# Proj. 3 Uncertainty Quantification

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#### **Project Description**

Salmon is a tool for measuring gene expression, and gives estimates of transcript abundances. Using 'bootstrapping' technique, we can measure the confidence in these estimates. However, this approach tends to underestimate the uncertainty with more transcripts falling out of the interval than expected.

Our goal is to filter out these failed transcripts, find common properties between them, and come up with a quality score based on those properties which measures our confidence in the estimates.

#### **Our Approaches**

- Parse the data files (poly\_truth.tsv, quant\_bootstrap.tsv, quant.sf);
- 2. Find range of confidence interval;
- 3. Filter out the failed transcripts;
- 4. Group data by true/failed transcripts, analyze common properties.

#### **Implementation Details**

#### Parse quant\_bootstraps.tsv

This file gives us the bootstrap data (200 rounds of sample taking).

Note: we are using DataFrame from the pandas package to easier do data analysis.

#### Parse poly\_truth.tsv

This file gives us the true count of each transcript.

#### Parse quant.sf

This file gives us some attributes of the transcripts.

```
quant_file = open(root_path + "quant.sf")
lines = quant_file.readlines()
quant_file.close()
quant = [line[:-1].split('\t') for line in lines]
```

We find and retrieve the intersecting transcript ids of poly\_truth and quant\_bootstraps, and sort each id's data by ascending order. There are transcripts in quant\_bootstraps that don't show up in poly\_truth, we'll deal with them later.

```
set_qb_id = set(df_quant_boot.columns)
set_pt_id = set(df_poly_truth.transcript_id)
intersect_ids = set_qb_id & set_pt_id
sort_qb = []
use_id = []
for id in intersect_ids:
    listed = list(df_quant_boot[id])
    listed.sort()
    use_id.append(id)
    sort_qb.append(listed)
sort_qb = list(map(list,zip(*sort_qb)))
```

#### Find confidence interval

Since we have already sorted each transcript id's data, we can find an empirical confidence interval of 95% by locating the numbers at index

```
(total_length) * 2.5% and (total_length) * 97.5%,
which would be the lower and upper bound.
```

```
df_poly_truth = df_poly_truth.set_index(['transcript_id'])
sum = len(sort_qb)
percent2dot5 = df_qb_sorted.loc[int(sum*0.025)-1]
percent97dot5 = df_qb_sorted.loc[int(sum*0.975)-1]
```

#### Find the failed transcripts

Compare the counts given by poly\_truth with the lower and upper bound we found earlier. If not in range we treat it as a failed transcript, else true.

```
true_id = []
false_id = []
for id in use_id:
    down = float(percent2dot5[id])
    up = float(percent97dot5[id])
    true_count = float(df_poly_truth.loc[id])
    if down < true_count < up:
        true_id.append(id)
    else:
        false_id.append(id)</pre>
```

We go back to deal with the 'diff' transcript ids we ignored earlier. The counts of these diff transcript ids are zero, and we assume that these are true transcripts.

```
true_id.extend(list(set_qb_id.difference(set_pt_id)))
all_id = true_id[:]
all_id.extend(false_id)
```

We add a label of 1 representing true transcripts and 0 representing failed transcripts for easy grouping later on.

## Common Properties of Failed Transcripts

We group the data by true and failed transcripts, and first observe the mean, std, max and min.

Observing the mean, the average TPM and NumReads of failed transcripts is a lot bigger than the true ones.

label	Length	EffectiveLength	ТРМ	NumReads	
0	2445.007350	2245.894343	52.556885	1784.370671	
1	1905.694197	1706.993431	0.841197	20.235059	

label	Length	Length EffectiveLength		NumReads	
0	2320.422083	2121.332135	41.485945	1378.079501	
1	1901.681056	1703.008206	0.154861	6.681882	

With the std, we find that failed transcripts tend to have a significantly larger variance of TPM and NumReads.

la	bel	Length EffectiveLength		ТРМ	NumReads	
	0	2380.728560	2380.669848	448.368497	15740.352178	
	1	2055.905694	2055.526264	45.873037	235.234059	

label	Length	EffectiveLength	ТРМ	NumReads 13629.239858	
0	2302.710811	2302.636007	396.200319		
1	2059.808430	2059.407366	9.489371	152.179457	

### Min values don't seem to have much to offer other than maybe a slightly bigger length.

label	Name	Length	EffectiveLength	ТРМ	NumReads
0	ENST00000000233	158	10.987	0.0	0.0
1	ENST00000000412	21	9.784	0.0	0.0

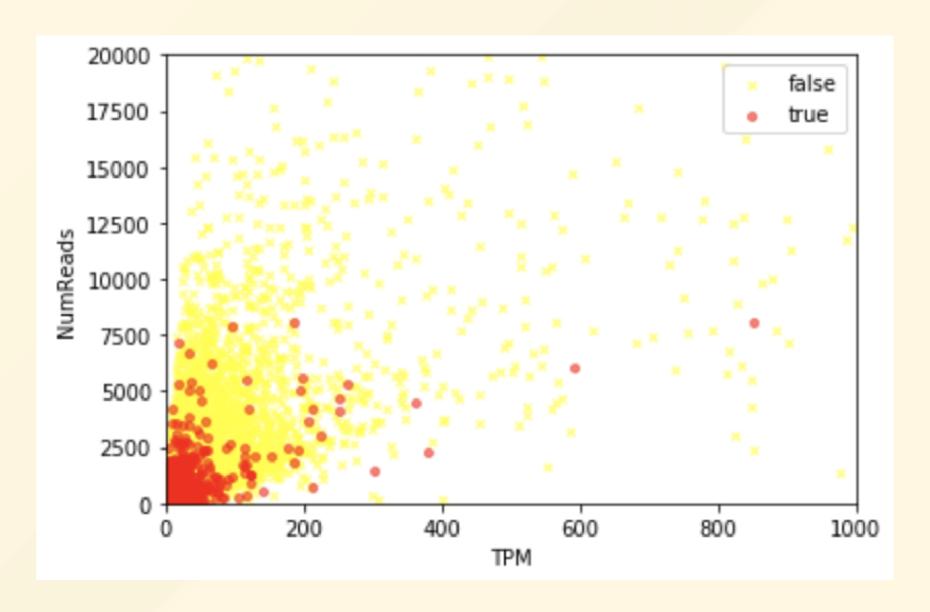
label	Name	Length	EffectiveLength	ТРМ	NumReads
0	ENST00000000233	82	10.861	0.0	0.0
1	ENST00000000412	21	9.784	0.0	0.0

With max values, the TPM and NumReads attributes seem interesting, which given the info from std, tells us that failed transcripts have a significantly wider range and variance of TPM and NumReads.

label	Name	Length	EffectiveLength	ТРМ	NumReads
0	ENST00000610278	101518	101318.991	23356.420222	1.109005e+06
1	ENST00000610279	109224	109024.991	10710.459004	3.769085e+04

label	Name	Length	EffectiveLength	ТРМ	NumReads
0	ENST00000610279	101518	101318.991	23356.420222	1.109005e+06
1	ENST00000610276	109224	109024.991	2435.717783	3.110509e+04

This looks more clear when we plot the distribution:



#### Classification Models

We used sklearn package for this.

First we gave **linear regression** a shot, and we ended up with:

mse= 0.24674520923824544 accuracy= 0.5450268817204301

The results were not ideal. Looks like it doesn't seem to be linear, so we tried another classification model.

#### With support vector regression, we achieved:

mse= 0.06162988735258758 accuracy= 0.8892588614393125

88.9% was a pretty high accuracy level, but we also managed to reach:

mse= 0.06162988735258758 accuracy= 0.9667338709677419

96.7% accuracy after we dropped the '0' count value transcripts from the 'true' group.