Backcross data from plants heterozygous for $\mathrm{L} 9_{2}$ and $\mathrm{A}_{1}$. In every combination listed in column one, the A allele is in the left-most chromosome. The first 8 entries are from Rhoades and Dempsey (1953).

Chromosome structure Linkage phase Heterozygous parents Lg A lg a
Lg a lg A Total Recomb. \% S.E.

| N/N C |  |  | 522 | 568 | 203 | 219 | 1512 | 27.8 | 1.16 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| In/N C | m | 3015 | 2142 | 24 | 84 | 5605 | 1.9 | . 18 |  |
| $\mathrm{In} / \mathrm{N}$ C | h | 1410 | 1215 | 6 | 8 | 2639 | 0.5 | . 14 |  |
| In/In C | m | 514 | 420 | 401 | 356 | 1691 | 44.8 | 1.21 |  |
| In/In R | m | 393 | 340 | 610 | 592 | 1935 | 37.9 | 1.11 |  |
| In/In R | h | 333 | 311 | 606 | 566 | 181 | 6 | 35.5 | 1.12 |
| In Df - Dp/In | R | ${ }_{\text {h }}$ | 823 | 68 | 1335 | 229 | 2455 | 36.3 | . 97 |
| In Df-Dp/In | R | m | 192 | 18 | 255 | 34 | 499 | 42.1 | 2.21 |
| N Df-Dp/N | R | 564 | 1045 | 2052 | 1031 | 4692 | 34.3 | . 69 |  |
| N Df-Dp/N | R | ${ }^{\text {h }}$ | 32 | 399 | 1773 | 86 | 2790 | 33.4 | . 89 |
| $\mathrm{N} / \mathrm{N}^{*} \mathrm{R}$ | m | 738 | 767 | 1522 | 1485 | 4512 | 33.4 | . 70 |  |
| In/In C | m | 1087 | 988 | 880 | 922 | 3877 | 46.5 | . 80 |  |
| In/In C | h | 350 | 385 | 341 | 334 | 1410 | 47.9 | 1.33 |  |
| In/In R | m | 307 | 365 | 509 | 475 | 1656 | 40.5 | 1.20 |  |
| In/In R | h | 302 | 311 | 422 | 462 | 1497 | 41.0 | 1.27 |  |

*The lgA chromosome was derived from a double exchange from an In/N plant.
The average unweighted percentage of recombination between $\operatorname{Lg}_{2}$ and $\mathrm{A}_{1}$ in $\mathrm{N} / \mathrm{N}$ plants is 30.6 while in the homozygous inversion plants where A is nearer the centromere and Lg is more distal the percentages of recombination varied from 35.5 to 47.9 with an unweighted mean of 42.0 percent. The difference in recombination percentages in $N / N$ and In/In plants can be accounted for by a centromere effect on crossing over. Since both Lg and A are in the inverted segment it can be argued, on the basis that closer proximity to the centromere results in a decrease in exchange frequency and conversely an increase in crossing ovor when further removed, that the physical distance of the proximal break of the inversion from Lg is less than the distance of the second break from A.

The Df-Dp chromosomes derived from In $3 a / N$ plants have varying portions of the proximal part of the long arm of chromosome 3 in duplicate. One of these Df-Dp chromosomes was tested to determine whether or not the $\mathrm{gl}_{6}$ locus, which is proximal to $\lg _{2}$, was included in the duplicated segment of the Df-Dp chromosome. The following crosses were made:


Sum $=1841$
Gl6-A recombination $=49.1 \%$
\% Gl6 in seedlings $=51.1$
\% A in seedlings $=31.2$
Since earlier studies showed that approximately $26 \%$ of Df-Dp ovules function, is [sic] clear from the $\mathrm{Gl}: \mathrm{gl}$ ratio that $\mathrm{Gl}_{6}$ is not included in the duplicated piece of 3 L for if it were there would be approximately 50 percent more Gl than gl seedlings and a $1: 1$ ratio was obtained. This argument is based on the following table. It is clear from the 49.1 percent recombination between $\mathrm{Gl}_{6}$ and A that at least one crossover occurred between $\mathrm{Gl}_{6}$ and the Df in all, or nearly all, megasporocytes so we can eliminate the products of no exchange bivalents and consider only single and double exchanges.


Type of exchange Gl not in duplication $\qquad$
Ngl N Gl Dp-Df gl Dp-Df Gl

| singles | 1 | 1 | 1 | 1 |
| :--- | :--- | :--- | :--- | :--- |
| 2 strand doubles | 2 | 0 | 0 | 2 |
| 3 strand doubles | 2 | 2 | 2 | 2 |
| 4 strand doubles | 0 | 2 | 2 | 0 |
|  | 5 | 5 | 5 | 5 |

If $\mathrm{Gl}_{6}$ is not in duplication a $1: 1$ ratio is expected. This was found, [sic]

If $\mathrm{Gl}_{6}$ were in duplication, a ratio of approximately $1.5: 1.0$ should occur. The observed ratio of Gl:gl deviates significantly from 1.5:1.0 but is very close to a 1:1.
2. Ears from crosses of $\mathrm{Bt}_{1} / \mathrm{bt}_{1}$ plants with $\mathrm{sh}_{3}$ showed a $1: 1$ segregation for shrunken kernel type. Although the phenotypes of $\mathrm{sh}_{3}$ and $\mathrm{bt}_{1}$ are quite dissimilar, they are allelic.
3. A new $\mathrm{gl}_{1}$ was found on chromosome 5 . Its linkage relations with $\mathrm{A}_{2}$ and $B t_{1}$ are shown below. Only colored seeds $\left(A_{2}\right)$ were used since another aleurone factor was segregating.


| $B t$ | $B t$ | $b t$ | $b t$ |
| :---: | :---: | :---: | :---: |
|  | 15 | 69 | 0 |$\quad \Sigma=1365$

Region (1) $\mathrm{gl}-\mathrm{A}_{2}=15 \div 1365=1.1 \%$
Region (2) $\mathrm{A}_{2}-\mathrm{Bt}=694 \div 1365=5.1 \%$

The order is: $\quad$| gl A | Bt |
| :---: | :---: |
| 1 | 5 |

On the basis of negative allelism tests with unplaced glossies, the new gl was designated $\mathrm{gl}_{17}$.
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