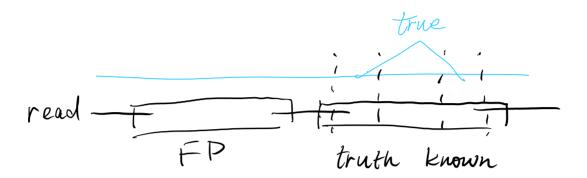
## Feb 10 2021 Meeting

Wednesday, February 10, 2021 10:35 AM

## 1. Separate FP cases and truth known cases

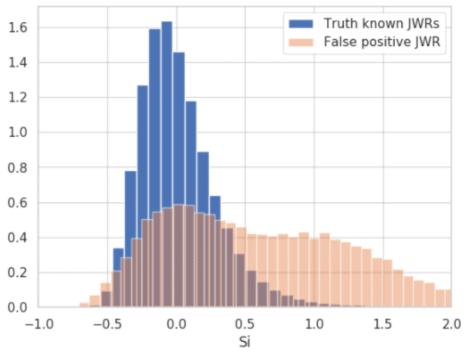


- For each reads, the true isoform is known so that the true splice site within each read can be known.
- In the analysis, junction within reads(JWR) a divided into 2 categories: False Positive and Truth known. For JWR that mapped within 10nt from both side of true splice junctions will be identified as Truth known JWR (613790 96.48%), otherwise False Positive JWR (22371 in total (3.52%)'
- Accuracy for Truth known JWR:

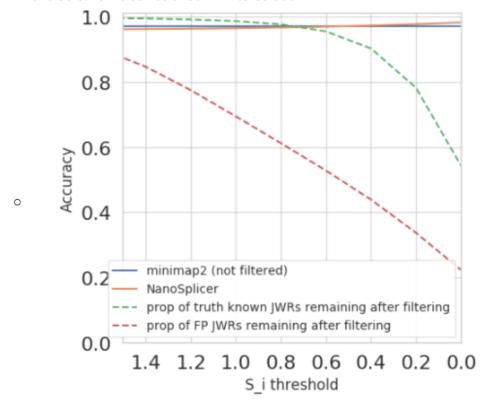
minimap accuracy: 97.2287%
NanoSplicer accuracy: 95.9815%
proportion of Junction within read identified by both softeware: 94.4238%

## 2. S\_i analysis

• Si distribution for False Positive JWR and Truth known JWR



- · Apply different Si threshold
  - Minimap2 result are not filtered, so that it's a flat line
  - Accuracy is calculated on Truth known JWRs only
  - The accuracy are the same at Si = 0.45, with 8.0% of truth known JWR filtered out, and 53.8% of False Positive JWR filtered out.



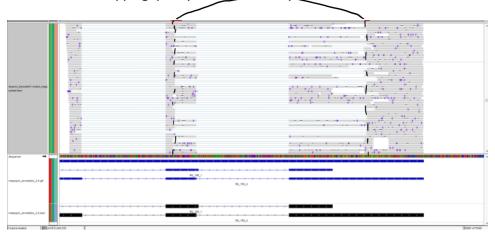
## 2.1 S\_i analysis (figure out the bimodal distribution of FP JWR

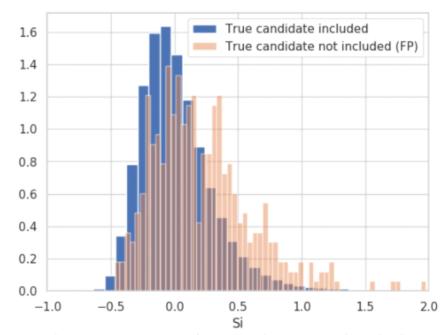
I manually checked some False positive JWR set and find some typical categories

• Junction mapping quality seem to be okay but not in annotation.

I manually checked some False positive JWK set and TING some typical categories

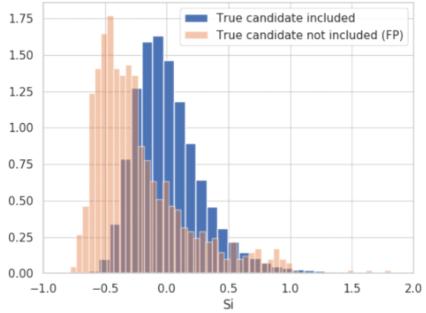
• Junction mapping quality seem to be okay but not in annotation.





• Guppy fail to trim poly-A: Poly-A tail doesn't exist in the fastq file for most of the reads, but we can still find some reads come with ploy A (planning to manually filter out them).





• False Positive introduced when minimap2 failed to identify small exons

