09. Milestone Project 2: SkimLit □□

In the previous notebook (<u>NLP fundamentals in TensorFlow (https://github.com/mrdbourke/tensorflow-deep-learning/blob/main/08 introduction to nlp in tensorflow.ipynb)</u>), we went through some fundamental natural lanuage processing concepts. The main ones being **tokenzation** (turning words into numbers) and **creating embeddings** (creating a numerical representation of words).

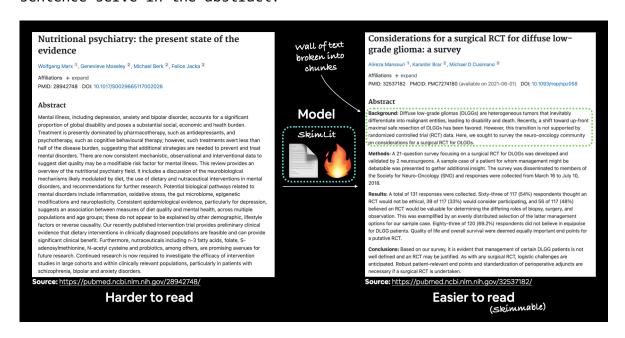
In this project, we're going to be putting what we've learned into practice.

More specificially, we're going to be replicating the deep learning model behind the 2017 paper <u>PubMed 200k RCT: a Dataset for Sequenctial Sentence Classification in Medical Abstracts</u> (https://arxiv.org/abs/1710.06071).

When it was released, the paper presented a new dataset called PubMed 200k RCT which consists of $\sim\!200,000$ labelled Randomized Controlled Trial (RCT) abstracts.

The goal of the dataset was to explore the ability for NLP models to classify sentences which appear in sequential order.

In other words, given the abstract of a RCT, what role does each sentence serve in the abstract?



Example inputs (harder to read abstract from PubMed (https://pubmed.ncbi.nlm.nih.gov/28942748/)) and outputs (easier to read abstract (https://pubmed.ncbi.nlm.nih.gov/32537182/)) of the model we're going to build. The model will take an abstract wall of text and predict the section label each sentence should have.

Model Input

For example, can we train an NLP model which takes the following input (note: the following sample has had all numerical symbols replaced with "@"):

To investigate the efficacy of @ weeks of daily low-dose oral prednisolone in improving pain , mobility , and systemic low-grade inflammation in the short term and whether the effect would be sustained at @ weeks in older adults with moderate to severe knee osteoarthritis (OA). A total of @ patients with primary knee OA were randomized @:@ ; @ received @ mg/day of prednisolone and @ received placebo for @ weeks. Outcome measures included pain reduction and improvement in function scores and systemic inflammation markers. Pain was assessed using the visual analog pain scale (@-@ mm). Secondary outcome measures included the Western Ontario and McMaster Universities Osteoarthritis Index scores , patient global assessment (PGA) of the severity of knee OA , and @-min walk distance (@MWD)., Serum levels of interleukin @ (IL-@) , IL-@ , tumor necrosis factor (TNF) - , and high-sensitivity Creactive protein (hsCRP) were measured. There was a clinically relevant reduction in the intervention group compared to the placebo group for knee pain , physical function , PGA , and @MWD at @ weeks. The mean difference between treatment arms (@%CI) was@(@-@@), p < @ ; @ (@-@ @) , p < @ ; @ (@-@ @) , p < @ ; and @ (@-@ @) , p < @ , respectively. Further , there was a clinically relevant reduction in the serum levels of IL-@ , IL-@ , TNF - , and hsCRP at @ weeks in the intervention group when compared to the placebo group. These differences remained significant at @ weeks. The Outcome Measures in Rheumatology Clinical Trials-Osteoarthritis Research Society International responder rate was @ % in the intervention group and @ % in the placebo group (p < @). Low-dose oral prednisolone had both a short-term and a longer sustained effect resulting in less knee pain , better physical function , and attenuation of systemic inflammation in older patients with knee OA (ClinicalTrials.gov identifier NCT@).

Model output

And returns the following output:

['###24293578\n',

'OBJECTIVE\tTo investigate the efficacy of @ weeks of daily lo w-dose oral prednisolone in improving pain , mobility , and sys temic low-grade inflammation in the short term and whether the effect would be sustained at @ weeks in older adults with moder ate to severe knee osteoarthritis (OA) .\n',

'METHODS\tA total of @ patients with primary knee OA were rand omized @:@ ; @ received @ mg/day of prednisolone and @ received placebo for @ weeks .\n',

'METHODS\tOutcome measures included pain reduction and improve ment in function scores and systemic inflammation markers .\n', 'METHODS\tPain was assessed using the visual analog pain scale (@-@ mm) .\n',

'METHODS\tSecondary outcome measures included the Western Onta rio and McMaster Universities Osteoarthritis Index scores , pat ient global assessment (PGA) of the severity of knee OA , and @-min walk distance (@MWD) .\n',

'METHODS\tSerum levels of interleukin @ (IL-@) , IL-@ , tumo r necrosis factor (TNF) - , and high-sensitivity C-reactive p rotein (hsCRP) were measured .\n',

'RESULTS\tThere was a clinically relevant reduction in the int ervention group compared to the placebo group for knee pain , p hysical function , PGA , and @MWD at @ weeks . \n' ,

'RESULTS\tThe mean difference between treatment arms (@ % CI) was @ (@-@ @) , p < @ ; @ (@-@ @) , p < @ ; @ (@-@ @) , p < @ , respectively .\n',

'RESULTS\tFurther , there was a clinically relevant reduction in the serum levels of IL-@ , IL-@ , TNF - , and hsCRP at @ wee ks in the intervention group when compared to the placebo group .\n',

'RESULTS\tThese differences remained significant at @ weeks .\n',

'RESULTS\tThe Outcome Measures in Rheumatology Clinical Trials -Osteoarthritis Research Society International responder rate w as @ % in the intervention group and @ % in the placebo group (p < @) .\n',

'CONCLUSIONS\tLow-dose oral prednisolone had both a short-term and a longer sustained effect resulting in less knee pain , bet ter physical function , and attenuation of systemic inflammatio n in older patients with knee OA (ClinicalTrials.gov identifier NCT@) .\n',

'\n']

Problem in a sentence

The number of RCT papers released is continuing to increase, those without structured abstracts can be hard to read and in turn slow down researchers moving through the literature.

Solution in a sentence

Create an NLP model to classify abstract sentences into the role they play (e.g. objective, methods, results, etc) to enable researchers to skim through the literature (hence $SkimLit \square\square$) and dive deeper when necessary.

☐ **Resources:** Before going through the code in this notebook, you might want to get a background of what we're going to be doing. To do so, spend an hour (or two) going through the following papers and then return to this notebook:

- Where our data is coming from: <u>PubMed 200k RCT: a Dataset for Sequential Sentence Classification in Medical Abstracts</u>
 (https://arxiv.org/abs/1710.06071)
- 2. Where our model is coming from: <u>Neural networks for joint sentence classification in medical paper abstracts</u>
 (https://arxiv.org/pdf/1612.05251.pdf).

What we're going to cover

Time to take what we've learned in the NLP fundmentals notebook and build our biggest NLP model yet:

- Downloading a text dataset (<u>PubMed RCT200k from GitHub</u> (https://github.com/Franck-Dernoncourt/pubmed-rct))
- Writing a preprocessing function to prepare our data for modelling
- Setting up a series of modelling experiments
 - Making a baseline (TF-IDF classifier)
 - Deep models with different combinations of: token embeddings, character embeddings, pretrained embeddings, positional embeddings
- Building our first multimodal model (taking multiple types of data inputs)
 - Replicating the model architecture from https://arxiv.org/pdf/1612.05251.pdf (https://arxiv.org/pdf/1612.05251.pdf)
- Find the most wrong predictions
- Making predictions on PubMed abstracts from the wild

How you should approach this notebook

You can read through the descriptions and the code (it should all run, except for the cells which error on purpose), but there's a better option.

Write all of the code yourself.

Yes. I'm serious. Create a new notebook, and rewrite each line by yourself. Investigate it, see if you can break it, why does it break?

You don't have to write the text descriptions but writing the code yourself is a great way to get hands-on experience.

Don't worry if you make mistakes, we all do. The way to get better and make less mistakes is to write more code.

☐ **Resource:** See the full set of course materials on GitHub: https://github.com/mrdbourke/tensorflow-deep-learning (<a href="https:

Confirm access to a GPU

Since we're going to be building deep learning models, let's make sure we have a GPU.

In Google Colab, you can set this up by going to Runtime -> Change runtime type -> Hardware accelerator -> GPU.

If you don't have access to a GPU, the models we're building here will likely take up to 10x longer to run.

```
In [ ]: # Check for GPU
!nvidia-smi -L
```

GPU 0: Tesla T4 (UUID: GPU-90b6bfd2-2dbc-6214-b3b0-835ecd7fd102)

Get data

Before we can start building a model, we've got to download the PubMed 200k RCT dataset.

In a phenomenal act of kindness, the authors of the paper have made the data they used for their research availably publically and for free in the form of .txt files <u>on GitHub (https://github.com/Franck-Dernoncourt/pubmed-rct)</u>.

We can copy them to our local directory using git clone https://github.com/Franck-Dernoncourt/pubmed-rct.

In []: !git clone https://github.com/Franck-Dernoncourt/pubmed-rct.git
!ls pubmed-rct

Cloning into 'pubmed-rct'...
remote: Enumerating objects: 33, done.
remote: Counting objects: 100% (3/3), done.
remote: Compressing objects: 100% (3/3), done.
remote: Total 33 (delta 0), reused 0 (delta 0), pack-reused 30
Unpacking objects: 100% (33/33), done.
PubMed_200k_RCT
PubMed_200k_RCT_numbers_replaced_with_at_sign
PubMed_20k_RCT
PubMed_20k_RCT_numbers_replaced_with_at_sign
README.md

Checking the contents of the downloaded repository, you can see there are four folders.

Each contains a different version of the PubMed 200k RCT dataset.

Looking at the <u>README file (https://github.com/Franck-Dernoncourt/pubmed-rct)</u> from the GitHub page, we get the following information:

- PubMed 20k is a subset of PubMed 200k. I.e., any abstract present in PubMed 20k is also present in PubMed 200k.
- PubMed_200k_RCT is the same as PubMed_200k_RCT_numbers_replaced_with_at_sign, except that in the latter all numbers had been replaced by @. (same for PubMed_20k_RCT vs.
 - PubMed_20k_RCT_numbers_replaced_with_at_sign).
- Since Github file size limit is 100 MiB, we had to compress PubMed_200k_RCT\train.7z and PubMed_200k_RCT_numbers_replaced_with_at_sign\train.zip. To uncompress train.7z, you may use 7-Zip on Windows, Keka on Mac OS X, or p7zip on Linux.

To begin with, the dataset we're going to be focused on is PubMed_20k_RCT_numbers_replaced_with_at_sign .

Why this one?

Rather than working with the whole 200k dataset, we'll keep our experiments quick by starting with a smaller subset. We could've chosen the dataset with numbers instead of having them replaced with @ but we didn't.

Let's check the file contents.

In []: # Check what files are in the PubMed_20K dataset
!ls pubmed-rct/PubMed_20k_RCT_numbers_replaced_with_at_sign

dev.txt test.txt train.txt

Beautiful, looks like we've got three separate text files:

- train.txt training samples.
- dev.txt dev is short for development set, which is another name for validation set (in our case, we'll be using and referring to this file as our validation set).
- test.txt test samples.

To save ourselves typing out the filepath to our target directory each time, let's turn it into a variable.

```
In [ ]: # Start by using the 20k dataset
data_dir = "pubmed-rct/PubMed_20k_RCT_numbers_replaced_with_at_sign/"
```

```
In [ ]: # Check all of the filenames in the target directory
import os
filenames = [data_dir + filename for filename in os.listdir(data_dir)]
filenames
```

Preprocess data

Okay, now we've downloaded some text data, do you think we're ready to model it?

Wait...

We've downloaded the data but we haven't even looked at it yet.

What's the motto for getting familiar with any new dataset?

I'll give you a clue, the word begins with " ν " and we say it three times.

```
Vibe, vibe, vibe?
```

Sort of... we've definitely got to the feel the vibe of our data.

```
Values, values?
```

Right again, we want to *see* lots of values but not quite what we're looking for.

```
Visualize, visualize, visualize?
```

Boom! That's it. To get familiar and understand how we have to prepare our data for our deep learning models, we've got to visualize it.

Because our data is in the form of text files, let's write some code to read each of the lines in a target file.

```
In []: # Create function to read the lines of a document
def get_lines(filename):
    """
    Reads filename (a text file) and returns the lines of text as a list.

Args:
    filename: a string containing the target filepath to read.

Returns:
    A list of strings with one string per line from the target filena
    For example:
    ["this is the first line of filename",
        "this is the second line of filename",
        "..."]

with open(filename, "r") as f:
    return f.readlines()
```

Alright, we've got a little function, get_lines() which takes the filepath of a text file, opens it, reads each of the lines and returns them.

Let's try it out on the training data (train.txt).

```
In [ ]: train_lines = get_lines(data_dir+"train.txt")
    train_lines[:20] # the whole first example of an abstract + a little mo
```

Out[7]: ['###24293578\n',

'OBJECTIVE\tTo investigate the efficacy of @ weeks of daily low-dose oral prednisolone in improving pain , mobility , and systemic low-grad e inflammation in the short term and whether the effect would be sustained at @ weeks in older adults with moderate to severe knee osteoarth ritis (OA) .\n',

'METHODS\tA total of @ patients with primary knee OA were randomized @:@ ; @ received @ mg/day of prednisolone and @ received placebo for @ weeks .\n',

'METHODS\tOutcome measures included pain reduction and improvement in function scores and systemic inflammation markers .\n',

'METHODS\tPain was assessed using the visual analog pain scale (@-@ mm) .\n',

'METHODS\tSecondary outcome measures included the Western Ontario and McMaster Universities Osteoarthritis Index scores , patient global ass essment (PGA) of the severity of knee OA , and @-min walk distance (@MWD) .\n',

'METHODS\tSerum levels of interleukin @ (IL-@) , IL-@ , tumor necro sis factor (TNF) - , and high-sensitivity C-reactive protein (hsCRP) were measured .\n',

'RESULTS\tThere was a clinically relevant reduction in the interventi on group compared to the placebo group for knee pain , physical functi on , PGA , and @MWD at @ weeks .\n',

'RESULTS\tThe mean difference between treatment arms (@ % CI) was @ (@-@ @) , p < @ ; @ (@-@ @) , p < @ ; and @ (@-@ @) , p < @ , respectively .\n',

'RESULTS\tFurther , there was a clinically relevant reduction in the serum levels of IL-@ , IL-@ , TNF - , and hsCRP at @ weeks in the intervention group when compared to the placebo group .\n',

'RESULTS\tThese differences remained significant at @ weeks .\n',

'RESULTS\tThe Outcome Measures in Rheumatology Clinical Trials-Osteoa rthritis Research Society International responder rate was @ % in the intervention group and @ % in the placebo group (p < @) .\n',

'CONCLUSIONS\tLow-dose oral prednisolone had both a short-term and a longer sustained effect resulting in less knee pain , better physical function , and attenuation of systemic inflammation in older patients with knee OA (ClinicalTrials.gov identifier NCT@) .\n',

'\n',

'###24854809\n',

'BACKGROUND\tEmotional eating is associated with overeating and the development of obesity .\n',

'BACKGROUND\tYet , empirical evidence for individual (trait) differ ences in emotional eating and cognitive mechanisms that contribute to eating during sad mood remain equivocal .\n',

'OBJECTIVE\tThe aim of this study was to test if attention bias for f ood moderates the effect of self-reported emotional eating during sad mood (vs neutral mood) on actual food intake .\n',

'OBJECTIVE\tIt was expected that emotional eating is predictive of el evated attention for food and higher food intake after an experimental ly induced sad mood and that attentional maintenance on food predicts food intake during a sad versus a neutral mood .\n',

'METHODS\tParticipants (N = @) were randomly assigned to one of the two experimental mood induction conditions (sad/neutral) .\n'l

Reading the lines from the training text file results in a list of strings containing different abstract samples, the sentences in a sample along with the role the sentence plays in the abstract.

The role of each sentence is prefixed at the start of each line separated by a tab (\t) and each sentence finishes with a new line (\t).

Different abstracts are separated by abstract ID's (lines beginning with ###) and newlines (\n).

Knowing this, it looks like we've got a couple of steps to do to get our samples ready to pass as training data to our future machine learning model.

Let's write a function to perform the following steps:

- Take a target file of abstract samples.
- Read the lines in the target file.
- For each line in the target file:
 - If the line begins with ### mark it as an abstract ID and the beginning of a new abstract.
 - Keep count of the number of lines in a sample.
 - If the line begins with \n mark it as the end of an abstract sample.
 - Keep count of the total lines in a sample.
 - Record the text before the \t as the label of the line.
 - Record the text after the \t as the text of the line.
- Return all of the lines in the target text file as a list of dictionaries containing the key/value pairs:
 - "line_number" the position of the line in the abstract (e.g.
 3).
 - "target" the role of the line in the abstract (e.g. OBJECTIVE).
 - "text" the text of the line in the abstract.
 - "total_lines" the total lines in an abstract sample (e.g. 14).
- Abstract ID's and newlines should be omitted from the returned preprocessed data.

Example returned preprocessed sample (a single line from an abstract):

```
[{'line_number': 0,
              'target': 'OBJECTIVE',
              'text'. 'to investigate the efficacy of @ weeks of daily low-
In []: def preprocess text with line numbers(filename):
          """Returns a list of dictionaries of abstract line data.
          Takes in filename, reads its contents and sorts through each line,
          extracting things like the target label, the text of the sentence,
          how many sentences are in the current abstract and what sentence numb
          the target line is.
          Args:
              filename: a string of the target text file to read and extract li
              from.
          Returns:
              A list of dictionaries each containing a line from an abstract,
              the lines label, the lines position in the abstract and the total
              of lines in the abstract where the line is from. For example:
              [{"target": 'CONCLUSION',
                "text": The study couldn't have gone better, turns out people a
                "line number": 8,
                "total_lines": 8}]
          input_lines = get_lines(filename) # get all lines from filename
          abstract lines = "" # create an empty abstract
          abstract_samples = [] # create an empty list of abstracts
          # Loop through each line in target file
          for line in input_lines:
            if line.startswith("###"): # check to see if line is an ID line
              abstract id = line
              abstract_lines = "" # reset abstract string
            elif line.isspace(): # check to see if line is a new line
              abstract_line_split = abstract_lines.splitlines() # split abstract
              # Iterate through each line in abstract and count them at the sam
              for abstract line number, abstract line in enumerate(abstract lin
                line_data = {} # create empty dict to store data from line
                target_text_split = abstract_line.split("\t") # split target la
                line_data["target"] = target_text_split[0] # get target label
                line_data["text"] = target_text_split[1].lower() # get target t
                line_data["line_number"] = abstract_line_number # what number 1
                line_data["total_lines"] = len(abstract_line_split) - 1 # how n
                abstract_samples.append(line_data) # add line data to abstract
            else: # if the above conditions aren't fulfilled, the line contains
              abstract_lines += line
          return abstract_samples
```

Beautiful! That's one good looking function. Let's use it to preprocess each of our RCT 20k datasets.

In []: # Get data from file and preprocess it %%time train_samples = preprocess_text_with_line_numbers(data_dir + "train.txt val_samples = preprocess_text_with_line_numbers(data_dir + "dev.txt") # test_samples = preprocess_text_with_line_numbers(data_dir + "test.txt") len(train_samples), len(val_samples), len(test_samples)

CPU times: user 450 ms, sys: 89.4 ms, total: 540 ms

Wall time: 540 ms

How do our training samples look?

In []: # Check the first abstract of our training data
train_samples[:14]

```
Out[10]: [{'line_number': 0,
            'target': 'OBJECTIVE',
            'text': 'to investigate the efficacy of @ weeks of daily low-dose or
         al prednisolone in improving pain , mobility , and systemic low-grade
         inflammation in the short term and whether the effect would be sustain
         ed at @ weeks in older adults with moderate to severe knee osteoarthri
         tis ( oa ) .',
            'total_lines': 11},
          {'line_number': 1,
            'target': 'METHODS',
            'text': 'a total of @ patients with primary knee oa were randomized
         @:@ ; @ received @ mg/day of prednisolone and @ received placebo for @
         weeks .',
           'total_lines': 11},
          {'line_number': 2,
            'target': 'METHODS',
            'text': 'outcome measures included pain reduction and improvement in
         function scores and systemic inflammation markers .',
            'total_lines': 11},
          {'line_number': 3,
            'target': 'METHODS',
            'text': 'pain was assessed using the visual analog pain scale ( @-@
         mm ) .',
           'total_lines': 11},
          {'line_number': 4,
            'target': 'METHODS',
            'text': 'secondary outcome measures included the western ontario and
         mcmaster universities osteoarthritis index scores , patient global ass
         essment ( pga ) of the severity of knee oa , and @-min walk distance (
         @mwd ) .',
            'total_lines': 11},
          {'line_number': 5,
            'target': 'METHODS',
            'text': 'serum levels of interleukin @ ( il-@ ) , il-@ , tumor necro
         sis factor (tnf) - , and high-sensitivity c-reactive protein (hscrp
         ) were measured .',
           'total_lines': 11},
          {'line_number': 6,
            'target': 'RESULTS',
            'text': 'there was a clinically relevant reduction in the interventi
         on group compared to the placebo group for knee pain , physical functi
         on , pga , and @mwd at @ weeks .',
            'total_lines': 11},
          {'line_number': 7,
            'target': 'RESULTS',
            'text': 'the mean difference between treatment arms ( @ % ci ) was @
         ( @-@ @ ) , p < @ ; @ ( @-@ @ ) , p < @ ; @ ( @-@ @ ) , p < @ ; and @
         ( @-@ @ ) , p < @ , respectively .',
           'total_lines': 11},
          {'line_number': 8,
            'target': 'RESULTS',
            'text': 'further , there was a clinically relevant reduction in the
         serum levels of il-@ , il-@ , tnf - , and hscrp at @ weeks in the inte
         rvention group when compared to the placebo group .',
           'total_lines': 11},
          {'line_number': 9,
            'target': 'RESULTS',
```

```
'text': 'these differences remained significant at @ weeks .',
  'total_lines': 11},
 {'line_number': 10,
  'target': 'RESULTS',
  'text': 'the outcome measures in rheumatology clinical trials-osteoa
rthritis research society international responder rate was @ % in the
intervention group and @ % in the placebo group ( p < @ ) .',
  'total_lines': 11},
 {'line_number': 11,
  'target': 'CONCLUSIONS',
  'text': 'low-dose oral prednisolone had both a short-term and a long
er sustained effect resulting in less knee pain , better physical func
tion , and attenuation of systemic inflammation in older patients with
knee oa ( clinicaltrials.gov identifier nct@ ) .',
  'total_lines': 11},
 {'line number': 0,
  'target': 'BACKGROUND',
  'text': 'emotional eating is associated with overeating and the deve
lopment of obesity .',
  'total_lines': 10},
 {'line_number': 1,
  'target': 'BACKGROUND',
  'text': 'yet , empirical evidence for individual ( trait ) differenc
es in emotional eating and cognitive mechanisms that contribute to eat
ing during sad mood remain equivocal .',
  'total_lines': 10}]
```

Fantastic! Looks like our preprocess_text_with_line_numbers() function worked great.

How about we turn our list of dictionaries into pandas DataFrame's so we visualize them better?

In []: import pandas as pd
 train_df = pd.DataFrame(train_samples)
 val_df = pd.DataFrame(val_samples)
 test_df = pd.DataFrame(test_samples)
 train_df.head(14)

Out[11]:		target	text	line_number	total_lines
	0	OBJECTIVE	to investigate the efficacy of @ weeks of dail	0	11
	1	METHODS	a total of @ patients with primary knee oa wer	1	11
	2	METHODS	outcome measures included pain reduction and i	2	11
	3	METHODS	pain was assessed using the visual analog pain	3	11
	4	METHODS	secondary outcome measures included the wester	4	11
	5	METHODS	serum levels of interleukin @ (il-@) , il-@	5	11
	6	RESULTS	there was a clinically relevant reduction in t	6	11
	7	RESULTS	the mean difference between treatment arms ($@$	7	11
9 10	RESULTS	further , there was a clinically relevant redu	8	11	
	9	RESULTS	these differences remained significant at @ we	9	11
	RESULTS	the outcome measures in rheumatology clinical	10	11	
	11	CONCLUSIONS	low-dose oral prednisolone had both a short-te	11	11
:	12	BACKGROUND	emotional eating is associated with overeating	0	10
	13	BACKGROUND	yet , empirical evidence for individual (trai	1	10

Now our data is in DataFrame form, we can perform some data analysis on it.

In []: # Distribution of labels in training data

train_df.target.value_counts()

Out[12]: METHODS 59353

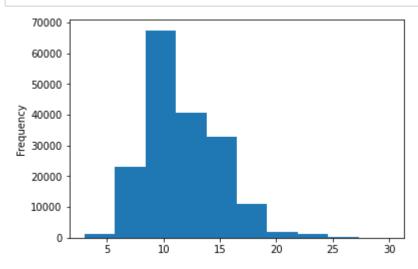
RESULTS 57953 CONCLUSIONS 27168 BACKGROUND 21727 OBJECTIVE 13839

Name: target, dtype: int64

Looks like sentences with the OBJECTIVE label are the least common.

How about we check the distribution of our abstract lengths?

In []: train_df.total_lines.plot.hist();



Okay, looks like most of the abstracts are around 7 to 15 sentences in length.

It's good to check these things out to make sure when we do train a model or test it on unseen samples, our results aren't outlandish.

Get lists of sentences

When we build our deep learning model, one of its main inputs will be a list of strings (the lines of an abstract).

We can get these easily from our DataFrames by calling the tolist() method on our "text" columns.

```
In [ ]: # Convert abstract text lines into lists
         train_sentences = train_df["text"].tolist()
         val_sentences = val_df["text"].tolist()
         test_sentences = test_df["text"].tolist()
         len(train_sentences), len(val_sentences), len(test_sentences)
Out[14]: (180040, 30212, 30135)
 In [ ]: | # View first 10 lines of training sentences
         train_sentences[:10]
Out[15]: ['to investigate the efficacy of @ weeks of daily low-dose oral predni
         solone in improving pain , mobility , and systemic low-grade inflammat
         ion in the short term and whether the effect would be sustained at @ w
         eeks in older adults with moderate to severe knee osteoarthritis ( oa
         ) .',
          'a total of @ patients with primary knee oa were randomized @:@ ; @ r
         eceived @ mg/day of prednisolone and @ received placebo for @ weeks
          'outcome measures included pain reduction and improvement in function
         scores and systemic inflammation markers .',
          'pain was assessed using the visual analog pain scale ( @-@ mm ) .',
          'secondary outcome measures included the western ontario and mcmaster
         universities osteoarthritis index scores , patient global assessment (
         pga ) of the severity of knee oa , and @-min walk distance ( @mwd )
          'serum levels of interleukin @ ( il-@ ) , il-@ , tumor necrosis facto
         r ( tnf ) - , and high-sensitivity c-reactive protein ( hscrp ) were m
         easured .',
          'there was a clinically relevant reduction in the intervention group
         compared to the placebo group for knee pain , physical function , pga
         , and @mwd at @ weeks .',
          'the mean difference between treatment arms ( @ % ci ) was @ ( @-@ @
         ) , p < @ ; @ ( @-@ @ ) , p < @ ; @ ( @-@ @ ) , p < @ ; and @ ( @-@ @
         ) , p < @ , respectively .',
          'further , there was a clinically relevant reduction in the serum lev
         els of il-@ , il-@ , tnf - , and hscrp at @ weeks in the intervention
         group when compared to the placebo group .',
          'these differences remained significant at @ weeks .']
```

Alright, we've separated our text samples. As you might've guessed, we'll have to write code to convert the text to numbers before we can use it with our machine learning models, we'll get to this soon.

Make numeric labels (ML models require numeric labels)

We're going to create one hot and label encoded labels.

We could get away with just making label encoded labels, however, TensorFlow's CategoricalCrossentropy loss function likes to have one hot encoded labels (this will enable us to use label smoothing later on).

To numerically encode labels we'll use Scikit-Learn's <u>OneHotEncoder</u> (https://scikit-

learn.org/stable/modules/generated/sklearn.preprocessing.OneHotEncoder.
and LabelEncoder _(http://scikit-

<u>learn.org/stable/modules/generated/sklearn.preprocessing.LabelEncoder.h</u> classes.

Label encode labels

```
In []: # Extract labels ("target" columns) and encode them into integers
    from sklearn.preprocessing import LabelEncoder
    label_encoder = LabelEncoder()
    train_labels_encoded = label_encoder.fit_transform(train_df["target"].t
    val_labels_encoded = label_encoder.transform(val_df["target"].to_numpy(
    test_labels_encoded = label_encoder.transform(test_df["target"].to_nump

# Check what training labels look like
    train_labels_encoded
```

Out[17]: array([3, 2, 2, ..., 4, 1, 1])

dtype=object))

Now we've trained an instance of LabelEncoder, we can get the class names and number of classes using the classes_ attribute.

```
In []: # Get class names and number of classes from LabelEncoder instance
    num_classes = len(label_encoder.classes_)
    class_names = label_encoder.classes_
    num_classes, class_names
Out[18]: (5, array(['BACKGROUND', 'CONCLUSIONS', 'METHODS', 'OBJECTIVE', 'RESUL TS'],
```

Creating a series of model experiments

We've proprocessed our data so now, in true machine learning fashion, it's time to setup a series of modelling experiments.

We'll start by creating a simple baseline model to obtain a score we'll try to beat by building more and more complex models as we move towards replicating the sequence model outlined in <u>Neural networks</u> <u>for joint sentence classification in medical paper abstracts</u> (https://arxiv.org/pdf/1612.05251.pdf).

For each model, we'll train it on the training data and evaluate it on the validation data.

Model 0: Getting a baseline

Our first model we'll be a TF-IDF Multinomial Naive Bayes as recommended by Scikit-Learn's machine learning map (https://scikit-learn.org/stable/tutorial/machine learning map/index.html).

To build it, we'll create a Scikit-Learn Pipeline which uses the TfidfVectorizer (https://scikit-

<u>learn.org/stable/modules/generated/sklearn.feature_extraction.text.Tfid</u> class to convert our abstract sentences to numbers using the TF-IDF (term frequency-inverse document frequecy) algorithm and then learns to classify our sentences using the <u>MultinomialNB_(https://scikit-learn.org/stable/modules/generated/sklearn.naive_bayes.MultinomialNB.ht</u> aglorithm.

Due to the speed of the Multinomial Naive Bayes algorithm, it trains very quickly.

We can evaluate our model's accuracy on the validation dataset using the score() method.

Out[20]: 0.7218323844829869

Nice! Looks like 72.1% accuracy will be the number to beat with our deeper models.

Now let's make some predictions with our baseline model to further evaluate it.

```
In [ ]: # Make predictions
baseline_preds = model_0.predict(val_sentences)
baseline_preds
```

```
Out[21]: array([4, 1, 3, ..., 4, 4, 1])
```

To evaluate our baseline's predictions, we'll import the calculate_results() function we created in the <u>previous notebook</u> https://github.com/mrdbourke/tensorflow-deep-learning/blob/main/08 introduction to nlp in tensorflow.ipynb) and added it to our helper_functions.py script https://github.com/mrdbourke/tensorflow-deep-learning/blob/main/extras/helper_functions.py) to compare them to the ground truth labels.

More specificially the calculate_results() function will help us obtain the following:

- Accuracy
- Precision
- Recall
- F1-score

Download helper functions script

Let's get our helper_functions.py script we've been using to store helper functions we've created in previous notebooks.

```
In []: # Download helper functions script
         !wget https://raw.githubusercontent.com/mrdbourke/tensorflow-deep-learn
         --2021-08-24 23:56:53-- https://raw.githubusercontent.com/mrdbourke/t
         ensorflow-deep-learning/main/extras/helper_functions.py (https://raw.g
         ithubusercontent.com/mrdbourke/tensorflow-deep-learning/main/extras/he
         lper_functions.py)
         Resolving raw.githubusercontent.com (raw.githubusercontent.com)... 18
         5.199.110.133, 185.199.111.133, 185.199.108.133, ...
         Connecting to raw.githubusercontent.com (raw.githubusercontent.com)|18
         5.199.110.133|:443... connected.
         HTTP request sent, awaiting response... 200 OK
         Length: 10246 (10K) [text/plain]
         Saving to: 'helper_functions.py'
         helper functions.py 100%[==========] 10.01K --.-KB/s
                                                                             in
         0s
         2021-08-24 23:56:53 (89.8 MB/s) - 'helper_functions.py' saved [10246/1
         02461
         Now we've got the helper functions script we can import the
         caculate_results() function and see how our baseline model went.
In [ ]: # Import calculate_results helper function
         from helper_functions import calculate_results
In [ ]: |# Calculate baseline results
         baseline_results = calculate_results(y_true=val_labels_encoded,
                                              y_pred=baseline_preds)
         baseline_results
Out[24]: {'accuracy': 72.1832384482987,
           'f1': 0.6989250353450294,
          'precision': 0.7186466952323352,
```

Preparing our data for deep sequence models

'recall': 0.7218323844829869}

Excellent! We've got a working baseline to try and improve upon.

But before we start building deeper models, we've got to create vectorization and embedding layers.

The vectorization layer will convert our text to numbers and the embedding layer will capture the relationships between those numbers.

To start creating our vectorization and embedding layers, we'll need to import the appropriate libraries (namely TensorFlow and NumPy).

```
In [ ]: import numpy as np
import tensorflow as tf
from tensorflow.keras import layers
```

Since we'll be turning our sentences into numbers, it's a good idea to figure out how many words are in each sentence.

When our model goes through our sentences, it works best when they're all the same length (this is important for creating batches of the same size tensors).

For example, if one sentence is eight words long and another is 29 words long, we want to pad the eight word sentence with zeros so it ends up being the same length as the 29 word sentence.

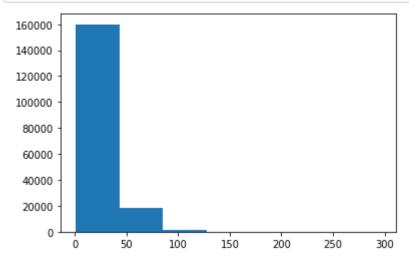
Let's write some code to find the average length of sentences in the training set.

```
In [ ]: # How long is each sentence on average?
    sent_lens = [len(sentence.split()) for sentence in train_sentences]
    avg_sent_len = np.mean(sent_lens)
    avg_sent_len # return average sentence length (in tokens)
```

Out[26]: 26.338269273494777

How about the distribution of sentence lengths?

```
In [ ]: # What's the distribution look like?
import matplotlib.pyplot as plt
plt.hist(sent_lens, bins=7);
```



Looks like the vast majority of sentences are between 0 and 50 tokens in length.

```
In [ ]: # How long of a sentence covers 95% of the lengths?
   output_seq_len = int(np.percentile(sent_lens, 95))
   output_seq_len
```

Out[28]: 55

Wonderful! It looks like 95% of the sentences in our training set have a length of 55 tokens or less.

When we create our tokenization layer, we'll use this value to turn all of our sentences into the same length. Meaning sentences with a length below 55 get padded with zeros and sentences with a length above 55 get truncated (words after 55 get cut off).

```
□ Question: Why 95%?
```

We could use the max sentence length of the sentences in the training set.

```
In [ ]: # Maximum sentence length in the training set
max(sent_lens)
```

Out[29]: 296

However, since hardly any sentences even come close to the max length, it would mean the majority of the data we pass to our model would be zeros (sinces all sentences below the max length would get padded with zeros).

□ **Note:** The steps we've gone through are good practice when working with a text corpus for a NLP problem. You want to know how long your samples are and what the distribution of them is. See section 4 Data Analysis of the PubMed 200k RCT paper
(https://arxiv.org/pdf/1710.06071.pdf) for further examples.

Create text vectorizer

Now we've got a little more information about our texts, let's create a way to turn it into numbers.

To do so, we'll use the <u>TextVectorization</u> (https://www.tensorflow.org/api docs/python/tf/keras/layers/experimenta layer from TensorFlow.

We'll keep all the parameters default except for max_tokens (the number of unique words in our dataset) and output_sequence_length (our desired output length for each vectorized sentence).

Section 3.2 of the <u>PubMed 200k RCT paper</u> (https://arxiv.org/pdf/1710.06071.pdf) states the vocabulary size of the PubMed 20k dataset as 68,000. So we'll use that as our max tokens parameter.

```
In [ ]: # How many words are in our vocabulary? (taken from 3.2 in https://arxi
max_tokens = 68000
```

And since discovered a sentence length of 55 covers 95% of the training sentences, we'll use that as our output_sequence_length parameter.

Great! Looks like our text_vectorizer is ready, let's adapt it to the training data (let it read the training data and figure out what number should represent what word) and then test it out.

```
In [ ]: # Adapt text vectorizer to training sentences
    text_vectorizer.adapt(train_sentences)
```

```
In [ ]: # Test out text vectorizer
import random
target_sentence = random.choice(train_sentences)
print(f"Text:\n{target_sentence}")
print(f"\nLength of text: {len(target_sentence.split())}")
print(f"\nVectorized text:\n{text_vectorizer([target_sentence])}")
```

Tovt.

http://www.clinicaltrials.gov (http://www.clinicaltrials.gov) .

Length of text: 2

Vectorized text: [[2243 0]]

Cool, we've now got a way to turn our sequences into numbers.

★ Exercise: Try running the cell above a dozen or so times. What do you notice about sequences with a length less than 55?

Using the <u>get_vocabulary()</u>
https://www.tensorflow.org/api_docs/python/tf/keras/layers/experimenta
method of our text_vectorizer we can find out a few different tidbits about our text.

```
In []: # How many words in our training vocabulary?
    rct_20k_text_vocab = text_vectorizer.get_vocabulary()
    print(f"Number of words in vocabulary: {len(rct_20k_text_vocab)}"),
    print(f"Most common words in the vocabulary: {rct_20k_text_vocab[:5]}")
    print(f"Least common words in the vocabulary: {rct_20k_text_vocab[-5:]}
```

Number of words in vocabulary: 64841 Most common words in the vocabulary: ['', '[UNK]', 'the', 'and', 'of'] Least common words in the vocabulary: ['aainduced', 'aaigroup', 'aache ner', 'aachen', 'aaacp']

And if we wanted to figure out the configuration of our text_vectorizer we can use the get_config() method.

```
In [ ]: # Get the config of our text vectorizer
    text_vectorizer.get_config()

Out[35]: {'batch_input_shape': (None,),
        'dtype': 'string',
        'max_tokons': 68000
```

'max_tokens': 68000,
'name': 'text_vectorization',
'ngrams': None,
'output_mode': 'int',
'output_sequence_length': 55,
'pad_to_max_tokens': False,
'split': 'whitespace',
'standardize': 'lower_and_strip_punctuation',
'trainable': True}

Create custom text embedding

Our token_vectorization layer maps the words in our text directly to numbers. However, this doesn't necessarily capture the relationships between those numbers.

To create a richer numerical representation of our text, we can use an **embedding**.

As our model learns (by going through many different examples of abstract sentences and their labels), it'll update its embedding to better represent the relationships between tokens in our corpus.

We can create a trainable embedding layer using TensorFlow's Embedding

(https://www.tensorflow.org/tutorials/text/word_embeddings) layer.

Once again, the main parameters we're concerned with here are the inputs and outputs of our Embedding layer.

The input_dim parameter defines the size of our vocabulary. And the output_dim parameter defines the dimension of the embedding output.

Once created, our embedding layer will take the integer outputs of our text_vectorization layer as inputs and convert them to feature vectors of size output_dim .

Let's see it in action.

```
In [ ]: # Create token embedding layer
        token_embed = layers.Embedding(input_dim=len(rct_20k_text_vocab), # ler
                                       output_dim=128, # Note: different embedd
                                       # Use masking to handle variable sequend
                                       mask zero=True,
                                       name="token_embedding")
        # Show example embedding
        print(f"Sentence before vectorization:\n{target_sentence}\n")
        vectorized_sentence = text_vectorizer([target_sentence])
        print(f"Sentence after vectorization (before embedding):\n{vectorized_s
        embedded_sentence = token_embed(vectorized_sentence)
        print(f"Sentence after embedding:\n{embedded_sentence}\n")
        print(f"Embedded sentence shape: {embedded_sentence.shape}")
        Sentence before vectorization:
        http://www.clinicaltrials.gov (http://www.clinicaltrials.gov) .
        Sentence after vectorization (before embedding):
        [[2243
                                                                    0
        0
             0
                                                                        0
                                 0
                                           0
                                                0
                                                     0
        0
             0
                  0
                       0
                            0
                                 0
                                      0
                                           0
                                                0
                                                    0
                                                         0
                                                                    0
                                                                        0
        0
                                      0
             0
                  0
                                 0
                                           0
                                                0
                                                    0
                                                         0
                                                                    0
                                                                        0]]
        Sentence after embedding:
        [[[ 0.0105269
                        0.04210378 0.04194726 ...
                                                   0.0111946
                                                                0.02379807
           -0.0433928 ]
          [-0.02391002 0.04098249 -0.0333933 ...
                                                   0.01311035 -0.02968328
           -0.047244721
          [-0.02391002 0.04098249 -0.0333933 ...
                                                   0.01311035 -0.02968328
           -0.047244721
          [-0.02391002  0.04098249  -0.0333933  ...
                                                   0.01311035 -0.02968328
           -0.047244721
          [-0.02391002 \quad 0.04098249 \quad -0.0333933 \quad \dots \quad 0.01311035 \quad -0.02968328
           -0.047244721
          -0.04724472]]]
```

Embedded sentence shape: (1, 55, 128)

Create datasets (as fast as possible)

We've gone through all the trouble of preprocessing our datasets to be used with a machine learning model, however, there are still a few steps we can use to make them work faster with our models.

Namely, the tf.data API provides methods which enable faster data loading.

☐ **Resource:** For best practices on data loading in TensorFlow, check out the following:

- tf.data: Build TensorFlow input pipelines
 (https://www.tensorflow.org/quide/data)
- <u>Better performance with the tf.data API</u> (https://www.tensorflow.org/quide/data performance)

The main steps we'll want to use with our data is to turn it into a PrefetchDataset of batches.

Doing so we'll ensure TensorFlow loads our data onto the GPU as fast as possible, in turn leading to faster training time.

To create a batched PrefetchDataset we can use the methods <u>batch()</u> (https://www.tensorflow.org/api_docs/python/tf/data/Dataset#batch) and prefetch()

(https://www.tensorflow.org/api_docs/python/tf/data/Dataset#prefetch),
the parameter tf.data.AUTOTUNE

(https://www.tensorflow.org/api docs/python/tf/data#AUTOTUNE)
also allow TensorFlow to determine the optimal amount of compute to

- In []: # Turn our data into TensorFlow Datasets
 train_dataset = tf.data.Dataset.from_tensor_slices((train_sentences, tr
 valid_dataset = tf.data.Dataset.from_tensor_slices((val_sentences, val_
 test_dataset = tf.data.Dataset.from_tensor_slices((test_sentences, test
 train_dataset
- - In []: # Take the TensorSliceDataset's and turn them into prefetched batches
 train_dataset = train_dataset.batch(32).prefetch(tf.data.AUTOTUNE)
 valid_dataset = valid_dataset.batch(32).prefetch(tf.data.AUTOTUNE)
 test_dataset = test_dataset.batch(32).prefetch(tf.data.AUTOTUNE)
 train_dataset

Model 1: Conv1D with token embeddings

Alright, we've now got a way to numerically represent our text and labels, time to build a series of deep models to try and improve upon our baseline.

All of our deep models will follow a similar structure:

Input (text) -> Tokenize -> Embedding -> Layers -> Output (labe
l probability)

The main component we'll be changing throughout is the Layers component. Because any modern deep NLP model requires text to be converted into an embedding before meaningful patterns can be discovered within.

The first model we're going to build is a 1-dimensional Convolutional Neural Network.

We're also going to be following the standard machine learning workflow of:

- Build model
- Train model

In []: # Get summary of Conv1D model model_1.summary()

Model: "model"

Layer (type)	Output Shape	Param #
<pre>input_1 (InputLayer)</pre>	[(None, 1)]	0
text_vectorization (TextVect	(None, 55)	0
token_embedding (Embedding)	(None, 55, 128)	8299648
conv1d (Conv1D)	(None, 55, 64)	41024
global_average_pooling1d (Gl	(None, 64)	0
dense (Dense)	(None, 5)	325

Total params: 8,340,997 Trainable params: 8,340,997 Non-trainable params: 0

Wonderful! We've got our first deep sequence model built and ready to go.

Checking out the model summary, you'll notice the majority of the trainable parameters are within the embedding layer. If we were to increase the size of the embedding (by increasing the output_dim parameter of the Embedding layer), the number of trainable parameters would increase dramatically.

It's time to fit our model to the training data but we're going to make a mindful change.

Since our training data contains nearly 200,000 sentences, fitting a deep model may take a while even with a GPU. So to keep our experiments swift, we're going to run them on a subset of the training dataset.

More specifically, we'll only use the first 10% of batches (about 18,000 samples) of the training set to train on and the first 10% of batches from the validation set to validate on.

□ **Note:** It's a standard practice in machine learning to test your models on smaller subsets of data first to make sure they work before scaling them to larger amounts of data. You should aim to run many smaller experiments rather than only a handful of large experiments. And since your time is limited, one of the best ways to run smaller experiments is to reduce the amount of data you're working with (10% of the full dataset is usually a good amount, as long as it covers a similar distribution).

```
In [ ]:
     # Fit the model
     model_1_history = model_1.fit(train_dataset,
                         steps_per_epoch=int(0.1 * len(train_datas
                         epochs=3,
                         validation_data=valid_dataset,
                         validation_steps=int(0.1 * len(valid_data
     Epoch 1/3
     accuracy: 0.6280 - val_loss: 0.6956 - val_accuracy: 0.7350
     Epoch 2/3
     - accuracy: 0.7528 - val_loss: 0.6353 - val_accuracy: 0.7666
     Epoch 3/3
     - accuracy: 0.7742 - val_loss: 0.6017 - val_accuracy: 0.7839
```

Brilliant! We've got our first trained deep sequence model, and it didn't take too long (and if we didn't prefetch our batched data, it would've taken longer).

Time to make some predictions with our model and then evaluate them.

```
In [ ]: # Evaluate on whole validation dataset (we only validated on 10% of bat
         model_1.evaluate(valid_dataset)
         - accuracy: 0.7845
Out[42]: [0.6039669513702393, 0.784489631652832]
 In [ ]: # Make predictions (our model outputs prediction probabilities for each
        model_1_pred_probs = model_1.predict(valid_dataset)
         model_1_pred_probs
Out[43]: array([[4.65810180e-01, 1.68030873e-01, 9.11973268e-02, 2.48495579e-0
         1,
                2.64659580e-021,
                [3.85841161e-01, 3.26967925e-01, 1.09655075e-02, 2.69699097e-0
         1,
                6.52625971e-03],
                [1.50063470e-01, 1.18270982e-02, 1.80714345e-03, 8.36278021e-0
         1,
                2.42987626e-051,
                [5.72754607e-06, 8.22585251e-04, 5.39675879e-04, 1.08408301e-0
         6,
                9.98630941e-011,
                [5.39277196e-02, 4.82838809e-01, 9.59893093e-02, 5.87528460e-0
         2,
                3.08491379e-011,
                [2.05477521e-01, 5.33861935e-01, 5.17199412e-02, 8.40113238e-0
         2,
                1.24929301e-01]], dtype=float32)
 In [ ]:
        # Convert pred probs to classes
        model_1_preds = tf.argmax(model_1_pred_probs, axis=1)
        model_1_preds
Out[44]: <tf.Tensor: shape=(30212,), dtype=int64, numpy=array([0, 0, 3, ..., 4,</pre>
         1, 1])>
        # Calculate model_1 results
 In [ ]:
         model_1_results = calculate_results(y_true=val_labels_encoded,
                                           y_pred=model_1_preds)
        model 1 results
Out[45]: {'accuracy': 78.44896067787634,
          'f1': 0.7822985872046467,
          'precision': 0.7814941086130403,
          'recall': 0.7844896067787634}
```

Model 2: Feature extraction with pretrained token embeddings

Training our own embeddings took a little while to run, slowing our experiments down.

Since we're moving towards replicating the model architecture in <u>Neural Networks for Joint Sentence Classification in Medical Paper Abstracts (https://arxiv.org/pdf/1612.05251.pdf)</u>, it mentions they used a <u>pretrained GloVe embedding (https://nlp.stanford.edu/projects/glove/)</u> as a way to initialise their token embeddings.

To emulate this, let's see what results we can get with the <u>pretrained Universal Sentence Encoder embeddings from TensorFlow Hub (https://tfhub.dev/google/universal-sentence-encoder/4)</u>.

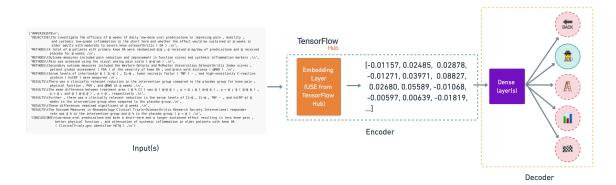
□ **Note:** We could use GloVe embeddings as per the paper but since we're working with TensorFlow, we'll use what's available from TensorFlow Hub (GloVe embeddings aren't). We'll save <u>using pretrained GloVe embeddings</u> (https://keras.io/examples/nlp/pretrained word embeddings/) as an extension.

The model structure will look like:

Inputs (string) -> Pretrained embeddings from TensorFlow Hub (U
niversal Sentence Encoder) -> Layers -> Output (prediction prob
abilities)

You'll notice the lack of tokenization layer we've used in a previous model. This is because the Universal Sentence Encoder (USE) takes care of tokenization for us.

This type of model is called transfer learning, or more specifically, **feature extraction transfer learning**. In other words, taking the patterns a model has learned elsewhere and applying it to our own problem.



The feature extractor model we're building using a pretrained embedding from TensorFlow Hub.

To download the pretrained USE into a layer we can use in our model, we can use the https://www.tensorflow.org/hub/api_docs/python/hub/KerasLayer) class.

We'll keep the pretrained embeddings frozen (by setting trainable=False) and add a trainable couple of layers on the top to tailor the model outputs to our own data.

 \square **Note:** Due to having to download a relatively large model (~916MB), the cell below may take a little while to run.

Beautiful, now our pretrained USE is downloaded and instantiated as a hub.KerasLayer instance, let's test it out on a random sentence.

```
In [ ]: # Test out the embedding on a random sentence
    random_training_sentence = random.choice(train_sentences)
    print(f"Random training sentence:\n{random_training_sentence}\n")
    use_embedded_sentence = tf_hub_embedding_layer([random_training_sentence])
    print(f"Sentence after embedding:\n{use_embedded_sentence[0][:30]} (truprint(f"Length of sentence embedding:\n{len(use_embedded_sentence[0])}"
```

Random training sentence:

data were collected from @,@ @th - and @th-grade students in these communities using anonymous cross-sectional surveys in @ and @ and analyzed in @.

```
Sentence after embedding:
[-0.03330918  0.06432742 -0.04426299 -0.03340627  0.01580565  0.042607 87
  0.05223562  0.02998829 -0.09009711  0.01004721 -0.01640431 -0.029705 98
  0.06300306  0.0500071 -0.02137795  0.01532154  0.00676913  0.041741 87
  0.00070422 -0.08601078  0.02839869  0.07599191 -0.05524107 -0.008624 89
  -0.04704431  0.05998407 -0.00028414 -0.01645767  0.05340267 -0.064738 02] (truncated output)...
```

Length of sentence embedding: 512

Nice! As we mentioned before the pretrained USE module from TensorFlow Hub takes care of tokenizing our text for us and outputs a 512 dimensional embedding vector.

Let's put together and compile a model using our tf_hub_embedding_layer.

Building and fitting an NLP feature extraction model from TensorFlow Hub

In []: # Get a summary of the model model_2.summary()

Model: "model_1"

Layer (type)	Output Shape	Param #
input_2 (InputLayer)	[(None,)]	0
universal_sentence_encoder ((None, 512)	256797824
dense_1 (Dense)	(None, 128)	65664
dense_2 (Dense)	(None, 5)	645

Total params: 256,864,133 Trainable params: 66,309

Non-trainable params: 256,797,824

Checking the summary of our model we can see there's a large number of total parameters, however, the majority of these are non-trainable. This is because we set training=False when we instatiated our USE feature extractor layer.

So when we train our model, only the top two output layers will be trained.

```
In [ ]: |# Fit feature extractor model for 3 epochs
        model_2.fit(train_dataset,
                  steps_per_epoch=int(0.1 * len(train_dataset)),
                  epochs=3,
                  validation_data=valid_dataset,
                  validation_steps=int(0.1 * len(valid_dataset)))
        Epoch 1/3
        562/562 [================ ] - 9s 12ms/step - loss: 0.9150
        - accuracy: 0.6498 - val_loss: 0.7939 - val_accuracy: 0.6895
        Epoch 2/3
        - accuracy: 0.7022 - val_loss: 0.7523 - val_accuracy: 0.7058
        Epoch 3/3
        - accuracy: 0.7127 - val_loss: 0.7367 - val_accuracy: 0.7138
Out[50]: <keras.callbacks.History at 0x7f1f82200850>
In [ ]: # Evaluate on whole validation dataset
        model_2.evaluate(valid_dataset)
        3 - accuracy: 0.7143
Out[51]: [0.7403141856193542, 0.7142525911331177]
        Since we aren't training our own custom embedding layer, training is
        much quicker.
        Let's make some predictions and evaluate our feature extraction
        model.
       # Make predictions with feature extraction model
In [ ]:
        model_2_pred_probs = model_2.predict(valid_dataset)
        model_2_pred_probs
Out[52]: array([[4.3034002e-01, 3.5648319e-01, 2.3842952e-03, 2.0301045e-01,
               7.7820425e-031,
              [3.5814181e-01, 4.8381555e-01, 3.4082821e-03, 1.5132000e-01,
               3.3143654e-03],
              [2.2659931e-01, 1.5397042e-01, 2.1248475e-02, 5.6063354e-01,
               3.7548285e-02],
              [1.7298853e-03, 5.6997794e-03, 5.2002735e-02, 8.3732005e-04,
               9.3973035e-011,
              [3.5631014e-03, 4.8854645e-02, 1.8445115e-01, 1.2047966e-03,
               7.6192635e-011,
              [1.7391451e-01, 3.0626863e-01, 4.5657125e-01, 5.7948320e-03,
               5.7450745e-02]], dtype=float32)
```

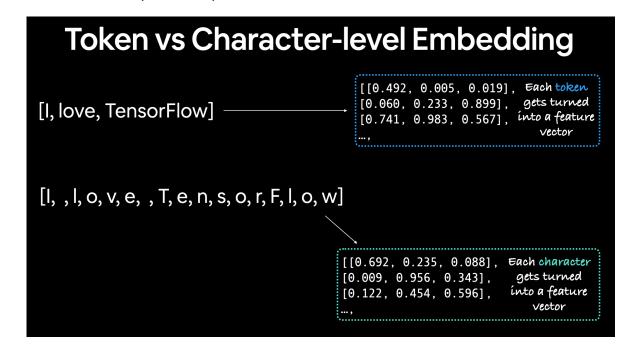
Model 3: Conv1D with character embeddings

Creating a character-level tokenizer

The <u>Neural Networks for Joint Sentence Classification in Medical</u>
<u>Paper Abstracts (https://arxiv.org/pdf/1612.05251.pdf)</u> paper mentions their model uses a hybrid of token and character embeddings.

We've built models with a custom token embedding and a pretrained token embedding, how about we build one using a character embedding?

The difference between a character and token embedding is that the **character embedding** is created using sequences split into characters (e.g. hello -> [h, e, 1, 1, o]) where as a **token embedding** is created on sequences split into tokens.



Token level embeddings split sequences into tokens (words) and embeddings each of them, character embeddings split sequences into characters and creates a feature vector for each.

We can create a character-level embedding by first vectorizing our sequences (after they've been split into characters) using the TextVectorization

(https://www.tensorflow.org/api docs/python/tf/keras/layers/experimenta
class and then passing those vectorized sequences through an
 Embedding

(https://www.tensorflow.org/api_docs/python/tf/keras/layers/Embedding)
layer.

Before we can vectorize our sequences on a character-level we'll need

```
In []: # Make function to split sentences into characters
def split_chars(text):
    return " ".join(list(text))

# Test splitting non-character-level sequence into characters
split_chars(random_training_sentence)
```

Out[55]: 'data were collected from @,@ @th - and @th-grade students in these communities using anonymous cross-sectional surveys in @ and @ and analyzed in @ .'

Great! Looks like our character-splitting function works. Let's create character-level datasets by splitting our sequence datasets into characters.

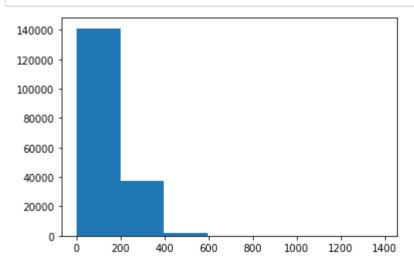
In []: # Split sequence-level data splits into character-level data splits
 train_chars = [split_chars(sentence) for sentence in train_sentences]
 val_chars = [split_chars(sentence) for sentence in val_sentences]
 test_chars = [split_chars(sentence) for sentence in test_sentences]
 print(train_chars[0])

investigate the efficacy o f of daily low-dose oral predniso lone in improving pain , mobility and systemic low-grade inflammation in the short term and whether ffect would be sustained weeks at @ older adults with moderate to se knee osteoarthritis (

To figure out how long our vectorized character sequences should be, let's check the distribution of our character sequence lengths.

Out[57]: 149.3662574983337

In []: # Check the distribution of our sequences at character-level import matplotlib.pyplot as plt plt.hist(char_lens, bins=7);



Okay, looks like most of our sequences are between 0 and 200 characters long.

Let's use NumPy's percentile to figure out what length covers 95% of our sequences.

```
In [ ]: # Find what character length covers 95% of sequences
    output_seq_char_len = int(np.percentile(char_lens, 95))
    output_seq_char_len
```

Out[59]: 290

Wonderful, now we know the sequence length which covers 95% of sequences, we'll use that in our TextVectorization layer as the output_sequence_length parameter.

□ **Note:** You can experiment here to figure out what the optimal output_sequence_length should be, perhaps using the mean results in as good results as using the 95% percentile.

We'll set max_tokens (the total number of different characters in our sequences) to 28, in other words, 26 letters of the alphabet + space + 00V (out of vocabulary or unknown) tokens.

Nice! Now we've adapted our char_vectorizer to our character-level sequences, let's check out some characteristics about it using the get_vocabulary()

(https://www.tensorflow.org/api_docs/python/tf/keras/layers/experimenta method.

4

We can also test it on random sequences of characters to make sure it's working.

```
In [ ]: # Test out character vectorizer
        random_train_chars = random.choice(train_chars)
        print(f"Charified text:\n{random_train_chars}")
        print(f"\nLength of chars: {len(random_train_chars.split())}")
        vectorized_chars = char_vectorizer([random_train_chars])
        print(f"\nVectorized chars:\n{vectorized_chars}")
        print(f"\nLength of vectorized chars: {len(vectorized_chars[0])}")
        Charified text:
            persistent
                                  question
                                                    i s
                                                          whether
        raprofessional home visitors
                                                                  might
        produce comparable effects
        Length of chars: 86
        Vectorized chars:
        [[ 5 14 2 8
                       9
                             9
                                3
                                   2
                                      6
                                         3 26 16
                                                  2
                                                     9
                                                        3
        2
                 2
                                               2
                                                                 5 12 13
           3 13
                    8 14
                          5
                                5 14
                                      8
                                         7 17
                                                  9
                                                     9
                                                           7
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        2
          21
              4
                 9
                   4
                      3
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                             8
                                9 15
                                      4 18 13
                                               3 14
                                                     8
                                                        7 10 16 11
                                                                    2 11
        14
                             2 17 17
                                      2 11
                                            3
           5
                 5 22 12
                          2
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                                0
                                                  0
        0
```

Length of vectorized chars: 290

0]]

You'll notice sequences with a length shorter than 290 (output_seq_char_length) get padded with zeros on the end, this ensures all sequences passed to our model are the same length.

Also, due to the standardize parameter of TextVectorization being "lower_and_strip_punctuation" and the split parameter being "whitespace" by default, symbols (such as @) and spaces are removed.

□ **Note:** If you didn't want punctuation to be removed (keep the @, % etc), you can create a custom standardization callable and pass it as the standardize parameter. See the <u>TextVectorization</u>

(https://www.tensorflow.org/api_docs/python/tf/keras/layers/exper

Creating a character-level embedding

We've got a way to vectorize our character-level sequences, now's time to create a character-level embedding.

Just like our custom token embedding, we can do so using the tensorflow.keras.layers.Embedding
(https://www.tensorflow.org/api_docs/python/tf/keras/layers/Embedding)
class.

Our character-level embedding layer requires an input dimension and output dimension.

The input dimension (input_dim) will be equal to the number of different characters in our char_vocab (28). And since we're following the structure of the model in Figure 1 of <u>Neural Networks for Joint Sentence Classification in Medical Paper Abstracts</u> (https://arxiv.org/pdf/1612.05251.pdf), the output dimension of the character embedding (output_dim) will be 25.

```
In [ ]: # Create char embedding layer
      char_embed = layers.Embedding(input_dim=NUM_CHAR_TOKENS, # number of di
                              output_dim=25, # embedding dimension of e
                              mask_zero=False, # don't use masks (this
                              name="char embed")
      # Test out character embedding layer
      print(f"Charified text (before vectorization and embedding):\n{random_t
      char_embed_example = char_embed(char_vectorizer([random_train_chars]))
      print(f"Embedded chars (after vectorization and embedding):\n{char_embedding}
      print(f"Character embedding shape: {char_embed_example.shape}")
      Charified text (before vectorization and embedding):
         persistent question is whether
      raprofessional home visitors
      produce comparable effects
      Embedded chars (after vectorization and embedding):
      [[[-0.01923175 -0.01720572 0.04752548 ... -0.00952251 -0.03900822
         -0.016066511
        [ 0.02746468 -0.0157405 -0.03408597 ... 0.00957409 -0.04426242
         0.025472671
        0.02328778]
        [-0.02053725  0.04947415  -0.03963646  ...  0.04780323  0.00831659
         -0.000332821
        -0.000332821
        -0.00033282111
```

Character embedding shape: (1, 290, 25)

Wonderful! Each of the characters in our sequences gets turned into a 25 dimension embedding.

Building a Conv1D model to fit on character embeddings

Now we've got a way to turn our character-level sequences into numbers (char_vectorizer) as well as numerically represent them as an embedding (char_embed) let's test how effective they are at encoding the information in our sequences by creating a character-level sequence model.

The model will have the same structure as our custom token embedding model (model_1) except it'll take character-level sequences as input instead of token-level sequences.

```
Input (character-level text) -> Tokenize -> Embedding -> Layers
(Conv1D, GlobalMaxPool1D) -> Output (label probability)
```

In []: # Check the summary of conv1d_char_model model_3.summary()

Model: "model_3_conv1D_char_embedding"

Layer (type)	Output Shape	Param #
input_3 (InputLayer)	[(None, 1)]	0
char_vectorizer (TextVectori	(None, 290)	0
char_embed (Embedding)	(None, 290, 25)	1750
conv1d_1 (Conv1D)	(None, 290, 64)	8064
global_max_pooling1d (Global	(None, 64)	0
dense_3 (Dense)	(None, 5)	325
Total params: 10 130		

Total params: 10,139 Trainable params: 10,139 Non-trainable params: 0

Before fitting our model on the data, we'll create char-level batched PrefetchedDataset's.

Just like our token-level sequence model, to save time with our experiments, we'll fit the character-level model on 10% of batches.

```
In []: # Fit the model on chars only
       model_3_history = model_3.fit(train_char_dataset,
                                 steps_per_epoch=int(0.1 * len(train_char_
                                 epochs=3,
                                 validation_data=val_char_dataset,
                                 validation_steps=int(0.1 * len(val_char_d
        Epoch 1/3
        - accuracy: 0.4940 - val_loss: 1.0555 - val_accuracy: 0.5864
        Epoch 2/3
        - accuracy: 0.5996 - val_loss: 0.9542 - val_accuracy: 0.6267
        Epoch 3/3
        - accuracy: 0.6410 - val_loss: 0.8712 - val_accuracy: 0.6722
In []: # Evaluate model 3 on whole validation char dataset
       model_3.evaluate(val_char_dataset)
       - accuracy: 0.6588
Out[69]: [0.8873457908630371, 0.6587779521942139]
       Nice! Looks like our character-level model is working, let's make
       some predictions with it and evaluate them.
In []: # Make predictions with character model only
       model_3_pred_probs = model_3.predict(val_char_dataset)
       model_3_pred_probs
Out[70]: array([[0.14757556, 0.40572175, 0.06463172, 0.33147973, 0.05059117],
              [0.33927342, 0.25103244, 0.01815611, 0.35482603, 0.03671199],
              [0.15584646, 0.11993653, 0.10841523, 0.5846738, 0.03112798],
              [0.02389161, 0.04591153, 0.07370146, 0.01870138, 0.83779407],
              [0.01393123, 0.10044182, 0.55801105, 0.05441127, 0.2732046],
              [0.28039306, 0.582124 , 0.04549843, 0.04313584, 0.04884864]],
             dtype=float32)
In [ ]: # Convert predictions to classes
       model_3_preds = tf.argmax(model_3_pred_probs, axis=1)
       model_3_preds
Out[71]: <tf.Tensor: shape=(30212,), dtype=int64, numpy=array([1, 3, 3, ..., 4,</pre>
       2, 1])>
```

'precision': 0.6545009464773592, 'recall': 0.6587779690189329}

Model 4: Combining pretrained token embeddings + character embeddings (hybrid embedding layer)

Alright, now things are going to get spicy.

In moving closer to build a model similar to the one in Figure 1 of <u>Neural Networks for Joint Sentence Classification in Medical Paper Abstracts (https://arxiv.org/pdf/1612.05251.pdf)</u>, it's time we tackled the hybrid token embedding layer they speak of.

This hybrid token embedding layer is a combination of token embeddings and character embeddings. In other words, they create a stacked embedding to represent sequences before passing them to the sequence label prediction layer.

So far we've built two models which have used token and characterlevel embeddings, however, these two models have used each of these embeddings exclusively.

To start replicating (or getting close to replicating) the model in Figure 1, we're going to go through the following steps:

- Create a token-level model (similar to model_1)
- 2. Create a character-level model (similar to model_3 with a slight modification to reflect the paper)
- 4. Build a series of output layers on top of 3 similar to Figure 1 and section 4.2 of <u>Neural Networks for Joint Sentence</u> <u>Classification in Medical Paper Abstracts</u> (https://arxiv.org/pdf/1612.05251.pdf)
- 5. Construct a model which takes token and character-level sequences as input and produces sequence label probabilities as output

 \triangleleft

```
In [ ]: # 1. Setup token inputs/model
        token_inputs = layers.Input(shape=[], dtype=tf.string, name="token_inpu")
        token_embeddings = tf_hub_embedding_layer(token_inputs)
        token output = layers.Dense(128, activation="relu")(token embeddings)
        token_model = tf.keras.Model(inputs=token_inputs,
                                      outputs=token_output)
        # 2. Setup char inputs/model
        char_inputs = layers.Input(shape=(1,), dtype=tf.string, name="char_inpu")
        char_vectors = char_vectorizer(char_inputs)
        char_embeddings = char_embed(char_vectors)
        char_bi_lstm = layers.Bidirectional(layers.LSTM(25))(char_embeddings) #
        char_model = tf.keras.Model(inputs=char_inputs,
                                     outputs=char bi lstm)
        # 3. Concatenate token and char inputs (create hybrid token embedding)
        token_char_concat = layers.Concatenate(name="token_char_hybrid")([token_char_hybrid"))
                                                                            char
        # 4. Create output layers - addition of dropout discussed in 4.2 of htt
        combined_dropout = layers.Dropout(0.5)(token_char_concat)
        combined_dense = layers.Dense(200, activation="relu")(combined_dropout)
        final_dropout = layers.Dropout(0.5)(combined_dense)
        output_layer = layers.Dense(num_classes, activation="softmax")(final_dr
        # 5. Construct model with char and token inputs
        model_4 = tf.keras.Model(inputs=[token_model.input, char_model.input],
                                  outputs=output_layer,
                                  name="model_4_token_and_char_embeddings")
```

Woah... There's a lot going on here, let's get a summary and plot our model to visualize what's happening.

In []: # Get summary of token and character model
 model_4.summary()

Layer (type) cted to	Output	Shape	Param #	Conne
char_input (InputLayer)	[(None	, 1)]	0	
token_input (InputLayer)	[(None	,)]	0	
<pre>char_vectorizer (TextVectorizat input[0][0]</pre>	(None,	290)	0	char_
universal_sentence_encoder (Ker _input[0][0]	(None,	512)	256797824	token
char_embed (Embedding) vectorizer[1][0]	(None,	290, 25)	1750	char_
dense_4 (Dense) rsal_sentence_encoder[1][0]	(None,	128)	65664	unive
bidirectional (Bidirectional) embed[1][0]	(None,	50)	10200	char_
token_char_hybrid (Concatenate) _4[0][0]	(None,	178)	0	dense bidir
ectional[0][0]				
dropout (Dropout) _char_hybrid[0][0]	(None,	178)	0	token
dense_5 (Dense) ut[0][0]	(None,	200)	35800	dropo
dropout_1 (Dropout) _5[0][0]	(None,	200)	0	dense
dense_6 (Dense) ut_1[0][0]	(None,	5)	1005	dropo
	=====:	========	========	=====

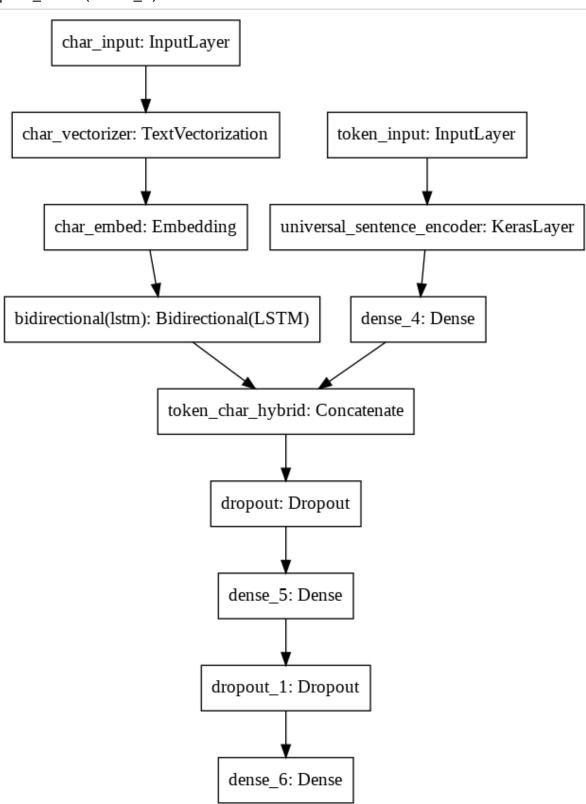
Total params: 256,912,243

Trainable params: 114,419

In []: # Plot hybrid token and character model

from tensorflow.keras.utils import plot_model
plot_model(model_4)

Out[75]:



Now that's a good looking model. Let's compile it just as we have the rest of our models.

```
Note: Section 4.2 of Neural Networks for Joint Sentence
Classification in Medical Paper Abstracts
(https://arxiv.org/pdf/1612.05251.pdf) mentions using the
SGD (stochastic gradient descent) optimizer, however, to
stay consistent with our other models, we're going to use
the Adam optimizer. As an exercise, you could try using
tf.keras.optimizers.SGD
(https://www.tensorflow.org/api docs/python/tf/keras/optimizers/S
instead of tf.keras.optimizers.Adam
(https://www.tensorflow.org/api docs/python/tf/keras/optimizers/A
and compare the results.
```

And again, to keep our experiments fast, we'll fit our token-character-hybrid model on 10% of training and validate on 10% of validation batches. However, the difference with this model is that it requires two inputs, token-level sequences and character-level sequences.

We can do this by create a tf.data.Dataset with a tuple as it's first input, for example:

• ((token_data, char_data), (label))

In []: |# Combine chars and tokens into a dataset

f.string, tf.string), tf.float64)>)

Let's see it in action.

Combining token and character data into a tf.data dataset

Fitting a model on token and character-level sequences

```
# Fit the model on tokens and chars
In [ ]:
        model_4_history = model_4.fit(train_char_token_dataset, # train on data
                                  steps_per_epoch=int(0.1 * len(train_char_
                                  epochs=3,
                                  validation_data=val_char_token_dataset,
                                  validation steps=int(0.1 * len(val char t
        Epoch 1/3
        4 - accuracy: 0.6159 - val_loss: 0.7859 - val_accuracy: 0.6898
        4 - accuracy: 0.6958 - val_loss: 0.7139 - val_accuracy: 0.7301
        Epoch 3/3
        9 - accuracy: 0.7057 - val_loss: 0.6826 - val_accuracy: 0.7410
In [ ]: |# Evaluate on the whole validation dataset
        model_4.evaluate(val_char_token_dataset)
        945/945 [============== ] - 20s 21ms/step - loss: 0.689
        9 - accuracy: 0.7362
Out[80]: [0.6899493336677551, 0.7362306118011475]
        Nice! Our token-character hybrid model has come to life!
        To make predictions with it, since it takes multiplie inputs, we can
        pass the predict() method a tuple of token-level sequences and
        character-level sequences.
        We can then evaluate the predictions as we've done before.
In [ ]:
        # Make predictions using the token-character model hybrid
        model_4_pred_probs = model_4.predict(val_char_token_dataset)
        model_4_pred_probs
Out[81]: array([[4.5224771e-01, 3.3035564e-01, 2.7360097e-03, 2.0863818e-01,
               6.0224836e-031,
              [2.7638477e-01, 5.4655772e-01, 3.0270391e-03, 1.7166287e-01,
               2.3676162e-03],
              [3.1192392e-01, 1.4092967e-01, 4.3908428e-02, 4.5166326e-01,
               5.1574774e-021,
              [6.1459269e-04, 5.8821058e-03, 4.2544805e-02, 1.3780949e-04,
               9.5082068e-011,
              [9.3976045e-03, 7.2958224e-02, 1.8798770e-01, 4.0314477e-03,
               7.2562498e-01],
              [2.6598454e-01, 3.5141158e-01, 2.8458366e-01, 3.5951469e-02,
               6.2068779e-02]], dtype=float32)
```

Model 5: Transfer Learning with pretrained token embeddings + character embeddings + positional embeddings

It seems like combining token embeddings and character embeddings gave our model a little performance boost.

But there's one more piece of the puzzle we can add in.

What if we engineered our own features into the model?

Meaning, what if we took our own knowledge about the data and encoded it in a numerical way to give our model more information about our samples?

The process of applying your own knowledge to build features as input to a model is called **feature engineering**.

Can you think of something important about the sequences we're trying to classify?

If you were to look at an abstract, would you expect the sentences to appear in order? Or does it make sense if they were to appear sequentially? For example, sequences labelled CONCLUSIONS at the beggining and sequences labelled OBJECTIVE at the end?

Abstracts typically come in a sequential order, such as:

```
OBJECTIVE ...METHODS ...METHODS ...METHODS ...RESULTS ...
```

• CONCLUSIONS ...

- BACKGROUND ...
- OBJECTIVE ...
- METHODS ...
- METHODS ...
- RESULTS ...
- RESULTS ...
- CONCLUSIONS ...
- CONCLUSIONS ...

Of course, we can't engineer the sequence labels themselves into the training data (we don't have these at test time), but we can encode the order of a set of sequences in an abstract.

For example,

- Sentence 1 of 10 ...
- Sentence 2 of 10 ...
- Sentence 3 of 10 ...
- Sentence 4 of 10 ...
- ...

You might've noticed this when we created our preprocess_text_with_line_numbers() function. When we read in a text file of abstracts, we counted the number of lines in an abstract as well as the number of each line itself.

Doing this led to the "line_number" and "total_lines" columns of our DataFrames.

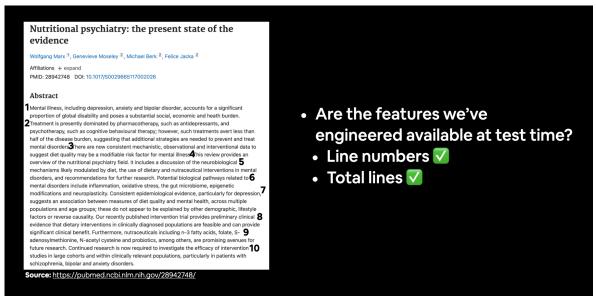
In []: # Inspect training dataframe train_df.head()

Out[84]:		target	text	line_number	total_lines
	0	OBJECTIVE	to investigate the efficacy of @ weeks of dail	0	11
	1	METHODS	a total of @ patients with primary knee oa wer	1	11
	2	METHODS	outcome measures included pain reduction and i	2	11
	3	METHODS	pain was assessed using the visual analog pain	3	11
	4	METHODS	secondary outcome measures included the wester	4	11

The "line_number" and "total_lines" columns are features which didn't necessarily come with the training data but can be passed to our model as a **positional embedding**. In other words, the positional

embedding is where the sentence appears in an abstract.

We can use these features because they will be available at test time.



Since abstracts typically have a sequential order about them (for example, background, objective, methods, results, conclusion), it makes sense to add the line number of where a particular sentence occurs to our model. The beautiful thing is, these features will be available at test time (we can just count the number of sentences in an abstract and the number of each one).

Meaning, if we were to predict the labels of sequences in an abstract our model had never seen, we could count the number of lines and the track the position of each individual line and pass it to our model.

Create positional embeddings

Okay, enough talk about positional embeddings, let's create them.

Since our "line_number" and "total_line" columns are already numerical, we could pass them as they are to our model.

But to avoid our model thinking a line with "line_number"=5 is five times greater than a line with "line_number"=1, we'll use one-hot-encoding to encode our "line_number" and "total_lines" features.

To do this, we can use the tf.one_hot
hot) utility.

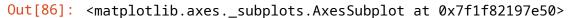
tf.one_hot returns a one-hot-encoded tensor. It accepts an array (or tensor) as input and the depth parameter determines the dimension of the returned tensor.

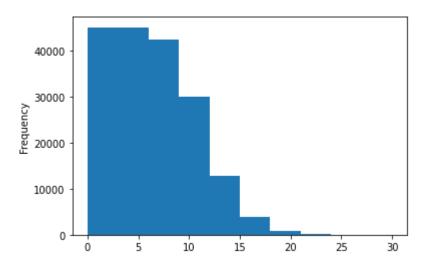
To figure out what we should set the depth parameter to, let's investigate the distribution of the "line_number" column.

□ **Note:** When it comes to one-hot-encoding our features,
Scikit-Learn's <u>OneHotEncoder (https://scikit-learn.org/stable/modules/generated/sklearn.preprocessing.OneHotEnclass</u> is another viable option here.

```
In [ ]: # How many different line numbers are there?
          train_df["line_number"].value_counts()
Out[85]:
          0
                15000
          1
                15000
          2
                15000
          3
                15000
          4
                14992
          5
                14949
          6
                14758
          7
                14279
          8
                13346
          9
                11981
          10
                10041
          11
                 7892
          12
                 5853
          13
                 4152
          14
                 2835
          15
                 1861
                 1188
          16
          17
                   751
          18
                   462
          19
                   286
          20
                   162
          21
                   101
          22
                    66
          23
                    33
          24
                    22
          25
                    14
          26
                     7
          27
                     4
          28
                     3
          29
                     1
          30
                     1
          Name: line_number, dtype: int64
```

```
In [ ]: # Check the distribution of "line_number" column
train_df.line_number.plot.hist()
```





Looking at the distribution of the "line_number" column, it looks like the majority of lines have a position of 15 or less.

Knowing this, let's set the depth parameter of tf.one_hot to 15.

In []: # Use TensorFlow to create one-hot-encoded tensors of our "line_number"
 train_line_numbers_one_hot = tf.one_hot(train_df["line_number"].to_nump
 val_line_numbers_one_hot = tf.one_hot(val_df["line_number"].to_numpy(),
 test_line_numbers_one_hot = tf.one_hot(test_df["line_number"].to_numpy()

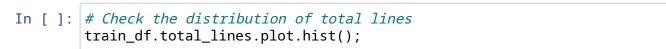
Setting the depth parameter of tf.one_hot to 15 means any sample with a "line_number" value of over 15 gets set to a tensor of all 0's, where as any sample with a "line_number" of under 15 gets turned into a tensor of all 0's but with a 1 at the index equal to the "line_number" value.

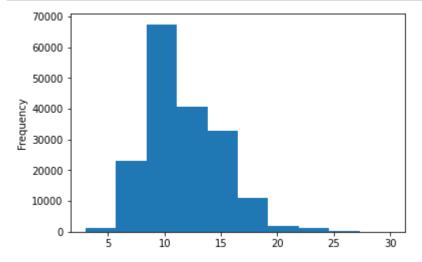
□ **Note:** We could create a one-hot tensor which has room for all of the potential values of "line_number" (depth=30), however, this would end up in a tensor of double the size of our current one (depth=15) where the vast majority of values are 0. Plus, only ~2,000/180,000 samples have a "line_number" value of over 15. So we would not be gaining much information about our data for doubling our feature space. This kind of problem is called the **curse of dimensionality**. However, since this we're working with deep models, it might be worth trying to throw as much information at the model as possible and seeing what happens. I'll leave exploring values of the depth parameter as an extension.

```
In [ ]: # Check one-hot encoded "line_number" feature samples
    train_line_numbers_one_hot.shape, train_line_numbers_one_hot[:20]
Out[88]:
    (TensorShape([180040, 15]), <tf.Tensor: shape=(20, 15), dtype=float32,
    numpy=
     [0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0., 0., 0., 0., 0.]
        [0., 0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0., 0., 0., 0.]
        [0., 0., 0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0., 0., 0.]
        [0., 0., 0., 0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0., 0.]
        [0., 0., 0., 0., 0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0.]
        [0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0., 0., 0., 0., 0.]
        [0., 0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0., 0., 0., 0.]
        [0., 0., 0., 0., 0., 0., 0., 1., 0., 0., 0., 0., 0., 0., 0.]],
        dtype=float32)>)
```

We can do the same as we've done for our "line_number" column witht he "total_lines" column. First, let's find an appropriate value for the depth parameter of tf.one_hot.

In []: # How many different numbers of lines are there? train_df["total_lines"].value_counts() Out[89]: 11 Name: total_lines, dtype: int64





Looking at the distribution of our "total_lines" column, a value of 20 looks like it covers the majority of samples.

```
In [ ]: # Check the coverage of a "total_lines" value of 20
     np.percentile(train_df.total_lines, 98) # a value of 20 covers 98% of s
Out[91]: 20.0
     4
     Beautiful! Plenty of converage. Let's one-hot-encode our
      "total lines" column just as we did our "line_number" column.
In [ ]: # Use TensorFlow to create one-hot-encoded tensors of our "total_lines"
     train_total_lines_one_hot = tf.one_hot(train_df["total_lines"].to_numpy
     val_total_lines_one_hot = tf.one_hot(val_df["total_lines"].to_numpy(),
     test_total_lines_one_hot = tf.one_hot(test_df["total_lines"].to_numpy()
     # Check shape and samples of total lines one-hot tensor
     train_total_lines_one_hot.shape, train_total_lines_one_hot[:10]
Out[92]: (TensorShape([180040, 20]), <tf.Tensor: shape=(10, 20), dtype=float32,</pre>
     numpy=
      0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.],
           0.,
           0., 0., 0., 0.1,
           0.,
           0., 0., 0., 0.]], dtype=float32)>)
```

Building a tribrid embedding model

Woohoo! Positional embedding tensors ready.

It's time to build the biggest model we've built yet. One which incorporates token embeddings, character embeddings and our newly crafted positional embeddings.

We'll be venturing into uncovered territory but there will be nothing here you haven't practiced before.

More specifically we're going to go through the following steps:

- Create a token-level model (similar to model_1)
- Create a character-level model (similar to model_3 with a slight modification to reflect the paper)
- 3. Create a "line_number" model (takes in one-hot-encoded
 "line_number" tensor and passes it through a non-linear layer)
- 4. Create a "total_lines" model (takes in one-hot-encoded
 "total_lines" tensor and passes it through a non-linear layer)
- 5. Combine (using <u>layers.Concatenate</u> (https://www.tensorflow.org/api docs/python/tf/keras/layers/Concater the outputs of 1 and 2 into a token-character-hybrid embedding and pass it series of output to Figure 1 and section 4.2 of Networks for Joint Sentence Classification in Medical Paper Abstracts (https://arxiv.org/pdf/1612.05251.pdf)
- 6. Combine (using <u>layers.Concatenate</u> (https://www.tensorflow.org/api_docs/python/tf/keras/layers/Concater the outputs of 3, 4 and 5 into a token-character-positional tribrid embedding
- 7. Create an output layer to accept the tribrid embedding and output predicted label probabilities
- 8. Combine the inputs of 1, 2, 3, 4 and outputs of 7 into a tf.keras.Model (https://www.tensorflow.org/api_docs/python/tf/keras/Model)

Woah! That's alot... but nothing we're not capable of. Let's code it.

```
In [ ]: # 1. Token inputs
        token_inputs = layers.Input(shape=[], dtype="string", name="token_input"
        token_embeddings = tf_hub_embedding_layer(token_inputs)
        token outputs = layers.Dense(128, activation="relu")(token embeddings)
        token_model = tf.keras.Model(inputs=token_inputs,
                                     outputs=token_outputs)
        # 2. Char inputs
        char_inputs = layers.Input(shape=(1,), dtype="string", name="char_input")
        char_vectors = char_vectorizer(char_inputs)
        char_embeddings = char_embed(char_vectors)
        char_bi_lstm = layers.Bidirectional(layers.LSTM(32))(char_embeddings)
        char model = tf.keras.Model(inputs=char_inputs,
                                    outputs=char_bi_lstm)
        # 3. Line numbers inputs
        line_number_inputs = layers.Input(shape=(15,), dtype=tf.int32, name="li")
        x = layers.Dense(32, activation="relu")(line_number_inputs)
        line number_model = tf.keras.Model(inputs=line_number_inputs,
                                            outputs=x)
        # 4. Total lines inputs
        total_lines_inputs = layers.Input(shape=(20,), dtype=tf.int32, name="to
        y = layers.Dense(32, activation="relu")(total_lines_inputs)
        total_line_model = tf.keras.Model(inputs=total_lines_inputs,
                                           outputs=v)
        # 5. Combine token and char embeddings into a hybrid embedding
        combined_embeddings = layers.Concatenate(name="token_char_hybrid_embedd")
        z = layers.Dense(256, activation="relu")(combined_embeddings)
        z = layers.Dropout(0.5)(z)
        # 6. Combine positional embeddings with combined token and char embeddi
        z = layers.Concatenate(name="token_char_positional_embedding")([line_nu
                                                                         total 1
                                                                         z])
        # 7. Create output layer
        output_layer = layers.Dense(5, activation="softmax", name="output_layer
        # 8. Put together model
        model 5 = tf.keras.Model(inputs=[line_number_model.input,
                                          total_line_model.input,
                                          token model.input,
                                          char_model.input],
                                  outputs=output_layer)
```

There's a lot going on here... let's visualize what's happening with a summary by plotting our model.

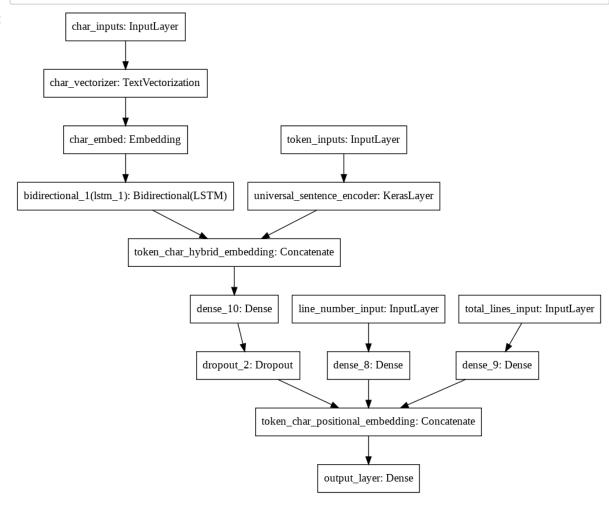
In []: # Get a summary of our token, char and positional embedding model
model_5.summary()

Layer (type) cted to	Output Shape	Param #	Conne
char_inputs (InputLayer)	[(None, 1)]	0	
<pre>char_vectorizer (TextVectorizat inputs[0][0]</pre>	(None, 290)	0	char_
token_inputs (InputLayer)	[(None,)]	0	
char_embed (Embedding) vectorizer[2][0]	(None, 290, 25)	1750	char_
universal_sentence_encoder (Ker _inputs[0][0]	(None, 512)	256797824	token
bidirectional_1 (Bidirectional) embed[2][0]	(None, 64)	14848	char_
token_char_hybrid_embedding (Corsal_sentence_encoder[2][0]	(None, 576)	0	unive bidir
ectional_1[0][0]			
line_number_input (InputLayer)	[(None, 15)]	0	
total_lines_input (InputLayer)	[(None, 20)]	0	
dense_10 (Dense) _char_hybrid_embedding[0][0]	(None, 256)	147712	token
dense_8 (Dense) number_input[0][0]	(None, 32)	512	line_
dense_9 (Dense) _lines_input[0][0]	(None, 32)	672	total
dropout_2 (Dropout) _10[0][0]	(None, 256)	0	dense

```
token_char_positional_embedding (None, 320)
                                                                   dense
_8[0][0]
                                                                   dense
_9[0][0]
                                                                   dropo
ut_2[0][0]
output_layer (Dense)
                                 (None, 5)
                                                       1605
                                                                   token
_char_positional_embedding[0
Total params: 256,964,923
Trainable params: 167,099
Non-trainable params: 256,797,824
from tensorflow.keras.utils import plot_model
```

In []: # Plot the token, char, positional embedding model plot_model(model_5)

Out [95]:



Visualizing the model makes it much easier to understand.

Essentially what we're doing is trying to encode as much information about our sequences as possible into various embeddings (the inputs to our model) so our model has the best chance to figure out what label belongs to a sequence (the outputs of our model).

You'll notice our model is looking very similar to the model shown in Figure 1 of <u>Neural Networks for Joint Sentence Classification in Medical Paper Abstracts (https://arxiv.org/pdf/1612.05251.pdf)</u>. However, a few differences still remain:

- We're using pretrained TensorFlow Hub token embeddings instead of GloVe emebddings.
- We're using a Dense layer on top of our token-character hybrid embeddings instead of a bi-LSTM layer.
- Section 3.1.3 of the paper mentions a label sequence optimization layer (which helps to make sure sequence labels come out in a respectable order) but it isn't shown in Figure 1. To makeup for the lack of this layer in our model, we've created the positional embeddings layers.
- Section 4.2 of the paper mentions the token and character embeddings are updated during training, our pretrained TensorFlow Hub embeddings remain frozen.
- The paper uses the <u>SGD</u>
 (https://www.tensorflow.org/api docs/python/tf/keras/optimizers/SGD)
 optimizer, we're going to stick with <u>Adam</u>
 (https://www.tensorflow.org/api docs/python/tf/keras/optimizers/Adam

```
In [ ]: # Check which layers of our model are trainable or not
for layer in model_5.layers:
    print(layer, layer.trainable)
```

<keras.engine.input_layer.InputLayer object at 0x7f1fa3d725d0> True <keras.layers.preprocessing.text_vectorization.TextVectorization objec</pre> t at 0x7f1ecec2d0d0> True <keras.engine.input layer.InputLayer object at 0x7f1fa3d22210> True <keras.layers.embeddings.Embedding object at 0x7f1ececbc8d0> True <tensorflow_hub.keras_layer.KerasLayer object at 0x7f1fa586e5d0> False <keras.layers.wrappers.Bidirectional object at 0x7f1fa3dff090> True <keras.layers.merge.Concatenate object at 0x7f1fb14effd0> True <keras.engine.input_layer.InputLayer object at 0x7f1fa3ec1890> True <keras.engine.input layer.InputLayer object at 0x7f1fa3ef9610> True <keras.layers.core.Dense object at 0x7f1fb14efbd0> True <keras.layers.core.Dense object at 0x7f1fa3d9b110> True <keras.layers.core.Dense object at 0x7f1fb14e4f10> True <keras.layers.core.Dropout object at 0x7f1fb14efed0> True <keras.layers.merge.Concatenate object at 0x7f1fb14e49d0> True <keras.layers.core.Dense object at 0x7f1fb14f3790> True

Now our model is constructed, let's compile it.

This time, we're going to introduce a new parameter to our loss function called label_smoothing. Label smoothing helps to regularize our model (prevent overfitting) by making sure it doesn't get too focused on applying one particular label to a sample.

For example, instead of having an output prediction of:

• [0.0, 0.0, 1.0, 0.0, 0.0] for a sample (the model is very confident the right label is index 2).

It's predictions will get smoothed to be something like:

• [0.01, 0.01, 0.096, 0.01, 0.01] giving a small activation to each of the other labels, in turn, hopefully improving generalization.

```
□ Resource: For more on label smoothing, see the great blog post by PyImageSearch, <u>Label smoothing with Keras</u>, <u>TensorFlow</u>, <u>and Deep Learning</u> (<a href="https://www.pyimagesearch.com/2019/12/30/label-smoothing-with-keras-tensorflow-and-deep-learning/">https://www.pyimagesearch.com/2019/12/30/label-smoothing-with-keras-tensorflow-and-deep-learning/</a>)
```

Create tribrid embedding datasets and fit tribrid model

Model compiled!

Again, to keep our experiments swift, let's fit on 20,000 examples for 3 epochs.

This time our model requires four feature inputs:

- Train line numbers one-hot tensor (train_line_numbers_one_hot)
- Train total lines one-hot tensor (train_total_lines_one_hot)
- Token-level sequences tensor (train_sentences)
- 4. Char-level sequences tensor (train_chars)

We can pass these as tuples to our tf.data.Dataset.from_tensor_slices() method to create appropriately shaped and batched PrefetchedDataset 's.

```
In [ ]: # Create training and validation datasets (all four kinds of inputs)
        train_pos_char_token_data = tf.data.Dataset.from_tensor_slices((train_l
                                                                 train t
                                                                 train s
                                                                 train_d
        train_pos_char_token_labels = tf.data.Dataset.from_tensor_slices(train_
        train_pos_char_token_dataset = tf.data.Dataset.zip((train_pos_char_toke
        train_pos_char_token_dataset = train_pos_char_token_dataset.batch(32).p
        # Validation dataset
        val_pos_char_token_data = tf.data.Dataset.from_tensor_slices((val_line))
                                                                val_total
                                                                val_sente
                                                                val chars
        val_pos_char_token_labels = tf.data.Dataset.from_tensor_slices(val_labe
        val_pos_char_token_dataset = tf.data.Dataset.zip((val_pos_char_token_da
        val_pos_char_token_dataset = val_pos_char_token_dataset.batch(32).prefe
        # Check input shapes
        train_pos_char_token_dataset, val_pos_char_token_dataset
Out[98]: (<PrefetchDataset shapes: (((None, 15), (None, 20), (None,)),</pre>
        (None, 5)), types: ((tf.float32, tf.float32, tf.string, tf.string), t
        f.float64)>,
         <PrefetchDataset shapes: (((None, 15), (None, 20), (None,), (None,)),
        (None, 5)), types: ((tf.float32, tf.float32, tf.string, tf.string), t
        f.float64)>)
        # Fit the token, char and positional embedding model
In [ ]:
        history_model_5 = model_5.fit(train_pos_char_token_dataset,
                                   steps_per_epoch=int(0.1 * len(train_pos_d
                                   epochs=3,
                                   validation_data=val_pos_char_token_datase
                                   validation_steps=int(0.1 * len(val_pos_ch
        Epoch 1/3
        3 - accuracy: 0.7260 - val_loss: 0.9930 - val_accuracy: 0.8002
        1 - accuracy: 0.8114 - val_loss: 0.9606 - val_accuracy: 0.8268
        Epoch 3/3
        7 - accuracy: 0.8180 - val_loss: 0.9493 - val_accuracy: 0.8271
        Tribrid model trained! Time to make some predictions with it and
```

evaluate them just as we've done before.

```
In []: # Make predictions with token-char-positional hybrid model
          model_5_pred_probs = model_5.predict(val_pos_char_token_dataset, verbos
          model_5_pred_probs
          945/945 [========== ] - 20s 20ms/step
Out[100]: array([[0.51536554, 0.10340027, 0.01223736, 0.34324795, 0.02574881],
                 [0.5037048 , 0.1263607 , 0.0476622 , 0.3120701 , 0.01020223],
                 [0.31137902, 0.10944027, 0.11880615, 0.3917513 , 0.06862326],
                 [0.04232275, 0.09047632, 0.04658423, 0.02905692, 0.7915597],
                 [0.03812133, 0.3116883 , 0.10215054, 0.02388792, 0.5241519 ],
                 [0.18210074, 0.5038779 , 0.18253621, 0.03620264, 0.0952825 ]],
                dtvpe=float32)
  In [ ]: # Turn prediction probabilities into prediction classes
          model_5_preds = tf.argmax(model_5_pred_probs, axis=1)
          model_5_preds
Out[101]: <tf.Tensor: shape=(30212,), dtype=int64, numpy=array([0, 0, 3, ..., 4,</pre>
          4, 1])>
  In [ ]: # Calculate results of token-char-positional hybrid model
          model_5_results = calculate_results(y_true=val_labels_encoded,
                                              y_pred=model_5_preds)
          model_5_results
Out[102]: {'accuracy': 82.6128690586522,
           'f1': 0.8250369638872138,
           'precision': 0.8244488224211757,
           'recall': 0.8261286905865219}
```

Compare model results

Far out, we've come a long way. From a baseline model to training a model containing three different kinds of embeddings.

Now it's time to compare each model's performance against each other.

We'll also be able to compare our model's to the <u>PubMed 200k RCT: a</u>
<u>Dataset for Sequential Sentence Classification in Medical Abstracts</u>
(https://arxiv.org/pdf/1710.06071.pdf) paper.

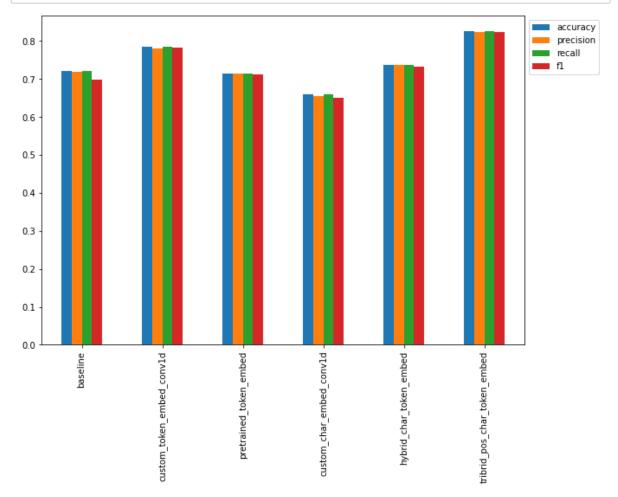
Since all of our model results are in dictionaries, let's combine them into a pandas DataFrame to visualize them.

Out[103]:

	accuracy	precision	recall	f1
baseline	72.183238	0.718647	0.721832	0.698925
<pre>custom_token_embed_conv1d</pre>	78.448961	0.781494	0.784490	0.782299
<pre>pretrained_token_embed</pre>	71.425261	0.714881	0.714253	0.711455
<pre>custom_char_embed_conv1d</pre>	65.877797	0.654501	0.658778	0.651686
hybrid_char_token_embed	73.623064	0.737075	0.736231	0.733561
tribrid_pos_char_token_embed	82.612869	0.824449	0.826129	0.825037

```
In [ ]: # Reduce the accuracy to same scale as other metrics
all_model_results["accuracy"] = all_model_results["accuracy"]/100
```

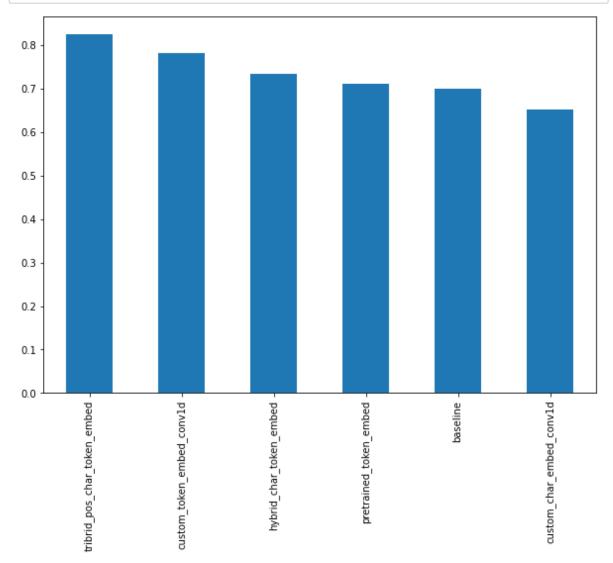
In []: # Plot and compare all of the model results
all_model_results.plot(kind="bar", figsize=(10, 7)).legend(bbox_to_anch)



Since the <u>PubMed 200k RCT: a Dataset for Sequential Sentence</u>
<u>Classification in Medical Abstracts</u>
(https://arxiv.org/pdf/1710.06071.pdf) paper compares their tested
model's F1-scores on the test dataset, let's take at our model's F1-scores.

I Note: We could've also made these comparisons in TensorBoard using the <u>TensorBoard</u> (https://www.tensorflow.org/api_docs/python/tf/keras/callbacks/Te callback during training.

In []: # Sort model results by f1-score
all_model_results.sort_values("f1", ascending=False)["f1"].plot(kind="b



Nice! Based on F1-scores, it looks like our tribrid embedding model performs the best by a fair margin.

Though, in comparison to the results reported in Table 3 of the <u>PubMed 200k RCT: a Dataset for Sequential Sentence Classification in Medical Abstracts (https://arxiv.org/pdf/1710.06071.pdf)</u> paper, our model's F1-score is still underperforming (the authors model achieves an F1-score of 90.0 on the 20k RCT dataset versus our F1-score of ~82.6).

There are some things to note about this difference:

- Our models (with an exception for the baseline) have been trained on $\sim 18,000$ (10% of batches) samples of sequences and labels rather than the full $\sim 180,000$ in the 20k RCT dataset.
 - This is often the case in machine learning experiments though, make sure training works on a smaller number of samples, then

upscale when needed (an extension to this project will be training a model on the full dataset).

• Our model's prediction performance levels have been evaluated on the validation dataset not the test dataset (we'll evaluate our

Save and load best performing model

4

Since we've been through a fair few experiments, it's a good idea to save our best performing model so we can reuse it without having to retrain it.

We can save our best performing model by calling the save()
save and serialize#the short and method on it.

In []: # Save best performing model to SavedModel format (default)
model_5.save("skimlit_tribrid_model") # model will be saved to path spe

WARNING:absl:Found untraced functions such as lstm_cell_4_layer_call_a nd_return_conditional_losses, lstm_cell_4_layer_call_fn, lstm_cell_5_l ayer_call_and_return_conditional_losses, lstm_cell_5_layer_call_fn, lstm_cell_4_layer_call_fn while saving (showing 5 of 10). These function s will not be directly callable after loading.

INFO:tensorflow:Assets written to: skimlit_tribrid_model/assets
INFO:tensorflow:Assets written to: skimlit_tribrid_model/assets

Optional: If you're using Google Colab, you might want to copy your saved model to Google Drive (or download it https://colab.research.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.research.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.research.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.research.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.research.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.research.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.research.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h https://colab.google.com/notebooks/io.ipynb#scrollTo=hauvGV4h <a href="https://colab.goog

In []: # Example of copying saved model from Google Colab to Drive (requires 0
!cp skim_lit_best_model -r /content/drive/MyDrive/tensorflow_course/s

Like all good cooking shows, we've got a pretrained model (exactly the same kind of model we built for model_5 <u>saved and stored on Google Storage</u>

(https://storage.googleapis.com/ztm tf course/skimlit/skimlit best mode

So to make sure we're all using the same model for evaluation, we'll download it and load it in.

And when loading in our model, since it uses a couple of custom
objects

(https://www.tensorflow.org/guide/keras/save and serialize#custom objec (our TensorFlow Hub layer and TextVectorization layer), we'll have to load it in by specifying them in the custom_objects parameter of tf.keras.models.load model()

(https://www.tensorflow.org/api docs/python/tf/keras/models/load model)

```
In [ ]:
        # Download pretrained model from Google Storage
        !wget https://storage.googleapis.com/ztm_tf_course/skimlit/skimlit_trib
        !mkdir skimlit_qs_model
        !unzip skimlit_tribrid_model.zip -d skimlit_qs_model
        --2021-08-25 00:03:10-- https://storage.googleapis.com/ztm_tf_course/
        skimlit/skimlit_tribrid_model.zip (https://storage.googleapis.com/ztm_
        tf_course/skimlit/skimlit_tribrid_model.zip)
        Resolving storage.googleapis.com (storage.googleapis.com)... 142.250.1
        41.128, 142.251.2.128, 74.125.137.128, ...
        Connecting to storage.googleapis.com (storage.googleapis.com)|142.250.
        141.128|:443... connected.
        HTTP request sent, awaiting response... 200 OK
        Length: 962561955 (918M) [application/zip]
        Saving to: 'skimlit_tribrid_model.zip'
        skimlit_tribrid_mod 100%[=============] 917.97M 40.9MB/s
                                                                             in
        12s
        2021-08-25 00:03:23 (74.5 MB/s) - 'skimlit_tribrid_model.zip' saved [9
        62561955/962561955]
        Archive: skimlit_tribrid_model.zip
           creating: skimlit_qs_model/skimlit_tribrid_model/
           creating: skimlit_gs_model/skimlit_tribrid_model/variables/
          inflating: skimlit_qs_model/skimlit_tribrid_model/variables/variable
        s.index
          inflating: skimlit_qs_model/skimlit_tribrid_model/variables/variable
        s.data-00000-of-00001
          inflating: skimlit_gs_model/skimlit_tribrid_model/keras_metadata.pb
          inflating: skimlit_gs_model/skimlit_tribrid_model/saved_model.pb
           creating: skimlit_qs_model/skimlit_tribrid_model/assets/
In [ ]: # Import TensorFlow model dependencies (if needed) - https://github.com
        import tensorflow_hub as hub
        import tensorflow as tf
        from tensorflow.keras.layers import TextVectorization
        model_path = "skimlit_qs_model/skimlit_tribrid_model/saved_model.pb"
        # Load downloaded model from Google Storage
        loaded_model = tf.keras.models.load_model(model_path)#,
                                                  # Note: with TensorFlow 2.5+
                                                  # (created when using model.s
                                                  # parameter. I'm leaving the
                                                   # custom_objects={"TextVector
                                                                     "KerasLayeı
```

Make predictions and evalaute them against the truth labels

To make sure our model saved and loaded correctly, let's make predictions with it, evaluate them and then compare them to the prediction results we calculated earlier.

```
In []: | # Make predictions with the loaded model on the validation set
         loaded_pred_probs = loaded_model.predict(val_pos_char_token_dataset, ve
         loaded_preds = tf.argmax(loaded_pred_probs, axis=1)
         loaded_preds[:10]
         Out[111]: <tf.Tensor: shape=(10,), dtype=int64, numpy=array([0, 0, 3, 2, 2, 4,</pre>
         4, 4, 4, 1])>
 In [ ]: # Evaluate loaded model's predictions
         loaded model results = calculate results(val labels encoded,
                                                 loaded preds)
         loaded_model_results
Out[112]: {'accuracy': 82.74526678141136,
           f1': 0.8264355957043299,
           'precision': 0.8258640600563426,
           'recall': 0.8274526678141136}
         Now let's compare our loaded model's predictions with the prediction
         results we obtained before saving our model.
 In []: # Compare loaded model results with original trained model results (she
         np.isclose(list(model_5_results.values()), list(loaded_model_results.va
Out[124]: array([ True, True, True, True])
          It's worth noting that loading in a SavedModel unfreezes all layers
          (makes them all trainable). So if you want to freeze any layers,
         you'll have to set their trainable attribute to False.
```

In []: # Check loaded model summary (note the number of trainable parameters)
loaded_model.summary()

Layer (type) cted to	Output Shape	Param #	Conne
char_inputs (InputLayer)	[(None, 1)]	0	
<pre>char_vectorizer (TextVectorizat inputs[0][0]</pre>	(None, None)	0	char_
token_inputs (InputLayer)	[(None,)]	0	
<pre>char_embed (Embedding) vectorizer[0][0]</pre>	(None, None, 25)	1750	char_
universal_sentence_encoder (Ker _inputs[0][0]	(None, 512)	256797824	token
bidirectional_1 (Bidirectional) embed[0][0]	(None, 64)	14848	char_
token_char_hybrid_embedding (Corsal_sentence_encoder[0][0]	(None, 576)	0	unive bidir
ectional_1[0][0]			21411
line_number_input (InputLayer)	[(None, 15)]	0	
total_lines_input (InputLayer)	[(None, 20)]	0	
dense_10 (Dense) _char_hybrid_embedding[0][0]	(None, 256)	147712	token
dense_8 (Dense) number_input[0][0]	(None, 32)	512	line_
dense_9 (Dense) _lines_input[0][0]	(None, 32)	672	total
dropout_2 (Dropout) _10[0][0]	(None, 256)	0	dense

```
token_char_positional_embedding (None, 320)
                                                  dense
_8[0][0]
                                                  dense
_9[0][0]
                                                  dropo
ut_2[0][0]
output_layer (Dense)
                        (None, 5)
                                        1605
                                                  token
char positional embedding[0
______
_____
Total params: 256,964,923
Trainable params: 167,099
Non-trainable params: 256,797,824
```

Evaluate model on test dataset

To make our model's performance more comparable with the results reported in Table 3 of the <u>PubMed 200k RCT: a Dataset for Sequential Sentence Classification in Medical Abstracts</u> (https://arxiv.org/pdf/1710.06071.pdf) paper, let's make predictions on the test dataset and evaluate them.

```
In [ ]: # Create test dataset batch and prefetched
         test_pos_char_token_data = tf.data.Dataset.from_tensor_slices((test_lin
                                                                      test_tot
                                                                      test_sen
                                                                      test_cha
         test_pos_char_token_labels = tf.data.Dataset.from_tensor_slices(test_la
         test pos char token dataset = tf.data.Dataset.zip((test pos char token
         test_pos_char_token_dataset = test_pos_char_token_dataset.batch(32).pre
          # Check shapes
         test_pos_char_token_dataset
Out[115]: <PrefetchDataset shapes: (((None, 15), (None, 20), (None,), (None,)),</pre>
          (None, 5)), types: ((tf.float32, tf.float32, tf.string, tf.string), t
         f.float64)>
 In [ ]: # Make predictions on the test dataset
         test_pred_probs = loaded_model.predict(test_pos_char_token_dataset,
                                               verbose=1)
         test_preds = tf.argmax(test_pred_probs, axis=1)
         test_preds[:10]
         Out[116]: <tf.Tensor: shape=(10,), dtype=int64, numpy=array([3, 3, 2, 2, 4, 4,</pre>
         4, 1, 4, 0])>
```

> It seems our best model (so far) still has some ways to go to match the performance of the results in the paper (their model gets 90.0 F1-score on the test dataset, where as ours gets ~82.1 F1-score).

However, as we discussed before our model has only been trained on 20,000 out of the total $\sim 180,000$ sequences in the RCT 20k dataset. We also haven't fine-tuned our pretrained embeddings (the paper fine-tunes GloVe embeddings). So there's a couple of extensions we could try to improve our results.

Find most wrong

One of the best ways to investigate where your model is going wrong (or potentially where your data is wrong) is to visualize the "most wrong" predictions.

The most wrong predictions are samples where the model has made a prediction with a high probability but has gotten it wrong (the model's prediction disagreess with the ground truth label).

Looking at the most wrong predictions can give us valuable information on how to improve further models or fix the labels in our data.

Let's write some code to help us visualize the most wrong predictions from the test dataset.

First we'll convert all of our integer-based test predictions into their string-based class names.

```
In [ ]: %%time
    # Get list of class names of test predictions
    test_pred_classes = [label_encoder.classes_[pred] for pred in test_pred
    test_pred_classes
```

CPU times: user 10.2 s, sys: 856 ms, total: 11.1 s Wall time: 9.42 s

Now we'll enrich our test DataFame with a few values:

- A "prediction" (string) column containing our model's prediction for a given sample.
- A "pred_prob" (float) column containing the model's maximum prediction probabiliy for a given sample.
- A "correct" (bool) column to indicate whether or not the model's prediction matches the sample's target label

```
In []: # Create prediction-enriched test dataframe
    test_df["prediction"] = test_pred_classes # create column with test pre
    test_df["pred_prob"] = tf.reduce_max(test_pred_probs, axis=1).numpy() #
    test_df["correct"] = test_df["prediction"] == test_df["target"] # creat
    test_df.head(20)
```

Out[119]:		target	text	line_number	total_lines	prediction	pred_prob	C)
	0	BACKGROUND	this study analyzed liver function	0	8	OBJECTIVE	0.513077	

0	BACKGROUND	this study analyzed liver function abnormaliti	0	8	OBJECTIVE	0.513077
1	RESULTS	a post hoc analysis was conducted with the use	1	8	OBJECTIVE	0.310540
2	RESULTS	liver function tests (lfts) were measured at	2	8	METHODS	0.801705
3	RESULTS	survival analyses were used to assess the asso	3	8	METHODS	0.627319
4	RESULTS	the percentage of patients with abnormal lfts	4	8	RESULTS	0.718288
5	RESULTS	when mean hemodynamic profiles were compared i	5	8	RESULTS	0.879730
6	RESULTS	multivariable analyses revealed that patients	6	8	RESULTS	0.548948
7	CONCLUSIONS	abnormal lfts are common in the adhf populatio	7	8	CONCLUSIONS	0.445276
8	CONCLUSIONS	elevated meld- xi scores are associated with po	8	8	RESULTS	0.529703
9	BACKGROUND	minimally invasive endovascular aneurysm repai	0	12	BACKGROUND	0.545452
10	BACKGROUND	the aim of this study was to analyse the cost	1	12	OBJECTIVE	0.495984

	target	text	line_number	total_lines	prediction	pred_prob	C
11	METHODS	resource use was determined from the amsterdam	2	12	METHODS	0.587782	
12	METHODS	the analysis was performed from a provider per	3	12	METHODS	0.852491	
13	METHODS	all costs were calculated as if all patients h	4	12	METHODS	0.573058	
14	RESULTS	a total of @ patients were randomized .	5	12	RESULTS	0.674374	
15	RESULTS	the @-day mortality rate was @ per cent after	6	12	RESULTS	0.664036	
16	RESULTS	at @months , the total mortality rate for evar	7	12	RESULTS	0.897093	
17	RESULTS	the mean cost difference between evar and or w	8	12	RESULTS	0.828620	
18	RESULTS	the incremental cost-effectiveness ratio per p	9	12	RESULTS	0.803249	
19	RESULTS	there was no significant difference in quality	10	12	RESULTS	0.729450	

Looking good! Having our data like this, makes it very easy to manipulate and view in different ways.

How about we sort our DataFrame to find the samples with the highest "pred_prob" and where the prediction was wrong ("correct" == False)?

```
In [ ]: # Find top 100 most wrong samples (note: 100 is an abitrary number, you
top_100_wrong = test_df[test_df["correct"] == False].sort_values("pred_
top_100_wrong
```

Out[120]: target text line_number total	tal_lines prediction pred_p	rob
---	-----------------------------	-----

					p	
16347	BACKGROUND	to evaluate the effects of the lactic acid bac	0	12	OBJECTIVE	0.944838
13874	CONCLUSIONS	symptom outcomes will be assessed and estimate	4	6	METHODS	0.941099
1221	RESULTS	data were collected prospectively for @ months	3	13	METHODS	0.928523
13598	METHODS	-@ % vs. fish : -@ % vs. fish + s : -@ % ; p <	6	9	RESULTS	0.918107
21382	OBJECTIVE	design , settings , participants , and interve	3	13	METHODS	0.918088
12269	RESULTS	patients received oral se tablets (@ mcg) or	4	10	METHODS	0.821220
9881	RESULTS	the primary outcome was bp control , and secon	4	11	METHODS	0.821166
1220	RESULTS	the group intervention consisted of @ weekly c	2	13	METHODS	0.821033
22105	RESULTS	we randomised @ statin treated cvd patients an	3	12	METHODS	0.820954
16840	RESULTS	the primary endpoint was a composite of cardio	3	12	METHODS	0.820538

Great (or not so great)! Now we've got a subset of our model's most wrong predictions, let's write some code to visualize them.

```
In []: # Investigate top wrong preds
for row in top_100_wrong[0:10].itertuples(): # adjust indexes to view c
_, target, text, line_number, total_lines, prediction, pred_prob, _ =
    print(f"Target: {target}, Pred: {prediction}, Prob: {pred_prob}, Line
    print(f"Text:\n{text}\n")
    print("----\n")
```

Target: BACKGROUND, Pred: OBJECTIVE, Prob: 0.9448384046554565, Line number: 0, Total lines: 12

Text:

to evaluate the effects of the lactic acid bacterium lactobacillus sal ivarius on caries risk factors .

Target: CONCLUSIONS, Pred: METHODS, Prob: 0.9410986304283142, Line number: 4, Total lines: 6

Text:

symptom outcomes will be assessed and estimates of cost-effectiveness made .

Target: RESULTS, Pred: METHODS, Prob: 0.9285234808921814, Line number: 3, Total lines: 13

Text:

data were collected prospectively for @ months beginning after complet ion of the first @ group clinic appointments (@ months post randomization) .

Target: METHODS, Pred: RESULTS, Prob: 0.9181066155433655, Line number: 6, Total lines: 9

Text:

-@ % vs. fish : -@ % vs. fish + s : -@ % ; p < @) but there were no s ignificant differences between groups .

Target: OBJECTIVE, Pred: METHODS, Prob: 0.9180881381034851, Line numbe
r: 3, Total lines: 13

Text:

design , settings , participants , and intervention : ten healthy , no rmal-weight men were studied in randomized , double-blind fashion , each receiving a @-minute intraduodenal infusion of l-trp at @ (total @ kcal) or @ (total @ kcal) kcal/min or saline (control) .

Target: METHODS, Pred: RESULTS, Prob: 0.9168640971183777, Line number: 5, Total lines: 7

Text:

at this time , an as@ response was achieved by @ (@ %) and @ (@ %) patients in groups @ and @ , respectively (p < @ for all) .

Target: RESULTS, Pred: METHODS, Prob: 0.9164432883262634, Line number: 3, Total lines: 16

Text:

a cluster randomised trial was implemented with @,@ children in @ gove rnment primary schools on the south coast of kenya in @-@ .

Target: BACKGROUND, Pred: OBJECTIVE, Prob: 0.914499819278717, Line number: 0, Total lines: 9

Text:

to compare the efficacy of the newcastle infant dialysis and ultrafilt ration system (nidus) with peritoneal dialysis (pd) and convention al haemodialysis (hd) in infants weighing < @ kg .

Target: RESULTS, Pred: METHODS, Prob: 0.9141932725906372, Line number: 4, Total lines: 13

Text:

baseline measures included sociodemographics , standardized anthropome trics , asthma control test (act) , gerd symptom assessment scale , pittsburgh sleep quality index , and berlin questionnaire for sleep ap nea .

Target: BACKGROUND, Pred: OBJECTIVE, Prob: 0.9109073877334595, Line number: 0, Total lines: 11

Text:

to assess the temporal patterns of late gastrointestinal (gi) and ge nitourinary (gu) radiotherapy toxicity and resolution rates in a ran domised controlled trial (all-ireland cooperative oncology research g roup @-@) assessing duration of neo-adjuvant (na) hormone therapy f or localised prostate cancer .

What do you notice about the most wrong predictions? Does the model make silly mistakes? Or are some of the labels incorrect/ambiguous (e.g. a line in an abstract could potentially be labelled OBJECTIVE or BACKGROUND and make sense).

A next step here would be if there are a fair few samples with inconsistent labels, you could go through your training dataset, update the labels and then retrain a model. The process of using a model to help improve/investigate your dataset's labels is often referred to as **active learning**.

Make example predictions

Okay, we've made some predictions on the test dataset, now's time to really test our model out.

To do so, we're going to get some data from the wild and see how our model performs.

In other words, were going to find an RCT abstract from PubMed, preprocess the text so it works with our model, then pass each sequence in the wild abstract through our model to see what label it predicts.

For an appropriate sample, we'll need to search PubMed for RCT's (randomized controlled trials) without abstracts which have been split up (on exploring PubMed you'll notice many of the abstracts are already preformatted into separate sections, this helps dramatically with readability).

Going through various PubMed studies, I managed to find the following unstructured abstract from <u>RCT of a manualized social treatment for high-functioning autism spectrum disorders</u>
(https://pubmed.ncbi.nlm.nih.gov/20232240/):

This RCT examined the efficacy of a manualized social intervention for children with HFASDs. Participants were randomly assigned to treatment or wait-list conditions. Treatment included instruction and therapeutic activities targeting social skills, face-emotion recognition, interest expansion, and interpretation of non-literal language. A response-cost program was applied to reduce problem behaviors and foster skills acquisition. Significant treatment effects were found for five of seven primary outcome measures (parent ratings and direct child measures). Secondary measures based on staff ratings (treatment group only) corroborated gains reported by parents. High levels of parent, child and staff satisfaction were reported, along with high levels of treatment fidelity. Standardized effect size estimates were primarily in the medium and large ranges and favored the treatment group.

Looking at the large chunk of text can seem quite intimidating. Now imagine you're a medical researcher trying to skim through the literature to find a study relevant to your work.

Sounds like quite the challenge right?

Enter SkimLit □□!

Let's see what our best model so far (model_5) makes of the above abstract.

But wait...

As you might've guessed the above abstract hasn't been formatted in the same structure as the data our model has been trained on. Therefore, before we can make a prediction on it, we need to preprocess it just as we have our other sequences.

More specifically, for each abstract, we'll need to:

- 1. Split it into sentences (lines).
- 2. Split it into characters.
- 3. Find the number of each line.
- 4. Find the total number of lines.

Starting with number 1, there are a couple of ways to split our abstracts into actual sentences. A simple one would be to use Python's in-built split() string method, splitting the abstract wherever a fullstop appears. However, can you imagine where this might go wrong?

Another more advanced option would be to leverage spaCy's
(https://spacy.io/)
(a very powerful NLP library) sentencizer
(https://spacy.io/usage/linguistic-features#sbd)
class. Which is an easy to use sentence splitter based on spaCy's English language model.

I've prepared some abstracts from PubMed RCT papers to try our model on, we can download them $\underline{\text{from GitHub}}$

(https://raw.githubusercontent.com/mrdbourke/tensorflow-deeplearning/main/extras/skimlit example abstracts.json).

```
In [ ]: import json
        # Download and open example abstracts (copy and pasted from PubMed)
        !wget https://raw.githubusercontent.com/mrdbourke/tensorflow-deep-learn
        with open("skimlit_example_abstracts.json", "r") as f:
          example_abstracts = json.load(f)
        example_abstracts
        --2021-08-25 00:08:37-- https://raw.githubusercontent.com/mrdbourke/t
        ensorflow-deep-learning/main/extras/skimlit_example_abstracts.json (ht
        tps://raw.githubusercontent.com/mrdbourke/tensorflow-deep-learning/mai
        n/extras/skimlit_example_abstracts.json)
        Resolving raw.githubusercontent.com (raw.githubusercontent.com)... 18
        5.199.110.133, 185.199.108.133, 185.199.109.133, ...
        Connecting to raw.githubusercontent.com (raw.githubusercontent.com)|18
        5.199.110.133|:443... connected.
        HTTP request sent, awaiting response... 200 OK
        Length: 6737 (6.6K) [text/plain]
        Saving to: 'skimlit_example_abstracts.json'
        skimlit example abs 100%[============] 6.58K --.-KB/s
                                                                            in
        0s
        2021-08-25 00:08:37 (82.5 MB/s) - 'skimlit_example_abstracts.json' sav
        ed [6737/6737]
        NameError
                                                  Traceback (most recent call
        last)
        <ipython-input-122-aa3c30151e9a> in <module>()
              4 with open("skimlit_example_abstracts.json", "r") as f:
                  example_abstracts = json.load(f)
        ---> 5
              7 example_abstracts
        NameError: name 'json' is not defined
In [ ]: # See what our example abstracts look like
        abstracts = pd.DataFrame(example_abstracts)
        abstracts
```

Now we've downloaded some example abstracts, let's see how one of them goes with our trained model.

First, we'll need to parse it using spaCy to turn it from a big chunk of text into sentences.

```
In []: # Create sentencizer - Source: https://spacy.io/usage/linguistic-featur
from spacy.lang.en import English
nlp = English() # setup English sentence parser
sentencizer = nlp.create_pipe("sentencizer") # create sentence splittin
nlp.add_pipe(sentencizer) # add sentence splitting pipeline object to s
doc = nlp(example_abstracts[0]["abstract"]) # create "doc" of parsed se
abstract_lines = [str(sent) for sent in list(doc.sents)] # return detec
abstract_lines
```

Beautiful! It looks like spaCy has split the sentences in the abstract correctly. However, it should be noted, there may be more complex abstracts which don't get split perfectly into separate sentences (such as the example in <u>Baclofen promotes alcohol abstinence in alcohol dependent cirrhotic patients with hepatitis C virus (HCV) infection (https://pubmed.ncbi.nlm.nih.gov/22244707/)</u>, in this case, more custom splitting techniques would have to be investigated.

Now our abstract has been split into sentences, how about we write some code to count line numbers as well as total lines.

To do so, we can leverage some of the functionality of our preprocess_text_with_line_numbers() function.

```
In []: # Get total number of lines
total_lines_in_sample = len(abstract_lines)

# Go through each line in abstract and create a list of dictionaries cd
sample_lines = []
for i, line in enumerate(abstract_lines):
    sample_dict = {}
    sample_dict["text"] = str(line)
    sample_dict["line_number"] = i
    sample_dict["total_lines"] = total_lines_in_sample - 1
    sample_lines.append(sample_dict)
sample_lines
```

Now we've got "line_number" and "total_lines" values, we can one-hot encode them with tf.one_hot just like we did with our training dataset (using the same values for the depth parameter).

```
In [ ]: # Get all line_number values from sample abstract
    test_abstract_line_numbers = [line["line_number"] for line in sample_li
    # One-hot encode to same depth as training data, so model accepts right
    test_abstract_line_numbers_one_hot = tf.one_hot(test_abstract_line_numb
    test_abstract_line_numbers_one_hot
```

In []: # Get all total_lines values from sample abstract
 test_abstract_total_lines = [line["total_lines"] for line in sample_lin
 # One-hot encode to same depth as training data, so model accepts right
 test_abstract_total_lines_one_hot = tf.one_hot(test_abstract_total_line
 test_abstract_total_lines_one_hot

We can also use our split_chars() function to split our abstract lines into characters.

In []: # Split abstract lines into characters
abstract_chars = [split_chars(sentence) for sentence in abstract_lines]
abstract_chars

Alright, now we've preprocessed our wild RCT abstract into all of the same features our model was trained on, we can pass these features to our model and make sequence label predictions!

In []: # Turn prediction probabilities into prediction classes
 test_abstract_preds = tf.argmax(test_abstract_pred_probs, axis=1)
 test_abstract_preds

Now we've got the predicted sequence label for each line in our sample abstract, let's write some code to visualize each sentence with its predicted label.

In []: # Turn prediction class integers into string class names
 test_abstract_pred_classes = [label_encoder.classes_[i] for i in test_a
 test_abstract_pred_classes

In []: # Visualize abstract lines and predicted sequence labels
for i, line in enumerate(abstract_lines):
 print(f"{test_abstract_pred_classes[i]}: {line}")

Nice! Isn't that much easier to read? I mean, it looks like our model's predictions could be improved, but how cool is that?

Imagine implementing our model to the backend of the PubMed website to format any unstructured RCT abstract on the site.

Or there could even be a browser extension, called "SkimLit" which would add structure (powered by our model) to any unstructured RCT abtract.

And if showed your medical researcher friend, and they thought the predictions weren't up to standard, there could be a button saying "is this label correct?... if not, what should it be?". That way the dataset, along with our model's future predictions, could be improved over time.

Of course, there are many more ways we could go to improve the model, the usuability, the preprocessing functionality (e.g. functionizing our sample abstract preprocessing pipeline) but I'll leave these for the exercises/extensions.

☐ **Question:** How can we be sure the results of our test example from the wild are truly *wild*? Is there something we should check about the sample we're testing on?

***** Exercises

- Train model_5 on all of the data in the training dataset for as many epochs until it stops improving. Since this might take a while, you might want to use:
- tf.keras.callbacks.ModelCheckpoint
 (https://www.tensorflow.org/api_docs/python/tf/keras/callbacks/Model
 to save the model's best weights only.
- tf.keras.callbacks.EarlyStopping
 (https://www.tensorflow.org/api docs/python/tf/keras/callbacks/Early
 to stop the model from training once the validation loss has
 stopped improving for ~3 epochs.
- 2. Checkout the <u>Keras guide on using pretrained GloVe embeddings</u> (<u>https://keras.io/examples/nlp/pretrained word embeddings/</u>). Can you get this working with one of our models?
- Hint: You'll want to incorporate it with a custom token Embedding Embedding layer.
- It's up to you whether or not you fine-tune the GloVe embeddings or leave them frozen.
- 3. Try replacing the TensorFlow Hub Universal Sentence Encoder pretrained embedding for the <u>TensorFlow Hub BERT PubMed expert</u> (<u>https://tfhub.dev/google/experts/bert/pubmed/2</u>) (a language model pretrained on PubMed texts) pretrained embedding. Does this effect results?
 - Note: Using the BERT PubMed expert pretrained embedding requires an extra preprocessing step for sequences (as detailed in the

TensorFlow Hub guide
(https://tfhub.dev/google/experts/bert/pubmed/2)).

- Does the BERT model beat the results mentioned in this paper? https://arxiv.org/pdf/1710.06071.pdf
 (https://arxiv.org/pdf/1710.06071.pdf)
- 4. What happens if you were to merge our line_number and total_lines features for each sequence? For example, created a X_of_Y feature instead? Does this effect model performance?
- Another example: line_number=1 and total_lines=11 turns into line_of_X=1_of_11.
- 5. Write a function (or series of functions) to take a sample abstract string, preprocess it (in the same way our model has been trained), make a prediction on each sequence in the abstract and return the abstract in the format:

PREDICTED_LABEL : SEQUENCE
 PREDICTED_LABEL : SEQUENCE
 PREDICTED_LABEL : SEQUENCE
 PREDICTED_LABEL : SEQUENCE

• . . .

You can find your own unstructured RCT abstract from PubMed or try this one from: <u>Baclofen promotes alcohol abstinence in</u> <u>alcohol dependent cirrhotic patients with hepatitis C virus</u> (HCV) infection (https://pubmed.ncbi.nlm.nih.gov/22244707/).

□ Extra-curriculum

- For more on working with text/spaCy, see spaCy's advanced NLP <a href="mailto:spaCy. If you're going to be working on production-level NLP problems, you'll probably end up using spaCy.
- For another look at how to approach a text classification problem like the one we've just gone through, I'd suggest going through Google's Machine Learning Course for text classification (https://developers.google.com/machine-learning/guides/text-classification).
- Since our dataset has imbalanced classes (as with many real-world datasets), so it might be worth looking into the TensorFlow guide for different methods to training a model with imbalanced classes (https://www.tensorflow.org/tutorials/structured data/imbalanced data