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CHROMOSOME COUNTS IN THE SECTION SIMIOLUS OF THE GENUS MIMULUS (SCROPHULARIACEAE). IV.

BARID B. MUKHERJEE AND ROBERT K. VICKERY, JR.

This report¹ on the determination of chromosome numbers in the section *Simiolus* of the genus *Mimulus* is part of a long range investigation into the evolution of species in *Mimulus* (Vickery, 1951). The chromosome numbers and configurations presented in this article indicate a lack of cytological differentiation between several of the currently accepted species (Pennell, 1951) of the section *Simiolus*. Also they reveal the presence of aneuploidy in different populations of two other species, and, lastly, they fill an important gap in the previously indicated (Mukherjee and Vickery, 1959) polyploid series that extends from North to South America.

Essentially the same method of bud fixation was employed as in the previous investigation (Mukherjee and Vickery, 1959), i.e., fixation in two parts absolute ethanol to one part glacial acetic acid saturated with ferric acetate, followed by staining of the anthers in iron-aceto-carmine. Work now in progress indicates that there may be possible improvements in this schedule. Each chromosome number determination is based on counts from an average of approximately eight pollen mother cells. Camera lucida drawings were made for three or four figures for each count and, in addition, photomicrographs were taken of many of the configurations. Herbarium specimens of each culture have been or will be deposited in the Garrett Herbarium of the University of Utah (UT).

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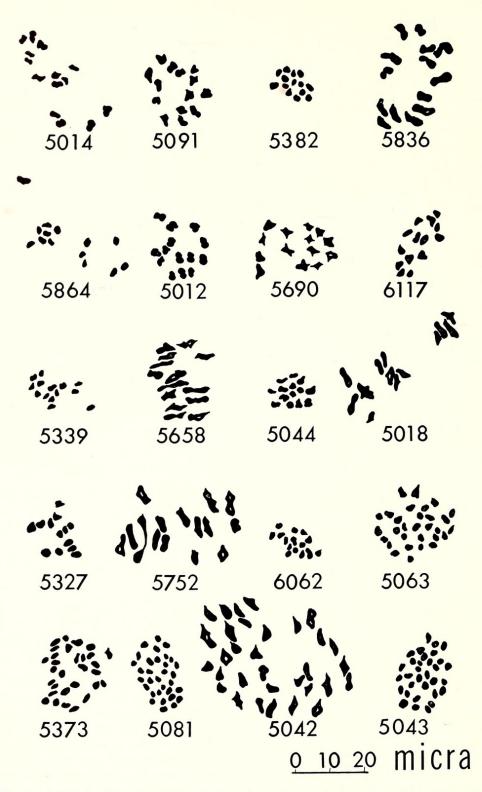


FIG. 1. Meiotic chromosomes of Mimulus: M. guttaus, 5014, 5091, 5382, 5836, 5864; M. tilingii var. tilingii, 5012, 5690, 6117; M. laciniatus, 5339; M. laxus, 5658; M. nasutus, 5044, 5018, 5327; M. platycalyx, 5752; M. glabratus var. utahensis, 6062; M. glabratus var. fremontii, 5063, 5373; M. tigrinus, 5081; M. luteus, 5042, 5043. All cells are in or near second metaphase except 5382, 5836, 5012, 5690, 5658, 5018, and 5752, which are in first metaphase. (Camera lucida drawings as reproduced, $= \times 840$).

A total of thirty-three cultures was studied during the present investigation (see table 1). They include representatives of ten species and varieties of the section Simiolus: Mimulus guttatus DC., M. tilingii Regel var. tilingii, M. laciniatus Gray, M. nasutus Greene, M. laxus Pennell, M. platycalyx Pennell, M. glabratus var. utahensis Pennell, M. glabratus var. fremontii (Benth.) Grant, M. tigrinus hort., and M. luteus L.

All fifteen cultures of *M*. guttatus were found to have n=14 chromosomes. The configurations were regular and similar to those previously observed for other cultures of *M. guttatus* (Vickery, 1955; Muhkerjee, Wiens, and Vickery, 1957; Mukherjee and Vickery, 1959) and, therefore, only a few of the camera lucida drawings of M. guttatus chromosomes were included in figure 1. The fifteen cultures examined represent much of the geographical range of M. guttatus (see table 1) and much of its morphological and physiological diversity as well. Morphologically the cultures differ from each other in the average height of the plants, shape of the leaves, amount and distribution of anthocyanin pigmentation, and in the size of the flowers. Physiologically they differ in growth rates, time and speed of flowering, and time and speed of maturing seeds. Cytologically the only detectable difference observed among the fifteen cultures was the presence of marked chromosome stickiness under the present fixation schedule in two of the annual races. Despite the wide range of morphological and physiological differences between the various cultures, they all exhibit apparently similar karyotypes.

The chromosome numbers of six cultures of M. tilingii var. tilingii from the Sierra Nevada (see table 1) were found to be n=14 as in M. guttatus. Mimulus tilingii var. tilingii is related to M. guttatus on the basis of morphology (Hitchcock, Cronquist, Ownbey and Thompson, 1959), but is separated from it by strong crossing barriers (Vickery, 1956). The chromosome configurations of the six cultures were regular and similar to those of M. guttatus and to our first M. tilingii var. tilingii count (Vickery, 1955), but differed in number from our more recent report of n=15 for a Utah population of M. tilingii var. tilingii (Mukherjee and Vickery, 1959). The populations studied represent much of the morphological diversity present in M. tilingii var. tilingii in the Sierra Nevada (table 1).

Mimulus laciniatus and M. laxus, species that are genetically closely related to M. guttatus (Vickery, 1956, and unpublished), were found to have n=14 chromosomes as does M. guttatus. The karyotypes of the three species are apparently indistinguishable. Mimulus laciniatus is morphologically strikingly different from M. guttatus, whereas M. laxus is closely similar. Probably M. laciniatus should be treated as a variety of M. guttatus, while M. laxus should be considered as synonymous with M. guttatus.

The chromosomes of three cultures from widely scattered populations (see table 1) of *M. nasutus* were counted. Two of the cultures have n=13 chromosomes and the third has n=14, which confirms the previous de-

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TABLE 1. CHROMOSOME COUNTS IN MIMULUS, SECTION SIMIOLUS

n=14 M. guttatus DC.

Barry Summit, Humboldt County, California, altitude 3,400 feet, Keck 6007 (5005).

Vosemite Junction (rocky creek), Tuolumne County, California, altitude 1300 feet, *Hiesey 560* (5006).

- Lee Vining Canyon, Mono County, California, altitude 8000 feet, Clausen 2039 (5014).
- Kern River, Kern County, California, altitude 1000 feet, L. Bean, April 16, 1949 (5085).

Botanic Garden strain, Hortus Cluj, Romania (5091).

Rio Santo Thomas, Baja California, Mexico, altitude ca. 20 feet, C. and L. Hubbs, spring 1950 (5382).

San Dimas Canyon, Los Angeles County, California, altitude 1500 feet, R.K. Vickery, Jr., September 29, 1950 (5678).

Hugh's Canyon, Salt Lake County, Utah, altitude 6000 feet, N. Chamberlain, spring, 1952 (5836).

Skagg's Springs, Sonoma County, California, altitude ca. 50 feet, R. Holm, spring, 1951 (5864).

Old Mine, Big Cottonwood Canyon, Salt Lake County, Utah, altitude 7650 feet, Vickery 683 (5961).

Neff Canyon, Salt Lake County, Utah, altitude 5500 feet, D. Wiens, September 6, 1956 (5995).

Moab, Grand County, Utah, altitude 4100 feet, Vickery 762 (6080).

Cane's Spring, San Juan County, Utah, altitude 5800 feet, Vickery 763 (6081).
Ledgemere, Big Cottonwood Canyon, Salt Lake County, Utah, altitude 5100 feet, Vickery 880 (6082).

East Creek, Morgan County, Utah, altitude 5700 feet, Vickery 883 (6083).

n=14 M. tilingii Regel var. tilingii

- Slate Creek (near Carnegie Transplant Garden), Mono County, California, altitude 10,000 feet, *Clausen 2075* (5012). (In flower at time 6120, 6121, 6122 were collected in bud.)
- Budd Lake, Tuolumne County, California, altitude 10,250 feet, C.W. Sharsmith, September 13, 1950 (5690).
- Tributary to Slate Creek (near Carnegie Transplant Garden), Mono County, California, altitude 10,050 feet, Vickery 1379 (6117).
- Slate Creek (near Carnegie Transplant Garden), Mono County, California, altitude 10,000 feet, *Vickery 1382* (6120). (Light green, large leaves.)

Same locality-Vickery 1383 (6121), dark green, medium sized leaves.

Same locality-Vickery 1384 (6122), dark green, small leaves.

n=14 M. laciniatus Gray

Lake Eleanor road, Tuolumne County, California, altitude 4200 feet, Vickery 179 (5339).

n=14 M. laxus Pennell

Yreka, Siskiyou County, California, altitude 3000-3200 feet, Pennell 26163 (5658).

n = 14 M. nasutus Greene

Hasting's Reservation, Monterey County, California, altitude 1500 feet, Stebbins 701 (5044).

n=13 M. nasutus Greene

San Augustine Pass, Dona Ana County, New Mexico, altitude 4500 feet, O. Norwell, October 30, 1946 (5018).

Wild Cat Creek, near Yosemite Junction, Tuolumne County, California, altitude 475 feet, Vickery 168 (5327). 1960]

n=15 M. platycalyx Pennell
Crystal Lakes Reservoir, San Mateo County, California, altitude 800 feet,
G. Oberlander, April, 1951 (5752).
n=15 M. glabratus var. utahensis Pennell
Johnson Pass, Tooele County, Utah, altitude 5800 feet, D. Wiens, October, 1956 (6062).
n=30 M. glabratus var. fremontii (Benth.) Grant
Whipple Mountains, San Bernardino County, California, collected April 21,
1940. U.C. 667,449 (5063).
Kakernot Springs, Alpine Creek, Brewster County, Texas, Cory 53186, May 18,
1946 (5373).
n=32 M. tigrinus hort.
Garden seed from the "Carlos Thays" Botanic Garden, Buenos Aires, Argentina
(5081).
n=32 M. luteus L.
Vicinity of Illapel, Coquimbo, Chile, altitude 6200 feet, Plant Introduction and
Exploration Division (U.S.D.A.) no. 144,535 (5042).
n=30+0, 1, or 2 M. luteus L.
Vicinity of Illapel, Coquimbo, Chile, altitude 2000 feet, Plant Introduction and
Exploration Division (U.S.D.A.) no. 144,536 (5043).

termination by G. L. Stebbins, Jr. (personal communictation) for the same culture. The n=13 cultures exhibit partial crossing barriers with M. guttatus, whereas the n=14 culture crosses readily with M. guttatus (Vickery, 1956, unpublished). The cytologic and crossing results suggest to us that M. nasutus as presently described (Grant, 1924, and Pennell, 1951) includes at least two different entities. The proper naming of these entities must await further investigation and a detailed study of the taxonomic literature and type specimens of M. nasutus and and its relatives.

Culture 5752, which was found to have n=15 chromosomes, was identified as *M. platycalyx* with some misgivings. Although the plants clearly exhibit Pennell's main key character of "fruiting calyces being fully as wide as long," yet the corolla throats are open and not filled by "a palate nearly closing orifice" (Pennell, 1951). Furthermore, Pennell had described *M. platycalyx* as occurring in the "southern Sierra Nevada from Mariposa to Tulare County, California," whereas the plants from which culture 5752 were grown came from the Crystal Lakes region of the outer Coast Ranges of California. However, even if this culture does not properly belong to *M. platycalyx*, it does represent an entity that is distinct from *M. guttatus* on the basis of morphology, crossing behavior (Vickery, 1956, in press), and cytology. Here again a sound taxonomic decision must await further critical study of this entity and the literature.

The culture of M. glabratus var. utahensis (6062) from the Stansbury Mountains near the Great Salt Lake had n=15 chromosomes as did culture 5265 from the population at Bicknell, Wayne County, Utah (Mukherjee, Wiens, and Vickery, 1957). In contrast, the population from the shore of Mono Lake at the western edge of the Great Basin has n=14 chromosomes (Vickery, 1955). Perhaps M. glabratus var. utahensis also

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consists of two morphologically similar but cytologically different entities as does M. nasutus, although corroborative crossing data is not yet available.

Two cultures of M. glabratus var. fremontii from southern California and Texas were found to have n=30 chromosomes, although some of the plants of culture 5063 were observed to have as few as n=26 chromosomes. This chromosome number fills an important gap in the polyploid series connecting the Great Basin form, M. glabratus var. utahensis, n=14 and n=15, with the South American M. glabratus var. parviflorus (Lindl.) Grant, n=45, and its ally M. pilosiusculus HBK., n=46 (Mukherjee and Vickery, 1959). Mimulus glabratus var. fremontii is approximately intermediate in appearance between the other two varieties. It is 5 to 15 centimeters in height, whereas M. glabratus var. utahensis varies from 10 to 50 centimeters, and M. glabratus var. parviflorus is nearly prostrate. The leaves and flowers of M. glabratus var. fremontii are smaller than those of M. glabratus var. utahensis but larger than those of M. glabratus var. parviflorus. A broader cytogenetic and taxonomic study of the Mimulus glabratus complex of species is now being undertaken.

The culture of *M. tigrinus* from the Botanic Garden of Buenos Aires, Argentina, was found to have n=32 chromosomes. This count agrees with the previous reports of Brozek (1932), Sugiura (Darlington and Wylie, 1955) and the authors (1959).

Chromosome counts obtained for two different cultures of M. luteus tend to support the previously indicated relationship of the horticultural species, M. tigrinus, to this wild species (Mukherjee and Vickery, 1959). Culture 5042 has n=32 chromosomes on the basis of two plants studied whereas culture 5043 was variable on the basis of the five or six plants examined. Of the twenty-three cells studied eleven had n=30 chromosomes, six had n=31 and six had n=32. The cause of the variability was not clear from the data obtained. Perhaps accessory chromosomes are involved. Tentatively, the chromosome number for this culture appears to be n=30+0, 1 or 2. The two cultures of M. luteus are morphologically similar, but are distinguishable on the basis of flower markings and the general growth habits of the plants.

In conclusion, this survey of chromosome numbers in section Simiolus has verified previously published counts for M. guttatus (n=14), M. tilingii var. tilingii (n=14), M. glabratus var. utahensis (n=15) and M. tigrinus (n=32). It has shown that two species that are genetically closely related to M. guttatus, M. laciniatus and M. laxus, have n=14 chromosomes also. In contrast, the culture, tentatively assigned to M. platycalyx, which is morphologically closely related to M. guttatus but genetically partially separated from it, has n=15 chromosomes. Mimulus nasutus appears to consist of two entities, one with n=14 chromosomes that is genetically closely related to M. guttatus, and the other with n=13chromosomes that is genetically partially isolated from M. guttatus and from the n=14 form of M. nasutus. Mimulus glabratus var. fremontii was found to have n=30 chromosomes, which neatly fills an important gap in the polyploid series of *M. glabratus* var. *utahensis*, n=15, to *M. glabratus* var. *parviflorus*, n=45. Lastly, one race of South American *M. luteus* has n=32 chromosomes, as had been previously reported for its horticultural derivatives, but the other race apparently has n=30+0, 1, or 2.

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FLOWERING RESPONSES IN PHACELIA SERICEA AND P. IDAHOENSIS¹

GEORGE W. GILLETT

In the study of variation in the *Phacelia sericea* complex [P. sericea (Graham) A. Gray subsp. sericea; P. idahoensis Henderson; and intermediates], experimental cultures of P. sericea subsp. sericea and of P. idahoensis could not be brought into flower under actual or simulated summer conditions. In these cultures, the daily photoperiod was extended by incandescent lights, when necessary, to between 16 and 20 hours. Later cultures were brought into flower, however, by simulating the fall conditions of the natural environment to the extent of materially reducing

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