Increased apomixis expression concurrent with genetic and epigenetic variation in a newly synthesized *Eragrostis curvula* polypl oid

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*Eragrostis curvula* includes biotypes reproducing through obligate and facultative apomixis or, rarely, full sexuality. We previously generated a “tetraploid-dihaploid-tetraploid” series of plants consisting of a tetraploid apomictic plant (T), a sexual dihaploid plant (D) and a tetraploid artificial colchiploid (C). Initially, plant C was nearly 100% sexual. However, its capacity to form non-reduced embryo sacs dramatically increased over a four year period (2003–2007) to reach levels of 85–90%. Here, we confirmed high rates of apomixis in plant C, and used AFLPs and MSAPs to characterize the genetic and epigenetic variation observed in this plant in 2007 as compared to 2003. Of the polymorphic sequences, some had no coding potential whereas others were homologous to retrotransposons and/or protein-coding-like sequences. Our results suggest that in this particular plant system increased apomixis expression is concurrent with genetic and epigenetic modifications, possibly involving transposable elements.

A pomixis in plants refers to a diverse group of developmental behaviors resulting in asexual reproduction through seeds⁵. Apomictic individuals bypass both meiotic reduction and egg cell fertilization to produce offspring that are genetic replicas of the maternal plant⁵. Few genotypes appear to be obligate for the apomictic trait, and those that do not totally exclude sexual processes are termed “facultative apomictic”, because they are able to produce seeds through both sexual and asexual means⁶. The line between apomixis and sexual reproduction is somewhat blurry, fueling the perception that the trait might have emerged from deregulation of sexuality rather than from the establishment of a new function⁴. Indeed, previous studies have shown that apomixis and sexual reproduction share key regulatory mechanisms⁵.

Given the established relationship between sexual and apomictic pathways, different models involving genetic and epigenetic mechanisms have been proposed to explain the occurrence of apomixis at the molecular level⁶. There is increasing evidence to support epigenetic control of apomixis⁷, including the observations that in most apomicts chronological and structural reproductive pattern variations occur in individual plants in response to environmental changes⁸,⁹. Variations in apomictic reproduction rates in response to environmental influences have been reported for several species and conditions⁷,⁸,¹⁰,¹¹.

Weeping lovegrass (*Eragrostis curvula* [Schrad.] Nees) is a perennial grass native to Southern Africa that displays a type of apomixis called pseudogamous diplospory¹². The megasporocyte of *E. curvula* undergoes two rounds of mitotic division to form a non-reduced tetranucleate embryo sac with an egg, two synergids, and one polar nucleus¹³. The *E. curvula* complex includes cytotypes with different ploidy levels (from 2x to 8x) that may undergo sexual reproduction, facultative apomixis or obligate apomixis¹⁴. Diploid (2n = 2x = 20) plants are sexual and rare¹⁵. Polyploids reproduce mainly by obligate apomixis, although both sexuality and facultative apomixis have also been reported¹⁴.

An euploid “back-and-forth” (4x - 2x - 4x) plant series was constructed displaying different ploidy levels and reproductive modes but sharing a common genetic background¹⁶. In order to construct the series, a fully sexual diploid plant D was generated from an apomictic tetraploid plant T, through inflorescence in vitro culture¹⁶. Seeds from a first generation diploid plant, derived from D after self-pollination, were treated with colchicine, leading to
the artificial generation of a tetraploid genotype named C. This colchloid initially produced variable progeny (as tested with RAPDs) and was subsequently classified as “highly sexual”. However, the possibility of a low level of apomixis expression was not discarded, because a plant with an increased ploidy level was detected among the progeny, probably originating from fertilization of an unreduced egg cell. Four years later, the reproductive mode of C was re-analyzed. Surprisingly, the plant exhibited a high proportion of apomictic events (around 85–90%, depending on the technique used to estimate the reproductive mode). Based on these results, Meier et al. proposed that the temporary expression of high levels of sexual reproduction observed in genotype C might be explained by the genomic stress caused by the in vitro culture and colchicine-induced chromosome duplication procedures.

The objective of the work reported here was to re-analyze the levels of apomixis in colchiploid plant C and characterize the range and nature of the genetic and epigenetic changes occurring during the period from 2003 to 2007, when the change in apomixis expression occurred, and then during the period 2007–2013.

Results

Analysis of reproductive mode. Plants were obtained as is shown in Fig. 1. As was mentioned above, the reproductive mode of plant C was analyzed in 2003 and 2007. An updated cytoembryological analysis of plant C reproductive mode was carried out in year 2009. A total of 233 ovaries at the correct stage were observed. Among these, 24 were found to be meiotic and 209 apomeiotic, corresponding to a potential sexuality level of 10%. Figure 2 shows the embryo sac development in the colchiploid plant C showing sexual (2a and 2c) and apomictic (2b and 2d) processes.

These results indicated that in 2009 the plant continued to display a level of apomixis as high as that in 2007. Therefore, the rate of apomixis in colchiploid plant C shifted from near 0% in 2003 (21) to 85–90% in 2007 (16), and remained at similar values in 2009 (89.7%).

Genetic and epigenetic structure of plants T and C. We searched for concurrent genetic and/or epigenetic modifications in plant C in 2003 and 2007, when the change in the rate of apomixis was observed. Genomic DNA samples extracted in 2003 and 2007 from plant C and the apomictic tetraploid control plant T were used to produce AFLP and MSAP profiles. Comparisons were performed between plants for the same year and between years for the same plants. A new MSAP analysis of both plants was conducted during the years 2007, 2011 and 2013.

Out of the 14 AFLP primer combinations used in samples from 2003 and 2007, five (5) showed no amplification, four (4) showed absence of amplification in at least one lane, and five (5) showed amplification in all lanes (E36-M37; E42-M36; E33-M38; E33-M37; E33-M33). The initial number of markers counted for each primer combination was 174, as follows: E36-M37: 44 markers; E42-M36: 32 markers; E33-M38: 35 markers; E33-M37: 31 markers; E33-M33: 32 markers. However, 39 individual bands showed inconsistencies in one of the technical replicates. Therefore, the corresponding markers were eliminated from analyses, rendering a total of 135 final markers with 74% (99) being monomorphic and 26% (36)
Three primer combinations did not amplify one or more lanes. In the first comparison (2003–2007), two did not amplify at all.

Full methylation of the external or both cytosines prevents digestion by both enzymes. HpaII will not cleave the DNA if either of the cytosines is fully hemi-methylated (single strand). On the contrary, MspI will not cleave the DNA if fully methylated (double-strand).

Cluster analysis was performed to estimate genomic similarity of the two plants (T and C) in 2003 and 2007. The results showed higher similarity in 2003 (90%) compared with 2007 (83%). This observation indicates that genome sequences diverged during this four-year period, since both plants have genetically changed mostly on the emergence of new bands (0/0/1/1 patterns) and three bands were associated with apomorphic expression (1/0/1/1 patterns).

To examine genome evolution in each particular plant during the period considered, band profiles were analyzed individually for each plant (Table 2). The total amount of variation over time was similar between plants (15–17%) (Table 2), with the appearance of new bands more frequent than the disappearance of former ones (Supplementary Table S2). This suggests the occurrence of de novo sequence insertion in the genome, because random point mutations would have led to both appearance and disappearance of bands at the same rate, while deletions would have led to most bands vanishing.

Cluster analysis was performed to estimate genomic similarity of the two plants (T and C) from 2003–2007. A similarity matrix was constructed from the MSAP data. Cluster analysis showed that after the four-year period, the plants underwent some changes in their genome (15–17%, see Table 2), the variation observed should be of epigenetic origin.

Polymorphisms arising between 2003 and 2007 were individually analyzed in T and C plants (Table 4). T polymorphic bands corresponding to demethylations were as frequent as those corresponding to methylations (8% demethylations vs. 7% methylations). Interestingly, demethylations mainly involved loci methylated at internal cytosines (12 out of 21) (see Supplementary Table S4). In C, by contrast, the time-variable polymorphic bands corresponded mostly to methylations (15% methylations vs. 5% demethylations). Transformations from internal-cytosine methylation to full methylation were the most frequent ones (See Supplementary Table S4).

The rate of methylation occurring in C was higher than that observed in T (15% vs. 7%). In genotype T, internal methylation to full methylation switches (I/M switches) represented 0.7% of the total analyzed loci (2 bands out of 266), whereas in genotype C they reached 6% (16 bands out of 266) (Supplementary Table S4). These results reveal that the I/M switch was 8.5 times more frequent in C than in T. It is important to note that I/M transitions could represent both genetic and/or epigenetic alterations, as pattern M reveals absence of bands after digestion with either enzyme (full methylation) or, alternatively, absence of the site due to sequence variation. However, since similar rates of sequence mutation were detected in T and C by AFLP (15–17%, see Table 2), the variation observed should be of epigenetic origin.

A similarity matrix was constructed from the MSAP data. Cluster analysis showed that after the four-year period, the plants underwent some changes in their genome (15–17%, see Table 2), the variation observed should be of epigenetic origin.

| Profile | T Plant | C Plant |
|---------|---------|---------|
| Absence of changes over time | 104 | 83% | 107 | 85% |
| Presence of changes over time | 22 | 17% | 19 | 15% |
| Total | 126 | 100% | 126 | 100% |

Table 2 | Variation in the AFLP profiles in each individual plant (T and C) from 2003–2007

Markers with missing data in at least one of the entries were eliminated from the matrix.

| Profiles                                      | Number of bands | Percentage (total) | Percentage (polymorphic) |
|-----------------------------------------------|-----------------|--------------------|--------------------------|
| Monomorphic patterns                          | 99              | 74%                |                          |
| Polymorphic patterns that were variable between plants and conserved over time (conservative variations) | 3               | 2%                  | 8%                       |
| Polymorphic patterns that were conserved between plants and variable over time                  | 7               | 5%                  | 20%                      |
| Polymorphic patterns that were conserved neither between plants nor over time                   | 26              | 19%                 | 72%                      |
| Subtotal polymorphic patterns                 | 36              | 26%                 | 100%                     |
| Total                                         | 135             | 100%                | 100%                     |

Table 1 | AFLP profiles produced from Tanganyika (T) and UNST1131 (C) from 2003–2007

Markers with missing data in at least one of the entries were eliminated from the matrix.
similarity between the plants increased from 71% to 82%. These observations indicate that cytosine methylation patterns in both plants changed over time, with a tendency to acquire the same epigenetic structure at least in some loci. Examples of AFLP and MSAP gels are shown in Fig. 4.

A new comparison between plants was performed between the years 2007–2011 and 2011–2013. Using six (6) primer combinations (HM4-E37, HM4-E40, HM6-E32, HM6-E40, HM7-E32 and HM7-E40) 26 out of 240 MSAP markers were polymorphic between years (Table 5). A new similarity matrix was constructed using these markers in the same way than before (2003–2007) (Fig. 3c). This analysis showed that the level of variation between plants along the years was lower than before (2003–2007) showing closer similarity between plants. It seems like with time both plants tend to stabilize their epigenomes and reach a similar methylation status. Plant C in the periods 2007–2011 and 2011–2013 showed one (1) methylation and one (1) de-methylation and three (3) methylations and six (6) de-methylations, respectively. In the same periods T showed nine (9) and six (6) methylations and two (2) and eight (8) de-methylations, respectively. Comparisons between plants showed that five (5) loci were methylated in T in 2011, but were not affected in C. The same five (5) loci were de-methylated in T in 2013. The rest of the polymorphic loci were affected in both plants in the following way: one (1) locus was de-methylated in 2011 and two (2) loci in 2013 in both plants T and C. Related to methylations, one (1) locus was methylated in T and C (2013), one (1) locus in T (2011) and the same locus in C two years later C (2013). Another locus experiences the same situation but in C (2011) and later in T (2013). Other methylations and de-methylations also occurred in independent loci.

Cloning and sequencing of genetic and epigenetic polymorphic bands. Some of the polymorphic bands obtained with AFLP or MSAP were cloned, sequenced, and compared with sequences in the NCBI non-redundant protein sequence (nr) and MIPS databases. The cloned bands were selected to represent different patterns of variation. Most of the AFLP sequences showed similarity to protein-coding genes in short segments (4 out of 5) but no significant similarities were found for the AFLP sequences when searching against the MIPS database (Supplementary Table S5). By contrast, nearly half of the MSAP sequences (3 out of 8) showed similarity to Gypsy or Copia retrotransposons (Supplementary Table S6). The presence of sequences with similarity to proteins was further analyzed (Supplementary Table S6). We isolated 27 additional polymorphic bands originated from AFLP from the same series. All these 27 sequences were also similar to retrotransposons, pseudogenes or retrotransposons + pseudogenes (unpublished).

Based on these results, it can be concluded that some of the variations detected using MSAP occurred in genomic regions showing similarity to mobile elements.

### Table 3 | MSAP profiles produced from Tanganyika (T) and UNST1131 (C) from 2003–2007

| Profiles | Number of bands | Percentage (total) | Percentage (polymorphic) |
|----------|-----------------|--------------------|--------------------------|
| Monomorphic patterns | 201 | 76% |
| Polymorphic patterns that were variable between plants and conserved over time | | |
| Demethylations (T vs C) | 5 |
| Methylations (T vs C) | 1 |
| Subtotal | 6 | 2% | 9% |
| Polymorphic patterns that were conserved between plants but varied over time (conservative variations) | | |
| Demethylations (2003 vs 2007) | 7 |
| Methylations (2003 vs 2007) | 14 |
| Subtotal | 21 | 8% | 32% |
| Polymorphic patterns that were conserved neither between plants nor over time | | |
| Demethylations | 12 |
| Methylations | 17 |
| Demethylations/Methylations | 9 |
| Subtotal | 38 | 14% | 59% |
| Subtotal polymorphic | 65 | 24% | 100% |
| Total bands | **266** | **100%** |
Survey of retrotransposons in cDNA libraries generated from *E. curvula* genotypes T and C. To analyse the presence of transcripts related to retrotransposons in the plants under study, we screened EST databases corresponding to inflorescence cDNA libraries derived from T and C in 2003, constructed shortly after the colchicine-induced formation of plant C.

The percentage of sequences showing similarity to retrotransposons (E value ≤ 1 e-08) in the different EST libraries showed that such transcripts were more abundant in plant C at that time (Fig. 5). The Gypsy subfamily comprised the most common class of retrotransposons found, and it was especially abundant in plant C.

Discussion

In this work we confirmed the occurrence of significant variation over time in the rate of asexual reproduction in a newly-synthesized tetraploid *E. curvula* plant. During the period from 2003 to 2007 the rate of asexual reproduction changed from nearly 0% to almost 90% in the colchloid plant C, as revealed by cytoembryological studies. This modification of the apomixis level was accompanied by the occurrence of genetic and epigenetic modifications. Genomic polymorphisms were also detected over the same period in the natural tetraploid apomictic plant T, which did not exhibit any variation in the apomixis rate. However, since apomixis is proposed to be governed by one or few loci, any genetic modifications, even when slight, could be relevant. On the other hand, cytosine methylation levels were higher in the colchloid plant C with respect to the control during the analyzed period. Epigenetic modifications occurring during the same interval appeared to be mostly specific, making the plants more similar. It is interesting to note that even when the level of epigenetic variation between plants C and T was comparable, a higher rate of methylation occurred in the colchloid plant C. Thus, our results suggest that an increase in apomixis rate occurred along with an increase in global genome methylation. Further experiments should be performed in order to test whether apomixis expression is functionally associated with the establishment of methylation in particular genomic areas.

In previous work, genetic characterization of the “back and forth” ploidy series used herein revealed high levels of genetic rearrangement derived from the chromosome number reduction followed by a duplication event. The transition from plant T (4x) to plant D (2x) involved the expected occurrence of band loss (due to allele elimination during dihaploidization) but also the unforeseen emergence of novel bands. Surprisingly, the restoration of the original ploidy level by colchloidization (transition from D to C), revealed recovery of many of the alleles lost during the transition from T to D. A

### Table 4 | Variation in the MSAP profiles in each individual plant (T and C) from 2003–2007

| Profile                        | T  | C  |
|-------------------------------|----|----|
| Absence of changes over time  | 227| 213|
| Methylations occurring over time | 18 | 39 |
| Demethylations occurring over time | 21 | 14 |
| Total bands                   | 266| 266|

### Figure 4 | Examples of MSAP and AFLP profiles

Left panel: MSAP amplicon produced with the HM5-E36 primer combination. 1 indicates samples obtained in 2003. 2 indicates samples obtained in 2007. M indicates digestion with Mspl. H indicates digestion with HpaII (e.g., sample T1M corresponds to genotype Tanganyika, year 2003, Mspl digestion). Different cytosine methylation patterns are indicated by arrows on the left. Right panels: AFLP amplicons produced with the M38-E33 and M38-E36 primer combinations. 1 indicates samples obtained in 2003. 2 indicates samples obtained in 2007 (e.g., sample T1 corresponds to genotype Tanganyika, year 2003). Different genetic patterns produced by primer combination M38-E36 are indicated by arrows on the left. Full length blots are presented in the Supplementary Information.
consequence of this restoration phenomenon was the establishment of remarkable genetic similarity between the T and C polyploids\(^2^0\). Similar results reporting non-Mendelian emergence of bands present in previous generations but not in the immediate ancestors has been reported by Song et al. (1995)\(^2^1\) for \textit{Brassica} synthetic allopolyploids. Meccia et al. (2007)\(^2^8\) proposed that plants belonging to the "back and forth" series had a genetic structure typical of their ploidy level, with the potential to be specifically restored via an unknown mechanism. Further analysis showed that the epigenetic landscape and the transcriptome of the plants appeared to be also characteristics of the ploidy\(^2^2\). The transition from T to D was characterized by a general modification in the epigenetic landscape, which was restored to a large degree during the transition from D to C\(^2^2\). These results as a whole indicate that there are specific genetic, epigenetic, and expression landscapes associated with a particular ploidy level. Moreover, the results presented here suggest that the epigenetic landscape restoration observed during the transition from D to C\(^2^2\) was still progressing during the period 2003–2007. A new comparison performed among years 2007, 2011 and 2013 showed a clear tendency to increase the similarity of both plants at this level. During these years the plants were still changing but were closer than in 2003. The corresponding similarity coefficients were 0.71 and 0.85 in 2003 and 2013 respectively.

A major point to be considered is the ploidy homogeneity of plant C, which was obtained after colchicine treatment of seeds from self-pollinated line D. The possibility of plant C of being a ploidy chimera was analyzed carefully by flow cytometry and chromosome counting in previous work\(^1^6\). Homogeneity was further demonstrated in order to register C as a new plant variety in the RNC (Registro Nacional de Cultivares, National Cultivars Database, Buenos Aires, Argentina) at INASE (Instituto Nacional de Semillas, National Institute of Seeds, Argentina) (Registration number: RC9193, 2006–2026). It is important to note that \textit{E. curvula} tetraploids and the diploid plant \textit{E. curvula} (experimental code UNST1131) are easy to differentiate at first sight in the greenhouse. Diploids show wider leaves with a typical pubescence on the base of the leaf adaxial surface and are taller than tetraploids\(^4^6\). None of the morphological characteristics of diploid plants were ever observed in the colchiloid plant C, which showed morphology that was essentially analogous to that of the other tetraploids. If line C was a chimera, we would have expected that at least parts of the plant would have displayed morphology characteristic of the diploids. Moreover, from 2002–2012, numerous vegetative tillers were separated from the original plant C in order to propagate the genotype. In many of the derived plants, ploidy levels were determined by flow cytometry and chromosome counting. In all cases, they were found to be tetraploid. In the period from 2002 to 2007 the meiotic behavior of plant C (male meiosis) was carefully analyzed in anthers originating from several panicles. A tetraploid configuration with normal meiosis was always observed. In 2007, reduced and unembryos embryos were always detected in the same inflorescences. Based on the above-mentioned considerations, the hypothesis of plant C being a chimera was discarded.

The modification of the apomixis rate in plant C occurred along with genetic and epigenetic modifications, the latter involving a higher rate of methylation when compared to control plant T. The variability detected involved retrotransposons and gene sequences, as revealed by cloning and sequencing of the polymorphic fragments. Moreover, plant C expressed more transcripts related to retrotransposons. Our results are in agreement with those reported by Ochogavia et al.\(^2^3\) who detected high rates of \textit{Gypsy} retrotransposon expression in tetraploid sexual genotypes of \textit{Paspalum notatum} when compared with apomictic genotypes of the same ploidy. Further work should analyze the role of these transposable elements in the transition from sexuality to apomixis.

| Profile       | 2007–2011 | 2011–2013 | 2007–2011 | 2011–2013 |
|---------------|-----------|-----------|-----------|-----------|
| Methylation   | 9         | 6         | 1         | 3         |
| Demethylation | 2         | 8         | 1         | 6         |

**Methods**

**Plant material.** An \textit{Eragrostis curvula} euploid “back-and-forth” plant series of different ploidy levels and reproductive modes was used in this work\(^9\). The series consisted of: 1) an obligate apomictic tetraploid plant T (cv. Tanganyika, \(2n = 4x = 40\)); 2) a diploid plant D (experimental code UNST1122, \(2n = 2x = 20\), fully sexual), generated from tetraploid T by inflorescence in vitro culture; and 3) a tetraploid plant C (experimental code UNST1131, \(2n = 4x = 40\), initially classified as highly sexual), obtained by colchicine duplication of R1 seeds of diploid genotype D (Fig. 1). DNA samples were collected from leaves of plants T and C in 2003, 2007, 2011 and 2013. Three additional adult plants belonging to the natural cv. Tanganyika, obtained from seeds, were used as controls in order to analyze the methylation differences among plants. The plants analyzed were grown under the same routine greenhouse conditions.

**Cytomorphological studies.** Inflorescences were collected at the beginning of anthesis, when it is possible to observe all of the embryo sac developmental stages, as described in Meier et al. (2011). Briefly, after fixation in FAA (50% ethanol, 5% acetic acid, 10% formaldehyde and distilled water), individual spikelets were dehydrated in a tertiary butyl alcohol series, embedded in paraplast (Johansen 1940), and then sectioned at 10 mm and stained with safranin-fast green. Observations were carried out with a Nikon Eclipse TE300 light transmission microscope (Tokyo, Japan).

**Amplified fragment length polymorphisms (AFLPs).** AFLP studies were performed on samples extracted from leaves, according to Vos et al.\(^7\). Genomic DNA (600 ng) samples were double-digested with the enzymes EcoRI and MseI. The resulting fragments were then ligated to EcoRI and MseI adaptors to produce template for further amplifications (primer and adaptor sequences are shown in Supplementary Table S7). PCR amplifications using technical duplicates were carried out with 14 AFLP primer combinations for the comparisons between years 2003 and 2007. PCR products were separated on 6% (w/v) denaturing polyacrylamide gels, silver-stained, and digitized for analysis. Bands with identical migration amplified from different plant/year samples were considered to represent a marker. The standard used to select a particular marker to be included in the analysis was that all representative bands should have produced consistent patterns (presence/absence) in both technical replicates. Variation of less than 5% between technical replicates was required for a primer combination to be included in the analysis. Genetic similarities between the samples were calculated using Jaccard’s coefficient, and UPGMA cluster analysis was performed using the NTSYS software package.

![Figure 5 | Contribution of retrotransposon sequences to cDNA libraries of \textit{E. curvula}. Each bar indicates the percentage of the retrotransposons in the library. Ec02: Library produced from Tanganyika (T) inflorescences, Ec04: Library produced from UNST1131 (C) inflorescences.](www.nature.com/scientificreports)
Methylation-sensitive amplified polymorphisms (MSAPs). Detection of cytosine methylation pattern modifications was carried out by performing MSAP studies on DNA samples extracted from leaves, according to Xu et al. The methylation-sensitive isoschizomers HpaII and MspI were selected as frequent-cutting enzymes, and EcoRI was chosen as a rare-cutting enzyme (primer and adaptor sequences are shown in Supplementary Table S7). PCR amplifications were carried out with 14 and 6 MSAP primer combinations (for comparisons between years 2003 and 2007 and among years 2007, 2011 and 2013, respectively), using technical duplicates. PCR products were separated on 6% (w/v) denaturing polyacrylamide gels, silver-stained, and digitized for analysis. Bands were counted only when they were present in both replicates. Variation of less than 5% between technical replicates was required for a primer combination to be included in the analysis. MSAP data derived from the four samples were converted into a binary matrix, with 0 representing monomorphic patterns (00 and 11) and 1 representing polymorphic patterns (01 and 10). Matrices were analyzed to determine similarity coefficients between pairs of individuals and for group clustering. Jaccard’s coefficient (J) was used as the similarity index, and UPGMA cluster analysis was performed using the NT SYS software package.

Isolation, cloning and sequencing of polymorphic DNA fragments. Bands of interest were moistened with distilled sterile water and excised from polyacrylamide gels using a scalpel, then cut into smaller sections and eluted with buffer solution 

Methylation-sensitive amplified polymorphisms (MSAPs) methylation patterns were carried out using the BLAST tools available at NCBI (National Center for Biotechnology Information, http://www.ncbi.nlm.nih.gov/) and MIPS at PlantsDB (http://mips.helmholtz-muenchen.de/plant/genomes.jsp).

In silico detection of retrotransposons in E. curvula cDNA libraries. The BLASTN algorithm was used to detect sequences potentially corresponding to retroelements in two cDNA E. curvula libraries constructed from inflorescences of genotypes T and C. The libraries were named as follows: Ec02 (Tanganyika, apomictic tetraploid, T) and Ec04 (UNST131, sexual colchihploid, C). The libraries were constructed in 2003.

Reference sequences used for search queries were obtained from a public non-redundant database (http://wheat.pw.usda.gov/ITMI/Repeats/). Matching sequences with E-value ≤ 1 e-08 were considered to be similar. Results were expressed as percentages for each library.

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