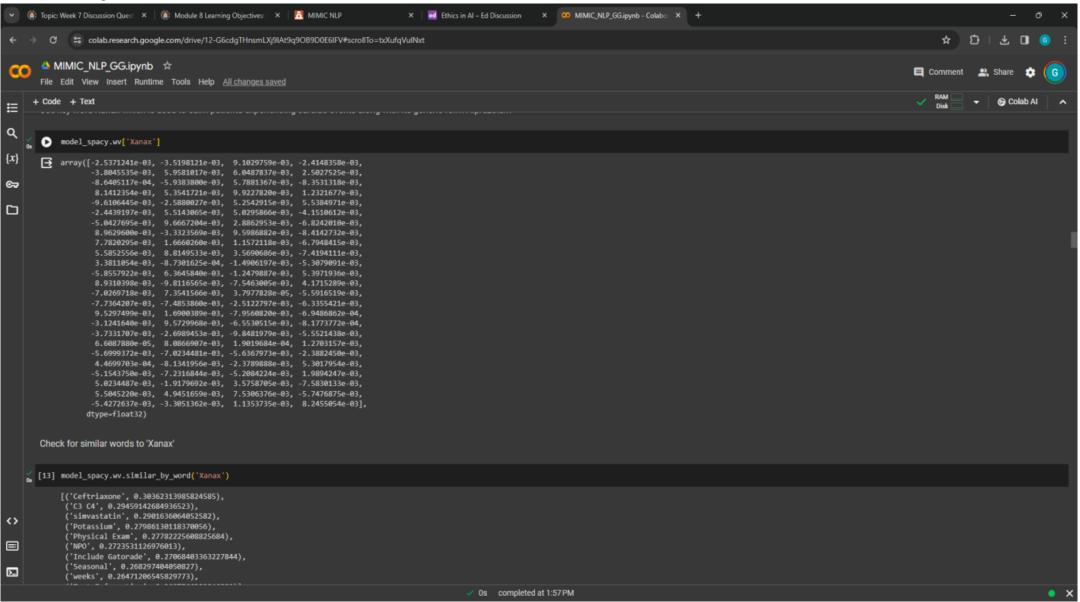
### Spacy Tutorial – Dependence Tree snapshot



#### Spacy Tutorial – Word2Vec

```
# Build corpus of all the entities extracted from the notes using spaCy model.
# The corpus is an array of arrays or list of lists where each of the nested lists corresponds to a note.
corpus=[]
for row in range(0, len(doc)):
 str tokens=[]
 tokens= nlp(doc[row]).ents
 for i in range(0, len(tokens)):
    str tokens.append(tokens[i].text)
  corpus.append(list(str tokens))
print(corpus)
#Get Word2Vec
import pandas as pd
pd.options.mode.chained assignment = None
import numpy as np
import re
import gensim
from gensim.models import Word2Vec
from sklearn.manifold import TSNE
import matplotlib.pyplot as plt
%matplotlib inline
model spacy = Word2Vec(corpus, min count=1)
#Check model vocabulary
model spacy.wv.key to index
#Use key word Xanax which is used to calm patients experiencing cardiac events along with its generic form Alprazolam
model spacy.wv['Xanax']
#Check for similar words to 'Xanax'
model spacy.wv.similar by word('Xanax')
#The output shows a number of words that have no similarity to "Xanax", likely due to its high degree of specificity in drug name.
```

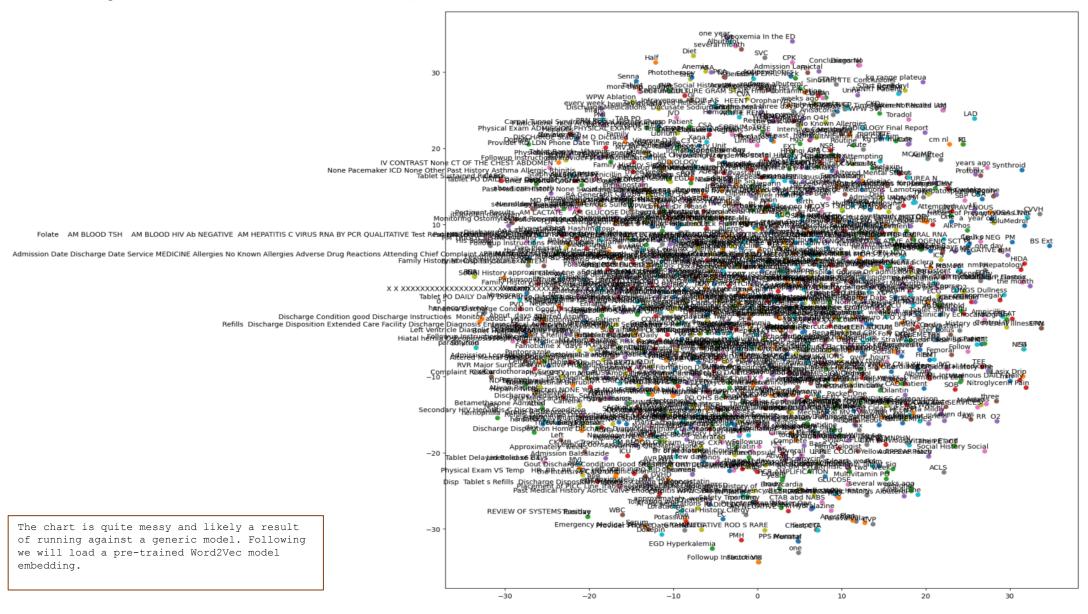
# Spacy Tutorial – Word2Vec snapshot



#### Spacy Tutorial – tSNE function and plotting

```
#Define the tSNE plot
def tsne plot(model, words, preTrained=False):
    "Creates and TSNE model and plots it"
    labels = []
    tokens = []
    for word in words:
      if preTrained:
          tokens.append(model[word])
          tokens.append(model.wv[word])
      labels.append(word)
    tokens = np.array(tokens)
    tsne model = TSNE(perplexity=30, early exaggeration=12, n components=2, init='pca', n iter=1000, random state=23)
    new values = tsne model.fit transform(tokens)
    x = []
    y = []
    for value in new values:
        x.append(value[0])
        y.append(value[1])
    plt.figure(figsize=(16, 16))
    for i in range(len(x)):
        plt.scatter(x[i],y[i])
       plt.annotate(labels[i],
                     xy=(x[i], y[i]),
                     xytext=(5, 2),
                     textcoords='offset points',
                     ha='right',
                     va='bottom')
    plt.show()
#Plot the corpus in a tSNE plot
vocabs = model spacy.wv.key to index.keys()
new v = np.array(list(vocabs))
tsne plot(model spacy, new v)
```

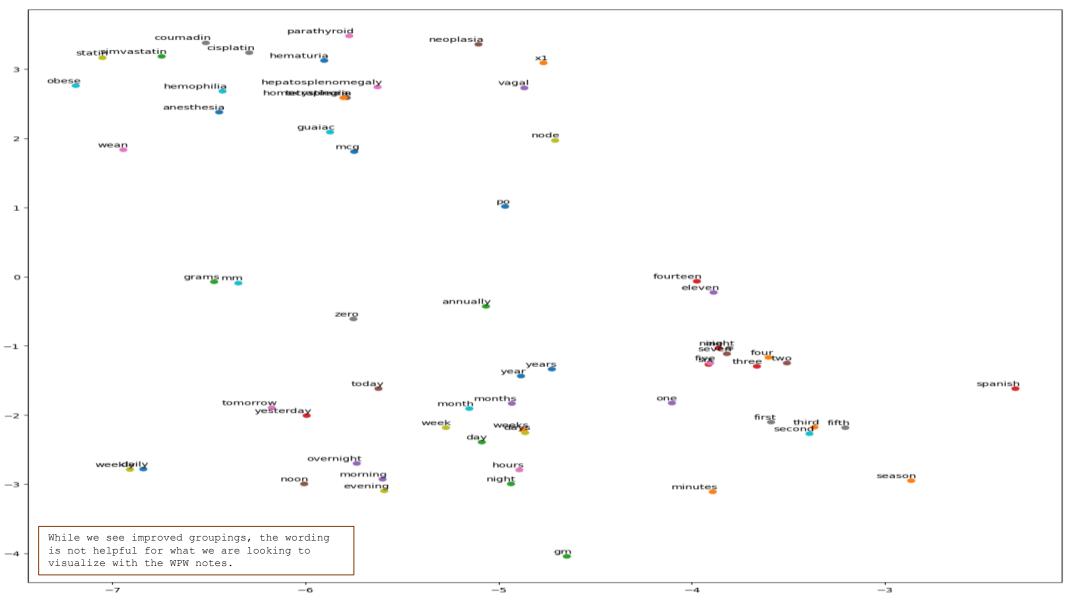
#### Spacy Tutorial – tSNE plot



### Spacy Tutorial – loading pre-trained model

```
# load pre-trained word2vec embeddings
import gensim
import gensim.downloader as api
info = api.info() # show info about available models/datasets
pretrained model= api.load("glove-wiki-gigaword-50") # download the model and return as object ready for use
#Check for words in the pre-trained model that are similar to 'cardiac' given WPW is a heart-related medical condition
pretrained model.most similar("cardiac")
#Generate a new corpus based on the pre-trained model
corpus in pretrained model = []
for word in vocabs:
 if word in pretrained model:
    corpus in pretrained model.append(word)
    print(word) #
#Generate a new tSNE plot using the pre-trained model
tsne plot(pretrained model, corpus in pretrained model, True)
```

## Spacy Tutorial – tSNE plot on pre-trained model



# SciSpacy Tutorial – loading and building notes for processing

```
#install SciSpacy
!pip install -U spacy
!pip install scispacy
#Load SciSpacy pre-trained models: using the spaCy NER model trained on BC5CDR corpus given it has the highest efficacy of the imported models
!pip install https://s3-us-west-2.amazonaws.com/ai2-s2-scispacy/releases/v0.4.0/en_ner_bc5cdr_md-0.4.0.tar.gz
import scispacy
import spacy
import en ner bc5cdr md
nlp = en ner bc5cdr md.load()
#Import condensed NOTEEVENTS file from the Spacy tutorial
# load condensed NOTEEVENTS file
import pandas as pd
import numpy as np
#connect to google drive
from google.colab import drive
drive.mount("/content/drive", force remount=True)
%cd drive/MyDrive
notes cardiac df = pd.read csv('wpw notes.tsv', low memory=False, sep='\t')#.set index('ROW ID')
```

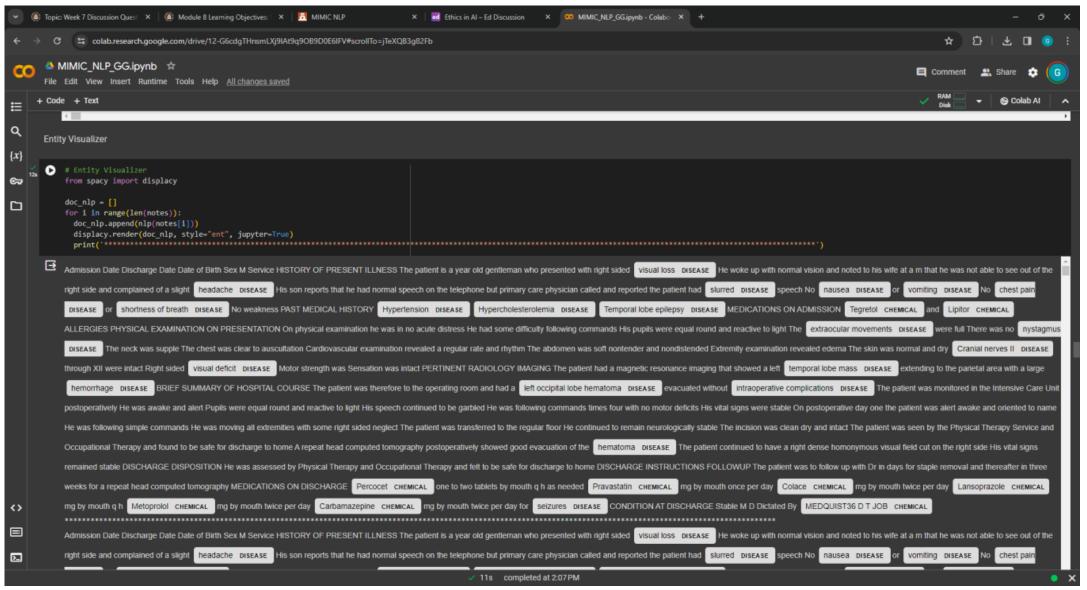
# SciSpacy Tutorial – loading and building notes for processing

```
#clean up the notes by removing excess punctuation, characters, and numbers
temp = []
def clean up(text series):
   return (text series
          .str.replace('\[\*\*[^\]]*\*\*\]','')
          .str.replace('<[^>]*>', '')
          .str.replace('[\W]+', ' ')
          .str.replace(' \d+', ' '))
temp = clean up(notes cardiac df['TEXT'])
temp.shape
doc scispacy = []
for i in range(len(temp)):
 doc scispacy.append(nlp(temp[i]))
 print(doc scispacy[-1])
 #save down cleaned-up version of the WPW notes
text file = open("clean wpw notes.txt", "w")
with open("clean wpw notes.txt", "w") as text file:
   for i in range(len(temp)):
     doc scispacy.append(nlp(temp[i]))
     print(doc scispacy[-1], file=text file)
text file.close
#Load cleaned-up WPW notes file: clean wpw notes.txt
from google.colab import files
uploaded = files.upload()
with open('clean wpw notes.txt', 'r') as fin:
 lines = fin.readlines()
 for line in lines:
   notes.append(line)
print(notes)
print(len(notes))
```

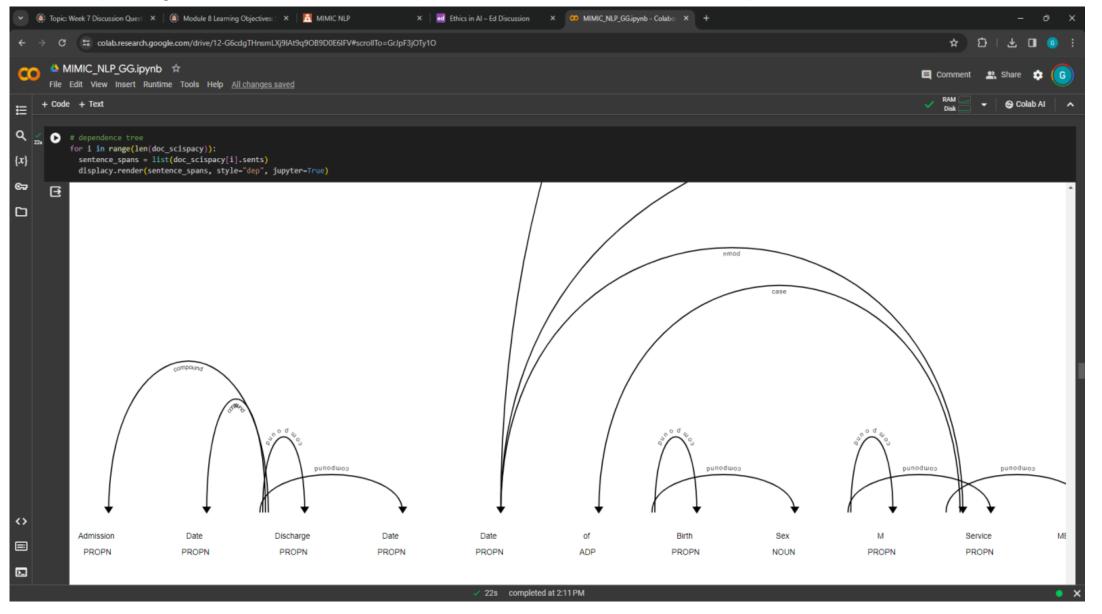
#### SciSpacy Tutorial – analyzing notes and visualization

```
#view notes
for i in range(len(notes)):
 print(notes[i])
#Get tokens for all patient notes
# Token
token without punct = []
for i in range(len(notes)):
 token without punct.append([token.orth for token in nlp(notes[i]) if not token.is punct or token.is space])
 print(token without punct[-1])
 # Entity Visualizer
from spacy import displacy
doc nlp = []
for i in range(len(notes)):
 doc nlp.append(nlp(notes[i]))
 displacy.render(doc nlp, style="ent", jupyter=True)
#Visualizing dependencies
# dependence tree
for i in range(len(doc scispacy)):
 sentence spans = list(doc scispacy[i].sents)
 displacy.render(sentence spans, style="dep", jupyter=True)
```

## SciSpacy Tutorial – Entity Visualizer snapshot



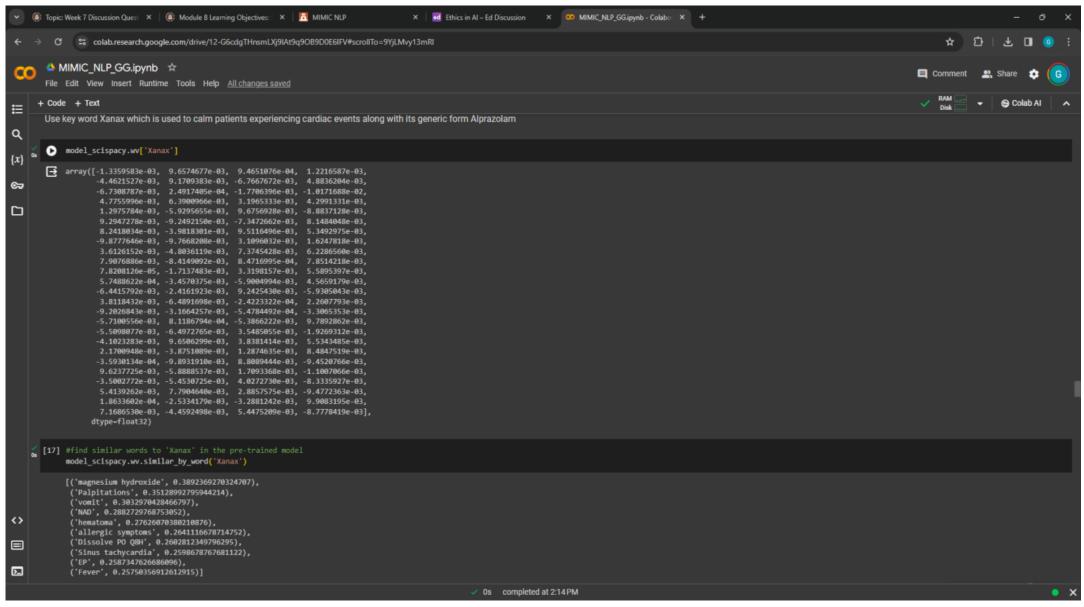
## SciSpacy Tutorial – Dependence Tree snapshot



#### SciSpacy Tutorial – loading notes and model, Word2Vec

```
#Build corpus of notes
# Build corpus of all the entities extracted from the notes using spaCy model.
# The corpus is an array of arrays or list of lists where each of the nested lists corresponds to a note.
corpus=[]
for row in range(0, len(notes)):
 str tokens=[]
  tokens= nlp(notes[row]).ents
 for i in range(0, len(tokens)):
   str tokens.append(tokens[i].text)
  corpus.append(list(str tokens))
print(corpus)
#import Word2Vec model
import gensim
import pandas as pd
pd.options.mode.chained assignment = None
import numpy as np
import re
from gensim.models import Word2Vec
from sklearn.manifold import TSNE
import matplotlib.pyplot as plt
%matplotlib inline
#Using SciSpacy and BC5CDR pre-trained model
import en ner bc5cdr md
nlp = en ner bc5cdr md.load()
#Define model
model scispacy = Word2Vec(corpus, min count=1)
#Use key word Xanax which is used to calm patients experiencing cardiac events along with its generic form Alprazolam
model scispacy.wv['Xanax']
#find similar words to 'Xanax in the pre-trained model
model scispacy.wv.similar by word('Xanax')
```

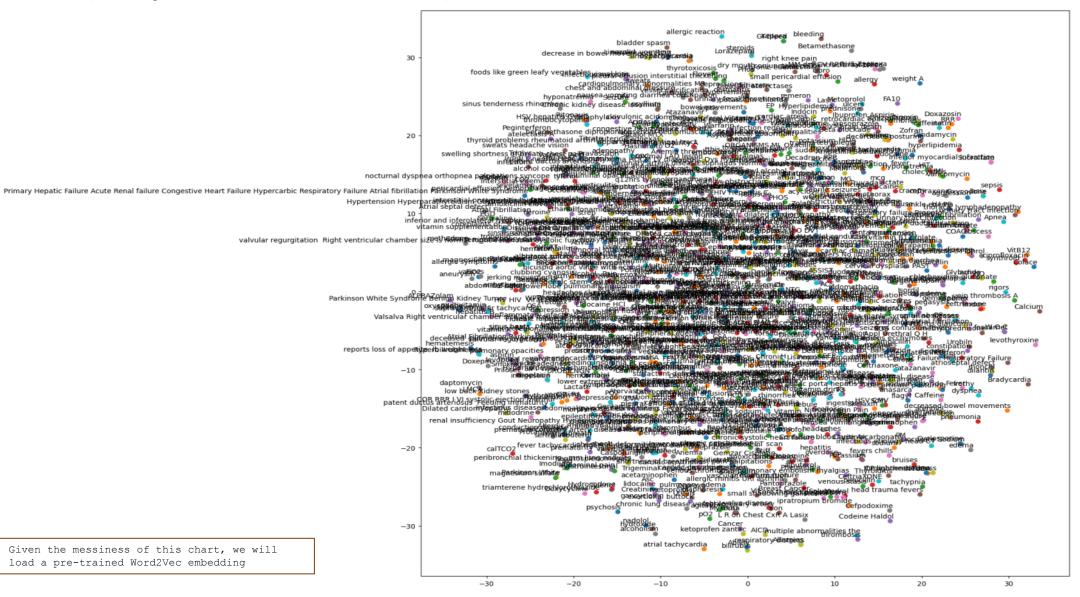
## SciSpacy Tutorial – Word2Vec snapshot



#### SciSpacy Tutorial – tSNE function and plotting

```
#Define the plot function
def tsne plot(model, words, preTrained=False):
    "Creates and TSNE model and plots it"
    labels = []
    tokens = []
    for word in words:
      if preTrained:
          tokens.append(model[word])
          tokens.append(model.wv[word])
      labels.append(word)
    tokens = np.array(tokens)
    tsne model = TSNE(perplexity=30, early exaggeration=12, n components=2, init='pca', n iter=1000, random state=23)
    new values = tsne model.fit transform(tokens)
    x = []
    y = []
    for value in new values:
        x.append(value[0])
       y.append(value[1])
    plt.figure(figsize=(16, 16))
    for i in range(len(x)):
        plt.scatter(x[i],y[i])
       plt.annotate(labels[i],
                     xy=(x[i], y[i]),
                     xytext=(5, 2),
                     textcoords='offset points',
                     ha='right',
                     va='bottom')
    plt.show()
#Plot the corpus in a TSNE plot
vocabs = model scispacy.wv.index to key # Access vocabulary using index to key
new v = np.array(list(vocabs))
tsne plot(model scispacy, new_v)
```

### SciSpacy Tutorial –tSNE plot



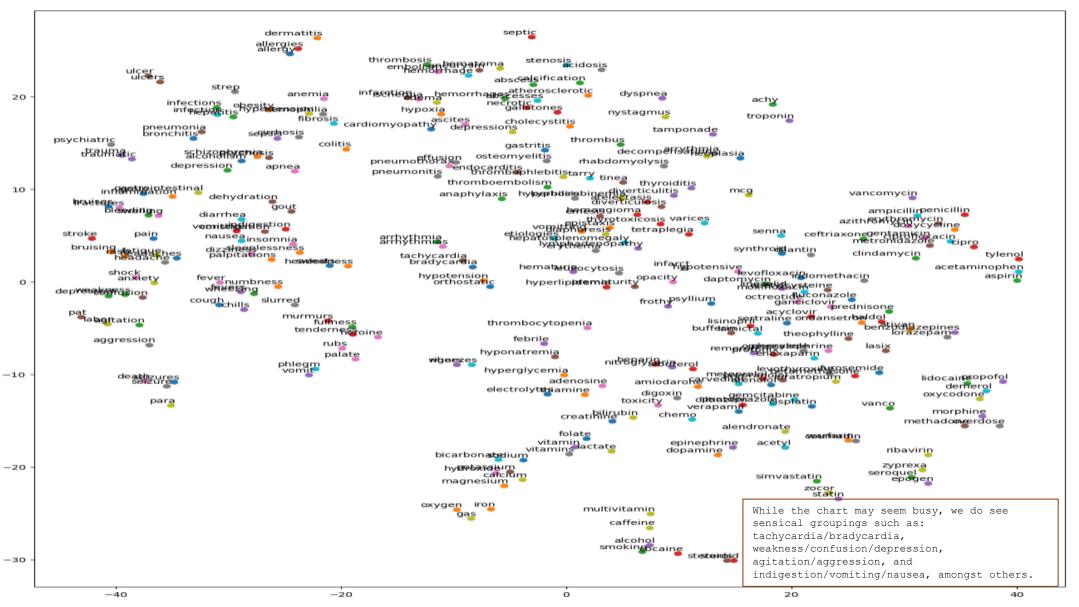
#### SciSpacy Tutorial – Loading pre-trained model

```
#Given the messiness of this chart, we will load a pre-trained Word2Vec embedding
# load pre-trained word2vec embeddings
import gensim
import gensim.downloader as api
info = api.info() # show info about available models/datasets
pretrained model= api.load("qlove-wiki-qiqaword-50") # download the model and return as object ready for use
#Since we are viewing patient notes regarding those diagnosed with WPW, we will search for words similar to tachycardia
pretrained model.most similar("tachycardia")
#Create a new corpus from the pre-trained model
new corpus in pretrained model = []
for word in new v:
   if word in pretrained model.key to index:
        new corpus in pretrained model.append(word)
        print(word) # Print out-of-vocabulary words
```

#### SciSpacy Tutorial – tSNE function and plotting

```
#Define the tSNE plot
import numpy as np
def tsne plot(model, words):
    "Creates a t-SNE model and plots it"
   labels = []
    tokens = []
    for word in words:
        if word in model:
            tokens.append(model[word])
            labels.append(word)
        else:
            print(f"Skipping '{word}' as it is not present in the model's vocabulary.")
    tsne model = TSNE(perplexity=11, early exaggeration=12, n components=2, init='pca', n iter=1000, random state=23)
    new values = tsne model.fit transform(np.array(tokens)) # Convert tokens to a NumPy array
    x = []
    y = []
    for value in new values:
        x.append(value[0])
        y.append(value[1])
    plt.figure(figsize=(16, 16))
    for i in range(len(x)):
        plt.scatter(x[i], y[i])
        plt.annotate(labels[i],
                     xy=(x[i], y[i]),
                     xytext=(5, 2),
                     textcoords='offset points',
                     ha='right',
                     va='bottom')
    plt.show()
#Generate the tSNE plot from the pre-trained model
tsne plot(pretrained model, new corpus in pretrained model)
```

### SciSpacy Tutorial -tSNE plot on pre-trained model



# ClinicalBERT Tutorial – loading the model, defining vocabulary

```
!pip install transformers
from transformers import AutoTokenizer, AutoModel, BertTokenizer, BertModel
import torch
bert model = BertModel.from pretrained('bert-base-uncased')
print(' Bert model is type:', type(bert model))
clinical model = AutoModel.from pretrained("emilyalsentzer/Bio ClinicalBERT")
print(' clinical model is type:', type(clinical model))
#ClinicalBERT vocabulary
bert tokenizer = BertTokenizer.from pretrained('bert-base-uncased')
clinical tokenizer = AutoTokenizer.from pretrained("emilyalsentzer/Bio ClinicalBERT")
blue tokenizer = AutoTokenizer.from pretrained('bionlp/bluebert pubmed mimic uncased L-12 H-768 A-12')
biobert tokenizer = AutoTokenizer.from pretrained('dmis-lab/biobert-base-cased-v1.2')
scibert tokenizer = AutoTokenizer.from pretrained('allenai/scibert scivocab uncased')
# blue tokenizer = AutoTokenizer.from pretrained('bionlp/bluebert pubmed mimic uncased L-24 H-1024 A-16')
print('clinical tokenizer is type:', type(clinical tokenizer))
```

# ClinicalBERT Tutorial – loading and analyzing tokens from notes

```
#using the notes from the Spacy and SciSpacy tutorials
#tokenization
for i in range(len(notes)):
  text = notes[i]
  bert tokens = bert tokenizer.tokenize(text)
  clinical tokens = clinical tokenizer.tokenize(text)
  bluebert tokens = blue tokenizer.tokenize(text)
  biobert tokens = biobert tokenizer.tokenize(text)
  # Pad out the clinical bert, bluebert list to be the same length.
  while len(clinical tokens) < len(bert tokens):</pre>
      clinical tokens.append("")
  while len(bluebert tokens) < len(bert tokens):</pre>
      bluebert tokens.append("")
  while len(biobert tokens) < len(bert tokens):</pre>
      biobert tokens.append("")
  # Label the columns.
  print('{:<12} {:<12} {:<12}'.format("BERT", "ClinicalBERT", "bluebert", "biobert"))</pre>
  print('{:<12} {:<12} {:<12}'.format("----", "------", "------", "------"))</pre>
  # Display the tokens.
  for tup in zip(bert tokens, clinical tokens, bluebert tokens, biobert tokens):
      print('{:<12} {:<12} {:<12}'.format(tup[0], tup[1], tup[2], tup[3]))</pre>
```

# ClinicalBERT Tutorial – loading and analyzing tokens from notes

```
#Comparing BERT to ClinicalBERT
# Use pandas just for table formatting.
import pandas as pd
# Some strange terms from the paper.
words = ['tachycardia',
         'cardiothoracic',
         'history of present illness'
# For each term...
for word in words:
    # Print it out
    print('\n\n', word, '\n')
    # Start a list of tokens for each model, with the first one being the model name.
    list a = ["BERT:"]
    list b = ["ClinicalBERT:"]
    # Run both tokenizers.
    list a.extend(bert tokenizer.tokenize(word))
    list b.extend( clinical tokenizer.tokenize(word))
    # Pad the lists to the same length.
    while len(list a) < len(list b):</pre>
        list a.append("")
    while len(list b) < len(list a):</pre>
        list b.append("")
    # Wrap them in a DataFrame to display a pretty table.
    df = pd.DataFrame([list a, list b])
    display(df)
#We see that in general, the two models have similar tokenizations on the selected words
```

# ClinicalBERT Tutorial – loading and analyzing tokens from notes

```
#The following is a dump of the vocabulary in ClinicalBERT and SciBERT
with open("vocabulary clinicalbert.txt", 'w') as f:
    # For each token in ClinicalBERT's vocabulary...
    for token in clinical tokenizer.vocab.keys():
        # Write it out, one per line.
        f.write(token + '\n')
with open("vocabulary scibert.txt", 'w') as f:
    # For each token in ClinicalBERT's vocabulary...
   for token in scibert tokenizer.vocab.keys():
        # Write it out, one per line.
        f.write(token + '\n')
```

```
import numpy as np
def get word indeces(tokenizer, text, word):
    Determines the index or indeces of the tokens corresponding to `word`
    within `text`. `word` can consist of multiple words, e.g., "cell biology".
    Determining the indeces is tricky because words can be broken into multiple
    tokens. I've solved this with a rather roundabout approach--I replace `word`
    with the correct number of `[MASK]` tokens, and then find these in the
    tokenized result.
    # Tokenize the 'word'--it may be broken into multiple tokens or subwords.
    word tokens = tokenizer.tokenize(word)
    # Create a sequence of `[MASK]` tokens to put in place of `word`.
   masks str = ' '.join(['[MASK]']*len(word tokens))
    # Replace the word with mask tokens.
    text masked = text.replace(word, masks str)
    print(text masked)
    # `encode` performs multiple functions:
    # 1. Tokenizes the text
    # 2. Maps the tokens to their IDs
    # 3. Adds the special [CLS] and [SEP] tokens.
    input ids = tokenizer.encode(text masked)
    print(input ids)
    # Use numpy's `where` function to find all indeces of the [MASK] token.
    mask token indeces = np.where(np.array(input ids) == tokenizer.mask token id)[0]
    return mask token indeces
```

```
#Select words to mask
words = ['abdomen', 'ablation', 'Acetaminophen', 'Alprazolam', 'aortic', 'artery', 'aspirin', 'Atrial', 'Cardiac', 'cardiothoracic', 'coronary',
'Coumadin', 'echocardiogram', 'edema', 'fibrillation', 'heart', 'heparin', 'Hypertension', 'Hypotension', ' 'palpitations', 'pericardial', 'pulmonary',
'tachycardia', 'ventricular', 'Warfarin', 'WPW', 'Xanax']
for i in range(len(notes)):
  text = notes[i]
  # [CLS]: 101; [SEP]: 102; [MASK]: 103; [PADDING]: 0
  print(clinical tokenizer.cls token id)
  print(clinical tokenizer.sep token id)
 print(clinical tokenizer.mask token id)
  print(clinical tokenizer.pad token id)
  print(get word indeces(clinical tokenizer, text, words[i]))
```

```
#Note that for visualization in this document, the function is split into two text
Boxes but is a single command in the Jupyter notebook.
#Get embeddings
def get embedding(b model, b tokenizer, text, word=''):
    Uses the provided model and tokenizer to produce an embedding for the
    provided `text`, and a "contextualized" embedding for `word`, if provided.
    # If a word is provided, figure out which tokens correspond to it.
    if not word == '':
        word indeces = get word indeces(b tokenizer, text, word)
    # Encode the text, adding the (required!) special tokens, and converting to
    # PyTorch tensors.
    encoded dict = b tokenizer.encode plus(
                                                   # Sentence to encode.
                        add special tokens = True, # Add '[CLS]' and '[SEP]'
                        return tensors = 'pt',  # Return pytorch tensors.
    input ids = encoded dict['input ids']
    b model.eval()
    # Run the text through the model and get the hidden states.
    bert outputs = b model(input ids)
```

```
# Run the text through BERT, and collect all of the hidden states produced
# from all 12 lavers.
with torch.no grad():
    outputs = b model(input ids,output hidden states=True )
    # Evaluating the model will return a different number of objects based on
    # how it's configured in the `from pretrained` call earlier. In this case,
    # because we set `output hidden states = True`, the third item will be the
    # hidden states from all layers. See the documentation for more details:
    # https://huggingface.co/transformers/model doc/bert.html#bertmodel
    hidden states = outputs[2]
# `hidden states` has shape [13 x 1 x <sentence length> x 768]
# Select the embeddings from the second to last layer.
# `token vecs` is a tensor with shape [<sent length> x 768]
token vecs = hidden states[-2][0]
# Calculate the average of all token vectors.
sentence embedding = torch.mean(token vecs, dim=0)
# Convert to numpy array.
sentence embedding = sentence embedding.detach().numpy()
# If `word` was provided, compute an embedding for those tokens.
if not word == '':
    # Take the average of the embeddings for the tokens in `word`.
    word embedding = torch.mean(token vecs[word indeces], dim=0)
    # Convert to numpy array.
    word embedding = word embedding.detach().numpy()
    return (sentence embedding, word embedding)
else:
    return sentence embedding
```

```
#Test out the get embeddings function
import nltk
nltk.download('stopwords')
nltk.download('punkt')
nltk.download('wordnet')
from nltk.corpus import stopwords
from nltk.tokenize import word tokenize
from nltk.stem import WordNetLemmatizer
import re
def clean text(text):
    # Tokenize the text into words
    words = text.split()
    # Remove special characters and convert to lowercase
    clean words = [word.lower() for word in words if word.isalnum()]
    # Remove stopwords
    stop words = set(stopwords.words("english"))
    filtered words = [word for word in clean words if word not in stop words]
    # Remove words with less than 4 characters and numbers. This is done in order to reduce noisy data and numbers dont contribute much in any NLP applications
    filtered words = [word for word in filtered words if len(word) >= 4 and not word.isdigit()]
    # Remove duplicate words
   cleaned text = " ".join(dict.fromkeys(filtered words)) # This is useful while plotting t-SNE plots
    return cleaned text
```

#### ClinicalBERT Tutorial – building notes for analysis

```
#in the first run, there was an issue with the tensor size so truncating notes to run
max length = 512
notes = [note[:max length] for note in notes]
words = [word[:max length] for word in words]
for i in range(len(notes)):
 text = notes[i]
  word = words[i] # words not recognized by the model will return nan, rest of the words will get embeddings
  text = clean text(text)
  clinical model.eval()
  # Get the embedding for the sentence, as well as an embedding for 'word'..
  (sen emb, word emb) = get embedding(clinical model, clinical tokenizer, text, word)
  print('Embedding sizes:')
  print(sen emb.shape)
  print(word emb.shape)
  print(sen emb)
  print(f'word embeddings for {word}')
  print(word emb)
```

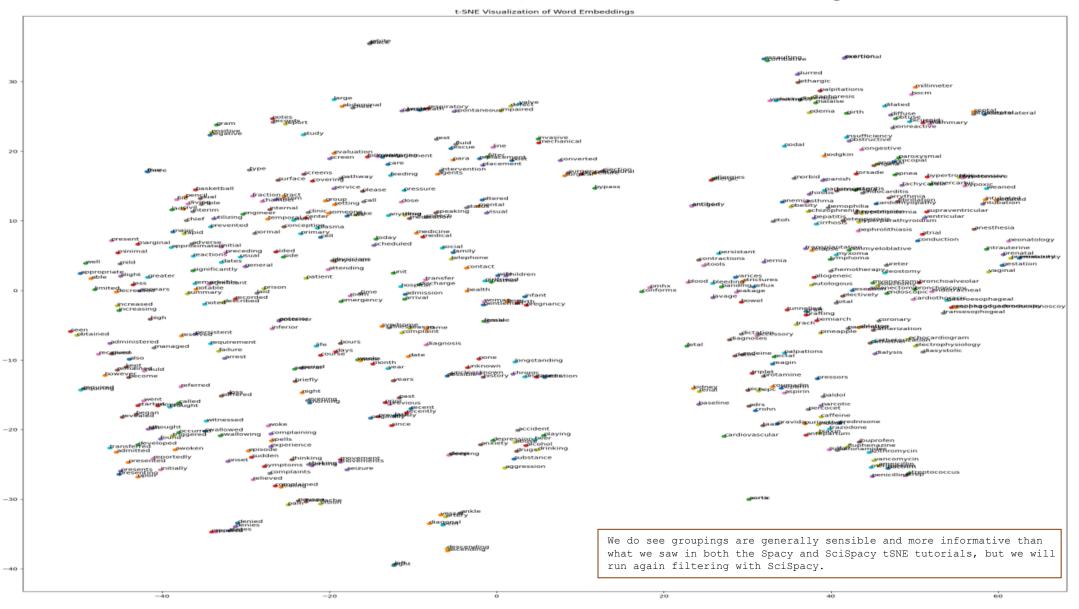
#### ClinicalBERT Tutorial – building notes for analysis

```
import scispacy
import spacy
nlp = spacy.load("en ner bc5cdr md")
# Build corpus of all the entities extracted from the notes using Spacy model.
# The corpus is an array of arrays or list of lists where each of the nested lists corresponds to a note.
for row in range(0, len(notes)):
 str tokens=[]
 tokens= nlp(notes[row]).ents
  for i in range(0, len(tokens)):
   str tokens.append(tokens[i].text)
 corpus.append(list(str tokens))
print(corpus)
#combining all notes to form a single note which will be easier to visualize in one graph
# Initialize an empty list to store the combined words
notes combined = []
# Iterate through the sublists and combine the words
for sublist in corpus:
    notes combined.extend(sublist)
print (notes combined)
# these notes contains all tokens that are extracted from the original notes using SciSpacy
notes combined = ' '.join(notes combined)
notes combined
# these are the original notes. No SciSpacy is used to filter tokens
all notes combined = '. '.join(notes)
all notes combined
```

#### ClinicalBERT Tutorial – visualization of all notes

```
# Visualization of all notes using ClinicalBert
import numpy as np
from sklearn.manifold import TSNE
import string
import matplotlib.pyplot as plt
from transformers import AutoModel, AutoTokenizer
# Load the BERT model and tokenizer
clinical model = AutoModel.from pretrained("emilyalsentzer/Bio ClinicalBERT")
clinical tokenizer = AutoTokenizer.from pretrained("emilyalsentzer/Bio ClinicalBERT")
clinical model.eval()
# Example input text
input text = clean text(all notes combined)
# Tokenize the input text using the BERT tokenizer
#input tokens = clinical tokenizer.tokenize(input text)
input tokens = input text.split()
# Initialize an empty list to store word embeddings
word embs = []
for token in input tokens:
    # Check if the token is a valid word
   if token not in string.punctuation:
        # Encode the token using the BERT model
        inputs = clinical tokenizer(token, return tensors="pt")
        with torch.no grad():
            outputs = clinical model(**inputs)
        token emb = outputs.last hidden state.mean(dim=1).squeeze().numpy()
        word embs.append(token emb)
# Perform t-SNE dimensionality reduction
tsne model = TSNE(n components=2, perplexity=10, random state=42)
word embs 2d = tsne model.fit transform(np.array(word embs))
# Create a scatter plot of the word embeddings in 2D space
plt.figure(figsize=(25, 25))
for i in range(len(word embs 2d)):
    plt.scatter(word embs 2d[i, 0], word embs 2d[i, 1])
   plt.annotate(input tokens[i], (word embs 2d[i, 0], word embs 2d[i, 1]))
plt.title("t-SNE Visualization of Word Embeddings")
plt.show()
```

### ClinicalBERT Tutorial – tSNE plot with no filtering



# ClinicalBERT Tutorial – visualization of notes filtered on SciSpacy

```
# Visualization of notes filtered with SciSpacy using ClinicalBert
import numpy as np
from sklearn.manifold import TSNE
import string
import matplotlib.pyplot as plt
from transformers import AutoModel, AutoTokenizer
# Load the BERT model and tokenizer
clinical model = AutoModel.from pretrained("emilyalsentzer/Bio ClinicalBERT")
clinical tokenizer = AutoTokenizer.from pretrained("emilyalsentzer/Bio ClinicalBERT")
clinical model.eval()
# Example input text
input text = clean text(notes combined)
# Tokenize the input text using the BERT tokenizer
#input tokens = clinical tokenizer.tokenize(input text)
input tokens = input text.split()
# Initialize an empty list to store word embeddings
word embs = []
for token in input tokens:
    # Check if the token is a valid word
   if token not in string.punctuation:
        # Encode the token using the BERT model
        inputs = clinical tokenizer(token, return tensors="pt")
        with torch.no grad():
            outputs = clinical model(**inputs)
        token emb = outputs.last hidden state.mean(dim=1).squeeze().numpy()
        word embs.append(token emb)
# Perform t-SNE dimensionality reduction
tsne model = TSNE(n components=2, perplexity=10, random state=42)
word embs 2d = tsne model.fit transform(np.array(word embs))
# Create a scatter plot of the word embeddings in 2D space
plt.figure(figsize=(25, 25))
for i in range(len(word embs 2d)):
    plt.scatter(word embs 2d[i, 0], word embs 2d[i, 1])
   plt.annotate(input_tokens[i], (word embs 2d[i, 0], word embs 2d[i, 1]))
plt.title("t-SNE Visualization of Word Embeddings")
plt.show()
```

# ClinicalBERT Tutorial – tSNE plot using SciSpacy filtering

