Cross Compatibility Analysis to Identify Suitable Parents of *Tagetes erecta* and *T. patula* for Heterotic Hybrid Breeding

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Abstract

Seven interspecific crosses of *Tagetes erecta* × *T. patula* were conducted in this study. The cross compatibility index, seed setting rate, germination rate, phenotype, ploidy level, heterosis over male parent and field performance were tested. The results showed that different cross compatibility indices of *Tagetes* vary greatly, from 10.44~114.41, which is far less than that of the intraspecific hybridization S-121 × I-506 index. The seed setting rate of each cross ranged from 15.67% to 49.0%, and the cross S-121 × TP-512 scored the highest seed setting rate, which was higher than that of the intraspecific cross S-121 × I-506 (35.85%). Compared with the male parent, F₁ has higher plant height, wider crown width, larger flower diameter and more ray florets and earlier bloom with two exceptions. The S-121 of *T. erecta* was diploid. All of the male parents of *T. patula* were tetraploids, and the combinations were triploids. The traits of plant height, plant width, flower size and number of ray florets showed positive heterosis over the male parents, and the days leading to flowering showed both positive and negative ones. Finally, S-121 × TP-379 and S-121 × TP-512 were selected as the two best combinations. This study confirmed that the two species of marigold could be crossed for better F₁ varieties with improved performance.

Keywords: heterosis; interspecific hybridization; *Tagetes erecta*; *T. patula*

Introduction

Both *Tagetes erecta* and *T. patula* are annual flowers in the Asteraceae family and are native to South America and Mexico (He et al., 2016). They can be grown under a broad range of climatic conditions and thus are cultivated worldwide. *Tagetes erecta*, often called the African marigold, has fully double flowers in various brilliant colors of variable height. It is of great importance for landscaping and can be used as bedding and as a potted plant which make a magnificent garden from spring to autumn. *Tagetes patula*, often called the French marigold, belongs to the genus *Tagetes* spp. The species is one of the most important annual ornamental plants in commercial cultivation (Ai et al., 2015; Cicevan et al., 2016). The French marigold is different from the African marigold in many ornamental traits. The former is smaller in flower size, dwarf and compact in the whole plant, and it has more flowers on each individual plant.

The common method of commercial seed production for marigolds is crosses by using male sterile lines as female parents for cross breeding purposes (Sreekala and Raghava, 2003; Namita et al., 2011; Ai et al., 2017). To date, an intraspecific cross has been developed to produce new varieties and seeds in *T. erecta* (Zhang et al., 2014). In this system, using male sterile lines of *T. erecta* as parents avoided the large manual work of emasculation to obtain F₁ hybrid seeds at a commercial production scale (He et al., 2009). Varieties of *T. patula* are inbreeding varieties and could be crossed with *T. erecta* as male parents (Li et al., 2005; Namita et al., 2009; Namita et al., 2011). F₁ hybrids of *T. erecta* × *T. patula* are often called triploid marigold and have a longer flowering duration (Namita et al., 2011). Few varieties of triploid marigold could be found in seed companies. One of the series was 'Zenith', including several colors. In ornamental plant breeding, interspecific hybridization is often used in combination as a system to

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Maintaining the concept of the importance and limited related research about marigolds, the objectives of this research were to: (1) create new crosses between two marigolds species to create new varieties with dwarf, compact, early flowering, double flower and vigorous growth traits, and (2) elucidate the heterosis over the male parents, and elucidate the difference between \textit{T. patula} and \textit{T. erecta} × \textit{T. patula}.

**Materials and Methods**

**Plant materials**

One male sterile line of \textit{T. erecta} (S-121) was designated as the female parent, while seven accessions of \textit{T. patula} (TP-508, TP-509, TP-510, TP-511, TP-512, TP-515 and TP-379) and one \textit{T. erecta} inbred line (I-506) were used as the male parents. The main traits of the parental lines are described in Table 1 and Fig. 1.

| Species          | Accession | Flower color | Flower type | Line          | Ploidy level | Source                              |
|------------------|-----------|--------------|-------------|---------------|--------------|-------------------------------------|
| \textit{T. erecta} | S-121     | Yellow       | Double (♂)  | Male sterile  | 2x           | Beijing Institute of Landscape Architecture |
|                  | I-506     | Orange       | Single (♀)  | Inbred lines  | 2x           |                                     |
| \textit{T. patula} | TP-508    | Yellow       | Anemone     | Inbred variety| 4x           | PanAmerican Seed Company            |
|                  | TP-509    | Orange       | Anemone     | Inbred variety| 4x           |                                     |
|                  | TP-510    | Yellow       | Crested     | Inbred variety| 4x           | PanAmerican Seed Company            |
|                  | TP-511    | Orange       | Crested     | Inbred variety| 4x           |                                     |
|                  | TP-512    | Red          | Anemone     | Inbred variety| 4x           | American Seed Company               |
|                  | TP-515    | Orange       | Double      | Inbred variety| 4x           | Company                             |
|                  | TP-379    | Yellow       | Anemone     | Inbred variety| 4x           |                                     |

Fig. 1. The phenotypic trait comparison of \textit{T. erecta}, \textit{T. patula} and their \textit{F}_{1} hybrids.
Artificial pollination
The study was conducted at the Beijing Institute of Landscape and Architecture, Beijing, China. The artificial pollination was conducted at the greenhouse in the autumn of 2015, and the field trial was conducted at the open experimental field in the spring of 2016. To ensure that the flowers of S-121 and the male parents engaged, the male sterile line of S-121 was sown in July 2nd (2015), and the rest of the materials were sown 18 days later. A total of 200-hole plugs were used for seed germination and the seedlings were transplanted into 12-hole plugs after another 4 week. The medium was the mixture of peat and vermiculite 4:1 which contains approximately 2g of 14-13-13 as base fertilizer per pot. Water soluble fertilizer, such as 14-0-14 and 20-10-20, was applied at approximately 50~100mg/L and watered and fertilized alternately. When the flower buds emerged, the water soluble fertilizer was applied at 10-30-20 every 15 days with the concentration approximately 200 mg/L.

The seven male lines were pinched for better branching and additional flowers. These plants bloomed approximately 60 days after transplantation. All of the crosses were created using artificial pollination. The crosses were carried out from September 28th to October 26th (2015) when their flowers were fully opened. Thirty male sterile plants and 144 plants of T. patula were used in this study in total. All of the pollen of each cross was collected on the same day, and pollination was started at the same time. For each cross, the pollen of the corresponding parent was collected on a sunny day morning after 10:00 AM. The flowers were bounced by hand, and the pollen dropped into a glass culture dish. The pollen was used to pollinate the male sterile plants of S-121 by a brush every other day and repeated 3~4 times to maintain enough pollen. More than 20 flowers were pollinated in each cross. The seeds were harvested and separately detailed in the gauze bags approximately 20 days later and stored at 4°C.

Field trial, data collection and statistical analysis
The field trials were conducted between February 16th and July 4th, 2016. The plants of the F<sub>1</sub> and parental lines were sown in February 16th and cultured under the same conditions as artificial pollination. The seven F<sub>1</sub> combinations and parents were evaluated in a randomized complete block design with three replicates, and each replicate contained 24 plants. The plants were transplanted into the open field on April 25th (2016) with an inter-row spacing of 40cm and grown under natural conditions. All the selected genotypes were treated uniformly with cultural practices for healthy growth and development.

Measurements of the ornamental traits were carried out from May 10th to June 20th, the time that all the plants were in full bloom. Five plants were randomly selected for each replicate. Ornamental traits, including plant height, width, flower color, flower size, flower type, number of ray florets and days leading to flowering were recorded. The mean and heterosis over male-parent (MP) values were calculated using the formula 100×(F<sub>1</sub>−MP)/MP. For different crosses, four to 33 capitulums were selected to calculate the distant hybridization cross compatibility index and seed setting rate. The distant hybridization cross compatibility index was calculated using the formula: seed number / capitulum number. Full seed number refers to the number of seeds which contain testa and embryo and developed from ovules. The seed setting rate was counted by the formula full seed number / flower number of one capitulum × 100%. The germination rate was counted using the formula germination number / total seed number × 100%. The cross compatibility, the seed number, seed setting rate, germination and heterosis were calculated using SPSS Statistics 17.0. In addition, the LSD (least significant difference) was used for multiple comparisons to determine the significant differences (P < 0.05) among the combinations for the mean values for each characteristic.

Chromosome counting and Flow cytometry analysis
The chromosomes were counted as previously described by Zhu (1982).

Accuri C6 flow cytometry (BD Company, the United States) was used to conduct a flow cytometry analysis. The tenders leaves (approx. 50 mg) were added to the precooled plate to which the precooled 1 mL lysate was added (LB01, Wu et al., 2016), and the leaves were quickly chopped with a sharp blade. In the process, the leaves should be soaked in the lysate. The supernatant of the prepared cell suspension was discarded. One hundred microliters of precooled lysis solution was added to 150 mL propidium iodide (PI) working solution and dye and placed at 4°C for 10min out of the light. Each sample was tested with 3 repeats, with at least 10000 cells with FL2-A channel meter. The CV (coefficient of variation) was less than 5%. The data was collected using CFlow Plus software (Wu et al., 2016).

Results and Discussion
Analysis of the cross compatibility index of T. erecta × T. patula
The cross compatibility index of the two Tagetes is shown in Table 2. The index of the seven crosses vary greatly, from 10.44~121.67; in particular, S-121 × TP-510 and S-121 × TP-511 display lower values of 19.93 and 10.44, respectively. In addition, the cross S-121 × TP-512 had the highest value among the interspecific crosses. The intraspecific hybridization S-121 × I-506 showed the highest cross compatibility index of 121.67.

When the value of the compatibility index is zero, it is proven that the parent cannot produce F<sub>1</sub> seed. Li et al. (2008) reported that the highest compatibility index was low with a mean value of 5.4 when the crosses were made using the diploid species of Dendranthema lavanduliföium, D. dichrum and D. nankingense. This study shows a high cross compatibility index and indicates that the crosses between the two species are compatible and there is no prefertilization barrier.

Analysis of seed setting rate and germination rate
The full seed numbers, total seed numbers per capitulum and seed setting rate of each cross are listed in Table 3. The number of full seeds varies greatly, from 36 to 122. S-121 × I-506 had the maximum number of full seeds with 122 and it has a significant difference with the crosses of S-121 × TP-508, S-121 × TP-510 and S-121 × TP-511. There is no significant difference in the total seed number.
Table 2. The cross compatibility index of T. erecta × T. patula

| Cross        | Seed number | Capitulum number | Cross compatibility index |
|--------------|-------------|------------------|--------------------------|
| S-121 × TP-379 | 1763        | 19               | 92.79                    |
| S-121 × TP-508 | 397         | 6                | 66.17                    |
| S-121 × TP-509 | 1674        | 16               | 104.63                   |
| S-121 × TP-510 | 279         | 14               | 19.93                    |
| S-121 × TP-511 | 94          | 9                | 40.34                    |
| S-121 × TP-512 | 1945        | 17               | 114.41                   |
| S-121 × TP-515 | 359         | 4                | 89.75                    |
| S-121 × I-506 | 4015        | 33               | 121.67                   |

Table 3. Seed setting rate and germination rate in different crosses of T. erecta × T. patula

| Cross        | Full seed number | Total seed number per capitulum | Seed setting rate | Germination rate |
|--------------|------------------|----------------------------------|-------------------|-----------------|
|              | Average          | Extreme                          | Average           | Extreme         |
| S-121 × TP-379 | 93 ab            | 45–136                           | 250 a             | 217–294         | 37.83 abc       | 84.0 af         |
| S-121 × TP-508 | 66 bcd           | 14–132                           | 213 a             | 167–290         | 29.33 bc        | 81.0 acg        |
| S-121 × TP-509 | 105 abc          | 37–173                           | 203 a             | 139–297         | 47.33 ac        | 67.0 bd         |
| S-121 × TP-510 | 37 d             | 23–58                            | 233 a             | 192–306         | 15.67 c         | 83.0 af         |
| S-121 × TP-511 | 36 d             | 22–48                            | 209 a             | 176–255         | 17.83 cd        | 83.0 af         |
| S-121 × TP-512 | 114 a            | 59–167                           | 231 a             | 149–298         | 49.0 e          | 77.0 g          |
| S-121 × TP-515 | 90 ab            | 60–112                           | 217 a             | 190–266         | 41.17 bc        | 65.0 d          |
| S-121 × I-506 | 122 a            | 69–180                           | 252 a             | 179–352         | 35.85 ac        | 92.0 e          |

The seed setting rate and germination rate in different crosses of T. erecta × T. patula

The seed setting rate varies from 15.67% to 49.0%. The cross S-121 × TP-512 shows the highest seed setting rate of 49.0%, and it had significant difference with S-121 × TP-508, S-121 × TP-510 and S-121 × TP-511. This is closely related to their genetic distance. Previous research suggested that T. patula was a close sibling species of T. erecta and T. patula might be an amphidiploid from T. erecta and T. tenuifolia or a species closely related to them (Towner, 1961). Some studies also proved this hypothesis (Zeng et al., 2010; He et al., 2016). Additional research about homologous chromosome pairing is suggested to conduct and observe if the synaptonemal complex could form.

The traits and heterosis of offspring resulting from cross-breeding

In whole plant type, the F1 plants of these seven combinations show no main stem, similar to their male parent varieties.

Fig. 1 shows the quantitative characteristics of the combinations and their parents. For the plant height, seven combinations are shorter than their female parents, but taller than their male parents (Fig. 1a). For plant width, the F1 hybrids are better than their male parents (Fig. 1b). For the flower size, all the combinations are larger than their male parents (Fig. 1c). For the trait of number of ray florets, pollination occurs.

This study shows that most of the interspecific crosses had a higher seed setting rate than the inner-specific ones. In many cases, the distant species often have different degrees of spatial and temporal isolation, different genetic material and information and inconsistent physiology, and they may cause difficulty in successfully hybridizing due to reproductive barriers (Deng et al., 2012). Reproductive barrier always exist in distant hybridization and render it unsuccessful (Deng et al., 2010). Better hybridization affinity can be achieved when the parents have a close genetic distance (Li et al., 2008). The two marigolds are two species and had reproductive isolation. However, they can produce seeds when crossed by artificial pollination; this indicates that the two species have a relatively close genetic relationship. Previous research suggested that T. patula was a close sibling species of T. erecta and T. patula might be an amphidiploid from T. erecta and T. tenuifolia or a species closely related to them (Towner, 1961). Some studies also proved this hypothesis (Zeng et al., 2010; He et al., 2016).

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all the F₁ offspring show more ray florets than their male parents (Fig. 1d). In addition, for the trait of days leading to flowering, the combinations bloom earlier than their male parents with the exception S-121 × TP-510 and S-121 × TP-511, which bloom 5 and 4 days later than their male parents, respectively (Fig. 1e).

For the quality characters, the traits of flower color, almost all of the combinations exhibit the same color with their male parents, except the cross S-121 × TP-512, whose flower color is orange (17A) and red (24A), with a gradual conversation from red color to orange along with blossom (Fig. 2).

As shown in Fig. 2, the female parent has no petals and pollen, and the male parents show different flower types including the anemone form (TP-379, TP-508, TP-509, and TP-512), crested (TP-510 and TP-511) and semidouble (TP-515). For the F₁ hybrids, the flower patterns of the crosses S-121 × TP-510 and S-121 × TP-511 are the same as that of their male parents. The crosses S-121 × TP-508, S-121 × TP-509, S-121 × TP-512, S-121 × TP-515 and S-121 × TP-379 display fully double flowers which differ from their male parents.

The ploidy level of each progeny material was estimated using flow cytometry analysis and chromosome counting.

Fig. 2. The flower traits, chromosome counting and flow cytometry analysis for the parent lines of *T. erecta* and *T. patula* and their F₁ hybrids
According to Fig. 2, the S-121 of *T. erecta* was diploid; all of the male parents of *T. patula* were tetraploids, and the combinations were triploids.

*T. erecta* is naturally diploid (2n=2x=24) while *T. patula* is allotetraploid (2n=4x=48) (Towner 1961; Chen et al., 1982; Qi et al., 2008; He et al., 2016). Thus, the chromosome number of the interspecific combinations of *T. erecta* and *T. patula* is 2n=3x=36 (Chen et al., 1982). Our results are consistent with the previous research.

The heterosis over male-parent for the five quantitative traits including plant height, plant width, flower size, number of ray florets and days leading to flowering were examined. The details are listed in Table 4. Most combinations had heterosis advantage compared with the male parent, with the exception of days leading to flowering. For plant height, the heterosis ranges from 13.6% to 60.4%, and the hybrid combination S-121 × TP-509 recorded the highest percentage of heterosis. The heterosis of the plant spread has a range of 2.9%~51.6%, and the cross S-121 × TP-379 obtained the highest heterosis of 51.6%, closely followed by the cross of S-121 × TP-509 (31.5%). The heterosis of flower size varied from 5.2% to 27.1%, and the heterosis of S-121 × TP-510 displayed the highest value. The highest heterosis for the ray floret number was attained by the cross S-121 × TP-379 (306.0%), which was closely followed by the cross of S-121 × TP-509 (198.2%). Most crosses have negative heterosis over their male parents for days leading to flowering with two exceptions.

**Field performance of the best crosses**

According to the main ornamental traits, including the flower pattern and color, flower number of individual plant and uniform population, both S-121 × TP-379 and S-121 × TP-512, are the best combinations. The performances are shown in Fig. 3. The two combinations show double flowers, compact plants and early flowering, which meet breeding target and market demands. S-121 × TP-379 have a relatively high seed setting rate (37.83%) and the highest germination rate of 84.0%, and with a mean of 26.2 cm in height, and 33.2 cm in plant width, double gold flower which is 6.3 cm in diameter and 54 days from sowing to flowering (Fig. 1, Fig. 3A, 3B).

S-121 × TP-512 have the highest seed setting rate of 49.0% and germination rate of 77.0%, with a height of 25 cm, spread of 36.4 cm, red and orange double flower, which is 6.7 cm in diameter, and 53 days from sowing to flowering (Fig. 1, Fig. 3C, 3D).

Table 4. Heterosis over the male-parent for the 5 ornamental traits in different crosses of *T. erecta* and *T. patula*

| Cross          | Plant height (% | Plant spread (% | Flower size (%) | Number of ray florets (%) | Days leading to flowering (%) |
|----------------|-----------------|-----------------|-----------------|--------------------------|-----------------------------|
| S-121 × TP-508 | 42.5 ad         | 2.9 ac          | 9.8 ad          | 148.9 a                  | -14.5 a                     |
| S-121 × TP-509 | 60.4 b          | 31.5 b          | 5.4 a           | 198.2 a                  | -11.7 a                     |
| S-121 × TP-510 | 42.5 abf        | 7.4 c           | 27.1 b          | 36.1 b                   | 9.8 b                       |
| S-121 × TP-511 | 41.9 df         | 14.4 abc        | 5.2 ac          | 45.6 b                   | 7.8 b                       |
| S-121 × TP-512 | 13.6 eg         | 12.7 abc        | 13.6 d          | 170.4 a                  | -5.4 c                      |
| S-121 × TP-515 | 25.1 cde        | 15.6 abc        | 6.6 ac          | 143.6 a                  | -5.7 cd                     |
| S-121 × TP-379 | 37.2 aef        | 51.6 d          | 6.8 ac          | 306.0 c                  | -1.8 d                      |

**Fig. 3. Individual and population performance of the selected combinations**
Conclusions

Hybridization between the male sterile line of the African marigold and inbred lines of French marigold was a good way to create better ornamental traits of marigold. According to the primary traits of compatibility, heterosis over male parents, and field performance, the crosses of S-121 × TP-379 and S-121 × TP-512 meet the breeding demand and were selected as the best combinations. S-121 × TP-515 has a relatively low germination rate, and further research is required to determine the reason. In addition, more studies about multyear and multisite trials of variety performances and seed production should be conducted for the future promotion of the selected crosses.

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References

Ai Y, He YH, Hu Y, Zhang Q, Pan C, Bao MZ (2014). Characterization of a novel male sterile mutant of Tagetes patula induced by heat shock. Euphytica 200(2):159-173.
Ai Y, Zhang CL, Sun YL, Wang WN, He YH, Bao MZ (2017). Characterization and functional analysis of five MADS-box B类 genes related to floral organ identification in Tagetes erecta. PLoS One 12(1):e0169777.
Ai Y, Zhang QH, Pan C, Zhang HY, Ma S, He YH, Bao MZ (2015). A study of heterosis, combining ability and heritability between two male sterile lines and ten inbred lines of Tagetes patula. Euphytica 203(2):349-366.
Badu A, Parnell D, Settlers A, Mihalke L, Sestras R (2012). Heterosis studies for response to Aphis fabae attack in Calendula. Bulletin of University of Agricultural Sciences and Veterinary Medicine Cluj-Napoca. Horticulture 69(1):40-47.
Chen JF, Lin YJ (1982). Chromosome pairing in interspecific hybrids of Tagetes patula and Tagetes erecta marigold. Cytologia 47(3-4):37-42.
Cicevan R, Al Hassan M, Sestras AF, Prahens J, Vicente O, Sestras RE, Boscaini M (2016). Screening for drought tolerance in cultivars of the ornamental genus Tagetes (Asteraceae). Peelf 4:e2133.
Deng YM, Teng NJ, Chen SM, Chen FD, Guan ZY, Song AP, Chang QS (2010). Reproductive barriers in the intergeneric hybridization between Chrysanthemum grandiflorum (Ramat) Kitam and Ajania przewalskii Poljak. (Asteraceae). Euphytica 174(1):41-50.
Deng YM, Ye XQ (2012). The prefertilization reproductive barriers and overcoming methods of horticultural crops distant hybridization. Acta Agriculturae Boreali-Sinica 27(5):81-86.
He YH, Ning GG, Sun YL, Qi YC, Bao MZ (2009). Identification of a SCAR marker linked to a recessive male sterile gene (TemA) and its application in breeding of marigold (Tagetes erecta). Plant Breeding 128(1):92-96.
He YH, Sun YL, Zheng RR, Ai Y, Cao Z, Bao MZ (2016). Induction of tetraploid male sterile Tagetes erecta by colchicine treatment and its application for interspecific hybridization. Horticultural Plant Journal 2(5):284-292.
Li FR, Zhang JC, Xu JR, Zhou JH (2005). Studies on the cross-breeding of Tagetes erecta L. × Tagetes patula L. and the sterility of hybrid. Inner Mongolia Agricultural University 26(2):51-54.
Li XL, Chen FD, Zhao HB (2008). Compatibility of interspecific cross in Dendranthema genus. Acta Horticultura Sinica 35(2):257-262.
Namita N, Singh KP, Bharadwaj C, Sharma TR, Sorah H, Raju DVS, Deshmukh RK (2011). Gene action and combining ability analysis for flower yield and its component traits in interspecific hybrids of marigold (Tagetes spp). Indian Journal of Agricultural Sciences 81(9):807-811.
Namita N, Singh KP, Bharadwaj CP, Prasad KV, Raju DVS (2009). Studies on character association and path analysis of quantitative traits among parental lines of marigold (Tagetes erecta and T. patula) and their interspecific F1 hybrids. Indian Journal of Horticulture 66(3):348-352.
Qi YC, Zhou GL, Gao Y (2008). Study on squash technique of root tip and analysis of chromosome karyotype in Tagetes erecta L. and their interspecific F1 hybrids. Indian Journal of Horticulture 66(3):348-352.
Sreekala C, Raghava SPS (2003). Exploitation of heterosis for carotenoid content in African marigold (Tagetes erecta L.) and its correlation with esterase polymorphism. Theoretical Applied Genetics 106(4):771-776.
Towner JW (1961). Cytogenetic studies on the origin of Tagetes patula. I. Meiosis and morphology of diploid and allotetraploid T. erecta × T. tenuifolia. American Journal of Botany 48(9):743-751.
Wu RH, Ge BB, Wang ML, Zhou Y, Feng H (2016). Estimation of genome size of fourteen Chinese old garden roses by flow cytometry. Journal of Beijing Forestry University 38(6):94-100.
Zeng L, Zhao LJ, Sun J, Zhao ZG, Yang F (2010). Analysis of genetic relatedness of genetic resources of Tagetes as revealed by ISSR. Scientia Agriculturae Sinica 43(1):215-222.
Zhang HJ, Dong AX, Wang T, Zhao LJ, Xin HB (2014). A New Cultivar of Marigold 'Jingmei 1'. Acta Horticultura Sinica 41(7):1521-1522.
Zhu C (1982). Plant chromosome and chromosome technology. Beijing Science Press.