# GYTOTAXONOMIC NOTES ON SOME GALIUM SPECIES 

E. KLIPHUIS<br>(Botanical Museum and Herbarium, Utrecht)

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It is well known that the chromosome number of a plant may be a character of importance to taxonomy; it can often give a better insight with respect to the place cf a taxon in the System. It is, therefore, particularly important to determine for any species the chromosome number of as many individuals from as many localities as possible. Publications of lists of documented chromosome numbers as presented by Löve in the IOPB reports in Taxon are very valuable and so are lists of chromosome numbers of plants from more restricted areas (Greenland: Jörgensen et al., 1958; Iceland: Löve and Löve, 1956; the Netherlands: Gadella and Kliphuis, 1963; Poland: Skalinska, 1950, Skalinska et al., 1957, 1959, 1961; Sweden, Skåne: L.övkvist, 1962).

These lists enable us to ascertain whether or not there are differences in chromosome numbers within a species and whether or not there is a relation with the geographical distribution or the ecological preference. Apart from the number, however, the size and shape of the chromosomes and, consequently, their 'portrait' may be of value. In the Angiospermae this character plays an important role almost exclusively in the Monocotyledones.

Differences in chromosome number within a species or species complex are sometimes, but certainly not always correlated with differences in morphology (Viola riviniana with $2 n=35,40,45,46$ and 47, Gadella, 1963; Listera ovata with $2 n=34,35,36$ and 38, Kliphuis, 1963). Even on the level of polyploidy (Galium verum with $2 n=22$ and 44, Lövkvist, 1962), we encounter this situation. The reverse situation also occurs: the species Galium mollugo L., collected in 91 different localities, always appeared to be a tetraploid ( $2 n=44$ ) but even though showing a great morphological variability which remains constant after several years of cultivation.

A careful analysis of the chromosome number of Galium aparine L., a species having a polyploid series with $2 n=22,44,66$ and, according to Fagerlind (1934), $2 n=88$, clearly showed that the level of polyploidy rather than the exact number of chromosomes is important.

The hexaploids generally have $2 n=64$, but within the same plant metaphase plates from the same roottip may have $2 n=61$, 62, 63, 65 and the normal hexaploid number $2 n=66$. However, we also found plants with 66 chromosomes in all countable metaphase plates of the same roottip.

The same phenomenon has been found in Cardamine pratensis L. (Berg, 1966), and in Symphytum officinale L. (Gadella and Kliphuis, in press).

Before arriving at a final conclusion it may be necessary to investigate other characters besides the chromosome number. The Dactylorchis maculata complex has been shown to have a diploid with $2 n=40$ as well as a tetraploid with $2 n=80$ chromosomes. Vermeulen (1947) separated the diploid from the tetraploid and described it as Dactylorchis fuchsii (Druce) Vermln. Not only cn account of those numbers but also because of morphological and ecological differences. Extensive biometric work by Heslop-Harrison (1948, 1951, 1954) on a considerable amount of material confirmed this conclusion.

Plants belonging to the Dactylorchis maculata complex occuring in the Netherlands were cultivated under identical conditions for several years. It was shown that differences between the diploid Dactylorchis fuchsii and the tetraploid Dactylorchis maculata became less clear, already after one year (Kliphuis, 1963).

From these examples it is clearly necessary to make cultivation experiments before expressing oneself definitely on such problems.

Galium hercynicum Weig. provides an example of a species in which morphological differences are correlated with the level of polyploidy. These differences remained constant during several years of cultivation. The diploid ( $2 n=22$ ) is distinguished from the tetraploid ( $2 n=44$ ) by a more fragile habit, more slender shoots, smaller flowers and leaves, and by the fact that it flowers about two weeks earlier than the tetraploid. Galium hercynicum is a common plant of Western Europe (southern Scandinavia, Denmark, Germany, Belgium, the Netherlands, France, northern Spain, Portugal and the British Isles).

The results of the measurements of the flowers are given in Table 1. In the first column: the investigated plants with the collection

Table 1.
(Measurements of the flowers)

| Flower 1963 in mm |  | $\boldsymbol{X}$ | $S D$ | $S E$ | $\mathcal{N}$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| No. plant: diploid$2 n=22$ |  |  |  |  |  |
| 106 | $\stackrel{L}{L}$ | 2.881 | 0.2047 | 0.0206 | 99 |
|  | W | 0.811 | 0.1082 | 0.0108 | 100 |
| 244 | $\stackrel{L}{L}$ | 3.038 | 0.2565 | 0.0257 | 100 |
|  | W | 0.931 | 0.0896 | 0.0090 | 100 |
| tetraploid$2 n=44$ |  |  |  |  |  |
| 386 | $\stackrel{L}{L}$ | 3.898 | 0.2224 | 0.0222 | 100 |
| 414 | $\stackrel{W}{L}$ | 0.953 3.992 | 0.1132 0.1212 | 0.0113 0.0121 | 100 100 |
|  | $\boldsymbol{W}$ | 1.021 | 0.0608 | 0.0061 | 100 |

CYtotaxonomic notes on some galium species
Table 2. (Measurements of the leaves)

| Leaf 1963 in mm |  |  | I |  |  |  | II |  |  |  | III |  |  |  | IV |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | $\boldsymbol{X}$ | $S D$ | SE | $\mathcal{N}$ | $\boldsymbol{X}$ | $S D$ | $S E$ | $N$ | $\boldsymbol{X}$ | SD | SE | $\mathcal{N}$ | $X$ | SD | SE | $\boldsymbol{N}$ |
| $\begin{aligned} & 106 \\ & \text { diploid } \\ & 2 n=22 \end{aligned}$ | 1962 | L | 3.332 | 0.655 | 0.12 | 30 | 3.552 | 0.747 | 0.144 | 27 | 1.350 | 0.698 | 0.174 | 16 | 3.640 | 0.408 | 0.091 | 20 |
|  |  | W | 0.582 | 0.265 | 0.045 | 33 | 1.028 | 0.131 | 0.025 | 27 | 0.690 | 0.397 | 0.099 | 16 | 1.288 | 0.222 | 0.049 | 20 |
|  | 1963 | $L$ | 4.432 | 0.800 | 0.080 | 100 | 4.200 | 0.498 | 0.050 | 100 | 2.656 | 0.635 | 0.090 | 50 | 4.112 | 0.824 | 0.117 | 50 |
|  |  | W | 0.681 | 0.175 | 0.017 | 100 | 0.975 | 0.154 | 0.015 | 100 | 0.933 | 0.069 | 0.021 | 50 | 1.312 | 0.219 | 0.031 | 50 |
|  | 1964 | L | 4.728 | 0.614 | 0.061 | 100 | 4.560 | 0.828 | 0.117 | 50 | 2.976 | 0.584 | 0.082 | 50 | 4.256 | 0.947 | 0.134 | 50 |
|  |  | W | 0.763 | 0.118 | 0.012 | 100 | 1.173 | 0.270 | 0.038 | 50 | 1.085 | 0.295 | 0.042 | 50 | 1.291 | 0.178 | 0.025 | 50 |
| 244 diploid$2 n=22$ | 1962 | $L$ | 4.713 | 1.016 | 0.126 | 64 | 4.880 | 1.286 | 0.181 | 50 | 2.984 | 0.723 | 0.218 | 11 | 4.512 | 1.272 | 0.127 | 100 |
|  |  | W | 0.764 | 0.302 | 0.039 | 60 | 1.139 | 0.258 | 0.036 | 50 |  |  |  | - | 1.564 | 0.204 | 0.022 | 90 |
|  | 1963 | L | 4.248 | 1.024 | 0.102 | 100 | 4.288 | 1.096 | 0.122 | 80 | 2.664 | 0.817 | 0.122 | 45 | 4.224 | 0.592 | 0.070 | 71 |
|  |  | W | 0.641 | 0.265 | 0.027 | 95 | 0.964 | 0.270 | 0.031 | 75 | 0.788 | 0.293 | 0.046 | 40 | 1.626 | 0.271 | 0.033 | 66 |
|  | 1964 | $L$ | 4.336 | 0.896 | 0.090 | 100 | 4.328 | 1.109 | 0.111 | 100 | 2.720 | 0.775 | 0.110 | 50 | 4.272 | 0.712 | 0.071 | 100 |
|  |  | W | 0.752 | 0.257 | 0.026 | 100 | 0.976 | 0.206 | 0.021 | 100 | 0.837 | 0.189 | 0.027 | 50 | 1.704 | 0.274 | 0.027 | 100 |
| 386 tetraploid $2 n=44$ | 1962 | $L$ | 5.912 | 1.144 | 0.114 | 100 | 6.880 | 1.688 | 0.169 | 100 | 4.526 | 1.016 | 0.110 | 70 | 5.312 | 1.066 | 0.107 | 100 |
|  |  | W | 0.841 | 0.324 | 0.032 | 100 | 1.448 | 0.191 | 0.019 | 100 | 1.520 | 0.372 | 0.044 | 70 | 2.326 | 0.284 | 0.028 | 100 |
|  | 1963 | L | 6.120 | 1.208 | 0.121 | 100 | 6.792 | 1.144 | 0.114 | 100 | 3.560 | 0.673 | 0.067 | 100 | 5.768 | 1.472 | 0.147 | 100 |
|  |  | W | 0.830 | 0.317 | 0.032 | 100 | 1.483 | 0.197 | 0.020 | 100 | 1.517 | 0.272 | 0.027 | 100 | 2.205 | 0.257 | 0.026 | 100 |
|  | 1964 | L | 6.064 | 1.392 | 0.139 | 100 | 7.864 | 1.568 | 0.157 | 100 | 4.680 | 0.801 | 0.127 | 40 | 5.328 | 1.546 | 0.218 | 50 |
|  |  | W | 0.839 | 0.084 | 0.008 | 100 | 1.636 | 0.215 | 0.022 | 100 | 1.582 | 0.233 | 0.039 | 40 | 2.059 | 0.256 | 0.026 | 100 |
| 414 tetraploid$2 n=44$ | 1962 | L | 5.128 | 0.917 | 0.092 | 100 | 7.072 | 1.046 | 0.105 | 100 | 3.400 | 0.869 | 0.087 | 100 | 4.480 | 1.212 | 0.169 | 50 |
|  |  | W | 0.937 | 0.179 | 0.018 | 100 | 1.754 | 0.202 | 0.020 | 100 | 1.478 | 0.226 | 0.032 | 50 | 1.822 | 0.299 | 0.042 | 50 |
|  | 1963 | $L$ | 5.680 | 1.488 | 0.016 | 90 | 6.720 | 1.480 | 0.171 | 75 | 3.264 | 0.422 | 0.060 | 50 | 5.424 | 1.216 | 0.122 | 100 |
|  |  | W | 0.993 | 0.297 | 0.031 | 90 | 1.734 | 0.493 | 0.057 | 75 | 1.654 | 0.349 | 0.049 | 50 | 2.435 | 0.322 | 0.032 | 100 |
|  | 1964 | L | 5.736 | 1.167 | 0.118 | 100 | 7.128 | 1.168 | 0.117 | 100 | 3.656 | 0.730 | 0.073 | 100 | 5.200 | 0.960 | 0.096 | 100 |
|  |  | W | 1.069 | 0.362 | 0.036 | 100 | 2.271 | 0.346 | 0.035 | 100 | 1.420 | 0.289 | 0.029 | 100 | 2.300 | 0.246 | 0.025 | 100 |

numbers, (the diploids: 106 and 244; the tetraploids: 386 and 414; localities: 106: Portugal, Botanical Garden Sacavem, 244: Portugal, Tras os Montes, Lisa do Alvoa; 386: the Netherlands, Zuid-Limburg, 414: the Netherlands, Duurswoude, prov. of Friesland). In the following columns is $L=$ length, $W=$ width both in $\mathrm{mm}, X=$ the mean of the length or width of the sample, $S D$ the standard deviation, $S E$ the standard error of the mean and $\mathcal{N}=$ the total number of the sample.

It is clear that there is a difference in length of the flowers between the diploid and tetraploid. Most striking, however, is the difference in size of the leaves. Three types of leaves can be distinguished on the flowering shoots: Type I, leaves on the upper third part of the shoot; Type II, those of the middle third part of the shoot, and Type III, the smallest leaves on the lowest part, mostly wilted. A fourth type is found on the non-flowering shoots.

The measurements of the leaves are given in Table 2. From these tables the differences between the diploid and tetraploid are evident. The same is true for the differences between the leave types mentioned before. It is also clear from the values given that these differences remain constant over the years.

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