Personalized Medicine: Redefining Cancer Treatment

<DataGeeks/>

AGENDA

- 1. Problem Definition
- 2. Dataset Explanation
- 3. Methodology
- 4. Results
- 5. Discussion

Problem Definition

- Develop a robust and accurate model to automate the process of identifying carcinogenic genetic mutations from medical text records.
- Could be analyzed as a text classification problem.
- Possible to eliminate the time consuming manual efforts.
- A chance to predict the cancer causing mutations early and treat the patient's tumor at a preliminary stage.

Dataset Explanation

- Four files in total training_variants, test_variants, training_text and test_text.
- Variants Files:
 - Information about genetic mutations
 - CSV file with 4 columns ID, Gene (gene where mutation is located), Variation (Amino acid change for this mutation), Class (1-9 class where this mutation is classified on)
 - The test_variants file does not contain the last column specifying the class.
- Text Files:
 - Clinical evidence (text) that human experts used to classify genetic mutations.
 - Double pipe (||) delimited file with 2 fields ID, Text (clinical evidence obtained from various research papers)
- Both sets of training and test files are linked using the ID field.
- Some of the test data is machine-generated to prevent hand labeling. Kaggle
 ignores the results from machine-generated samples in the final result.

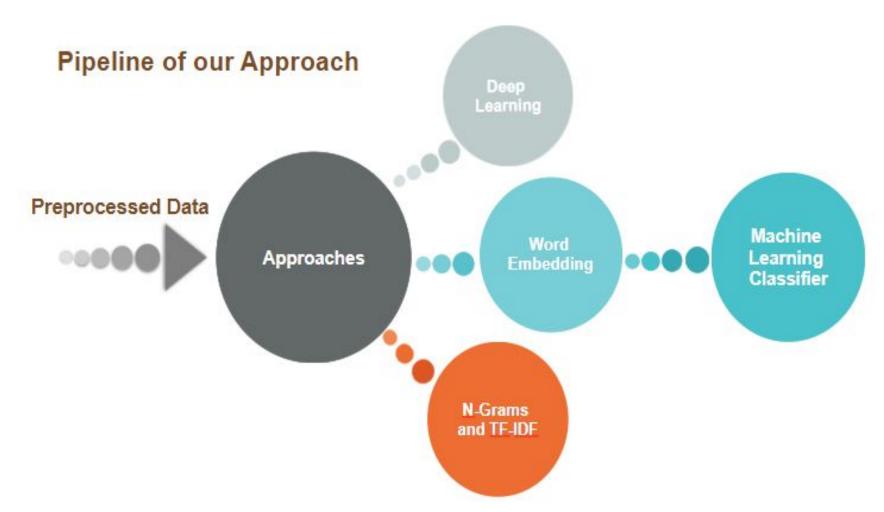
Dataset Glimpse

Training Variants:

ID ‡	Gene [‡]	Variation	Class
0	FAM58A	Truncating Mutations	1
1	CBL	W802*	2
2	CBL	Q249E	2
3	CBL	N454D	3
4	CBL	L399V	4

Training Text:

ID ‡	txt ÷
0	Cyclin-dependent kinases (CDKs) regulate a variety of fu
1	Abstract Background Non-small cell lung cancer (NSCLC)
2	Abstract Background Non-small cell lung cancer (NSCLC)
3	Recent evidence has demonstrated that acquired unipar
4	Oncogenic mutations in the monomeric Casitas B-lineag



Feature Engineering

TF-IDF

- Text Data: Unigrams and Bigrams on word level.
- Gene and Variants Encoding: 1-10 character level n-grams
- Label Encoding for Gene and Variants

Word Embeddings

- Glove vectors trained on general domain web data.
- Word2vec trained on PubMed medical text.
- Word2vec trained on our data.

Deep Learning

- Long Short Term Memory Networks
- Hierarchical LSTMs
- Convolutional Neural Networks

Results

$$logloss = -\frac{1}{N} \sum_{i=1}^{N} \sum_{j=1}^{M} y_{i,j} \log(p_{i,j})$$

Algorithm	Private Score	Public Score
Hierarchical LSTMs	3.18	1.30
CNN	2.50	2.42
TFIDF + SVD + Gradient Boosting	2.41	2.11
Glove + Logistic Regression	2.80	1.44
Word2Vec (PubMed) + Gradient Boosting	2.45	1.35
Word2Vec (Training Data) + Gradient Boosting	1.98	1.22

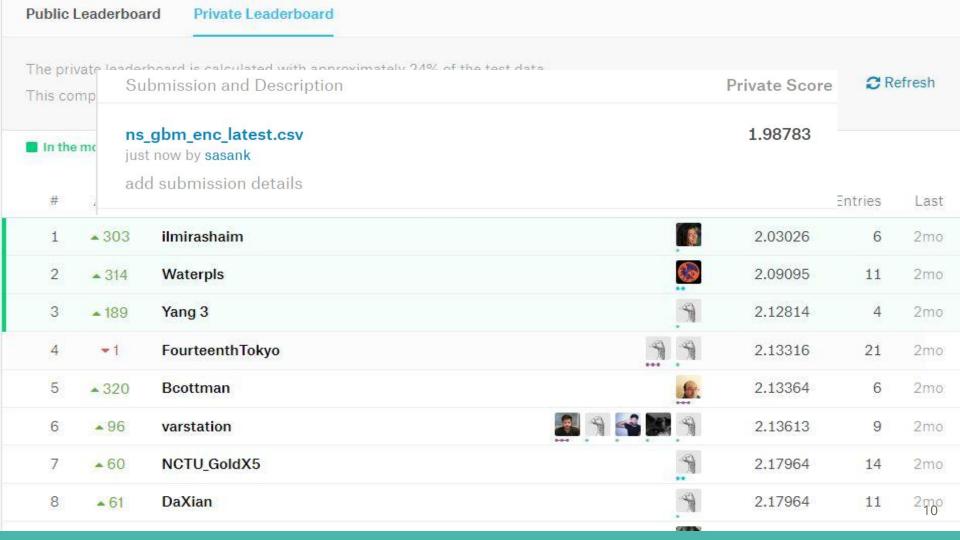
Learned Embeddings

Mutation Cosine Distance

substitution	0.269
variant	0.297
alteration	0.319
mutational	0.369
polymorphism	0.389

Disease Cosine Distance

malignancy	0.260
tumor	0.395
illness	0.411
disorder	0.414
seizures	0.415



Discussion

- Deep Learning based approaches performed poorly → Overfitting the training data.
- 2. Glove < Word2vec (Pubmed) < Word2vec trained on the dataset. Embeddings trained on the dataset performed better than pretrained word2vec embeddings and Glove Embeddings.
- 3. Take home: Learning word vector embeddings on training data is extremely effective. Boosting techniques significantly improve the performance of the classifiers.
- 4. Visualizing the Embeddings: http://projector.tensorflow.org/

Thank You!!