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Conference Paper

The Korea *Brassica* Genome Project: A glimpse of the *Brassica* genome based on comparative genome analysis with *Arabidopsis*

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Abstract

A complete genome sequence provides unlimited information in the sequenced organism as well as in related taxa. According to the guidance of the Multinational Brassica Genome Project (MBGP), the Korea Brassica Genome Project (KBGP) is sequencing chromosome 1 (cytogenetically oriented chromosome #1) of *Brassica rapa*. We have selected 48 seed BACs on chromosome 1 using EST genetic markers and FISH analyses. Among them, 30 BAC clones have been sequenced and 18 are on the way. Comparative genome analyses of the EST sequences and sequenced BAC clones from *Brassica* chromosome 1 revealed their homeologous partner regions on the *Arabidopsis* genome and a syntenic comparative map between *Brassica* chromosome 1 and *Arabidopsis* chromosomes. *In silico* chromosome walking and clone validation have been successfully applied to extending sequence contigs based on the comparative map and BAC end sequences. In addition, we have defined the (peri)centromeric heterochromatin blocks with centromeric tandem repeats, rDNA and centromeric retrotransposons. In-depth sequence analyses of five homeologous BAC clones and an *Arabidopsis* chromosomal region reveal overall co-linearity, with 82% sequence similarity. The data indicate that the *Brassica* genome has undergone triplication and subsequent gene losses after the divergence of *Arabidopsis* and *Brassica*. Based on in-depth comparative genome analyses, we propose a comparative genomics approach for conquering the *Brassica* genome. In 2005 we intend to construct an integrated physical map, including sequence information from 500 BAC clones and integration of fingerprinting data and end sequence data of more than 100 000 BAC clones. The sequences have been submitted to GenBank with accession numbers: 10 204 BAC ends of the KBrH library (CW978640–CW988843); KBrH138P04, AC155338; KBrH117N09, AC155337; KBrH097M21, AC155348; KBrH093K03, AC155347; KBrH081N08, AC155346; KBrH080L24, AC155345; KBrH077A05, AC155343; KBrH020D15, AC155340; KBrH015H17, AC155339; KBrH001H24, AC155335; KBrH080A08, AC155344; KBrH004D11, AC155341; KBrH117M18, AC146875; KBrH052O08, AC155342. Copyright © 2005 John Wiley & Sons, Ltd.

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Introduction

The *Arabidopsis* genome has been sequenced completely by an international consortium (the Arabidopsis Genome Initiative, 2000). *Arabidopsis* and *Brassica* diverged 14.5–20.4 million years ago from a common ancestor (Bowers *et al*., 2003). Comparative genetic mapping has revealed co-linear chromosome segments (Kowalski *et al*., 1994; Lagercrantz *et al*., 1996; Paterson *et al*., 2000, 2001; Schmidt *et al*., 2001) in the family Brassicaceae and linkage arrangements between *Arabidopsis* and *B. oleracea* (Lukens *et al*., 2003). The genomes of *Brassica* species have duplicated, perhaps triplicated, counterparts of the corresponding homeologous segments of *Arabidopsis* (O'Neill and Bancroft, 2000; Rana *et al*., 2004).

Brassica is one of the core genera in the family *Brassicaceae*. Six *Brassica* species are cultivated worldwide; three diploids: *B. rapa* (AA, $2n = 20$), *B. nigra* (BB, $2n = 16$) and *B. oleracea* (CC, $2n = 18$), and three amphidiploids (allotetraploids): *B. juncea* (AABB, $2n = 36$), *B. napus* (AACC, $2n = 38$) and *B. carinata* (BBCC, $2n = 34$) (U. 1935). The species *B. rapa* (syn. *campestris*), with 529 Mb per haploid genome equivalent (Johnston *et al*., 2005), was prioritized for sequencing by a multinational collaboration. The Multinational *Brassica* Genome Project (MBGP) and *Brassica rapa* Genome Sequencing Project (BrGSP) are aiming to completely sequence the genome of *Brassica rapa* inbred line 'Chiifu' (**http://www.brassicagenome.org**; **http://www.brassica-rapa.org**). Korea launched the Korea *Brassica* Genome Project (KBGP) for complete sequencing of the cytogenetic chromosome 1 using BAC-by-BAC shotgun sequencing. In-depth comparative sequence analyses of the sequenced *B. rapa* BAC clones revealed overall co-linearity with a homeologous region of the *Arabidopsis* genome. Comparative sequence analyses suggest that we can use the *Arabidopsis* genome as a backbone for *in silico* clone validation of seed BAC clones and physical mapping as in the report of Love *et al*., 2004.

Here we propose an efficient clone validation method for selecting chromosome-specific seed BACs using comparative physical mapping and BAC end sequences. In 2005, KBGP aims to sequence 500 BAC clones that correspond to the majority of *Arabidopsis* euchromatin regions. The 500 BACs will be distributed and mapped on *B. rapa* chromosomes through sequence tagged site (STS) or simple sequence repeat (SSR) markers. BAC end sequences of 100 000 BACs (STC) and fingerprinting polymorphism-based BAC contigs (FPC) will be available soon. Hence, the sequence and map information of 500 BACs can be integrated with STCs and FPCs, resulting in an integrated physical map. The integrated physical map will provide a high resolution genome wide comparative map with *Arabidopsis* and will be supplied to MBGP to accelerate the *Brassica* genome sequencing.

Materials and methods

DNA sequencing

Shotgun sequencing libraries were constructed in pCUGIblu31 for average insert size of 3 kb (Kim *et al*., 2004; Yang *et al*., 2004; Yang *et al*., 2005). BigDye terminators chemistry v3.0 (ABI) was used for the reactions. The sequences were analysed using ABI3730 automatic DNA sequencers (ABI). Base-calling was performed automatically using phred, and vector sequences were removed by CROSS_MATCH (Ewing and Green 1998; Ewing *et al*., 1998). High quality, vector-trimmed sequences were thus used for the sequence assembly of each BAC clone, using phrap and consed (Gordon *et al*., 1998).

Sequence analysis

Pairwise sequence comparison was conducted using PipMaker (Schwartz *et al*., 2000) and BLAST2 analysis (**http://www.ncbi.nlm.nih.gov/BLAST/**). MegaBLAST against the *Arabidopsis* chromosome database and BLAST-nr were used as needed (**http://www.ncbi.nlm.nih.gov/BLAST/**). Gene annotation was achieved using several web based gene prediction programs, e.g. FGENE-SH *Arabidopsis* (**http://www.softberry.com/berry.phtml**) and GeneMark *Arabidopsis* (**http://opal.biology.gatech.edu/GeneMark/eukhmm.cgi**). Repeats were identified using Repeatmasker (**http://ftp.genome.washington.edu/RM/webrepeatmaskerhelp.html**).

Fluorescence *in situ* hybridization

Our FISH protocol was adapted from Lim *et al*., (2001, 2005a) with minor modifications. FISH signals were pseudo-coloured and further improved for optimal brightness and contrast with Adobe Photoshop image processing software.

Results and discussion

Overview of *Brassica rapa* genome structure

A genetic map of *Brassica rapa*, using segregating doubled haploid lines of Chiifu and Kenshin, covering 1046 cM with 494 markers on 10 linkage groups, was constructed with 895 DNA markers, AFLP, PCR-RFLP, ESTP, CAPS and SSR (**http://www.brassicagenome.org**). We have constructed another EST-RFLP genetic map of *B. rapa* using 478 tissue-specific cDNA clones consisting of 176 cDNAs from immature flowers, 252 cDNAs from anthers and 50 from dark-grown seedlings of *B. rapa* ssp. *pekinensis* cv. Jangwon. This molecular map covered 3412 cM on 10 linkage groups. Aligning RFLP marker sequences on the counterpart *Arabidopsis* chromosomes shows syntenic co-linearity, resulting in a highly informative comparative genetic map (Kim, 2001). The karyotypes of *B. rapa* chromosomes were studied previously (Fukui *et al*., 1998; Snowdon *et al*., 2002; Koo *et al*, 2004). We further characterized chromosomes in detail using fluorescence *in situ* hybridization (FISH) using repetitive DNAs, such as 45S rDNA, 5S rDNA, centromeric repeats (CentBr) and centromere-specific retrotransposons (Lim *et al*., 2005a). The cytogenetic chromosomes were integrated with genetic maps by painting with chromosome-specific BAC clones identified by unique EST clones from each linkage group (LG1–LG10) (Lim *et al*., 2005b). The cytogenetic chromosome numbers, our linkage groups (LG1–LG10) and the international standard linkage numbers (R1–R10) (Lombard and Delourme, 2001) will be integrated soon.

We have sequenced four BAC clones that form the counterpart of an *Arabidopsis* chromosomal region (chromosome 5: 3.1–3.2 Mb) containing flowering locus C (FLC). Comparisons of the sequenced *Brassica* BAC clones with the homeologous regions of *Arabidopsis* showed overall co-linearity with 81% sequence similarity. The average sequence similarity between *Brassica* BACs is 82% with exceptionally high similarity (97%) of two clones, 117M18 and 52O08, representing two regions that have recently been duplicated. The co-linear 125 kb *Arabidopsis* sequence was reduced by up to 40% by deletions of DNA segments in *Brassica* BAC clones (Table 1). Among 36 genes in the 125 kb of *Arabidopsis* sequence, only 24, 17, 13, and 13 homologues remained in the common sequence of each BAC clone, 80A08, 4D11, 52O08 and 117M18, respectively. Only four genes remain in all four BAC clones, with 77–96% similarity in amino acid sequences. Newly emerged (or inserted)

Table 1. Comparison of four homologous *Brassica* BAC clones and its counterpart *Arabidopsis* sequence

	Homologous *Brassica* BAC clones				
Subject	**80A08**	**4D11**	**52O08**	**117M18**	***Arabidopsis* chrom. 5**
Insert size (bp)	110 219	106 476	153 587	132 883	—
Common sequence (begin)	1	15 001	58 254	18 227	3 134 987
Common sequence (end)	110 219	89 318	115 292	70 502	3 258 842
Total length of common sequence (bp)	110 217	74 316	57 037	52 274	123 855[b]
Aligned nucleotide (bp)[a]	67 412	47 325	33 870	31 161	
Internal deletion or substitution (bp)	42 795	26 991	23 167	21 113	
Co-linearity index[b]	0.9	0.6	0.5	0.4	1.0
Alignment index[c]	0.61	0.64	0.59	0.60	
Homology (*Arabidopsis* vs. Others) (%)	81.0	81.1	81.7	81.3	100.0
Homology (117M18 vs. Others) (%)	82.3	81.8	98.1	100.0	82.3

[a] A total of nucleotides that show significant sequence similarity with co-linear *Arabidopsis* sequence.
[b] represents genome expanding or reducing in *Brassica* BAC clones compared to the co-linear *Arabidopsis* sequence (= Total length of common sequence of *Brassica*/Common sequence of *Arabidopsis*.
[c] represents the significantly homeologous region in the common sequence (= Aligned nucleotide/Total length of common sequence.

genes including transposons are detected six, three, two and one times in each BAC clone, respectively. The data support previous reports (O'Neill and Bancroft, 2000; Rana *et al*., 2004) and provide in depth information about how triplicated *Brassica* genome sequences are modified after divergence with *Arabidopsis* at around 20 million years ago (Bowers *et al*., 2003).

Pericentromeric heterochromatin blocks in the *Brassica rapa* genome

The centromeric region of *Brassica* is occupied by 176 bp tandem repeats (Harrison and Heslop-Harrison, 1995). The 176 bp centromeric repeat of *Brassica* (named CentBr) occurred in 30% of our BAC end sequences (10 204 BAC ends of the KBrH library; GenBank accession numbers CW978 640–CW988 843) as tandem arrays, indicating that the CentBr is a major component of the *B. rapa* centromere. The CentBr sequences are subdivided into two classes, named CentBr1 and CentBr2, based on sequence similarity (82–84% between two classes and over 92% between members in each class). CentBr1 and CentBr2 occupy the centromeres of eight and two chromosomes, respectively (Lim *et al*., 2005a).

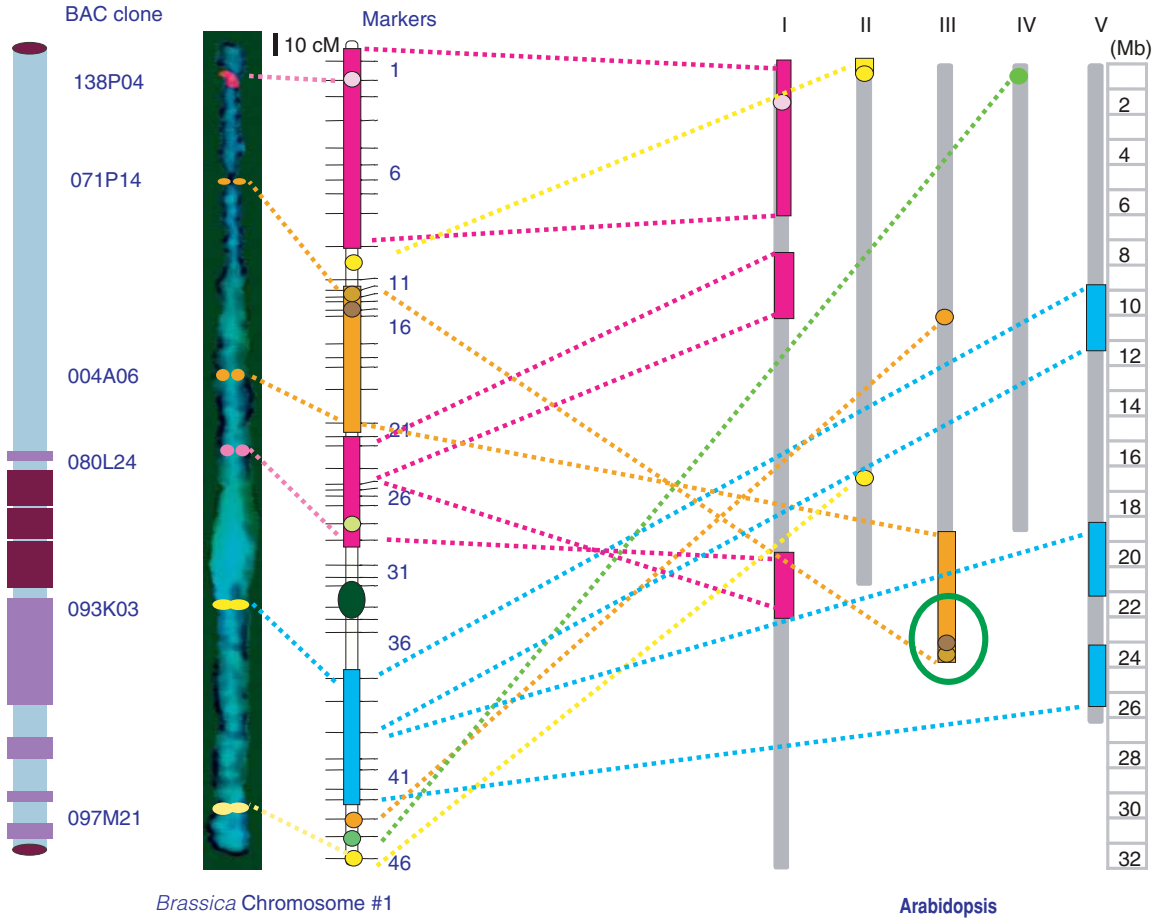

Figure 1. Comparative map of *B. rapa* chromosome 1 and *Arabidopsis* chromosomes based on sequence similarity of EST markers and sequenced BAC clones. The far left of the figure represents the features of chromosome 1, BAC clones were selected by filter hybridization using mapped EST markers and their actual chromosomal locations were confirmed by FISH analyses, using metaphase or pachytene phase chromosomes (left). The cartoon at the left of pachytene chromosome represents the features of chromosome 1, showing pericentromeric heterochromatin and heterochromatin (brown and purple boxes, respectively), based on numerous inspections by DAPI staining and FISH analyses using repetitive elements. The linkage groups containing 46 markers and corresponding elements of chromosome 1 are represented and the syntenic regions are represented on *Arabidopsis* chromosomes (right)

We have sequenced two centromeric BAC clones, KBrH015B20 (102 kb) and KBrH001P13 (17 kb), containing centromeric tandem repeats for increased understanding of major elements in the (peri)centromeric region of the *Brassica* genome. Careful sequence analysis revealed several families of centromere-specific retrotransposons of *Brassica* (CRB). Among these, two long terminal

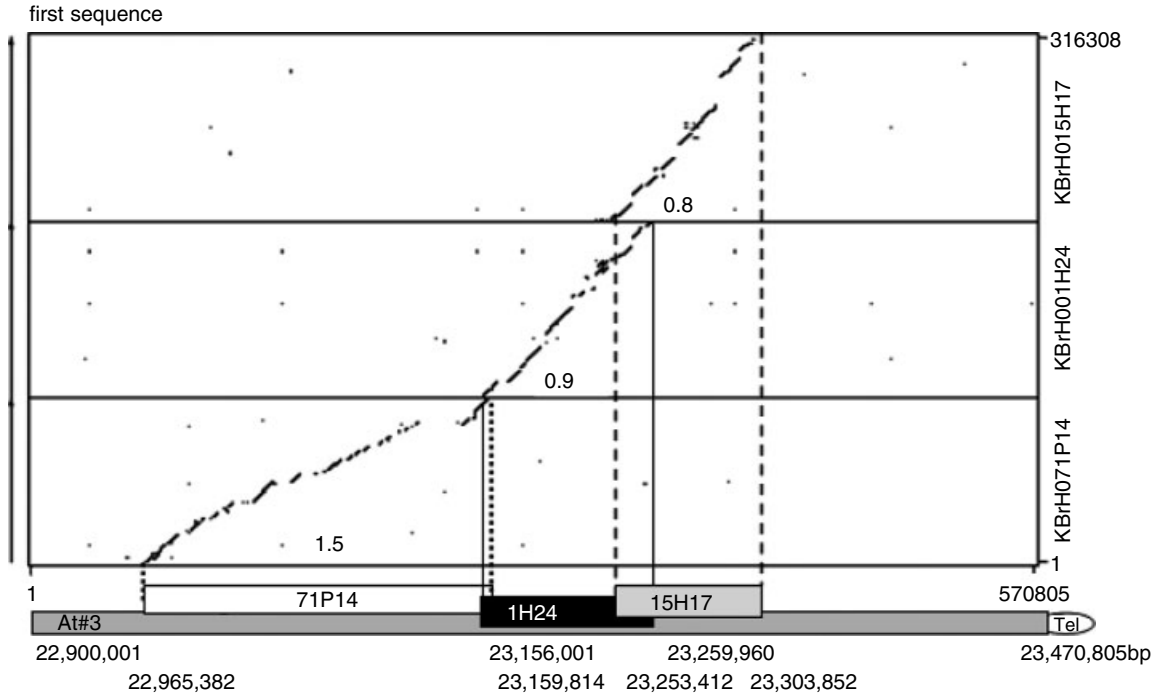

Figure 2. Dot-plot analysis of three contiguous sequenced BAC clones from *B. rapa* chromosome 1 and the counterpart region of *Arabidopsis* (chromosome 3, 22.9–23.3 Mb). The region is marked as a green circle in Figure 1. The beginning and ending nucleotide of the counterpart *Arabidopsis* sequence for the three *Brassica* BAC clones are represented as numericals under the figure. The co-linear index (= The length of co-linear *Arabidopsis* sequence/Co-linear *Brassica* BAC sequence which is in accordance with the slope) of each BAC clone is represented in the dot plot

Table 2. *Brassica* BAC clones sequenced and the counterpart *Arabidopsis* sequence

***Brassica* BAC Clone**		**Co-linear Index**	**Counterpart sequence in *Arabidopsis***			
Name	**Length**	***Arabidopsis*/*Brassica***	**Length**	**Begin**	**End**	**Chrom. No**
01H24	118 144	0.9	103 420	23 156 540	23 259 960	3
15H17	110 885	0.8	85 302	23 224 319	23 309 621	3
20D15	143 633	0.8	119 117	8 047 497	8 166 614	1
77A05	113 253	1.1	125 641	19 236 565	19 362 206	5
80L24	115 119	4.0	459 225	17 701 122	18 160 347	1
97M21	131 063	2.4	314 999	10 677 001	10 992 000	3
117N09	125 390	1.7	218 929	17 478 001	17 696 930	2
138P04	137 697	1.3	173 013	2 128 977	2 301 990	1
4D11	106 476	1.7	184 272	3 092 144	3 276 416	5
80A08	110 038	1.1	121 376	3 137 466	3 258 842	5
Average insert size (bp)	121 170	1.6	190 529			
STD	±12 682	1.0	116 235			

repeat (LTR) retrotransposons, a Ty3-gypsy-like one (PCRB; 9135 bp with 2047bp LTR) and a Ty1-copia-like one (CRB; 6010 bp with 597 bp LTR) predominantly occupied each BAC sequence. FISH analyses revealed that the CRB is a major component of the centromere of all chromosomes and the PCRB is a major component of the large pericentromeric heterochromatin regions of three chromosomes. Based on the BAC end sequence information and FISH analyses, we assume these four (peri)centromeric repeats occupy just over 40% of the *Brassica* genome. Since heterochromatin blocks are hard to sequence, we will focus on sequencing of euchromatin regions, which probably constitute less than 60% of the *Brassica rapa* genome.

Progress in sequencing chromosome 1 in Korea

The Korea *Brassica* Genome Project (KBGP) is aiming to complete the sequencing of cytogenetic chromosome 1 using three BAC libraries, KBrH (*Hin*dIII), KBrB (*Bam*HI), and KBrS (*Sau*3AI), of *B. rapa* ssp. *pekinensis* inbred line 'Chiifu'. Physical mapping is on-going by fingerprinting of the KBrH and KBrB libraries (**http://www.brassicagenome.org; http://www.brassica-rapa.org**). Anchoring the fingerprint polymorphism contigs (FPC) on the chromosome remains an obstacle to overcome for physical mapping and clone validation. We have selected 48 seed BACs on chromosome 1 through screening with EST markers and confirmation by FISH analyses. Among them, 30 BAC clones were sequenced and they show co-linearity with the counterpart homeologous region of *Arabidopsis*, with about 82% sequence similarity (Table 1).

The comparative analyses of the EST sequences mapped on chromosome 1 with their homeologous partner regions of *Arabidopsis* revealed counterparts in the *Arabidopsis* genome (Figure 1). The sequenced BAC clones show overall co-linearity with a counterpart *Arabidopsis* chromosomal region which was expected, based on the

BAC End Pairing	BAC Ends	Direction	Length	Matching			Span (bp)	Arabidopsis Match Region			GenBank No.	E Value
	KBrB092L06R	**+**	**792**	**121**	**-**	**764**	**207,171**	**441,816**	**-**	**442,459**	NC_003070.5	**1.00E-119**
	KBrB078D07F	+	660	228	-	495	200,497	449,221	-	449,488	NC_003070.5	3.00E-78
	KBrB069J20F	+	852	229	-	500	55,796	449,221	-	449,492	NC_003070.5	1.00E-99
	KBrB072O20R	+	561	242	-	513	162,673	449,221	-	449,492	NC_003070.5	4.00E-95
	KBrB064C19F	+	367	188	-	277	438,409	456,834	-	456,923	NC_003070.5	3.00E-27
	KBrH014E03F	+	767	677	-	754	175,318	486,300	-	486,377	NC_003070.5	2.00E-08
	KBrB006N24R	+	752	227	-	472	409,867	489,141	-	489,386	NC_003070.5	3.00E-41
	KBrB069J20R	-	342	1	-	126		505,017	-	504,892	NC_003070.5	2.00E-46
	KBrS006E09F	+	792	1	-	87	41,204	567,509	-	567,597	NC_003070.5	4.00E-19
	KBrS006E09R	-	686	197	-	514		608,713	-	608,396	NC_003070.5	1.00E-120
	KBrB072O20F	-	346	1	-	222		611,894	-	611,674	NC_003070.5	7.00E-25
	KBrB092L16R	+	665	159	-	372	120,321	612,015	-	612,234	NC_003070.5	4.00E-31
	KBrB001E05F	+	698	8	-	687	107,536	618,404	-	619,098	NC_003070.5	1.00E-147
	KBrB030F10R	**+**	**736**	**73**	**-**	**123**	**129,058**	**624,089**	**-**	**624,139**	NC_003070.5	**0.0001**
	KBrH008B01F	+	794	150	-	497	85,749	638,255	-	638,602	NC_003070.5	5.00E-83
	KBrB092L06F	**-**	**845**	**783**	**-**	**845**		**648,987**	**-**	**648,925**	NC_003070.5	**2.00E-15**
	KBrB078D07R	-	824	277	-	642		649,718	-	649,360	NC_003070.5	3.00E-72
	KBrB049O03F	+	802	161	-	201	105,990	651,701	-	651,741	NC_003070.5	6.00E-12
	KBrH014E03R	-	849	35	-	281		661,618	-	661,372	NC_003070.5	3.00E-44
	KBrH006J03R	+	711	145	-	221	163,565	666,344	-	666,420	NC_003070.5	5.00E-12
	KBrS011G09F	+	796	450	-	605	87,764	667,143	-	667,298	NC_003070.5	2.00E-30
	KBrS016E11R	+	699	167	-	275	120,558	681,105	-	681,213	NC_003070.5	6.00E-24
	KBrS005M08R	+	464	2	-	206	128,380	681,713	-	681,917	NC_003070.5	1.00E-33
	KBrB021O04F	+	838	643	-	822	266,085	685,555	-	685,734	NC_003070.5	1.00E-37
	KBrB069F04F	+	850	644	-	823	266,085	685,555	-	685,734	NC_003070.5	1.00E-37
	KBrB042P04R	+	532	1	-	189	283,170	709,677	-	709,862	NC_003070.5	1.00E-58
	KBrS005C04R	+	719	1	-	719	306,547	716,206	-	716,924	NC_003070.5	0
	KBrH008B01R	-	908	657	-	905		724,004	-	723,756	NC_003070.5	2.00E-64
	KBrB001E05R	-	896	435	-	828		725,940	-	725,548	NC_003070.5	1.00E-93
	KBrB056O02F	+	821	126	-	223	86,208	728,279	-	728,373	NC_003070.5	4.00E-13
	KBrB092L16F	-	942	6	-	540		732,336	-	731,793	NC_003070.5	1.00E-115
	KBrB089B13R	**+**	**839**	**1**	**-**	**242**	**341,507**	**732,356**	**-**	**732,597**	NC_003070.5	**3.00E-60**
	KBrB030F10F	**-**	**858**	**769**	**-**	**844**		**753,147**	**-**	**753,072**	NC_003070.5	**7.00E-21**
	KBrS011G09R	-	721	269	-	712		754,907	-	754,465	NC_003070.5	1.00E-140
	KBrB049O03R	-	683	54	-	196		757,691	-	757,549	NC_003070.5	3.00E-44

Figure 3. *In silico* allocation of *Brassica* BAC clones on *Arabidopsis* chromosomes. The beginning part of *Arabidopsis* chromosome 1 is represented. BAC clones are aligned on *Arabidopsis* chromosomes based on significant and directional matches of both ends within a 30–500 kb interval. The forward and reverse ends are marked as grey bars (left). An example of the minimum tiling path of the three BAC clones are boxed

comparative map (Figure 2). Based on the comparative physical map and micro-co-linearity between the *Brassica* and *Arabidopsis* sequences, we have proposed an efficient and novel clone validation method for sequencing in advance of the complete physical map. The *Brassica* BAC clones were allocated to *Arabidopsis* chromosomes by *in silico* allocation based on unique, significant (<1E-6), and directional matches: one BAC end is forward and the other end is the reverse, with a complement match within a 30–500 kb interval. BAC-FISH and STS mapping using BAC end sequences on the counterpart *Arabidopsis* chromosomal region showed the real locations of the BAC clones on the chromosomes. At least one in three BAC clones is mapped onto the expected region of chromosome 1 due to the triplicated nature of the *Brassica* genome. All the sequenced BAC clones provide a further starting point for selection of seed BAC clones for extending the sequence.

Integrated physical mapping

Successful clone validation based on *in silico* allocation to counterparts of chromosome 1 suggests a novel strategy for integrated physical mapping, using comparative mapping of BAC ends onto *Arabidopsis* chromosomes. The integrated physical mapping strategy encompasses *in silico* allocation of *B. rapa* BAC clones to the counterpart locations of *Arabidopsis* chromosomes, based on significant BLAST matches. A *Brassica* BAC clone (average size 120 kb) covers an average of 190 kb *Arabidopsis* sequence based on a co-linearity index of 1.6 (= co-linear *Arabidopsis* sequence/*Brassica* BAC nucleotide) (Table 2). We have analysed 91 000 BAC end sequences (Table 3). Among them, a total of 45 232 BAC end sequences (50%) show significant sequence similarity with unique *Arabidopsis* sequences, and a total of 4317 BAC clones (9.5%) are allocated on *Arabidopsis* chromosomes by significant matching with both ends within 30–500 kb interval (Table 3). These 4317 clones span 93 Mb of *Arabidopsis* euchromatin regions, representing 78.2% of the total *Arabidopsis* genome. A total of 26 Mb remain as unconvered gaps: among these 9.4 Mb (3.1 Mb, 1.8 Mb, 2.4 Mb, and 1.0 Mb from *Arabidopsis* chromosomes 1, 2, 3, 4, and 5, respectively) might be from euchromatin gaps at 116 sites ranged from over 20–585 498 bp, except for the 16.6 Mb of pericentromeric heterochromatin gaps. A single *Brassica* BAC clone spans an average of 147 kb (±74 kb) *Arabidopsis* sequence (Figure 3). A total of 500

Table 3. *Brassica* rapa BAC end sequence and the results of blast analyses against *Arabidopsis* chromosomes

	Unique hits (<1E-6)			BAC clones with both ends		
Library	Query No.	Hit No.	%	Pairhit No.	Clone No.	%[a]
KBrH (*Hin*dIII)	10 204	4195	41.1	1162	581	11.4
KBrB (*Bam*HI)	72 343	36 833	50.9	6908	3454	9.5
KBrS (*Sau*3AI)	8632	4204	48.7	564	282	6.5
Total	91 179	45 232	49.6	8634	4317	9.5

[a] 100* (Pairhit No./Query No.). Clone numbers represent the numbers of BAC clones allocated on *Arabidopsis* chromosome by directional hitting with both ends within 30–500 kb interval.

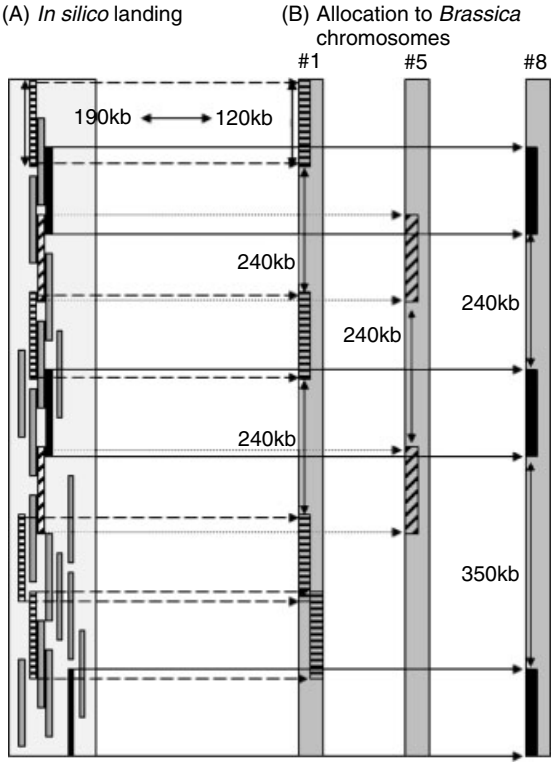

Figure 4. Schematic representation of the *in silico* landing on *Arabidopsis* chromosome and estimated real position on *Brassica* chromosomes. Minimum tiled BACs on *in silico* comparative allocation will be scattered onto three *Brassica* chromosomes. If the minimum tiled BACs are sequenced and mapped, fewer than 240 kb physical gaps will remain between BACs in each chromosome

BACs with an average 120 kb of insert will cover around 80 Mb of the euchromatin regions of the *Arabidopsis* genome (almost all of the euchromatin). The 500 BACs will be scattered into the triplicated regions on *Brassica* chromosomes (e.g. Figure 4). The actual chromosomal location of a sequenced BAC can be mapped on the genetic map through SSR or STS-PCR using its sequence information. Recently, we have selected the minimum tiled 629 *Brassica* BAC clones spanning 86 Mb of *Arabidopsis* from the *in silico* allocation (data is available at our website: **www.brassica-rapa.org**). Each BAC clone will be mapped on the *Brassica* chromosomes by STS mapping and FISH analyses. About 75 Mb from gene rich euchromatin regions of *Brassica* will be obtained from sequencing of the 629 BACs (average insert 120 kb) that may be distributed into 10 *B. rapa* chromosomes (average 60 BACs for each chromosome) with an average 240 kb gap (Figure 4). All the sequenced BAC clones will be provided to MBGP and used as a starting point for the selection of seed BAC clones extending to the flanking sides with minimum overlap based on sequence tagged connectors (STC). The results will provide in depth information about the comparative genomics between *Brassica* and *Arabidopsis*.

Complete sequencing of *Brassica rapa* will give great opportunities to increase our understanding of the evolution of the polyploidized genome and of agricultural aspects, especially for breeding and molecular farming, through finding novel or useful genes, not only in *B. rapa* but also in other important crops in the genus *Brassica*.

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