



Assessing diversity and clonal variation of Australia's grapevine germplasm: Curating the FALCON-Unzip Chardonnay *de novo* genome assembly

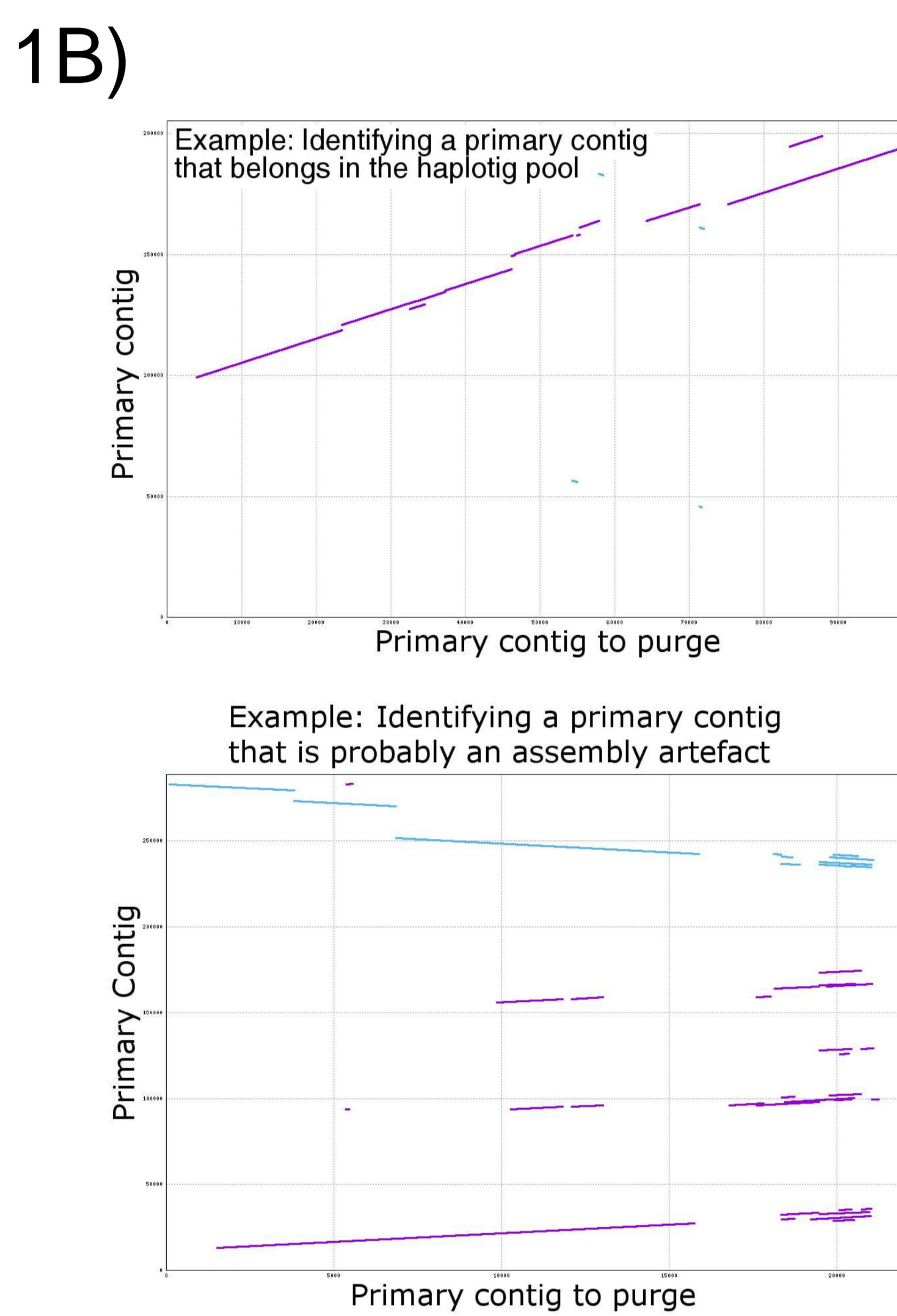
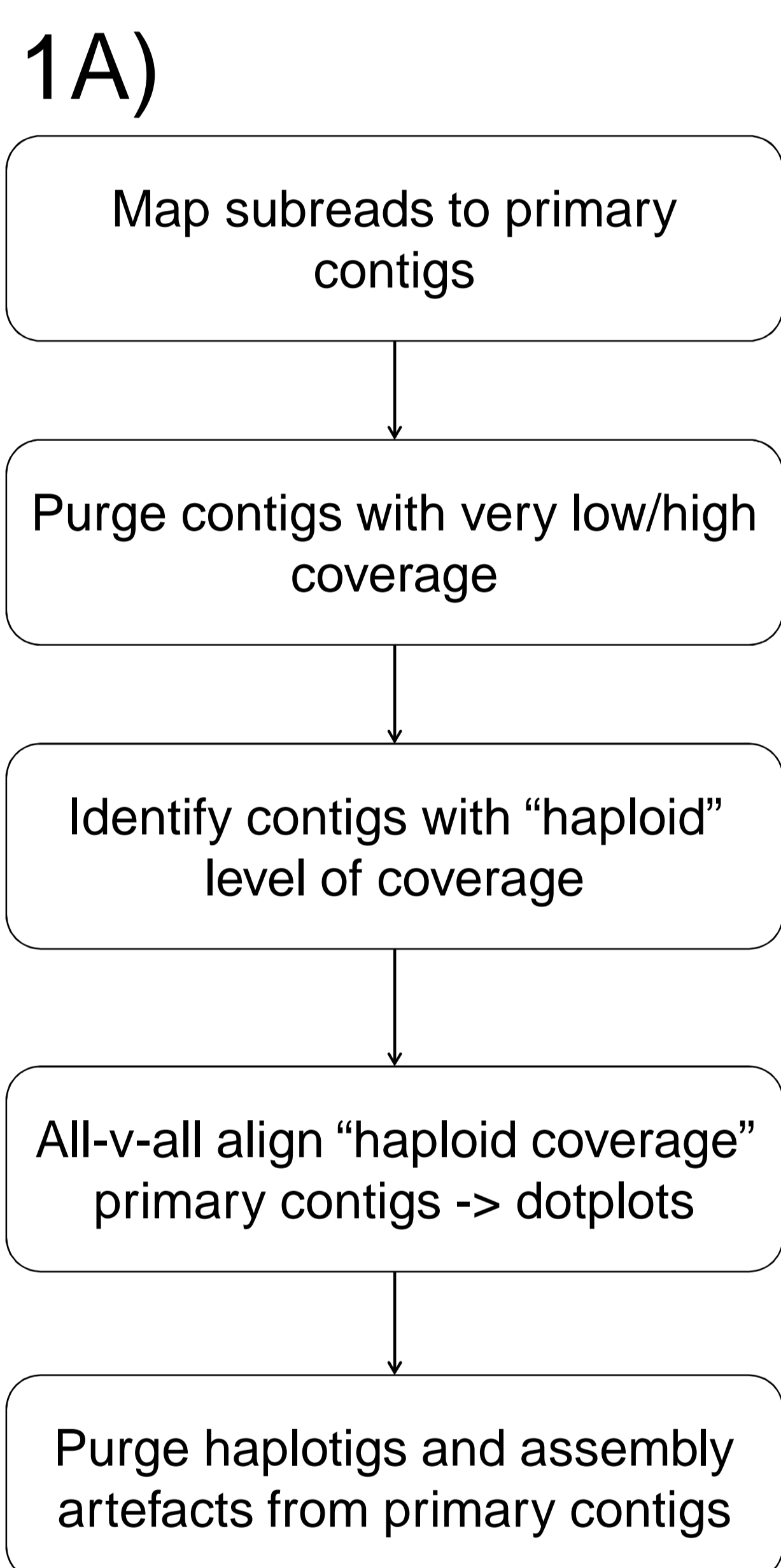
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Background: Until recently only two genome assemblies were publically available for grapevine—both *Vitis vinifera* L. Cv. Pinot Noir (PN). The best available PN genome assembly (Jaillon et al. 2007) is not representative of the genome complexity that is typical of wine-grape cultivars in the field and it is highly fragmented. To assess the genetic complexities of Chardonnay grapevine, assembly of a new *de novo* reference genome was needed. Here we describe a draft assembly using PacBio SMRT sequencing data and PacBio's new phased diploid genome assembler FALCON-Unzip (Chin et al. 2016).

I'm unzipped, now what do I do? 1A) Genome curation pipeline overview 1B) Identifying haplotigs and assembly graph artefacts with dotplots 1C) Assembly stats, comparison and improvement 1D) Validation with BUSCO (Simão et al. 2015)



1C) Assembly stats, comparison and improvement

	Pinot Noir (2007)	Chardonnay genome assembly	
		raw	curated
Size (Mb):	486.197	590.740	494.390
Contigs:	14 634	1 803	978
N50 (bp):	102 851	743 564	922 070
Largest Contig (bp):	653 287	6 339 470	6 347 624

1D) Validation with BUSCO (Simão et al. 2015)

	Chardonnay genome assembly	
	raw	curated
Complete BUSCOs:	892	892
Fragmented BUSCOs:	20	23
Missing BUSCOs:	44	41
Predicted completeness:	~ 95.4 %	~ 95.8 %

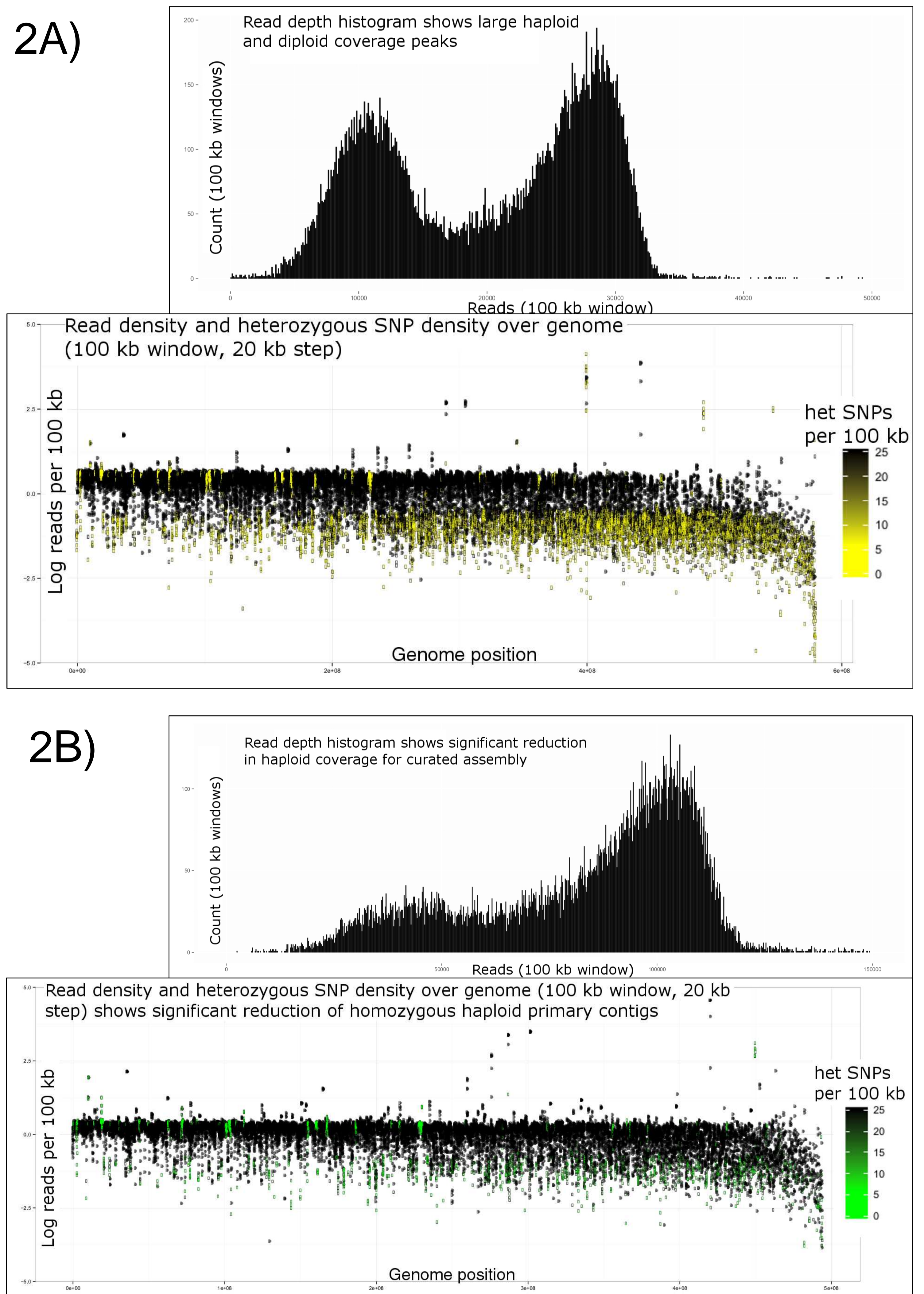
References:

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Read coverage and SNP density used to validate curation. Paired-end illumina reads mapped to primary contigs and heterozygous SNPs called. **2A) raw assembly 2B) curated assembly**



Conclusions:

- Long read sequencing enables highly contiguous phased genome assemblies
- FALCON-Unzip assemblies can be readily curated (even for problematic genomes)
- Phasing aids in genome curation

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The Australian Wine Research Institute