

## Resource Allocation

## How much resources do I need?

### ARTICLE

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### Effect of sequence depth and length in long-read assembly of the maize inbred NC358

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## What do I get with different resources?

**Table 1 Summary statistics for NC358 assemblies.**

Experiment <sup>a</sup>	21k_20x	21k_30x	21k_40x	21k_50x	21k_60x	21k_75x	11k_50x	16k_50x
Subreads size (Gb)	45.62	68.16	91.01	113.89	136.80	171.08	113.63	113.60
Subread coverage	20x	30x	40x	50x	60x	75x	50x	50x
Max read length (kb)	89.6	103.3	103.3	103.3	103.3	103.3	88.3	69.8
Subread N25 (kb)	30.1	30.1	30.1	30.1	30.1	30.1	14.5	21.6
Subread N50 (kb)	21.2	21.2	21.2	21.2	21.2	21.2	11.1	16.8
Corrected reads (Gb)	25.11	48.13	66.05	82.96	88.93	100.90	79.26	80.22
Corrected coverage	11x	21x	29x	37x	39x	44x	35x	35x
Corrected read N50 (kb)	18.42	17.13	17.10	17.25	18.80	20.05	10.37	14.48
Contig number	10,563	2015	641	407	360	327	5683	1036
Contig total (Gb)	1.60	2.11	2.12	2.12	2.13	2.13	2.10	2.12
Longest contig (Mb)	1.06	11.50	47.89	76.00	79.68	78.40	4.37	21.45
Contig N50 (Mb)	0.18	1.82	7.48	16.27	22.12	24.54	0.56	4.24
Longest scaffold (Mb)	198.5	198.7	237.1	237.2	237.1	237.3	205.4	237.6
Superscaffold N50 (Mb)	95.3	96.9	99.2	98.5	99.4	99.2	98.5	99.4
Assembled (%) <sup>b</sup>	70.4%	92.8%	93.3%	93.3%	93.7%	93.7%	92.4%	93.2%
Assembly gaps (%)	24.50%	0.90%	0.43%	0.34%	0.31%	0.31%	2.01%	0.48%
Effective assembly size (Gb) <sup>c</sup>	133	167	170	172	174	175	168	170
Optical map conflict <sup>d</sup>	594	125	56	31	22	21	386	107
Complete BUSCOs <sup>e</sup>	68.0%	95.5%	96.5%	96.4%	96.2%	96.3%	95.7%	96.7%
LTR Assembly Index (LAI)	12.2	19.8	20.4	20.2	20.4	20.6	19.1	21.0
Falcon CPU hour	1563	4162	6363	10,693	12,386	32,950	9721	9224
Falcon RAM (Gb)	75	75	75	75	75	75	75	75
Canu CPU hour	1860	4036	5959	7914	8849	11,520	6400	7174
Canu RAM (Gb)	61	112	149	177	201	120	183	174

<sup>a</sup>Each dataset was assembled only once with the Falcon-Canu hybrid approach (see Methods).

<sup>b</sup>Calculated based on total contig size and the estimated genome size of 2.2724 Gb.

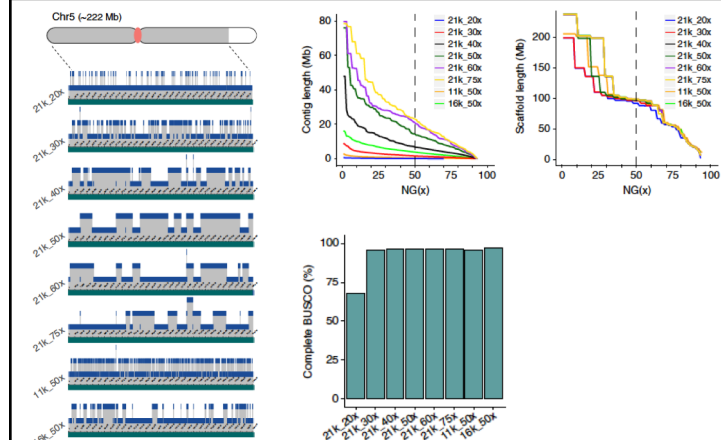
<sup>c</sup>Sum of unique 150-mers.

<sup>d</sup>The optical map was generated using the Direct Label and Stain (DLS) approach with enzyme DLE-1.

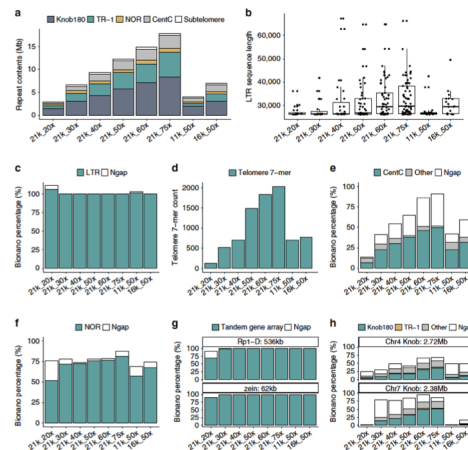
<sup>e</sup>Non-polished assemblies were used to calculate Benchmarking Universal Single-Copy Orthologs (BUSCO) scores.

CPU central processing.

## What do I get with different resources?



## What do I get with different resources?



## How much would it cost for sequencing for a short read assembly in maize?

- Assume you need ~250x depth
- Genome size is ~2.3Gb = 2,300,000,000bp
- Pricing at <https://genomics.umn.edu>
- What machine/chemistry will you use?
- How much will this cost in the least expensive scenario?

## How much would it cost for sequencing for a PacBio assembly in maize?

- Assume you need ~20x depth with HiFi reads
- Genome size is ~2.3Gb = 2,300,000,000bp
- Pricing at <https://genomics.umn.edu>