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► To cite this version:

Tingting Zhu, Le Wang, H el ene Rimbart, Juan Rodriguez, Karin Deal, et al.. Optical maps refine the bread wheat *Triticum aestivum* cv. Chinese Spring genome assembly. *Plant Journal*, 2021, 10.1111/tpj.15289 . hal-03285883

HAL Id: hal-03285883

<https://hal.inrae.fr/hal-03285883>

Submitted on 5 Nov 2021

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RESOURCE

Optical maps refine the bread wheat *Triticum aestivum* cv. Chinese Spring genome assembly

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Received 17 November 2020; revised 12 April 2021; accepted 19 April 2021; published online 24 April 2021.

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SUMMARY

Until recently, achieving a reference-quality genome sequence for bread wheat was long thought beyond the limits of genome sequencing and assembly technology, primarily due to the large genome size and > 80% repetitive sequence content. The release of the chromosome scale 14.5-Gb IWGSC RefSeq v1.0 genome sequence of bread wheat cv. Chinese Spring (CS) was, therefore, a milestone. Here, we used a direct label and stain (DLS) optical map of the CS genome together with a prior nick, label, repair and stain (NLRS) optical map, and sequence contigs assembled with Pacific Biosciences long reads, to refine the v1.0 assembly. Inconsistencies between the sequence and maps were reconciled and gaps were closed. Gap filling and anchoring of 279 unplaced scaffolds increased the total length of pseudomolecules by 168 Mb (excluding Ns). Positions and orientations were corrected for 233 and 354 scaffolds, respectively, representing 10% of the genome sequence. The accuracy of the remaining 90% of the assembly was validated. As a result of the increased contiguity, the numbers of transposable elements (TEs) and intact TEs have increased in IWGSC RefSeq v2.1 compared with v1.0. In total, 98% of the gene models identified in v1.0 were mapped onto this new assembly through development of a dedicated approach implemented in the MAGAAT pipeline. The numbers of high-confidence genes on pseudomolecules have increased from 105 319 to 105 534. The reconciled assembly enhances the utility of the sequence for genetic mapping, comparative genomics, gene annotation and isolation, and more general studies on the biology of wheat.

Keywords: direct label and stain, pseudomolecule, transposable element, gene collinearity, Hi-C.

INTRODUCTION

Wheat provides about one-fifth of the calories and proteins consumed by humans, and is annually planted on an area larger than any other crop. About 95% of the wheat area is planted with bread wheat (*Triticum aestivum*, $2n = 6x = 42$, genome formula AABBDD), which is used for bread, noodles, pastry, cookies and related products; and the remaining 5% is planted with durum wheat (*Triticum durum*

ssp. durum, $2n = 4x = 28$, genome formula AABB), which is used for pasta (Dubcovsky and Dvorak, 2007).

Wheat yields have not been increasing at a rate sufficient to meet anticipated demands (Ray *et al.*, 2013). The development of genomic resources, including a reference-quality genome sequence, is critical for accelerating genetic improvement of wheat. Due to polyploidy, the large sizes of the wheat subgenomes, and the high proportion of repeated sequences (85% of the genome; Wicker *et al.*,

2018), the assembly of a reference-quality genome sequence for a polyploid wheat species was only recently possible as a result of improvements in sequencing and sequence assembly technologies. The first reference-quality polyploid wheat genome sequence was reported for wild emmer wheat (*T. turgidum* ssp. *dicoccoides*, $2n = 4x = 28$, genome formula AABB) acc. 'Zavitan' (Avni *et al.*, 2017), and was followed by the publication of the reference sequences for bread wheat cv. Chinese Spring (CS; IWGSC, 2018) and durum wheat cv. Svevo (Maccaferri *et al.*, 2019).

The wild emmer genome sequence (WEW_v1.0) was assembled from whole-genome shotgun (WGS) reads by NRGene Inc. using their DeNovoMAGIC assembler (Avni *et al.*, 2017). Subsequent alignment of the WEW_v1.0 pseudomolecules to wheat genome-wide optical maps revealed some inconsistencies resulting from mis-orientation or positioning of scaffolds (Dvorak *et al.*, 2018). Using optical maps of Zavitan, about 10% of the WEW_v1.0 assembly was revised in the WEW_v2.0 assembly (Zhu *et al.*, 2019a).

An Illumina/DeNovoMAGIC assembly was also the core resource of the CS reference genome sequence (IWGSC RefSeq v1.0). This assembly was validated, enriched and improved using numerous independent resources (e.g. physical maps and mapped sequence tags) produced between 2008 (Paux *et al.*, 2008) and 2018 by the International Wheat Genome Sequencing Consortium (IWGSC). Genome-wide optical maps were not available for validation of IWGSC RefSeq v1.0, nor were they used in the validations of other CS genome assemblies, such as the assembly built from a combination of Illumina and Pacific Biosciences (PacBio) reads (*Triticum* 3.1; Zimin *et al.*, 2017) recently updated to *Triticum* 4.0 (Alonge *et al.*, 2020).

The first optical map built for a Triticeae species employed a chemistry utilizing a single-strand restriction endonuclease to nick DNA (Hastie *et al.*, 2013; Luo *et al.*, 2017). The nicks were enzymatically repaired and labeled, and the DNA molecules were counterstained for imaging. This chemistry was consequently called nick, label, repair and stain (NLRS). The first genome-wide optical map for the CS genome was built with the NLRS chemistry (Huo *et al.*, 2018). A drawback of this chemistry is that chance clusters of nick sites produce fragile regions in DNA molecules that are prone to breaking during sample preparation. This limits the length of optical contigs that can be built with this chemistry. A recently developed proprietary alternative, direct labeling and staining (DSL) chemistry (Bionano Genomics, San Diego, CA, USA), labels enzyme recognition sites directly without nicking DNA, enabling much longer optical contigs to be assembled.

Here we report a new release of the IWGSC reference genome assembly, namely, IWGSC RefSeq v2.1. A CS optical map based on the DLS chemistry was constructed and used together with the previously constructed NLRS map (Huo *et al.*, 2018) to correct and improve the IWGSC

RefSeq v1.0 assembly. In addition, gaps were closed using contigs built from WGS PacBio SMRT CS long reads (Zimin *et al.*, 2017). The transposable elements (TEs) in the resulting assembly IWGSC RefSeq v2.1 were reannotated, and gene annotation was updated by transferring the previously known gene models (v1.1) using a fine-tuned, dedicated strategy implemented in the Marker-Assisted Gene Annotation Transfer for Triticeae (MAGATT; <https://forge.mia.inra.fr/umr-gdec/magatt>) pipeline.

RESULTS AND DISCUSSION

Comparison of parental CS stock sequences

Chinese Spring is a bread wheat landrace (Liu *et al.*, 2018), and different CS stocks may be polymorphic. It was, therefore, important to determine the extent to which the stock used for the construction of the IWGSC RefSeq v1.0 genome sequence and the stock DV418, used to generate Illumina and PacBio reads for gap filling, differed. Approximately 117 Gb ($\sim 8 \times$ coverage) of DV418 Illumina reads were mapped to the IWGSC RefSeq v1.0 sequence. A total of 25 720 SNPs and 7959 one-base indels were detected. The average diversity between the two stocks was 2.39 polymorphic sites per 1 Mb, which is lower than the rate of sequencing error allowed for a gold-standard assembly (10 per 1 Mb). This low level of detected variation supported the use of the PacBio contigs of DV418 in gap filling.

Construction of the DLS optical map

Images of DLE-1-labeled CS DNA molecules > 150 kb and equivalent to about $136 \times$ genome coverage were collected and assembled into a genome-wide DLS optical map. The map contained 726 contigs with an N50 of 55.01 Mb and a total length of 14.41 Gb (Table 1). The DLS map was more contiguous than the CS NLRS optical map containing 11 727 contigs (N50 = 1.69 Mb; Huo *et al.*, 2018; Table 1).

Comparison of IWGSC RefSeq v1.0 with the DLS map

The 21 IWGSC RefSeq v1.0 pseudomolecules were aligned on the DLS optical map. Some alignments revealed

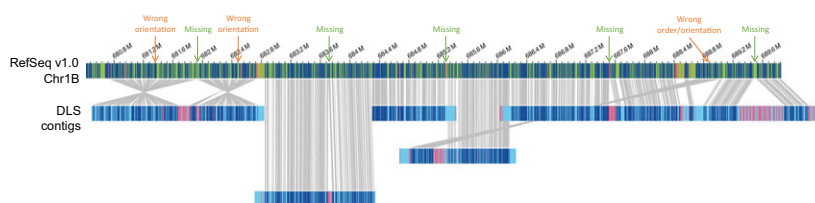
Table 1 Comparison of genome-wide optical maps of the CS genome built with the DLS and NLRS chemistries

Sample	DLS	NLRS
Enzyme	DLE-1	Nt. <i>BspQI</i>
Molecule size N50 (kb)	306	284
Molecule min. length (kb)	150	180
Molecule total length (Gb)	2096	1927
Genome coverage (\times)	136	113
Map contigs (no.)	726	11 727
Map total length (Gb)	14.41	14.15
Map contig length N50 (Mb)	55.01	1.69
Max. contig length (Mb)	331.43	10.52

DLS, direct label and stain; NLRS, nick, label, repair and stain.

Figure 1. Alignment of IWGSC RefSeq v1.0 with the direct label and stain (DLS) optical map.

The alignment of the distal region (from 680 to 690 Mb of the pseudomolecule) of Chr1B of RefSeq v1.0 (green box) to DLS map contigs (blue boxes). Ambiguous sequences, including missing sequences (pale green), mis-orientated scaffolds and mis-ordered scaffolds (orange), were observed.



discrepancies (Figure 1). Alignments on an independently constructed NLRS optical map were used to determine if a discrepancy was due to an error in the sequence or the DLS map. Most of the discrepancies in the sequence were mis-ordered or mis-oriented scaffolds within pseudomolecules, and fewer were caused by errors in the assembly of scaffolds. Discrepancies of 10 kb or less, if present, were not detected as they were below optical map resolution.

Pseudomolecule reconstruction

To correct ordering and orientation problems, pseudomolecule scaffolds were combined with unplaced scaffolds to produce a dataset of 138 546 scaffolds (scf_v0 in Table 2 and Figure 2) that was aligned to the DLS and NLRS optical maps to identify scaffolds with errors in their assembly. The alignments identified 81 conflicts within 72 scaffolds covering 429.408 Mb. The chimeric scaffolds were corrected, thereby increasing the number of scaffolds from 138 546 to 138 634, and decreasing their N50 from 6.871 to 6.775 Mb (scf_v1 in Table 2 and Figure 2).

Super-scaffolds were rebuilt from scf_v1 scaffolds with the aid of the DLS map. The total length of the resulting super-scaffolds was 14.712 Gb with an N50 of 64.014 Mb (scf_v2 in Table 2 and Figure 2). The scf_v2 scaffolds were then aligned to the NLRS optical map, which facilitated additional super-scaffolding of the sequences. The N50 of the resulting super-scaffolds (scf_v3 in Figure 2) reached 72.092 Mb, with the largest recording a length of 364.575 Mb (Table 2).

The scf_v3 super-scaffolds were ordered and oriented through alignment on high-density genetic maps. A total of 673 super-scaffolds (14.311 Gb) sharing two or more markers with genetic maps were anchored on the 21 wheat chromosomes to produce pseudomolecules_v1.1. A

comparison with the pseudomolecules of the IWGSC RefSeq v1.0 assembly showed that 279 more scaffolds (74.960 Mb) from unallocated scaffolds (ChrUn) were now anchored on the chromosomes, 354 scaffolds (469.834 Mb) were re-oriented, and 233 scaffolds (394.868 Mb) were moved into new locations (Table S3).

Gap closing and final refinement of the pseudomolecules

During the scaffold correction stage, the contigs within scaffolds remained largely the same as in pseudomolecules_v1.0, retaining gaps filled with Ns. In addition, there were 4021 gaps of unknown sizes in pseudomolecules_v1.0 (Table 3). Pseudomolecules_v1.1 acquired most of the intra-scaffold gaps and had 647 gaps of unknown sizes (Table 3). To close gaps in pseudomolecules_v1.1, 97 809 PacBio contigs equaling 12.939 Gb and N50 of 264 143 bp were polished with CS DV418 Illumina reads. Polishing corrected 7 221 194 (0.056%) base errors in the PacBio contigs. The PacBio contigs were aligned on the optical maps, and 96 363 of the contigs totaling 12.291 Gb were validated. The validated PacBio contigs were used for gap closure by aligning them on pseudomolecules_v1.1 and substituting N bases with actual nucleotide sequences. Of the 527 170 gaps in pseudomolecules_v1.1, 343 566, including 12 gaps of unknown length (Table 3), were closed to generate pseudomolecules_v2.0. In addition, 91 820 unplaced scaffolds with a total length of 351.583 Mb were assigned to ChrUn. The pseudomolecules_v2.0 along with ChrUn constituted the intermediate IWGSC RefSeq v2.0 assembly.

Comparison of IWGSC RefSeq v2.0 with IWGSC RefSeq v1.0 detected SNPs and single-base indels in 12 382 coding sequences (CDSs) from 9416 genes. Assuming that the differences were introduced into the assembly with the PacBio reads, the sequences harboring these CDSs were

Table 2 Scaffolding steps using optical maps

	scf_v0	Chimeras resolved (scf_v1)	Scaffolding using DLS map (scf_v2)	Scaffolding using NLRS map (scf_v3)
Scaffolds (no.)	138 546	138 634	134 957	134 925
Max. length (bp)	45 793 851	45 793 851	364 574 574	364 574 574
Total length (bp)	14 533 405 144	14 533 405 058	14 711 983 205	14 710 380 608
Sequence N50 (bp)	6 870 518	6 774 542	64 014 232	72 091 519
N%	1.80	1.80	2.99	3.00

DLS, direct label and stain; NLRS, nick, label, repair and stain.

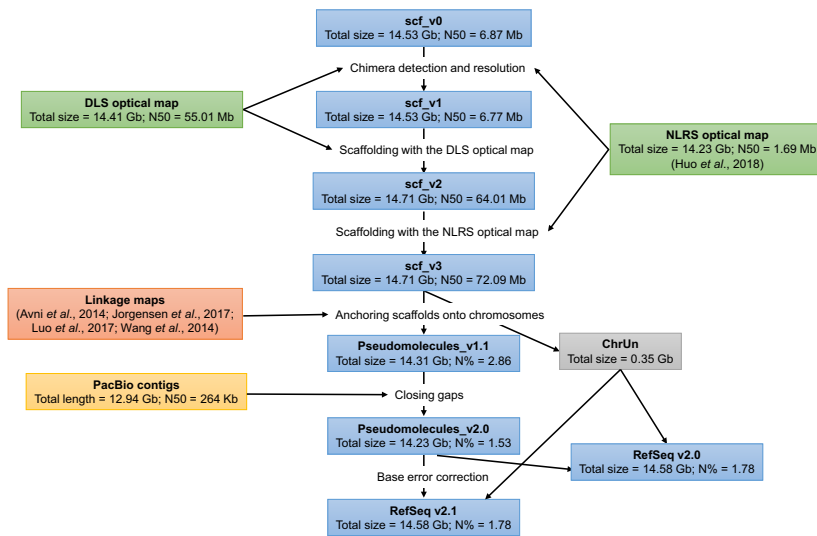


Figure 2. Overview of the strategy for reconstructing the IWGSC RefSeq v2.1 assembly. N% refers to the number of N bases placed into gaps in the assembly.

replaced with corresponding contigs from IWGSC RefSeq v1.0. This final refinement produced the final genome assembly IWGSC RefSeq v2.1 (Figure 3; Table 4).

Gene reannotation

Evaluation of the new sequence assembly showed that the number of differences between IWGSC RefSeq v1.0 and IWGSC RefSeq v2.1, primarily arising from gap filling, was too great to allow for a simple correlation between the two versions to generate new gene coordinates. Gene sequence alignments were, therefore, re-computed to map the gene annotation to the new assembly.

Table 3 Numbers of gaps of unknown sizes in pseudomolecules_v1.0 to v2.0

Chromosome	v2.0	v1.1	v1.0
Chr1A	13	13	145
Chr2A	18	19	171
Chr3A	7	7	186
Chr4A	34	34	236
Chr5A	13	13	151
Chr6A	11	11	158
Chr7A	15	15	196
Chr1B	45	45	191
Chr2B	38	38	273
Chr3B	48	48	237
Chr4B	59	63	204
Chr5B	54	55	250
Chr6B	61	62	276
Chr7B	58	58	281
Chr1D	16	16	133
Chr2D	16	17	167
Chr3D	19	20	159
Chr4D	24	26	115
Chr5D	27	28	170
Chr6D	27	27	121
Chr7D	32	32	201
Total	635	647	4021

Before transferring the gene annotation onto the new genome assembly, it was updated to IWGSC Annotation v1.2 by integrating a set of 117 novel genes and 81 microRNAs (miRNAs), many of which had been manually curated by the wheat community. The new annotation release containing 108 010 HC and 161 535 LC gene models (Table S1) was used to annotate IWGSC RefSeq v2.1.

Mapping a query gene on the whole wheat genome often leads to spurious alignments because of the high number of homologous gene copies. In order to mitigate this problem, the interval that harbored a gene to be mapped was first identified using Insertion Site-Based Polymorphisms (ISBP) markers, which mark junctions between neighboring TEs that are present in both the IWGSC RefSeq v1.0 and IWGSC RefSeq v2.1 assemblies. ISBPs are 150-mers representing a unique sequence shaped by the junction of a TE and its insertion site (De Oliveira *et al.*, 2020; Paux *et al.*, 2010; Rimbart *et al.*, 2018). Because ISBPs can be derived at each TE extremity, they are extremely abundant and well adapted for identifying corresponding loci across different assembly versions. In wheat, almost all genes are closely flanked by a pair of unique ISBPs.

A total of 5 394 172 ISBPs were designed on IWGSC RefSeq v1.0 and mapped to IWGSC RefSeq v2.1 (Table S1). Over 90% of these ISBPs (4 908 316) mapped fully and uniquely with no mismatches, giving an average density of one ISBP every 3 kb. An ISBP-flanked interval on IWGSC RefSeq v2.1 was assigned for 264 876 (98.3%) of the 269 545 gene models, corresponding to 107 877 high-confidence (HC) and 161 668 low-confidence (LC) gene models. They represented 207 575 intervals containing between 1 and 4 genes, with 83% of the intervals containing a single gene (Table S1). The gene intervals spanned 668, 683 and 512 Mb of the A-, B- and D-subgenome sequences, respectively, and 136 Mb of ChrUn scaffolds. The average size of an interval was 9.6 kb. A total of

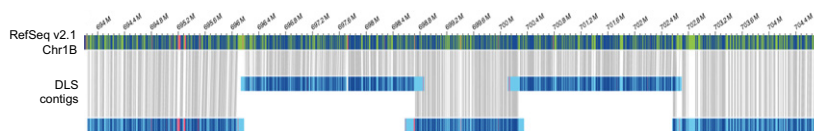


Figure 3. IWGSC RefSeq corrected with the direct label and stain (DLS) optical map.

Alignments of the region of the Chr1B pseudomolecule corresponding to that shown in Figure 1 (green box) to the DLS map contigs (blue boxes) show that most of the ambiguous regions have been resolved in this region of the IWGSC RefSeq v2.1 assembly.

Table 4 Comparison of the total and effective lengths (excluding Ns) of IWGSC RefSeq v2.1 pseudomolecules with those of IWGSC RefSeq v1.0

Chr ID	RefSeq v2.1			RefSeq v1.0			N%
	Length (bp)	Effective length (bp)	N%	Length (bp)	Effective length (bp)	N%	
Chr1A	598 660 471	590 447 024	1.37	594 102 056	585 682 930	1.42	
Chr2A	787 782 082	777 846 456	1.26	780 798 557	769 534 243	1.44	
Chr3A	754 128 162	743 748 734	1.38	750 843 639	739 934 526	1.45	
Chr4A	754 227 511	741 420 734	1.70	744 588 157	733 095 901	1.54	
Chr5A	713 360 525	704 968 565	1.18	709 773 743	699 527 711	1.44	
Chr6A	622 669 697	614 336 433	1.34	618 079 260	609 021 784	1.47	
Chr7A	744 491 536	733 447 483	1.48	736 706 236	725 041 360	1.58	
Chr1B	700 547 350	688 076 938	1.78	689 851 870	678 654 964	1.62	
Chr2B	812 755 788	798 522 160	1.75	801 256 715	789 089 512	1.52	
Chr3B	851 934 019	831 834 594	2.36	830 829 764	817 347 104	1.62	
Chr4B	673 810 255	665 179 646	1.28	673 617 499	663 297 446	1.53	
Chr5B	714 805 278	704 718 176	1.41	713 149 757	701 419 691	1.64	
Chr6B	731 188 232	717 245 234	1.91	720 988 478	709 167 234	1.64	
Chr7B	764 081 788	749 505 598	1.91	750 620 385	737 902 257	1.69	
Chr1D	498 638 509	491 298 183	1.47	495 453 186	486 750 394	1.76	
Chr2D	656 544 405	647 090 398	1.44	651 852 609	640 662 955	1.72	
Chr3D	619 618 552	611 768 249	1.27	615 552 423	604 864 457	1.74	
Chr4D	518 332 611	511 996 999	1.22	509 857 067	501 652 949	1.61	
Chr5D	569 951 140	563 314 725	1.16	566 080 677	555 817 680	1.81	
Chr6D	495 380 293	488 354 375	1.42	473 592 718	465 469 117	1.72	
Chr7D	642 921 167	633 773 979	1.42	638 686 055	626 564 746	1.90	
ChrUn	351 582 993	308 528 982	12.25	480 980 714	431 079 926	10.37	
Total	14 576 989 303	14 316 999 506	1.78	14 547 261 565	14 271 578 887	1.90	

241 201 (90%) genes with identical sequence between the two assemblies were unambiguously mapped, and 22 578 (8.5%) genes exhibited nucleotide differences (mismatches and/or indels) that affected only introns or UTRs. Together, they represented 263 779 (98%) genes from IWGSC RefSeq v1.0. Of these, 31 520 genes (12%) would not have transferred accurately if GMAP against the whole IWGSC RefSeq v2.1 had been used, demonstrating the importance of reducing the genome complexity before mapping as implemented in our pipeline MAGATT (<https://forgemia.inra.fr/umr-gdec/magatt>).

Of the remaining 2% of IWGSC RefSeq v1.0 genes, 2974 (1%) were aligned with mismatches/indels in the CDSs (or only partially aligned in rare cases), meaning that their sequence has changed in IWGSC RefSeq v2.1 compared

with v1.0. The remaining 2792 IWGSC RefSeq v1.0 genes (1%) could not be identified in IWGSC RefSeq v2.1. A total of 1258 genes previously on the ChrUn scaffolds were assigned to chromosomes in IWGSC RefSeq v2.1.

All genes were renamed to more accurately reflect the position of the gene and the annotation. For example, in a new name *TraesCS1A03G000000[LC]*, 1A and 03 specify the location of a low-confidence gene on chromosome 1A and the third annotation version, respectively.

In total, the new release IWGSC RefSeq Annotation v2.1 contains 266 753 genes comprising 106 913 HC genes and 159 840 LC genes (Table S1). Altogether, 105 534 HC and 155 624 LC genes were located on the pseudomolecules, and 1379 HC and 4216 LC genes were located on scaffolds assigned to ChrUn (Table S1).

Table 5 Numbers of collinear genes and syntenic blocks between subgenomes in the pseudomolecules of IWGSC RefSeq v1.0 and IWGSC RefSeq v2.1

Subgenome comparison	Collinear genes (no.)			Syntenic blocks (no.)		
	v1.0	v2.1	Difference (v2.1-v1.0)	v1.0	v2.1	Difference (v2.1-v1.0)
A and B	41 999	42 479	480	546	484	-62
A and D	44 280	45 008	728	581	500	-81
B and D	43 704	44 484	780	510	442	-68

Gene collinearity in homoeologous pseudomolecules

The refinement of the pseudomolecules was expected to improve gene collinearity between homoeologous pseudomolecules. The numbers of collinear genes and syntenic blocks (at least five genes were required to call a synteny block) between the A-, B- and D-subgenome homoeologous pseudomolecules were counted in the IWGSC RefSeq v2.1 and IWGSC RefSeq v1.0 assemblies to assess this expectation. In the pairwise subgenome comparisons, the number of collinear genes was greater ($P = 0.019$, two-tailed paired t -test, $N = 3$) in IWGSC RefSeq v2.1 than in IWGSC RefSeq v1.0 (Table 5). Concomitantly, there was a reduced fragmentation of syntenic blocks ($P = 0.006$, two-tailed paired t -test, $N = 3$), indicating higher collinearity between homoeologous chromosomes in the IWGSC RefSeq v2.1 assembly than in IWGSC RefSeq v1.0 (Table 5).

De novo TE annotation

De novo annotation of TEs with CLARITE (Daron *et al.*, 2014) annotated 4 199 592 TEs belonging to 506 families. The total length of TEs increased from 11 921 309 743 to 12 092 094 168 bp. As a result of the increase in pseudomolecule length in IWGSC RefSeq v2.1 compared with v1.0 (Table 6), the percentage of the CS genome accounted for by TEs (85.0%) in IWGSC RefSeq v2.1 was nearly identical to that reported for IWGSC RefSeq v1.0 (84.7%). Because the same approach, tools and criteria were used, the percentages of the genome represented by TEs and by individual TE super-families and families were very similar in the IWGSC RefSeq v1.0 (Wicker *et al.*, 2018) and v2.1 assemblies (Table 6). ISBPs can be used as a measure of similarity of the TE content (De Oliveira *et al.*, 2020; Rimbart *et al.*, 2018). Overall, 99.9% of ISBPs was shared by the two genome sequences.

Table 6 Comparison of the TE content in IWGSC RefSeq v1.0 and IWGSC RefSeq v2.1, and among subgenomes of IWGSC RefSeq v2.1

	IWGSC RefSeq v1.0	IWGSC RefSeq v2.1	IWGSC RefSeq v2.1 subgenomes		
			A	B	D
Pseudomolecules (bp)	14 066 280 851	14 225 829 371	4 975 319 984	5 249 122 710	4 001 386 677
Repeats (bp)	11 921 309 743	12 092 094 168	4 283 220 399	4 455 614 668	3 353 259 101
TEs (%) ^a	84.75	85.00	86.09	84.88	83.80
Class 1 ^b	67.6	66.9	71.0	66.8	62.2
<i>Gypsy</i> (RLG)	46.7	46.1	48.6	46.2	41.0
<i>Copia</i> (RLC)	16.7	16.5	19.4	16.0	16.3
Unclass. LTR RTs	3.24	3.26	2.21	3.47	3.74
LINEs (RIX)	0.9	1.05	0.81	1.12	1.09
SINEs (SIX)	0.01	0.01	0.01	0.01	0.01
Class 2 ^c	16.5	17.0	14.4	17.0	20.7
<i>CACTA</i> (DTC)	15.5	15.9	13.2	15.9	19.5
Mutator (DTM)	0.38	0.44	0.35	0.43	0.55
Unclass. with TIRs (DTX)	0.21	0.24	0.24	0.24	0.25
Harbinger (DTH)	0.16	0.18	0.17	0.17	0.19
Mariner (DTT)	0.16	0.17	0.15	0.17	0.19
Unclass. class 2 (DXX)	0.06	0.06	0.26	0.07	0.06
hAT (DTA)	0.006	0.009	0.009	0.011	0.008
Helitrons (DHH)	0.004	0.01	0.011	0.010	0.010
Unclass. repeats	0.68	0.95	0.78	0.93	0.88

RT, retrotransposon; TE, transposable element; TIR, terminal inverted repeats.

^a3-letter TE codes in parentheses, as defined in Wicker *et al.* (2007).

^bRNA transposons.

^cDNA transposons.

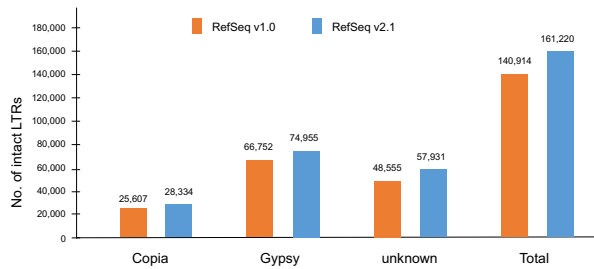


Figure 4. Numbers of intact LTR retrotransposons (RTs) in IWGSC RefSeq v1.0 and IWGSC RefSeq v2.1.

In IWGSC RefSeq v2.1, TEs accounted for similar percentages of the total lengths of the A-, B- and D-subgenome pseudomolecules (Table 6). The largest quantity of TEs was in the B-subgenome and the smallest was in the D-subgenome, mirroring the total lengths of pseudomolecules in the three subgenomes (Table 6). The percentages of subgenome lengths represented by individual super-families and families were similar among the subgenomes (Table 6), although in the D-subgenome the *Gypsy* super-family was under-represented and the *CACTA* super-family was over-represented relative to their percentages in the A- and B-subgenomes.

Gap closure improved the contiguity of TE sequences in IWGSC RefSeq v2.1 compared with v1.0. This was apparent from the numbers of intact LTR retrotransposons (RTs). There were more intact *Copia*, *Gypsy* and unknown LTR RTs ($P < 0.0001$ in each LTR RT class, two-tailed t -test with Bonferroni correction, $N = 21$) in IWGSC RefSeq v2.1 than in v1.0 (Figure 4; Table S4).

Comparisons of IWGSC RefSeq v2.1 with CS BAC sequences

Comparison of a genome sequence assembly with sequences of bacterial artificial chromosomes (BACs) assembled from reads produced with Sanger sequencing technology has been used in the past to assess quality of a genome sequence assembly (Luo *et al.*, 2017). The IWGSC RefSeq v2.1 sequence was compared with sequences of 15 CS BACs sequenced to high quality. Fourteen of these BACs could be mapped onto the corresponding pseudomolecules with mean identity $> 99.5\%$ and coverage $\geq 99.99\%$. This sequence identity was comparable to that of the 99.75% identity reported for an analogous comparison made with the *Ae. tauschii* Aet v4.0 genome sequence assembly (Luo *et al.*, 2017). Clone GU817319.1 was for an unknown reason exceptional; it mapped with only 98.85% identity and 89.92% coverage to the IWGSC RefSeq v2.1 sequence (Table S2).

Chromosome conformation capture sequencing (Hi-C) analysis

Hi-C was used to compare the structural integrity of the IWGSC RefSeq v1.0 and 2.1 genome assemblies at the

chromosome scale level. Hi-C contact matrices (Figure S1) showed the expected diagonal–antidiagonal matrix indicative of the Rab1 configuration of interphase nuclei (Mascher *et al.*, 2017). Contact matrices of the IWGSC RefSeq v1.0 assembly (Figure S2) indicated misassemblies in pseudomolecules Chr2A, Chr3A, Chr4A, Chr2B, Chr7B and Chr3D, which were corrected in the IWGSC RefSeq v2.1 assembly. A single discrepancy remained in the pericentromeric region of chromosome 4A. Conflicting signals of centromeric histone H3 (CENH3) localization from chromatin immunoprecipitation sequencing was reported for CS chromosome 4A (IWGSC, 2018). Whether heterogeneity in CS seed sources or some other factor is the cause of these signals is not clear.

Comparisons with the Triticum 4.0 assembly

IWGSC RefSeq v2.1 was compared with the recent CS assembly Triticum 4.0 (Alonge *et al.*, 2020), which was produced from CS DV418 sequence data with the assistance of IWGSC RefSeq v1.0. An analysis of a 10-Mb interval on chromosome 1A (Chr1A) provides an example of differences between the assemblies (Figure 5). In that region, the DLS optical map indicated that the IWGSC RefSeq v1.0 sequence harbored a 650-kb inversion and 602 kb of missing sequence (Figure 5a). Both the inversion and missing sequence also appeared in the Triticum 4.0 assembly (Figure 5b), but both were rectified in RefSeq v2.1 by changing the orientation of the 650-kb segment and inserting Ns equivalent to 602 kb, respectively (Figure 5c). There were only three DLE-1 restriction sites in this 602-kb interval (Figure 5a–c), while there should have been about 90 sites based on the genome average. The unusual structure of this 602-kb region is most likely due to the presence of repeats, which failed to assemble well in this region of Chr1A. In addition to the two rearrangements incorporated from IWGSC RefSeq v1.0, approximately 1.75 Mb of extra sequence was present in Triticum 4.0 in this interval, including a 250-kb sequence that our analysis assigned to Chr7A (indicated by the orange arrow in Figure 5b).

Aligning the three CS sequences on the DLS optical map indicated that many of the ambiguities in sequence order, sequence orientation and chromosome location were transferred from IWGSC RefSeq v1.0 to Triticum 4.0, while additional conflicting arrangements were introduced by the Triticum 4.0 assembly process itself (Table S5). In addition to conflicts in sequence arrangements, about 0.31 Gb of sequences from other chromosomes appear misassigned to pseudomolecules in Triticum 4.0, about 0.98 Gb of the sequence was missing (Table S5), and about 1.54 Gb of the Triticum 4.0 sequence could not be aligned with the DLS optical map. This may partially account for the total length of the Triticum 4.0 pseudomolecules exceeding that of the IWGSC RefSeq v2.1 by 6%.

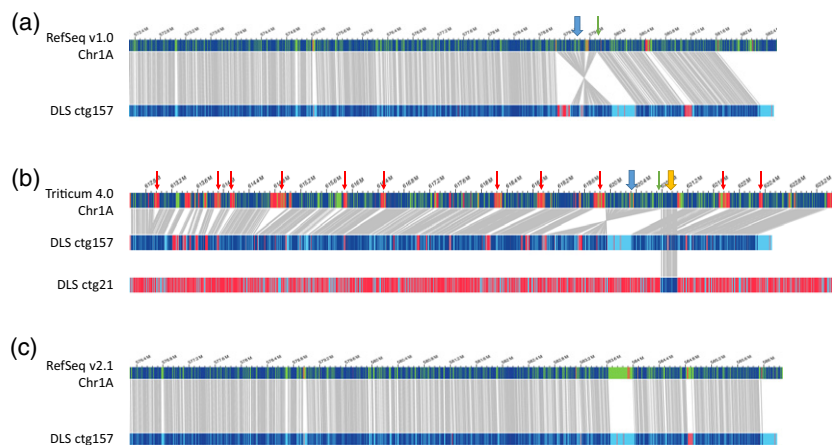


Figure 5. Alignments of a 10-Mb interval of Chr1A pseudomolecules in IWGSC RefSeq v1.0 (a), Triticum 4.0 (b) and IWGSC RefSeq v2.1 (c) with the DLS optical contigs.

Indicated are extra sequences (red arrows), missing sequences (green arrows), inverted sequences (blue arrows) and sequences from other chromosomes (orange arrow).

Concluding remarks

The deployment of optical maps plays two main roles in the assembly of a genome. First, an optical map represents an independent means for validating a sequence assembly. The deployment of the two CS optical maps validated about 90% of the IWGSC RefSeq v1.0 assembly (IWGSC, 2018), confirming that the assembly was indeed of high quality. Second, as the lengths of DLS optical contigs greatly exceed the lengths of sequence scaffolds, a DLS optical map is a valuable tool for scaffolding and super-scaffolding during sequence assembly. The deployment of the CS optical maps anchored 279 previously unplaced scaffolds (ChrUn) onto the pseudomolecules, reoriented 354 scaffolds and relocated 233 scaffolds, thus guiding the revision of approximately 10% of the genome sequence. Our experience illustrates the value of using multiple genome-wide optical maps to guide sequence assembly of complex genomes.

Closing gaps in the IWGSC RefSeq v1.0 sequence was another refinement of the assembly. The combined effect of anchoring 279 unplaced scaffolds (74.960 Mb) and gap filling with PacBio long-read sequences increased the length (excluding Ns) of the pseudomolecules by approximately 168 Mb (1.19%), from 13.840 Gb in IWGSC RefSeq v1.0 to 14.008 Gb in IWGSC RefSeq v2.1. Closing gaps also increased the number of intact LTR RTs by 13% as well as the overall number of identified TEs. It increased the number of HC genes on the pseudomolecules from 105 319 in Annotation v1.2 to 105 534 HC genes in Annotation v2.1, and reduced the total number of genes on unassigned scaffolds from 9449 in Annotation v1.2 to 5595 in Annotation v2.1 (Table S1).

Compared with IWGSC RefSeq v1.0, the gene space in the IWGSC RefSeq v2.1 pseudomolecules more faithfully reflects the divergence of the wheat subgenomes from a common ancestral genome in that a greater number of

collinear genes and a reduced fragmentation of syntenic blocks were found in the IWGSC RefSeq v2.1 pseudomolecules. These refinements of both the TE and gene space argue for the inclusion of long reads into a reference-genome sequence assembly. Importantly, we found that it was best to use PacBio contigs validated by optical maps to close gaps. This highlights the importance of combining multiple independent sources of information to produce a consensus that is less likely to suffer from issues related to a particular technology.

Revisions present in the IWGSC RefSeq v2.1 will enhance the utility of the genome sequence in breeding and research applications. Correcting scaffold order and orientation is critical for comparative studies and genetic mapping of quantitative trait loci, as well as any other approaches relying on linkage disequilibrium such as Genomic Prediction. The assignment of previously unplaced scaffolds into correct locations of the pseudomolecule and gap closure allows the genomic regions underpinning trait associations to be more fully studied, and facilitates a more complete gene annotation. Sizing gaps of previously unknown sizes allows local scale to be assessed, and the cross-validation of sequence with optical maps enables the identification of remaining problem regions.

Despite the refinements of the IWGSC RefSeq v2.1 assembly reported here, it is evident that more remains to be done. This study corrects most of the macro-scale assembly issues that were present in v1.0. Issues that are more difficult to resolve remain in localized regions and will require integration of new data types, especially in areas where the resolution of optical maps was inadequate to detect or resolve discrepancies. Also, some small scaffolds could not be aligned on the optical maps and were discarded. Some optical map contigs and aligned scaffolds may have been incorporated into the pseudomolecule either arbitrarily or were kept as in IWGSC RefSeq v1.0,

because of a lack of markers or ambiguity in genetic maps. Moreover, 351.583 Mb of assembled scaffolds remain unplaced and assigned to ChrUn. The sequences of the centromeric regions, rRNA loci, telomeric and subtelomeric regions, regions containing satellite DNA, and other hard-to-assemble regions remain largely unanchored and/or collapsed. The successful assembly of centromeric and telomeric regions of a walnut hybrid (~1.3 Gb genome size) with a combination of short- and long-read sequencing technologies and optical maps (Zhu *et al.*, 2019b) suggests that with these and other technologies the IWGSC sequence of the CS genome can be further refined. The IWGSC RefSeq v2.1 assembly we are now providing is a major advance for applied and basic applications.

EXPERIMENTAL PROCEDURES

Plants

Triticum aestivum cv. Chinese Spring DV418 was used to construct the DLS optical map and to produce the WGS PacBio SMRT long reads. DV418 is a doubled haploid of a CS accession provided by D.R. Knott, University of Saskatchewan, Saskatoon, Canada.

Comparison of the parental CS stocks

Illumina reads of CS stock DV418 (Zimin *et al.*, 2017) were downloaded from NCBI. These reads were mapped to the IWGSC RefSeq v1.0 genome assembly (IWGSC, 2018) using BWA-MEM (Li and Durbin, 2010) with default settings. After removing polymerase chain reaction duplicates, variant calling was performed and filtered with $QC \geq 20$ and depth ≥ 3 using SAMtools (Li *et al.*, 2009) and BCFtools (Narasimhan *et al.*, 2016). Only homozygous sites were kept and analyzed.

Construction of the CS DLS optical map

Seeds of CS were germinated and grown in the dark, and young leaves were collected from these seedlings. Ultra-high-molecular-weight (uHMW) DNA was isolated using the Plant DNA Isolation Kit (Bionano Genomics, San Diego, CA, USA). The uHMW DNA molecules were labeled with the DLE-1 enzyme (Bionano Genomics) and were then stained with the Bionano Prep™ DLS Kit (Bionano Genomics). A consensus optical map was *de novo* assembled with the Assembler tool of the Bionano Solve v3.3 package using significance cutoffs of $P < 1 \times 10^{-11}$ to generate draft consensus contigs, $P < 1 \times 10^{-12}$ for draft consensus contig extension, and $P < 1 \times 10^{-16}$ for final merging of the draft consensus contigs. A recipe of 'non-haplotype', 'noES' and 'noCut' was chosen. The initial optical map was then checked for potential chimeric contigs. Problematic regions were manually analyzed by altering contig assembly stringency, and by interrogating the NLRs CS optical map and DNA sequences.

Reconstruction of pseudomolecules

The stepwise process of pseudomolecule reconstruction is summarized in Figure 2.

Chimera resolution. The 21 pseudomolecules of the IWGSC RefSeq v1.0 genome assembly (IWGSC, 2018) were split into individual scaffolds at positions comprised of 100 Ns, which

represented gaps of unknown lengths between scaffolds. The scaffolds were combined with unplaced scaffolds (ChrUn) of the IWGSC RefSeq v1.0 assembly. All scaffolds were validated by aligning them on the CS DLS optical map using RefAligner tool of the Bionano Solve v3.3 package (Bionano Genomics), with an initial alignment cutoff of $P < 1 \times 10^{-10}$. Alignments in which a scaffold disagreed with the DLS optical map were aligned to the CS NLRs optical map (Huo *et al.*, 2018) to determine if the sequence or the DLS contig was incorrectly assembled. Incorrectly assembled scaffolds were disjoined.

Super-scaffolding. Conflict-free scaffolds were super-scaffolded using the Hybrid Scaffold pipeline of Bionano Solve v3.3 package (Bionano Genomics), with an alignment cutoff of $P < 1 \times 10^{-10}$. Gaps were filled with the number of Ns corresponding to the estimated length in bp between restriction sites flanking a gap on the DLS optical map. The NLRs optical map was used to further super-scaffold the sequences, which produced the final super-scaffolds.

Anchoring of super-scaffolds onto the chromosomes. High-density genetic maps of hexaploid wheat (Wang *et al.*, 2014), wild emmer (Avni *et al.*, 2014; Jorgensen *et al.*, 2017) and *Ae. tauschii* acc. AL8/78 (Luo *et al.*, 2017) were used to determine the order and orientation of super-scaffolds on each chromosome. The flow-sorted chromosome arm DNA (Chromosome Survey Sequencing) sequences (IWGSC, 2014) were used to confirm chromosome arm and subgenome allocation of super-scaffolds. Super-scaffolds were then linked with 100 Ns and anchored onto the 21 CS chromosomes.

Additional gap closure with PacBio contigs. Approximately 545 Gb of raw reads (~36 × coverage) of CS DV418 (NCBI SRX2994550) generated with PacBio sequencing technology were assembled with Falcon (Zimin *et al.*, 2017). To polish the PacBio contigs further, raw CS DV418 Illumina reads (NCBI SRX2994097) were downloaded from NCBI and filtered using Sickle v1.33 (Joshi and Fass, 2011) with default parameters. The clean Illumina reads were mapped on to the PacBio contigs using BWA v0.7.16 (Li and Durbin, 2010) and unique alignments were kept. 'SNP' or 'INDEL' calls were corrected with Pilon v1.18 (Walker *et al.*, 2014). These PacBio contigs were validated with the CS DLS and NLRs optical maps. Only those contigs that agreed with both maps were used for gap closing. The validated contigs were then aligned to pseudomolecules_v1.1 using BLAST. When a region in pseudomolecules_v1.1 aligned unambiguously with a PacBio contig and contained gap(s), it was replaced by the corresponding PacBio contig; thus, N bases in the pseudomolecules were filled with the actual nucleotide sequence, which resulted in Pseudomolecules_v2.0. The ChrUn super-scaffolds were created by linking adjacent unanchored scaffolds in a random order with 100 Ns. The pseudomolecules_v2.0 along with ChrUn constituted the IWGSC RefSeq v2.0 assembly.

Final refinement of the RefSeq v2.0 assembly. Primary transcripts of 133 744 HC genes (IWGSC, 2018) were mapped to RefSeq v2.0 to obtain new coordinates. Alignments of CDSs showing mis-matches and/or indels were labeled as 'problematic' and as needed to be corrected because of containing nucleotide errors due to the integration of PacBio contigs into the new assembly. Original contigs from the IWGSC RefSeq v1.0 harboring the 'problematic' CDSs were aligned against the RefSeq v2.0 using minimap2 (Li, 2018) with the default parameters and these regions in IWGSC RefSeq v2.0 were reverted to their original version (v1.0),

leading to a final genome assembly IWGSC RefSeq v2.1 (Figure 2).

TE annotation

Intact LTR RTs were identified with the LTR_finder (Xu and Wang, 2007). A non-redundant LTR library was generated. LTRs were annotated by searches with RepeatMasker 4.0.9 (<http://www.repeatmasker.org>) against the non-redundant LTR library, Repbase (Bao *et al.*, 2015) and TREP (Wicker *et al.*, 2002) databases.

To annotate all TEs, *de novo* TE models were generated using CLARITE and the curated wheat TE library ClariTeRep (Daron *et al.*, 2014) as described previously (IWGSC, 2018; Wicker *et al.*, 2018). CLARITE uses RepeatMasker for raw similarity search, to merge adjacent TE fragments into complete TE models, and eventually identifies nested insertions. The TE content similarity between v1.0 and v2.1 was compared by searching for overlaps with BEDTools intersect (Quinlan and Hall, 2010) between v1.0 ISBPs hits on v2.1 (from BAM file) and annotated v2.1 TEs (.gff3 file). Statistics and metrics for TE families were produced using bash command lines.

Gene annotation

Integration of manually curated genes (Annotation v1.2). Before performing a gene transfer onto the new genome assembly, a set of 198 genes manually curated by experts and communicated in 2020 to the IWGSC RefSeq Annotation group was integrated. A method described in IWGSC (2018) was applied to update gene annotation, which led to include an additional 117 novel genes and to correct 81 miRNAs. We generated a new release: Annotation v1.2, containing 108 010 HC and 161 535 LC gene models, which was used for gene transfer on the new assembly.

Transfer of curated genes on RefSeq v2.1. A Snakemake pipeline called MAGATT (Marker-Assisted Gene Annotation Transfer for Triticeae) was developed to transfer gene annotation from the IWGSC Annotation v1.2 onto the IWGSC RefSeq v2.1 assembly.

The pipeline performs this transfer automatically. First, 150-bp tags corresponding to ISBP markers in IWGSC RefSeq v1.0 were designed as described in De Oliveira *et al.* (2020), and mapped on IWGSC RefSeq v2.1 using BWA (Li and Durbin, 2010). Only ISBPs that were fully mapped with no mismatch and with a mapping quality of at least 30 (i.e. uniquely mapped) were selected. For each gene, the pipeline identified the closest 5' and 3' ISBPs in order to predict the coordinates of the smallest target interval harboring a query gene of IWGSC RefSeq v2.1. If there was no 5' or 3' ISBPs, the start/end coordinates of the chromosome were used as borders. Only intervals between 500 bp and 10 Mb were considered. When such an interval was defined, BLAT (Kent, 2002) was used to align the genomic sequence of each query gene (from transcription start site to termination site) on the extracted sequence of the target interval. If a full-length perfect match was observed (100% identity over 100% of the gene length, including Ns with BLAT option '-extendThroughN' between the two assembly versions, or if a full-length match with only mismatches was observed (but no indels), the pipeline calculated the positions of all gene-related features (mRNAs, exons, CDSs and UTRs) on IWGSC RefSeq v2.1. If indels were observed (that may occur because of the gap-filling step), the pipeline ran GMAP to perform a spliced alignment of the query mRNA on the target interval and to retrieve new feature positions. Finally, if no ISBP-based interval

was found to encompass a gene, MAGAAT ran GMAP on the whole genome assembly. Details can be found in Table S1.

Assessing the quality of IWGSC RefSeq v2.1

The sequences of 15 CS BACs (Table S2) assembled from Sanger reads were downloaded from the NCBI. The BACs originated from chromosomes 3A, 3B, 5A and 5D. They were mapped on the Chr3A, Chr3B, Chr5A and Chr5D pseudomolecules of IWGSC RefSeq v2.1 using BLAST, and identity and coverage were recorded.

Orthologous genes and syntenic blocks

Primary transcripts of 133 744 HC genes (IWGSC, 2018) were mapped to the pseudomolecules of IWGSC RefSeq v2.1 to obtain new coordinates. The all versus all bidirectional BLAST (Altschul *et al.*, 1990), with default parameters, was performed among protein sequences of the corresponding primary transcripts to identify putative orthologous genes. The top five hits based on identity and coverage of each HC gene were used to identify syntenic blocks among subgenomes using MCScanX (Wang *et al.*, 2012) with default settings.

Chromatin architecture analysis (Hi-C)

Chromosome conformation capture (Hi-C) sequencing data (IWGSC, 2018) were retrieved from ENA accession PRJEB25248. Hi-C reads were mapped with the TRITEX pipeline (Monat *et al.*, 2019). The IWGSC RefSeq v1.0 and v2.1 assemblies were digested *in silico* with *Hind*III using EMBOSS restrict (Rice *et al.*, 2000). Reads were trimmed at junction sites (AAGCTAGCTT) with cutdapt (Martin, 2011) and aligned to the assemblies with Minimap2 (Li, 2018). Alignment records were converted to Binary Sequence Alignment/Map (BAM) format with SAMtools (Li *et al.*, 2009) and sorted with Novosort (<http://www.novocraft.com/products/novosort/>). A list of Hi-C links was extracted from Hi-C alignments using BEDTools (Quinlan and Hall, 2010) and TRITEX scripts. A table of Hi-C links was imported to the R statistical environment (R Core Team, 2018) and contact matrices were plotted with TRITEX scripts.

ACKNOWLEDGEMENTS

The authors are grateful to the Mésocentre Clermont Auvergne University (France) and AuBi platform for providing computing and storage resources. The authors thank Dr Patrick E. McGuire for valuable suggestions and assistance with drafting the manuscript. This work was supported by the US National Science Foundation grant IOS-1929053 (JD and M-CL) and USDA Agricultural Research Service CRIS project 2030-21430-014-00-D (YQG). RDO was funded by a grant from the French Government managed by the Research National Agency (ANR) under the Investment for the Future program (BreedWheat project ANR-10-BTBR-03).

AUTHOR CONTRIBUTIONS

M-CL, JD, TZ and LW conceived the project. TZ, JCR, KRD and M-CL generated optical maps. LW performed comparative analysis of the two CS stocks. TZ and M-CL resolved chimera, and performed super-scaffolding and pseudomolecule construction. LW and TZ closed gaps and corrected base errors. HR and FC re-annotated genes, and RDO annotated TEs. LW, GK-G, JT, RA, YQG, HR, FC, JR, KE and MM contributed to evaluation and validation of the assembly. TZ, LW, FC, RDO, RA, M-CL and JD wrote the

first draft of the paper. All authors edited and approved the final draft.

CONFLICT OF INTEREST

The authors declare that they have no competing interest.

DATA AVAILABILITY

The IWGSC RefSeq v2.1 assembly and annotations are available at the IWGSC data repository hosted by URGI-INRAE (<https://wheat-urgi.versailles.inra.fr/Seq-Repository/Assemblies>) and NCBI under project PRJNA669381. The DLS optical map is available at <https://doi.org/10.25338/B8K917>.

SUPPORTING INFORMATION

Additional Supporting Information may be found in the online version of this article.

Figure S1. Intra-chromosomal Hi-C contact matrices of the IWGSC RefSeq v2.1.

Figure S2. Intra-chromosomal Hi-C contact matrices of the IWGSC RefSeq v1.0.

Table S1. Gene reannotation.

Table S2. Evaluation of IWGSC RefSeq v2.1 using random BAC assemblies.

Table S3. Detailed changes in IWGSC RefSeq v2.1 related to IWGSC RefSeq v1.0.

Table S4. Numbers of intact LTR retrotransposons in the pseudomolecules of IWGSC RefSeq v2.1 and IWGSC RefSeq v1.0.

Table S5. Comparison among IWGSC RefSeq v1.0, IWGSC RefSeq v2.1, and *Triticum* 4.0.

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