

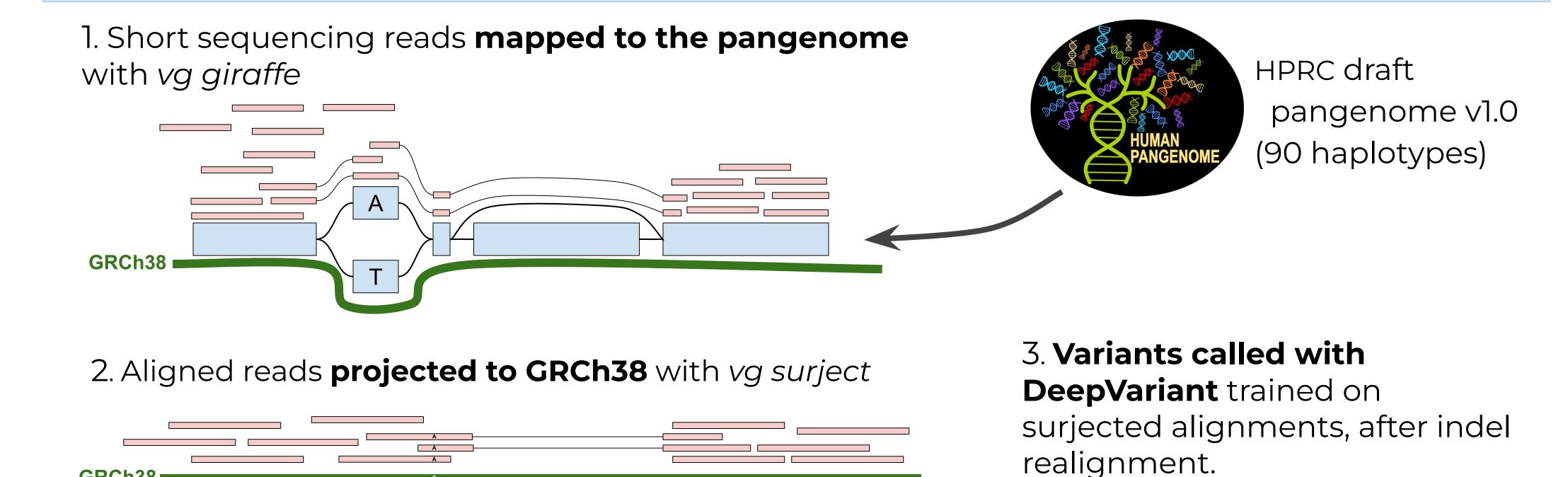
Accurate short variant calling from sequencing data with pangenomes and DeepVariant

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Background

- Accurate identification of genomic variants is critical for genetic studies and for clinical genetic testing
- Some genomics regions and variants are still challenging to detect with the standard techniques.
- The Human Pangenome Reference Consortium (HPRC) aims at producing high-quality phased de novo assemblies for more than 300 diverse individuals, and building a comprehensive and representative human pangenome

Methods



The HPRC pangenome improves the mapping of short reads and, as a result, the identification of short variants.

regions in

GRCh38.

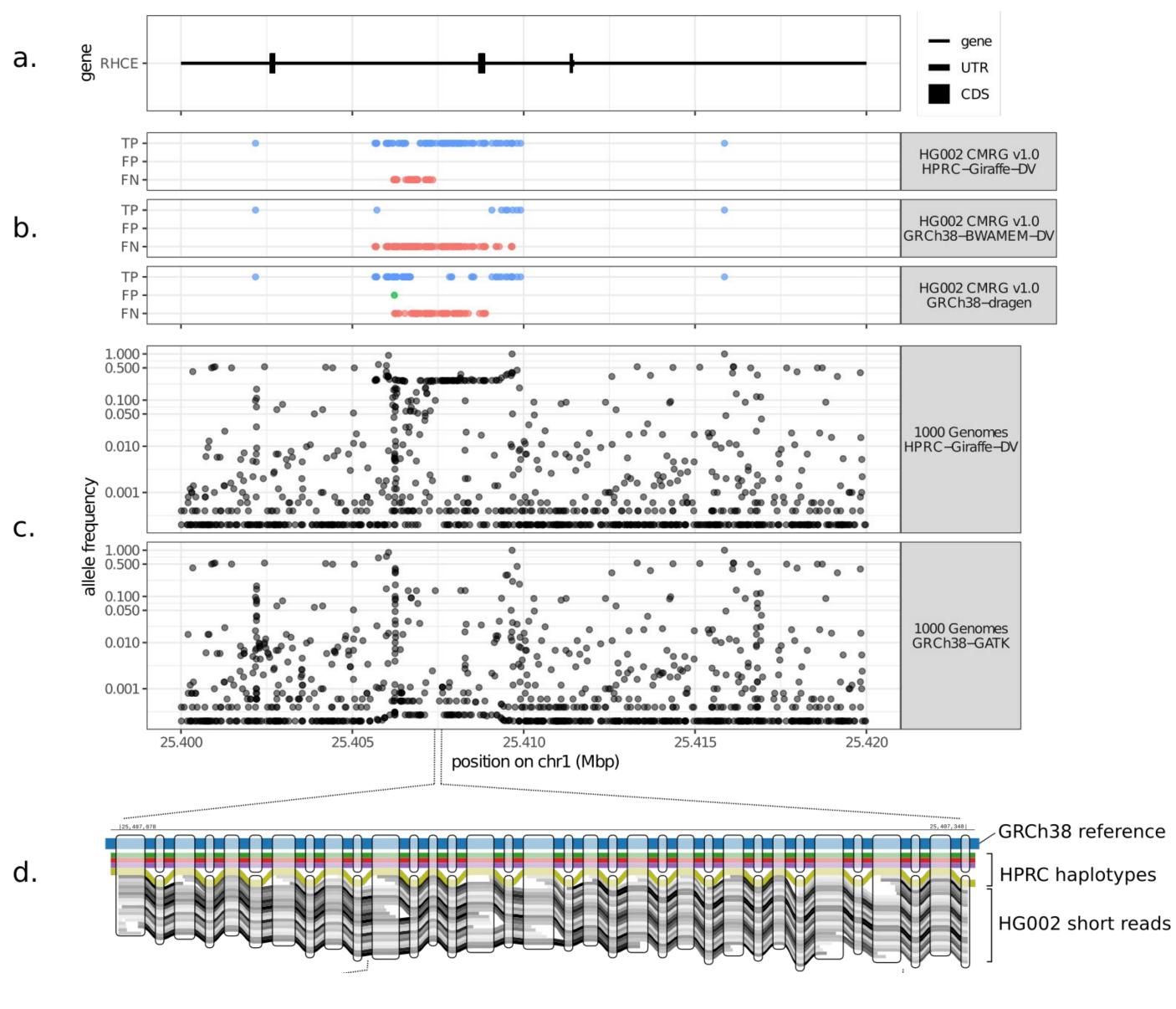
Evaluation against the Genome in a Bottle (GIAB) and Challenging Medically-Relevant Genes (CMRG) truthsets. CMRG v1.0 GIAB v4.2.1 HG005 -Evaluation of the HG002 GRCh38-based Evaluation of the HG002 CHM13-based HPRC pangenome HG001 0.980 0.975 0.9970 0.9960 0.9965 *excluding false duplications and collapsed Giraffe on HPRC-GRCh38 and DeepVariant BWAMEM on GRCh38 and DeepVariant

In RHCE (a), our approach can correctly call more variants in a region with gene conversion (b).

Giraffe on HPRC-CHM13 and DeepVariant

We could derive better allele frequencies from short read datasets in this region (c).

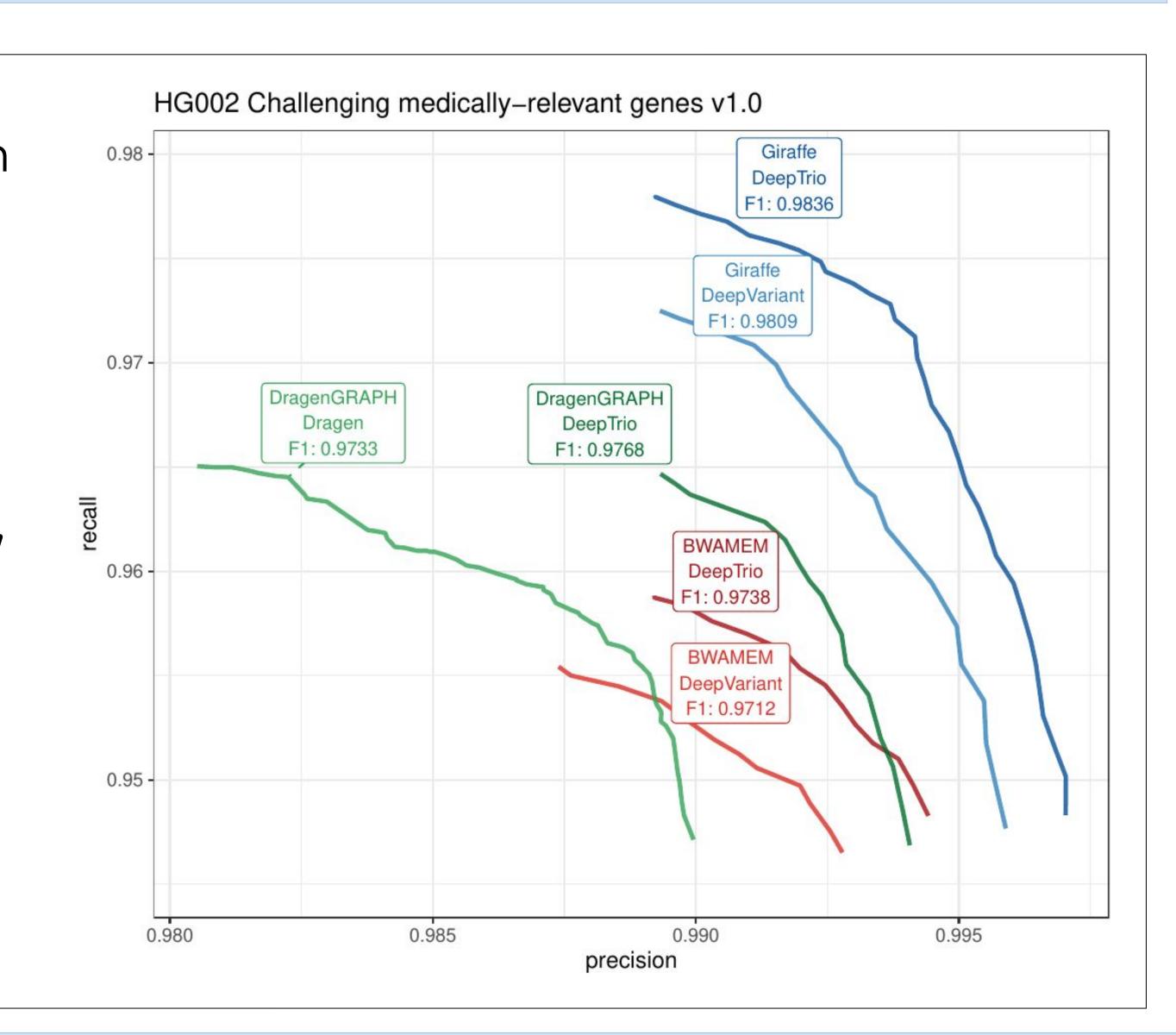
That's because the pangenome contains the gene-converted allele which ensure that reads are mapped to the correct location (d).



DragenGRAPH on GRCh38+ and Dragen

Best performance with DeepTrio trio-based calling

In challenging regions, single-sample pangenomic calling performed better than trio-based standard approaches



Conclusion

- Our pangenomic approach results in about 34% less errors, and calls short variants in a larger fraction of the genome, including in more challenging regions.
- Resources:
- A Draft Human Pangenome Reference. bioRxiv 2022 DOI:10.1101/2022.07.09.499321
- HPRC pangenomes: https://github.com/human-pangenomics/hpp_pangenome_resources
- Workflows (WDL) DOI: 10.5281/zenodo.6655968

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