

# Microsatellites for the genus *Cucurbita* and an SSR-based genetic linkage map of *Cucurbita pepo* L.

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**Abstract** Until recently, only a few microsatellites have been available for *Cucurbita*, thus their development is highly desirable. The Austrian oil-pumpkin variety Gleisdorfer Ölkürbis (*C. pepo* subsp. *pepo*) and the *C. moschata* cultivar Soler (Puerto Rico) were used for SSR development. SSR-enriched partial genomic libraries were established and 2,400 clones were sequenced. Of these 1,058 (44%) contained an SSR at least four repeats long. Primers were designed for 532 SSRs; 500 primer pairs produced fragments of expected size. Of these, 405 (81%) amplified polymorphic fragments in a set of 12 genotypes: three *C. moschata*, one *C. ecuadorensis*, and eight *C. pepo* representing all eight cultivar groups. On an average, *C. pepo* and *C. moschata* produced 3.3 alleles per primer pair, showing high inter-species transferability. There were 187 SSR markers detecting polymorphism between the USA oil-pumpkin variety “Lady Godiva” (O5) and the Italian crookneck variety “Bianco Friulano” (CN), which are the parents of our previous F<sub>2</sub> mapping population. It has been used to construct the first published *C. pepo* map, containing mainly RAPD and AFLP markers. Now the updated

map comprises 178 SSRs, 244 AFLPs, 230 RAPDs, five SCARs, and two morphological traits (*h* and *B*). It contains 20 linkage groups with a map density of 2.9 cM. The observed genome coverage (Co) is 86.8%.

## Introduction

The genus *Cucurbita* ( $2n = 2x = 40$ ) contains some of the earliest domesticated plant species (Smith 1997; Piperno and Stothert 2003). Today, three of them, *C. pepo* L., *C. moschata* Duchesne, and *C. maxima* Duchesne, are economically important crops worldwide (Robinson and Decker-Walters 1997; Loy 2004). Pumpkin seed oil and pumpkin seed extract are increasingly esteemed for their excellent nutritional quality and medicinal value, especially in the prevention and treatment of benign prostate hyperplasia (Blumenthal et al. 1998; Kreuter 2000; Schmidlin and Kreuter 2003). The recent increase in the popularity of pumpkin has stimulated the breeding of new cultivars. A devastating epidemic of Zucchini Yellow Mosaic Virus (ZYMV) in 1997, destroying half of the pumpkin harvest in Austria, made development of resistant varieties crucial (Pachner and Lelley 2004). A genetic map facilitates marker-assisted selection (MAS) speeding up the breeding process, it is crucial for future structural and functional genomic studies in *Cucurbita*. Until now only three genetic maps of *Cucurbita* have been constructed: two maps from inter-specific crosses between *C. pepo* and *C. moschata* (Lee et al. 1995; Brown and Myers 2002) and the third from an intraspecific cross of *C. pepo* (Zraidi et al. 2007). These maps contain only RAPD and AFLP markers. So far, only 27 microsatellites were available for *Cucurbita* (Stift et al. 2004). Therefore, it was highly desirable to

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develop a large number of microsatellite markers for constructing an SSR-based map.

Microsatellites, also referred to as simple sequence repeats (SSRs), are 1–6 bp long, tandemly repeated sequences. Since microsatellites are abundant in the genome, mostly co-dominant and hyperallelic, they are ideal tools for a broad application in basic and applied plant biology (e.g., Kashi et al. 1997; Gupta and Varshney 2000; Li et al. 2002; Anderson and Lübberstedt 2003; Varshney et al. 2005; Torada et al. 2006; Saha et al. 2006). ESTs are important sources for SSR development in many plants. Unfortunately, only 120 *Cucurbita* ESTs are available in the gene bank, nine of them including an SSR sequence. Only one of these is suitable for primer design. Nevertheless, partial genomic libraries, enriched for SSRs, can also provide an efficient source for microsatellite development.

SSR flanking sequences are often conserved among species and even genera (Katzir et al. 1996; Gaitán-Solís et al. 2002; Saha et al. 2006; Downey and Iezzoni 2000; Yamamoto et al. 2001; Kuleung et al. 2004). Successful SSR transfer between species within a genus has been reported in several genera, for example, *Olea* (Rallo et al. 2003), *Limonium* (Palop et al. 2000), *Prunus* (Wuensch and Hormaza 2002; Dirlewanger et al. 2002), *Vitis* (Di Gaspero et al. 2000), *Pinus* (González-Martínez et al. 2004), *Triticum* (Adonina et al. 2005) and *Cucumis* (Danin-Poleg et al. 2000, 2001; Ritschel et al. 2004). Of the first 27 *Cucurbita* SSRs developed from *C. pepo* (Stift et al. 2004), most were transferable to *C. moschata*, *C. maxima*, and *C. ecuadorensis*. A relatively large set of such highly transferable SSR markers will facilitate the development of an integrated or composite *Cucurbita* map by merging different maps (Song et al. 2004; Gonzalo et al. 2005). The transferability of SSR markers also will allow comparative genetic studies within this genus and may further elucidate the evolution of the different species in the genus. It might help to shed more light on the hypothetical paleopolyploid origin of the genus (Weeden and Robinson 1986). In this paper we report on the development of a large number of *Cucurbita* SSR markers and their transferability among species. We have updated our previous *C. pepo* map, containing only RAPD and AFLP markers (Zraidi et al. 2007), with 178 SSR and 105 new AFLP markers.

## Materials and methods

### Plant material and DNA extraction

For microsatellite development genomic DNA was isolated from the Austrian oil-pumpkin variety Gleisdorfer Ölkürbis (*C. pepo*) and the *C. moschata* cultivar Soler (Puerto Rico). For assessing microsatellite polymorphism

and transferability across species three *C. moschata*, one *C. ecuadorensis*, and eight *C. pepo* genotypes representing the eight cultivar groups (Paris 1986) were used (Table 1).

For mapping the previous Pumpkin–Crookneck population was used (Zraidi et al. 2007), consisting of 92 F<sub>2</sub> plants. It was derived from an inter-subspecific cross (*C. pepo* subsp. *pepo* × *C. pepo* subsp. *ovifera*) between the USA oil-pumpkin variety Lady Godiva (O5, ♀) and the Italian crookneck variety Bianco Friulano (CN, ♂). DNA was isolated from fresh young leaves using Promega's Wizard Genomic DNA Purification Kit (Promega Corp., Madison, USA). DNA concentration was determined by the GenQuant RNA/DNA Calculator (Amersham Biosciences Europe, Germany) according to manufacturer's protocol.

### Development of SSR-enriched partial genomic insert libraries

The methods used for constructing SSR-enriched libraries were based on the protocols reported by Connell et al. (1998) and Kumpatla et al. (2004). Five microgram of genomic DNA were nebulized (Invitrogen Corp., CA, USA) for 2 min and 15 s, in an ice-cold water bath at 33 psi. These conditions for nebulization produced fragments between 300 and 600 bp in length. Following nebulization, single-strand 5' or 3' overhangs were end-polished by mung bean nuclease (MBNase) (New England Biolabs, MA, USA). The excess MBNase was removed by Millipore Micropure-EZ Enzyme Remover columns (Millipore, MA, USA), and blunt-end adaptors were ligated onto the end-polished DNA. The adaptors were prepared by mixing equal

**Table 1** Genotypes used for assessing microsatellite polymorphism and transferability across species

No.	Species	Genotype	Fruit shape	Origin
1	<i>C. moschata</i>	Nigerian Local	Globular	Nigeria
2		Waltham Butternut	Cylindrical	USA
3		Zhou <sup>a</sup>	Flattened	China
4	<i>C. ecuadorensis</i>	Grif 9446 USDA	Globular	Ecuador
5	<i>C. pepo</i>	True French	Zucchini	GB
6		Striato di Italia	Cocozelle	Israel
7		General Patton	Straightneck	USA
8		Bianco Friulano	Crookneck	GB
9		Lady Godiva	Oil pumpkin	USA
10		Early White Bush	Scallop	France
11		Tay Belle	Acorn	USA
12		Alba	Vegetable Marrow	USA

<sup>a</sup> Zhou is a Chinese hull-less *C. moschata* genotype, first described by Zhou (1987), named here for the discoverer

amounts of AP-11 oligo (5'-CTCTTGCTTAGATCTGG-ACTA-3') and AP-12 oligo (5'-TAGTCCAGATCTAAGC AAGAGCACA-3'), heating them at 95°C for 5 min, and then slowly cooling the solution to room temperature. The adaptor-ligated DNA fragments were amplified with AP11 primer. The products were hybridized with the biotinylated SSR oligos (GA)13, (CA)13, (CAT)8, (CCA)8, (CAA)8, and (GAA)8. Hybridized fragments were captured by streptavidin-coated magnetic beads (Dyna) and residual biotinylated SSR oligos removed with Microcon YM-100 columns (Millipore). Captured fragments were amplified with AP11 primers and PCR products were cleaned up using the Promega Purification Kit. The purified PCR products were ligated into pSTBlue-1 AccepTor Vector, and then transferred to NovaBlue Single™ Competent Cells (Novagen, Merck Bioscience, Nottingham, GB). The transformed cells were cultured in LB agar with 70 µg/ml X-gal, 80 µM IPTG, 50 µg/ml Carbenicillin, and 15 µg/ml Tetracyclin for 15–18 h, after which white clones were picked and cultured in LB broth with 50 µg/ml Carbenicillin and 15 µg/ml Tetracyclin. Fifty microliter bacteria liquid was incubated at 99°C for 5 min to lyse the bacteria and destroy the DNase. PCR was performed with the lysed bacteria liquid (colony PCR). The vector specific primers used were as follows:

T7-fwd: 5'-TAATACGACTCACTATAGGG-3'  
SP6-rev: 5'-ATTTAGGTGACACTATAG-3'

Products of colony PCR were run on 1.5% agarose gel at 90 V for 20 min to screen for inserts longer than 300 bp. PCR products with appropriate size were purified by Exonuclease (ExoI) and Shrimp alkaline phosphatase (SAP) (Fermentas Inc., Ontario, Canada). DYEnamic ET Dye Terminator Kit (GE Healthcare, Uppsala, Sweden) was used for sequencing PCR according to manufacturer's protocol. Sequencing PCR products were purified with Sephadex G50 (GE Healthcare) in a Multiscreen HTS 96 well plate (Millipore). Sequencing was performed on a MegaBace 1000 DNA analysis system (GE Healthcare).

#### Microsatellite identification and primer design

Microsatellite detection was made by the SSRIT program (<http://www.gramene.org/gramene/searches/ssrtool>). Only sequences containing at least 4 di-, tri-, tetra-, or penta-nucleotide repeats were selected. To identify duplicates sequence comparisons were made using Stand-alone BLAST (2.2.12). Primers were designed with the online software Primer3 ([http://frodo.wi.mit.edu/cgi-bin/primer3/primer3\\_www.cgi](http://frodo.wi.mit.edu/cgi-bin/primer3/primer3_www.cgi)) using the following parameters: (1) minimum of seven di-nucleotide or five tri-nucleotide repeats, (2) primer length 18–27 nt, 20 nt considered to be optimal, (3) PCR product size 80–300 bp, (4) optimal

annealing temperature 60°C, (5) GC contents from 20 to 80% with 50% as optimum. Primers were synthesized by Invitrogen Corp. CA, USA.

#### PCR amplification and detection of SSR loci

SSR primers were first tested using genomic DNA of the respective species. Primers producing fragments were used for polymorphism test. The total volume of the PCR reaction mixture was 10 µl, and contained 27 ng DNA, 2 mM dNTPs, 0.25 units Taq DNA polymerase, and 1× reaction buffer containing 1.5 mM MgCl<sub>2</sub> (Gene craft, Germany), 0.25 pmoles forward primer with an M13 tail added to its 5' end (5'-CCCAGTCACGACGTTG-3'), 2.5 pmoles reverse primer, and 2.25 pmoles fluorescent-labeled M13 tail (FAM, HEX, synthesized by MWG, Ebersburg, D). PCR was performed as follows: denaturation at 95°C for 2 min, followed by seven cycles of 45 s at 94°C, 45 s at 68°C (with each cycle the annealing temp decreases 2°C), and of 60 s at 72°C. Products were subsequently amplified for 30 cycles at 94°C for 45 s, 54°C for 45 s, and 72°C for 60 s, with a final extension at 72°C for 5 min. When primers did not work in this PCR program final annealing temperature was reduced to 48°C. PCR products were separated using 10% polyacrylamide gels, 1× TBE buffer in a C.B.S. electrophoresis chamber (C.B.S. Scientific Inc., Del Mar, CA, USA). Electrophoresis conditions were set at constant 400 V and 10°C for 2 h. Gels were scanned by Typhoon (GE Healthcare, Uppsala, Sweden) in fluorescent mode.

Polymorphic index content (PIC) of an SSR locus was calculated as described by Smith et al. (1997), based on the set of 12 genotypes (Table 1):

$$\text{PIC} = 1 - \sum_{i=1}^n f_i^2,$$

where  $f_i$  is the frequency of the  $i$ th allele, and  $n$  is the total number of alleles for that locus.

#### Genotyping of AFLP markers

A set of 25 EcoRI and MseI selective primer combinations were used to produce AFLP markers for the Pumpkin–Crookneck population (O5 × CN), following the protocol of Hartl et al. (1999) and Buerstmayr et al. (2002).

#### Map construction

Segregation of each locus was calculated using a chi-square test ( $P = 0.05$ ). The markers, which showed highly significant segregation distortion ( $P < 0.001$ ) from the expected 1:2:1 (co-dominant) or 3:1 (dominant) ratio, were excluded from map construction. JoinMap® version 3.0

(Stam 1993; Van Ooijen and Voorrips 2001) was used for linkage analysis and map calculations. Linkage groups were determined using a LOD threshold of 3.0. Map construction was performed, using the Kosambi mapping function with JoinMap parameter settings as follows: Rec = 0.4, LOD = 1.0, Jump = 5. Some of the markers were removed, when they showed insufficient linkage or conflict with others. SSR, SCAR markers and two morphological traits were used to establish a skeleton map first. Then this map was used as “fixed order” to incorporate the AFLPs and RAPDs of the published *C. pepo* map (Zraidi et al. 2007). The new AFLP markers, produced in this study, were also added to this map. The genetic linkage map was drawn by MapChart software (Voorrips 2002).

#### Estimation of genome length and map coverage

Genome length ( $G_e$ ) was estimated by “moment estimator”,  $G_e = M(M - 1)X/K$  (Hulbert et al. 1988), where  $M$  is the number of linked markers,  $X$  is the maximum observed map distance between the locus pairs above a threshold LOD  $Z$  (Chakravarti et al. 1991), and  $K$  is the number of locus pairs having LOD values at or above  $Z$ . The value used for  $Z$  was LOD 3. The values of  $X$  and  $K$  were obtained from JoinMap using Kosambi mapping functions. The 95% confidence interval ( $I$ ) of the estimation of genome length was calculated using the equation  $I = G_e / (1 \pm 1.96/k^{-1/2})$  (Gerber and Rodolphe 1994). The observed genome coverage ( $C_o$ ) was defined as the ratio of the observed length of the map ( $G_{or}$ ) and the estimated genome length  $G_e$  (Cervera et al. 2001).

## Results

#### Development and characterization of microsatellites

In *C. pepo* a total of 1,056 clones were sequenced, of which 448 (42.4%) contained an SSR sequence of  $\geq 4$  repeats. The remaining 608 clones either contained SSRs with  $< 4$  repeats or were cryptic (SSR-similar) sequences. In 59 cases (5.6%) primers could not be designed due to insufficient length of SSR-flanking sequences. One hundred duplicate clones (9.5%) were removed. Ultimately, primers were designed for 230 unique SSR sequences (51.3%) having at least seven repeats of a di-nucleotide, or five repeats of a tri-nucleotide motif (Table 2). In *C. moschata* 1,344 clones were sequenced, of which 610 clones (45.4%) contained SSRs of  $\geq 4$  repeats. In 35 clones (2.6%) the SSR flanking sequences were too short to design primers. Twenty duplicate clones (1.5%) were removed. Finally, primers were designed for 342 unique SSR sequences (56%) with at least seven repeats of di- or five repeats of

**Table 2** Efficiency of SSR marker development in *C. pepo*

SSR development steps	Attrition	Total	%
Sequenced clones	0	1,056	100
SSR containing clones	1	448	42.4
Clones containing SSRs too close to one end	2	59	5.6 (13.2 of 1)
Primer pairs designed and tested	3	230	21.8 (51.3 of 1)
Working SSRs	4	193	18.3 (83.9 of 3)
Polymorphic SSRs	5	155	14.7 (80.3 of 4)

tri-nucleotide motifs (Table 3). In addition, sequence comparison revealed a cluster of 30 clones, with similar sequences containing 7–18 repeats of the GA/TC motif. Only two of them originated from *C. moschata*, the rest from *C. pepo*.

Microsatellites were classified by motif and structure. The majority of microsatellites, 68.3% of *C. pepo* and 74.6% of *C. moschata*, represented perfect repeats. In both species the majority of di-nucleotide motif was GA/TC, the predominant tri-nucleotide motif was GAA/CTT. Along with the targeted SSR motifs, for which biotinylated oligos have been used, a number of additional motifs were obtained and used for marker production. The average repeat number for perfect GA/TC motif was higher in *C. pepo* (17.8) than in *C. moschata* (10.4). In both species the repeat numbers of most GA/TC SSRs ranged from 7 to 11 (55.6% in *C. pepo* and 69% in *C. moschata*). However, while in *C. pepo* 25% of the perfect GA/TC SSRs showed repeat numbers  $\geq 20$ , in *C. moschata* only 3.9% were  $\geq 20$ .

#### Assessment of SSR polymorphism and transferability to other species

All 230 primer pairs developed from *C. pepo* were initially tested in the Austrian oil-pumpkin variety Gleisdorfer Ölkürbis (*C. pepo*). Of these, 193 (84%) gave a discrete PCR product with expected size and 19 (8.3%) yielded a complex banding pattern or a smear. The remaining 18 (7.8%) provided no product. All 193 working SSRs were subsequently tested in the 12 genotypes listed in Table 1,

**Table 3** Efficiency of SSR marker development in *C. moschata*

SSR development steps	Attrition	Total	%
Sequenced clones	0	1,344	100
SSR containing clones	1	610	45.4
Clones containing SSRs too close to one end	2	35	2.6 (5.7 of 1)
Primer pairs designed and tested	3	342	25.4 (56.0 of 1)
Working SSRs	4	307	22.8 (89.8 of 3)
Polymorphic SSRs	5	250	18.6 (81.4 of 4)

and 155 of them (80.3%) showed polymorphism with 2–9 alleles and an average PIC value of 0.62 (Table 2). All 342 primer pairs developed from *C. moschata* were initially evaluated in the cultivar Soler (Puerto Rico), and 307 (89.8%) amplified PCR products with expected size. Of these, 250 (81.4%) were polymorphic in the 12 genotypes with 2–7 alleles and an average PIC value of 0.65 (Table 3). In both species there were 3.3 alleles per locus on average. In total, 500 new SSR markers were developed, of which 405 showed polymorphism in the 12 genotypes.

Transferability of microsatellites across the three species is summarized in Table 4. Altogether 362 SSRs, that is, 76.2% of *C. pepo* and 70% of *C. moschata*, could be transferred to *C. equadorensis*, producing a unique allele in most cases. Transferability between *C. pepo* and *C. moschata* was even higher. Of the *C. pepo* SSRs 88.1% were transferable to *C. moschata* and 30.1% of them produced multiple alleles in three *C. moschata* genotypes. Of the *C. moschata* SSRs 87.3% could be used for *C. pepo* as well and 50.5% of them produced multiple alleles in the eight *C. pepo* genotypes.

#### An SSR-based map of *C. pepo*

Altogether 187 SSR markers, 94 developed from *C. pepo* and 93 from *C. moschata*, were polymorphic between the parents O5 and CN. Seven SSRs showed distorted segregation, but six of them could be mapped. As a first step, these SSR markers along with six SCARs and two morphological traits (*h* for hull-less seed and *B* for bush growth habit) were used to construct the skeleton map. This map finally comprises 172 SSRs, two morphological traits and three SCARs. It spans 1008.6 cM and contains 25 linkage groups (not shown).

Taking the skeleton map as fixed order, the *C. pepo* map, published by Zraidi et al. (2007), was updated. Altogether, the new map contains 659 loci: 178 SSR, 244 AFLP, including 105 produced in this experiment, 230 RAPD, five SCAR markers, and two morphological traits (*h* and *B*) (S1). A detailed description of the updated map is given in Table 5. Figure 1 shows the map with only the SSR markers. AFLP and RAPD markers are not shown in this map, but map distances occupied by them are included into the distance between the respective two SSR markers. For example, both ends of LGp11 without markers

represent the positions of AFLPs and RAPDs; or in LGp8, between SCAR\_1476 and CMTm66, the distance is longer than 50 cM because of the hidden AFLP and RAPD markers between them. The complete map is available as supplementary information (S1 in supplement). Out of the five SCAR markers, two were converted from co-dominant AFLP markers, and three were derived from RAPD markers. The map spans 1,936 cM with 20 linkage groups ranging in length from 24 cM in LGp19 to 224.8 cM in LGp3. The total number of markers in individual linkage groups varies from 11 to 53, and the marker density varies from 2.0 to 4.7 cM. On average, one linkage group covers 96.8 cM and contains 33 markers, that is, there was one marker in every 2.9 cM (Table 5). The distribution of SSR markers in the individual linkage groups ranges from 2 to 15 (Fig. 1; Table 5). The estimated genome length ( $G_e$ ) of *C. pepo*, according to the results of JoinMap, is 2,230.9 cM with the confidence interval between 2,175.6 and 2,288.1 cM. The observed genome map coverage ( $C_o$ ) is 86.8%. The corresponding linkage groups in the published map and in the updated map of this paper are listed in Table 5. The updated map is based on the marker order of the skeleton map as built by JoinMap. The published *C. pepo* map was created using Mapmaker. In the updated map there were two new linkage groups (LGp18 and LGp19) and each of LGp10, LGp11, LGp14, LGp15, and LGp20 included two linkage groups from the previous published map.

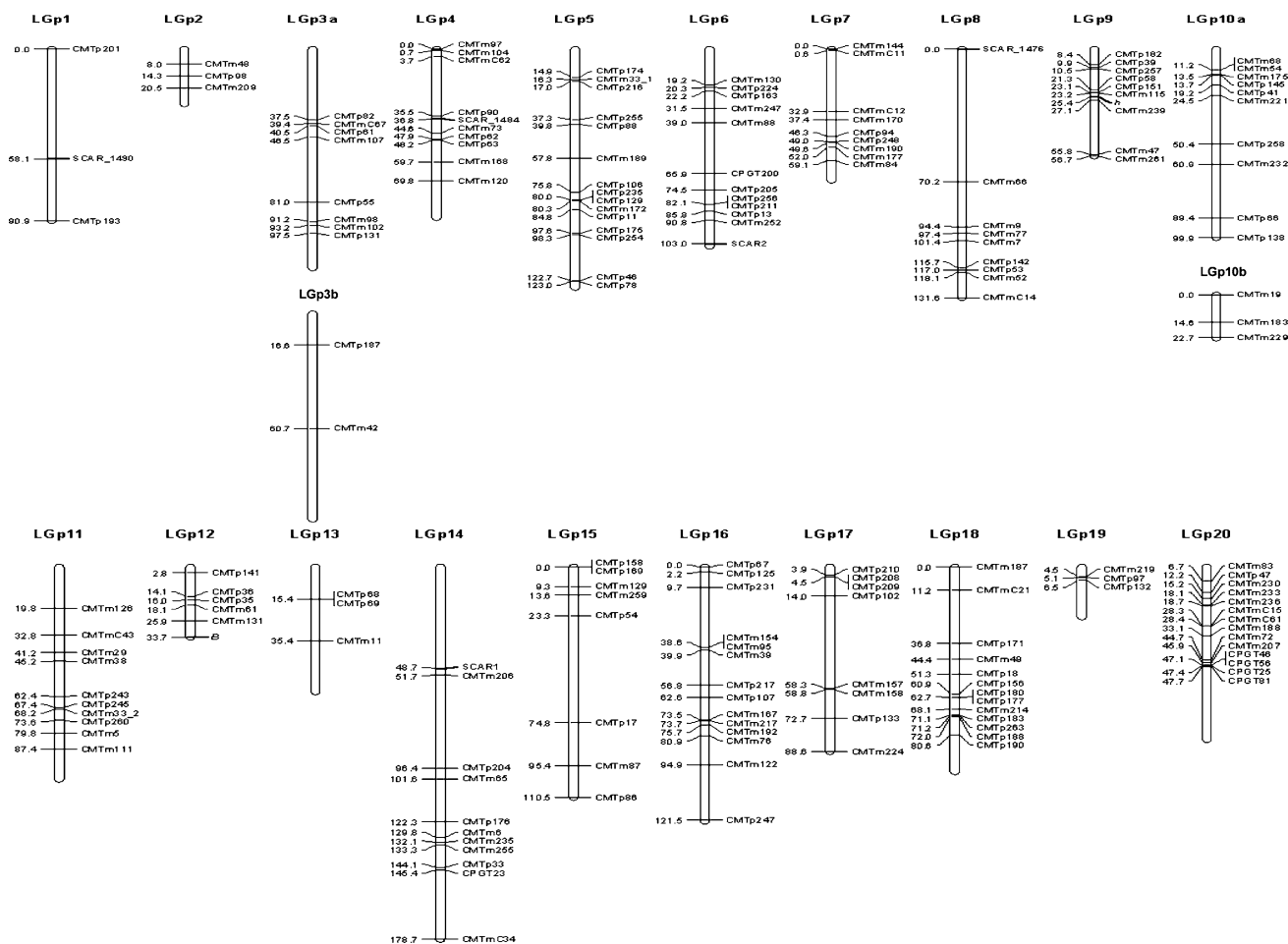
## Discussion

### Efficiency of SSR development from genomic libraries

Different enrichment technologies significantly increased the efficiency of microsatellite marker development (Kijas et al. 1994; Edwards et al. 1996; Zane et al. 2002). In non-enriched libraries the frequency of fragments containing SSRs may be as low as 0.7%, as was reported by Chiba et al. (2003) for melon. In the process of SSR development, Squirrell et al. (2003) defined the successive loss of sequenced fragments and designed primers, until arriving at a final number of “working SSRs” producing discrete bands of expected size, as “attrition rate”. This is an erosion of the originally high number of sequenced clones to a

**Table 4** Transferability of microsatellites across the 3 species: *C. pepo*, *C. moschata*, *C. equadorensis*

Origin	Total working SSRs	Transferable to <i>C. equadorensis</i>	Transferable to <i>C. pepo</i>		Transferable to <i>C. moschata</i>	
			Total	Polymorphic SSRs	Total	Polymorphic SSRs
<i>C. pepo</i>	193	147 (76.2%)	–	–	170 (88.1%)	58 (30.1%)
<i>C. moschata</i>	307	215 (70.0%)	268 (87.3%)	155 (50.5%)	–	–



**Fig. 1** A genetic linkage map of *C. pepo* showing only SSR, SCAR markers and the two morphological traits. Linkage groups (LGp) are numbered consecutively, subgroups are indicated by **a** and **b**. SCAR

markers are *bold*, and morphological traits are *bold and italic*. Cumulative marker distances are given in cM on the left side of the linkage groups

small final number of working SSRs. The reasons for erosion are (1) the absence of a useful SSR sequence in the clone, (2) duplications, (3) SSR sequences too close to one end of the fragment, and (4) primer pairs that produce either no fragment, or a complex banding pattern, or a smear. In our case this attrition rate was 80.2%, that is, approximately every fifth sequenced clone contained a working SSR. In different *Brassica* species, using enzyme digestion followed by colony hybridization for enrichment, Lowe et al. (2004) achieved similar enrichment efficiencies. After digesting genomic DNA with restriction enzymes in tall fescue (*Festuca arundinacea* Schreb.), Saha et al. (2006) reported 70% of clones containing SSR motifs. However, only every tenth of the sequenced clones produced a working SSR. One major reason for the large difference in efficiency to produce working SSRs in *Brassica* and *Cucurbita*, in comparison to tall fescue, may be due to genome complexity. The size of the *Cucurbita* genome is 1C ~ 0.5 pg DNA, and that of diploid *Brassica* is slightly higher, 1C between 0.6 and 0.7 pg DNA. In

contrast, the genome of tall fescue has 6.1 pg DNA and contains a large amount of repetitive sequences.

#### Characterization of microsatellites

In plants, the most frequently found SSR motif appears to be GA/TC (Lagercrantz et al. 1993; Li et al. 2002; Morgante et al. 2002; Saha et al. 2006). The genus *Cucurbita* seems to be no exception; 279, that is, 55.8%, of the 500 working *Cucurbita* SSRs have the GA/TC motif, but only 6 the CA/GT motif. The GA/TC microsatellite was found most frequently in the 5' flanking regions of genes in plants (Morgante et al. 2002; Zhang et al. 2004; Li et al. 2004). Song et al. (2005) found SSR markers within 5 cM to known genes in a much higher frequency than would be expected. Thus, GA/TC microsatellites may have a high probability to be linked to important traits.

A much higher proportion of perfect GA/TC SSRs having  $\geq 20$  repeat units (up to 69) was found in *C. pepo*

**Table 5** Description of the 20 linkage groups (LG) of the updated *C. pepo* map

Linkage groups	Map length (cM)	No. of markers	Marker density (cM/marker)	Maker types				Morphological traits	Corresponding linkage groups in the published map
				AFLP	RAPD	SSR	SCAR		
LG p1	90.9	29	3.1	12	14	2	1	–	LGpc1
LG p2	29.2	11	2.7	6	2	3	–	–	LGpc2
LG p3	224.8	48	4.7	17	21	10	–	–	LGpc3
LG p4	89.4	34	2.6	14	10	9	1	–	LGpc4
LG p5	126.3	43	2.9	19	9	15	–	–	LGpc5b
LG p6	104.5	38	2.8	13	13	11	1	–	LGpc6
LG p7	69.6	32	2.2	13	10	9	–	–	LGpc7
LG p8	131.6	42	3.1	16	16	9	1	–	LGpc8
LG p9	56.7	28	2.0	11	7	9	–	<i>h</i>	LGpc9
LG p10	122.6	53	2.3	22	18	13	–	–	LGpc10; LGpc18
LG p11	101.9	37	2.8	11	16	10	–	–	LGpc11; LGpc5a
LG p12	33.7	12	2.8	2	4	5	–	<i>B</i>	LGpc12
LG p13	59.9	16	3.7	6	7	3	–	–	LGpc13
LG p14	178.7	48	3.7	17	20	10	1	–	LGpc14; LGpc12
LG p15	110.5	36	3.1	19	9	8	–	–	LGpc15; LGpc19
LG p16	112	37	3.0	11	12	14	–	–	LGpc16
LG p17	88.6	25	3.5	7	10	8	–	–	LGpc17
LG p18	98.3	41	2.4	14	14	13	–	–	–
LG p19	24	11	2.2	4	4	3	–	–	–
LGp20	82.8	38	2.2	10	14	14	–	–	LGpc20; LGpc21
Total	1,936	659	2.9	244	230	178	5	2	–

than in *C. moschata*. A further difference between the two species was the presence of a cluster of highly similar sequences with the GA/TC repeat in *C. pepo*, which was found only twice in *C. moschata*. Primer pairs designed for 18 members of this cluster produced a smear, suggesting that this sequence was highly repeated in *C. pepo*, but occurred in a much lower frequency in *C. moschata*. These structural differences between the genomes of the two species might have arisen by an accelerated evolution of *C. pepo* after their separation from a common ancestor. The difference could be related, as Linda Wessel-Beaver suggests (personal communication, 2006), to their different use after domestication, before leaving the continent of origin. *C. pepo* has always been consumed immature, thus selection pressure for earliness might have been strong. On the other hand, *C. moschata* has been predominantly used as mature fruit, requiring as long as 4 months for ripening. During early cultivation period fruits could even be “stored” in the field and picked as needed. Thus, under tropical conditions in the region where both species originated, *C. pepo* might have undergone two cycles of selection per year whereas *C. moschata* had only one. Mutations could accumulate in *C. pepo* faster than in *C. moschata*.

#### The transferability of microsatellites across species

Within the Cucurbitaceae family, SSR markers developed for the genus *Cucumis* are generally transferable between cucumber and melon (Danin-Poleg et al. 2000, 2001; Ritschel et al. 2004). However, these markers have a low transferability to *Cucurbita* (Katzir et al. 1996). To assess genetic relationships in *C. pepo*, Paris et al. (2003) attempted to apply cucumber and melon SSR markers, described by Katzir et al. (1996) and Danin-Poleg et al. (2001). Seven melon and cucumber SSRs were used; they could, however, only be interpreted as dominant markers. Previously, 102 *Cucumis*-SSR primers developed by Fazio et al. (2002) were tested for polymorphism in four *C. pepo* genotypes for mapping (Zraidi et al. 2007). Only a few of these primer pairs generated a band, and none were polymorphic in the tested materials. These results suggest that SSRs are not easily transferable between genera within the Cucurbitaceae.

The genomic SSRs developed from *C. pepo* and *C. moschata* show a high (~90%) inter-species transferability. The SSR flanking sequences used for primer design are highly conserved, representing orthologous loci in these two species. This suggests a close phylogenetic relationship

between the two species, confirming the findings of Sanjur et al. (2002) and Stift et al. (2004). Of the 500 working SSRs, 405 were polymorphic in the set of 12 *Cucurbita* genotypes. Most likely, SSR markers could be applied to more species within the *Cucurbita* genus. There were 50.5% SSRs of *C. moschata* showing polymorphism in *C. pepo*, whereas only 30.1% of *C. pepo* SSRs appeared polymorphic in *C. moschata*. This can, however, also be due to the lower number of tested *C. moschata* (three) than *C. pepo* (eight) genotypes. The higher percentage of *C. pepo* microsatellites, which could be transferred to *C. ecuadorensis* in comparison to *C. moschata* (Table 4), implies a closer genetic relationship between *C. pepo* and *C. ecuadorensis* than between *C. moschata* and *C. ecuadorensis*, which is in line with findings of Sanjur et al. (2002). The transferability of SSRs will be useful for comparative mapping and evolution studies within the genus *Cucurbita*.

### *C. pepo* map

The first published *C. pepo* map, based on the F<sub>2</sub> population of O5 × CN, was composed solely of AFLP and RAPD markers (Zraidi et al. 2007). Such markers are useful for map construction of minor crops and in case of limited financial resources. They have the disadvantage of not being transferable to other populations, which greatly limits their application. The reason to update this published map was to create a more saturated and anchored linkage map for *C. pepo*, using the newly developed SSR markers. First an SSR skeleton map and then a high-density map were constructed. The high information content and transferability of SSRs are decisive for comparative alignment and integration of related maps. AFLP and RAPD markers filled the gaps where SSR markers were poorly represented and often facilitated linkage between unlinked SSR markers. Finally, only nine of the 187 polymorphic SSRs remained unlinked. The present *C. pepo* map contains 659 loci in total. Marker density thereby increased from 6.9 cM in the published map to 2.9 cM in the present map. However, marker density differs among linkage groups, and there are still two groups (LGp3 and LGp10) containing subgroups, which so far could not be linked. The SSR markers are distributed among all 20 linkage groups, which probably represent the 20 chromosomes of the haploid genome of *C. pepo*.

Twenty-one SSRs in this map (11.2%) have a null allele in one of the mapping parents. SSR markers are generally co-dominant, but the occurrence of null alleles in some genotypes has frequently been reported, for example, by Stachel et al. (2000) in wheat and Billotte et al. (2005) in oil palm (*Elaeis guineensis* Jacq). Only one of these 21 SSRs remained unlinked in the map.

Along with SSRs and two gene loci, six co-dominant SCARs were used to establish the skeleton map. Three (SCAR\_1476, SCAR\_1484, and SCAR\_1490) were converted from RAPD markers and three (SCAR1, SCAR2, and SCAR3) from co-dominant AFLPs by separately cutting out and sequencing the two co-dominant bands. Though AFLP and RAPD markers in most cases are dominant, co-dominant alleles can frequently be detected in both as well. In the previous *C. pepo* map out of 125 AFLP markers 12 were co-dominant (Zraidi et al. 2007), in this study 17 out of the 105 new AFLP markers turned out to be co-dominant.

The low rate of SSRs with distorted segregation (3.7%) was similar to the one found with AFLPs and RAPDs, when creating the first *C. pepo* map (Zraidi et al. 2007). The SSRs with distorted segregation are all located in different linkage groups. The low level of segregation distortion between O5 (“Lady Godiva”) and CN (the Italian crookneck variety “Bianco Friulano”), found with all three marker types, indicates regular meiosis and very low pre- or post-zygotic selection between the two parents.

Using the same mapping population, the present map, made by JoinMap and containing 659 loci, covers 1,936 cM. Thus it is much shorter than the previous map (2,234 cM) published by Zraidi et al. (2007), which contained only 323 markers, but was constructed by MAPMAKER. Comparable results were reported by Senior et al. (1997) and Vuylsteke et al. (1999) mapping the maize genome. They observed that, despite the larger number of markers, when using JoinMap the lengths of the individual linkage groups were shorter than those constructed by MAPMAKER. The reduction of map length using JoinMap has frequently been reported (Castiglioni et al. 1998; Bradeen et al. 2001; Gosselin et al. 2002; Tani et al. 2003; Pradhan et al. 2003). The inter-specific *Cucurbita* map, constructed by Brown and Myers (2002) using MAPMAKER, spanning 1,954 cM, comprises only 148 RAPD markers. These observed discrepancies could be due to the difference in the algorithms used by the two mapping softwares. The maximum-likelihood method used by MAPMAKER assumes the absence of crossover interference. However, when interference is present, JoinMap correctly produces shorter maps, even though both programs use the Kosambi mapping function (Stam 1993). Due to this difference, JoinMap will usually produce shorter maps than MAPMAKER, even using the same data set and same settings (Liebhard et al. 2003). As a consequence, genome length ( $G_c$ ) estimate by the methods of Hulbert et al. (1988) is also influenced by mapping software. In a pedigree of Sugi (*Cryptomeria japonica*)  $G_c$  was 2,168.5 cM according to MAPMAKER and 1,632.4 cM according to JoinMap (Tani et al. 2003). However, the observed map coverage ( $C_o$ ) is less affected



by the mapping software, because  $C_o$  is defined as the ratio of the observed length of the map ( $G_{or}$ ) and the estimated genome length ( $G_e$ ) (Cervera et al. 2001). The  $C_o$  of 86.8%, as calculated in this work, is very high.

Distribution of SSR markers on the current *C. pepo* map is non-random (Fig. 1). Their numbers range from 2 to 15 per LG. There are four LGs (LGp1, LGp2, LGp13 and LGp19) with less than five SSRs. SSR clustering or uneven distribution are usually found in linkage maps of plant species, for example, tomato (Broun and Tanksley 1996; Areshchenkova and Ganal 1999), Italian ryegrass (Hirata et al. 2006), rice (McCouch et al. 2002), barley (Ramsay et al. 1999), conifer (Elsik and Williams 2001), and sunflower (Tang et al. 2002). There are three possible explanations for the uneven distribution of SSR markers:

- (1) Cardle et al. (2000) and Morgante et al. (2002) reported that microsatellites are preferentially associated with the nonrepetitive fraction of plant genomes. In *Arabidopsis thaliana*, AG/CT SSRs occur significantly more frequently in transcribed regions and 5'UTRs than in introns and 3'UTRs (Morgante et al. 2002). Moreover, sequence analysis of DNA contigs in wheat and barley has shown the presence of clusters of closely linked genes forming gene islands that are separated by large stretches of repetitive DNA (Wicker et al. 2001, 2005; Varshney et al. 2006).
- (2) Homozygous regions between mapping parents may lead to gaps and insufficient coverage in the map.
- (3) The development of SSR markers from genomic DNA affects their distribution on the linkage maps. Ramsay et al. (1999) reported that SSRs isolated from genomic DNA libraries tend to concentrate in retrotransposons and dispersed repetitive-element DNA.

All the SSRs in this research were developed from genomic libraries. When more ESTs will be available in *Cucurbita* they might improve the distribution of SSRs in the current map.

In Austria, *C. pepo* has been cultivated for centuries primarily for seed oil production (Teppner 2000). A special feature of Austrian oil-pumpkin varieties is the completely hull-less seed coat (Teppner 2000; Zraidi et al. 2003), which is controlled by a recessive major gene, *h* (hull-less seed) (Schöniger 1950). Four SSR loci were found 1.5–3.6 cM from the hull-less locus (*h*) in LGp9. SSR markers closely linked to *h* would in the future be useful for map-based cloning of the gene responsible for testa lignification in *Cucurbita*. Cultivars with bush growth habit would increase the productivity in limited planting areas, so this is an important economical trait. The SSR marker CMTp131 in linkage group LGp12 was 7.8 cM from the bush locus (*B*). In a back-cross breeding program for more than one

trait, for example, combining hull-less seeds and bush growth habit, this marker could be very useful.

A saturated linkage map with entire genome coverage allows broad applications of molecular markers in plant breeding. Increasing marker density can easily be accomplished by using different mapping populations that segregate for important agronomic traits and harbor enough polymorphic SSR markers. Due to the good transferability between species and high reproducibility of SSR markers, they can easily be applied for map integration across different *Cucurbita* species, whose genetic variation can be utilized for breeding. The transferability of *Cucurbita* SSRs to *Cucumis* still awaits investigation.

In conclusion, we developed 500 microsatellite markers from SSR-enriched genomic libraries. They show a high transferability among species within the genus *Cucurbita*. The first *C. pepo* consensus map (Zraidi et al. 2007) has been updated by 178 SSR and new AFLP markers. The updated map contains 20 linkage groups with a map density of 2.9 cM. Four SSRs were detected to be linked to locus *h* (hull-less seed trait) and one SSR to locus *B* (bush growth habit). The primer sequences for the 500 SSRs and the updated *C. pepo* map with 659 loci are available as supplementary information.

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