AN ANALYSIS OF X-RAY INDUCED CHROMOSOMAL ABERRATIONS IN TRADESCANTIA

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The cause and nature of chromosomal alterations has become an important genetic problem since it has been shown that both mutation and speciation often are associated with structural alterations of the chromosomes (DOBZHANSKY 1937). The frequency of spontaneous structural changes is too low in most organisms to permit a statistical analysis of types or frequencies of alterations, although recently Giles (1940) has been able to analyze a considerable number of natural chromosomal aberrations in Tradescantia hybrids. In most respects these natural aberrations resemble those induced by X-rays, and although the spontaneous breaks are not caused by radiation, the same secondary factors seem to be involved in producing the various types and frequencies of aberrations. The X-ray induced aberrations can be produced in large numbers under controlled conditions, permitting adequate statistical analyses of types of aberrations and their frequency in relation to dosage, temperature, time of exposure, and radiation intensity.

The microspores of diploid species of Tradescantia provide excellent material for an analysis of X-ray induced chromosomal aberrations. The six chromosomes are large and the nuclear cycle can be timed rather accurately. During the summer months the microspore nucleus remains in the resting stage for about five days after the microspore is formed following meiosis. At about 30 hours before metaphase the chromosomes begin splitting, and at 23 to 24 hours before metaphase all chromosomes are split to form the sister chromatids. Thus the nuclear cycle from microspore formation to metaphase requires about six days. During the winter months the nuclear cycle is nearly twice as long and the prophase stage may begin at about 60-70 hours before metaphase. For an analysis of chromatid breaks induced at prophase the cells were fixed 24 hours after raying. Microspores fixed five days after raying were irradiated in the resting stage. Preparations were made at 48 hours after raying during the winter months, and 30 hours after raying during the summer months, for the analysis of very early prophase stages. The analyses of dosage, the time factor, and temperature effects were based on experiments done during the winter months. The division figures obtained provide a true random sample of aberrations because there is no chance of elimination of chromosomes or chromosome fragments.

The microspore cells were smeared, fixed in alcohol-acetic for a few
minutes, and stained with aceto-carmine. Several hundred division figures usually were obtained from each flower bud, and preparations with less than about a hundred metaphase or anaphase figures were discarded. Since there are six chromosomes in each microspore each preparation permitted the analysis of from 600 to 2000 chromosomes. The results to be presented are based on an analysis of more than 300,000 chromosomes.

The statistical treatment of the data presents a difficult problem. Most of the individual values given in the various tables are based on the average frequencies on three slides, and the total number of chromosomes is from 3,000 to 6,000. Neither the number of slides nor the number of chromosomes provides the correct value of \( n \) for a critical analysis of variation. Accordingly we have taken the deviation from the mean, in percent of the mean, for each series of observations involving three or more slides each, and have obtained the standard deviation and probable error from these figures. The value of \( n \) was taken as the number of series times \( n - 1 \) slides examined for each series. The generalized probable error is found to be about 10 percent of the mean. Many of the experiments were repeated several times with comparable results.

**TYPES OF CHROMOSOMAL ABERRATIONS**

Two general types of chromosomal aberrations are induced by X-rays, namely, chromatid breaks, and chromosome breaks. The chromatid breaks are induced at prophase when each chromosome consists of two sister chromatids, while the chromosome breaks are those induced during the resting stage when the chromosomes are in the form of single threads. Each of these general types of aberrations may be further classified into two groups: the aberrations caused by a single break which are referred to as one-hit breaks, and aberrations involving breaks in two different chromosomes or different loci of the same chromosomes which are referred to as two-hit aberrations. The origin and nature of these four classes of chromosomal aberrations is shown diagramatically in figure 1.

The one-hit chromatid breaks may involve one or both chromatids. The terminal deletion of one of the two chromatids (Plate I, figs. E and F) results in a chromatid fragment which is usually not included in either daughter nucleus at telophase. Often, however, the fragment appears to be separated from the centric arm only by an achromatic lesion, suggesting a partial breakage or imperfect reunion. Although these one-hit terminal deletions are rather frequent they cannot be scored accurately and have not been included in the analysis of X-ray breaks. The most frequent type of one-hit chromatid aberration is a break in both chromatids at the same locus followed by lateral fusion to produce a dicentric chromatid and an acentric U-shaped fragment (Plate I, B). At anaphase the dicentric chro-
matid forms a single bridge and the fragment tends to straighten out (Plate I, C). Such fragments are rarely included in the daughter nuclei. The ends of the fragment are the normal ends of sister chromatids (cf. figure 1).

The two-hit chromatid aberrations are of various types. The most frequent types are fusions between broken chromatids of two different chromosomes. These consist of reciprocal chromatid exchanges, producing a pseudo-crossover type of configuration (Plate I, D), or the broken ends may fuse to form a chromatid bridge between two chromosomes accompanied by an acentric fragment (Plate I, E). Other types have been found but they are rare. Occasionally the corresponding arms of a single chromosome are of unequal length at anaphase (Plate I, F) indicating an intercalary duplication such as Kauffmann and Bate (1938) have described in Drosophila. Fusions between breaks in each of the two arms of a single chromosome have produced a few chromatid rings, and in rare cases an acentric ring is produced from two breaks in a single arm.

<table>
<thead>
<tr>
<th>Chromosome aberrations</th>
<th>Chromatid aberrations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Resting</td>
<td>Pre-metaphase</td>
</tr>
<tr>
<td></td>
<td>Prophase</td>
</tr>
<tr>
<td>One Hit</td>
<td></td>
</tr>
<tr>
<td>Two Hit</td>
<td></td>
</tr>
</tbody>
</table>

**Figure 1.**—The types of chromosomal aberrations induced by X-rays at prophase and at the resting stage. Only the types which can be recognized at metaphase or anaphase have been included, since they are the types used in the various analyses.
The chromosome aberrations include both one-hit and two-hit types. The one-hit aberrations include both intercalary and terminal deficiencies. The loss of a terminal segment of a chromosome results in a shortened arm with no fusion of the ends of sister chromatids, and the acentric fragment consists of two separate chromatids at metaphase (Plate I, G). The small intercalary deletions presumably are deficiencies produced by breaks in adjacent relic spirals during the resting stage (figure 1). These deleted fragments are small (Plate I, G and N) and have not been included in the analysis of chromosome breaks. Most of these deficiencies have been found to be two-hit aberrations (Rick, unpublished observations).

The two-hit chromosome aberrations include reciprocal interchange or fusion between different chromosomes and fusion or exchange between the arms of the same chromosome. Occasionally a relatively large intercalary deletion is produced resulting in an acentric ring fragment. Reciprocal interchange between arms of different chromosomes can be detected only when the exchange is unequal (Plate I, J), and such exchanges are infrequent. Fusions between broken ends of two chromosomes result in a dicentric chromosome and an acentric fragment (Plate I, H and I).

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**DESCRIPTION OF PLATE I**

Photographs of Tradescantia microspore chromosomes showing the types of aberrations induced by X-rays. Compare with diagrams in figure 1.

**FIGURE A.**—Untreated,—six normal chromosomes at metaphase.

**FIGURE B.**—A one-hit chromatid aberration induced at prophase. Lateral fusions have occurred between the ends of the broken sister chromatids to produce a shortened arm with terminal fusion of sister chromatids and a U-shaped acentric fragment. Metaphase.

**FIGURE C.**—The same at anaphase showing the chromatid bridge and partially straightened U-shaped fragment.

**FIGURE D.**—A two-hit chromatid aberration. Reciprocal interchange between chromatids of different chromosomes.

**FIGURE E.**—The same, but with chromatids fused to form a chromatid bridge between chromosomes. Also a one-hit chromatid aberration—terminal deletion of only one of the two sister chromatids.

**FIGURE F.**—An intercalary chromatid duplication and deficiency involving the sister chromatids of one arm. Results in unequal arms at anaphase. Also a single chromatid terminal deletion.

**FIGURE G.**—Two one-hit chromosome aberrations. One is a terminal deletion of most of one arm, and the other a very small interstitial deletion.

**FIGURE H.**—A two-hit chromosome aberration. A dicentric chromosome and the accompanying fragment. Note relational coiling between centromeres.

**FIGURE I.**—Two two-hit chromosome aberrations. A dicentric and a locked ring with their accompanying fragments.

**FIGURE J.**—Unequal interchange between two chromosomes.

**FIGURES K-N.**—Behavior of dicentric chromosomes at anaphase. Also a terminal chromosome deletion and an interstitial deletion in figure 14.

**FIGURE O.**—A continuous ring chromosome at anaphase. Also somatic non-disjunction of two anaphase chromosomes.
broken ends of the terminal deletions fuse to produce a fragment consisting of parts of two chromosomes. The ends of the fragment are the normal ends of the two arms. The relational coiling between centromeres of the dicentric chromosome persists to metaphase and at anaphase the dicentric chromatids may separate freely, form an X-shaped bridge, or interlock (Plate I, K-N). Fusion between breaks in the two arms of the same chromosome results in a ring chromosome and an acentric fragment (Plate I, I). The ends of the fragment are the normal ends of the two arms. These ring chromosomes may separate freely at anaphase, they may be locked, or they may form a single large ring (Plate I, O). Occasionally a ring chromosome is locked around its rod fragment.

Combinations of chromosome and chromatid breaks may occur in the same cell or even in the same chromosome, but they are very rare. The individual chromosomes must split very rapidly at prophase (Sax and Mather 1939).

Other types of chromatid and chromosome aberrations undoubtedly occur, but inversions, transpositions or intra-chromosomal translocations, and simple intercalary translocations, either cannot be detected, or cannot be distinguished from reciprocal translocations. The types of aberrations included in the subsequent analyses consist of the one-hit chromatid breaks involving both chromatids, two-hit chromatid breaks which produce exchange or dicentric chromatids, and the two-hit chromosome breaks resulting in dicentric and ring chromosomes. These constitute most of the visible chromosomal alterations induced by X-ray doses used in the following experiments.

FREQUENCIES OF TYPES OF ABERRATIONS IN RELATION TO SPATIAL ARRANGEMENT OF THE CHROMOSOMES

The relative frequencies of one-hit and two-hit aberrations vary with the X-ray dosage and will be considered later. The various types of two-hit aberrations, both chromosome and chromatid, appear in the same relative frequencies at different dosages, even though the actual frequencies increase exponentially with dosage. The relative frequencies of the two-hit aberrations provide evidence regarding the conditions and limitations of fusion between broken ends of chromatids and chromosomes. The frequencies of types of chromatid and chromosome alterations are shown in table I. The dicentric chromatids are more than twice as frequent as the exchange chromatids. The exchange types cannot ordinarily be recognized at anaphase but most of these observations were made at metaphase so that the 2:1 ratio is approximately correct. The ratio of dicentric and ring chromatids is about 13 to 1, but for the chromosome aberrations the ratio of dicentrics and rings is 3.3 to 1.
The loci of breaks

Usually the exchange breaks are at similar loci in respect to the centromeres of the two chromosomes. An analysis of a random sample of dicentric and exchange chromatid aberrations showed 33 with breaks in both chromatids at about the same relative loci and only six which involved breaks at clearly different loci in the two chromosomes. A similar condition must obtain for chromosome breaks because unequal translocations are seldom observed.

The distribution of X-ray induced breaks must be at random among the chromosomes and at random for the loci of any one chromosome. But the

<table>
<thead>
<tr>
<th>TOTAL CHROMOSOMES</th>
<th>ONE HIT</th>
<th>TWO-HIT BREAKS</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>DELETION</td>
<td>EXCHANGE</td>
</tr>
<tr>
<td>9906</td>
<td>468</td>
<td>182</td>
</tr>
</tbody>
</table>

Frequencies of types of chromosome aberrations; 24 hours, 40–200 r.

<table>
<thead>
<tr>
<th>TWO-HIT ABERRATIONS</th>
<th>DICENTRICS</th>
<th>RINGS</th>
<th>RATIO</th>
</tr>
</thead>
<tbody>
<tr>
<td>10,344</td>
<td>7,936</td>
<td>2,408</td>
<td>3.3:1</td>
</tr>
</tbody>
</table>

aberrations resulting from these breaks are not at random for the various loci of the chromosome arms. The frequency of both chromosome and chromatid breaks which result in aberrations is greatest at the proximal end of the chromosome arm. The distribution of such chromatid and chromosome breaks is shown in graphic form in figure 2. The data were derived from an analysis of 340 chromosome aberrations and 334 chromatid aberrations (Sax and Mather 1939). The loci of breaks were divided into five groups; group one including those in the proximal fifth of the chromosome arm and the subsequent groups including the more distal loci. The difference in distribution of chromatid and chromosome breaks is statistically significant according to the $\chi^2$ test, but technical difficulties in determining breakage points in dicentric chromosomes may account for this difference.

The relation between X-ray dosage and frequency of chromosomal alterations

The relation between the X-ray dosage and mutation rate in Drosophila has been shown to be linear by a number of investigators (Timoféeff-
Recent work on both Drosophila and Tradescantia shows that the relation between X-ray dose and chromosomal aberrations is not linear. Oliver (1932), using genetic methods, was the first to show an exponential increase in chromosomal alterations with increased dosage in Drosophila. Direct cytological analysis of the salivary gland chromosomes of Drosophila also shows an exponential increase in chromosome changes with dosage (Bauer, Demerec and Kaufmann 1938). Muller (1938) also refers to experiments which indicate that the frequency of both translocations and inversions varies as the 3/2 power of the X-ray dose. The frequency of chromosome aberrations in Tradescantia plotted against dosage also shows an exponential curve (Sax 1938).

![Diagram](image)

**Figure 2.** Loci of chromosome and chromatid breaks involved in X-ray induced aberrations. Locus 1 includes the first fifth of the chromosome arm next to the centromere and locus 5 the distal fifth. If such breaks were at random 20 percent of all breaks should be found at each of the 5 loci. Data from Sax and Mather (1939). ○ = chromatid. ● = chromosome.

In all these experiments the frequency of chromosome aberrations increased approximately as the 3/2 power of the dosage. If each aberration is dependent upon two breaks, as is probably true in all these experiments, the frequency of aberrations should increase as the square of the dosage. In all these previous experiments the dosage was increased by increasing the time of exposure. Recently it has been found that the frequency of two-hit chromosomal aberrations is dependent upon the time-intensity factor, and that a high intensity of dosage is more effective in producing aberrations than the same dose given slowly (Sax 1939). Accordingly we should vary the dosage by varying the intensity rather than by varying the time of exposure.

In the following experiments on the relation between X-ray dosage and frequency of chromosomal aberrations, the dosage was varied by changing...
the distance between the target and the flower buds according to the inverse square law, so that variation in dosage was given without changing the exposure time. The minimum distance used was 24 cm, but even at this distance there is some variation in dosage, due to slight differences in the positions of the young buds in the inflorescences. At greater distances this experimental error is decreased. The source of the X-rays was a Coolidge tube with a tungsten target, operated at 10 ma, 134 K.V. with no screen.

The results of increased dosage on frequency of chromosomal aberrations induced at the resting stage are shown in table 2. The frequency of these two-hit aberrations increases as the square of the dosage. The equation for the curve is percent breaks, B, equals \((D/107)^{2.0}\), where D is the dosage in r units and 107 is a constant. The values of the exponent and constant were derived by plotting the logarithm of the dosage against the logarithm of the percent of effective breaks and obtaining the values from the graphs.

The low frequency of one-hit chromosome breaks induced at the resting stage does not permit an adequate comparison of the effects of varying dosage on one-hit and two-hit chromosome aberrations, so for this comparison we have used one-hit and two-hit chromatid aberrations induced at prophase. The frequency of chromatid breaks is much higher than that of chromosome breaks and it was not possible to analyze accurately the chromatid breaks induced at a dosage of much over 200 r. On the other hand it was possible to obtain an appreciable number of chromatid aberrations at 10 r. The effect of varying dosage on both one-hit and two-hit chromatid aberrations is shown in table 3. The frequency of the two-hit chromatid aberrations increases approximately as the square of the dosage. The equation for the curve is percent \(B = (D/67)^{1.9}\). The one-hit aberrations, however, show an approximately linear relation to dosage, the equation being percent \(B = (D/45)^{1.1}\). The deviations from the expected value

Table 2

<table>
<thead>
<tr>
<th>DOSE IN R UNITS</th>
<th>TOTAL CHROMOSOMES</th>
<th>CHROMOSOME ABERRATIONS*</th>
<th>%B</th>
<th>(D/107)^2</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>DICENTRIC</td>
<td>RING</td>
<td>TOTAL</td>
</tr>
<tr>
<td>100</td>
<td>1800</td>
<td>5</td>
<td>1</td>
<td>6</td>
</tr>
<tr>
<td>200</td>
<td>1710</td>
<td>21</td>
<td>8</td>
<td>29</td>
</tr>
<tr>
<td>300</td>
<td>1484</td>
<td>48</td>
<td>13</td>
<td>61</td>
</tr>
<tr>
<td>400</td>
<td>1902</td>
<td>104</td>
<td>28</td>
<td>132</td>
</tr>
<tr>
<td>500</td>
<td>1530</td>
<td>134</td>
<td>25</td>
<td>159</td>
</tr>
</tbody>
</table>

* Each aberration involves two breaks.
of the exponents may be due to experimental error since the data in this
table were derived from three different series of experiments extending
over a period of several months. It is possible that environmental condi-
tions or the output of the X-ray tube may have varied sufficiently so that

Table 3

The relation between X-ray dosage and one-hit and two-hit chromatid breaks; time constant.

<table>
<thead>
<tr>
<th>DOSE IN R UNITS</th>
<th>TOTAL CHROMOSOMES</th>
<th>CHROMATID ABERRATION BREAKS</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>1-HIT</td>
</tr>
<tr>
<td>10</td>
<td>6330</td>
<td>14</td>
</tr>
<tr>
<td>20</td>
<td>6930</td>
<td>24</td>
</tr>
<tr>
<td>40</td>
<td>8610</td>
<td>76</td>
</tr>
<tr>
<td>80</td>
<td>7368</td>
<td>148</td>
</tr>
<tr>
<td>120</td>
<td>4902</td>
<td>141</td>
</tr>
<tr>
<td>160</td>
<td>8292</td>
<td>375</td>
</tr>
<tr>
<td>200</td>
<td>8508</td>
<td>472</td>
</tr>
</tbody>
</table>

the data from different experiments were not strictly comparable. It is
also possible that some of the one-hit types of aberrations were produced
by two adjacent independent hits. In general, however, it is apparent that
"one-hit" aberrations show a linear response to dosage, while the "two-hit"
aberrations vary as the square of the dosage, when the exposure time is
held constant.

The effect of increased dosage on the frequencies of the various types of
aberrations is shown in the form of curves in figure 3. The curves for the
chromatid aberrations have been extrapolated beyond 200 r for comparison
with the two-hit chromosome aberration curve. The frequency of the two-
hit chromatid breaks is considerably greater than that of the two-hit
chromosome breaks. The two frequencies are not entirely comparable,
however, because the reciprocal interchanges can be detected when in-
duced at prophase, but only rarely when induced at the resting stage.

The relative frequencies of one-hit and two-hit chromosomal aberrations
vary not only with dosage, but also with the time of prophase development
at the time of irradiation. At very early prophase the exchange or fusions
between chromatids of different chromosomes are much less frequent in
relation to the number of terminal deletions than at later prophase stages.
The relative frequencies of these one-hit and two-hit chromatid aberrations
and the two-hit chromosome aberrations induced at early prophase, 48
hours before metaphase, are shown in table 4. The theoretical frequency
of two-hit chromatid breaks is based on the relative frequencies of one-hit
and two-hit aberrations produced at mid-prophase, while the theoretical
frequency of the chromosomal aberrations is derived from the equation
FIGURE 3.—The relation between the frequencies of chromosome and chromatid aberrations and the X-ray dosage in r units. Time of exposure constant. The curves for the one-hit and two-hit chromatid breaks have been extrapolated beyond 200 r. The equations for the curves are:

1. one-hit chromatid breaks, percent $B = (D/45)^{1.1}$,
2. two-hit chromatid breaks, percent $B = (D/67)^{1.9}$,
3. two-hit chromosome breaks, percent $B = (D/107)^{2.0}$ and
4. total chromatid breaks, percent $B = (D/37)^{1.4}$.

percent $B = (D/107)^2$, assuming that all chromosomes are single. The data show that most of the chromosomes were not yet split at this time. The aberrations induced in the split chromosomes show a much lower frequency of two-hit aberrations than would be expected at mid-prophase.

**TABLE 4**
Frequencies of one- and two-hit chromatid breaks induced at very early prophase; 48 hours.

<table>
<thead>
<tr>
<th>DOSE</th>
<th>TOTAL CHROMOSOMES</th>
<th>CHROMATID</th>
<th>CHROMOSOME</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>ONE-HIT %</td>
<td>TWO-HIT %</td>
<td>THEOR.</td>
</tr>
<tr>
<td>160 r</td>
<td>3276</td>
<td>26</td>
<td>0.8</td>
</tr>
<tr>
<td>320 r</td>
<td>3030</td>
<td>46</td>
<td>1.5</td>
</tr>
</tbody>
</table>
An analysis of the time-intensity factor in the X-ray production of various types of chromosomal aberrations has shown that the two hits necessary to produce an alteration involving two chromosomes need not occur simultaneously, but may occur over a period of time as long as an hour (Sax 1939). The time-intensity factor was varied by giving equal doses at different intensities, or by giving the same intensity of radiation over various periods of time by intermittent exposures. The intensity of radiation was varied by placing the inflorescences at various distances from the target according to the inverse square law. The results of the time factor at different X-ray intensities are shown in graphic form (figure 4). The dosage was 320 r and the intensity varied from 320 r/m to 20 r/m according to the time of exposure. The curve is based on analysis of 10,620 chromosomes. The chromosomes which were given 320 r in one minute show twice as many aberrations as those which were given 320 r in sixteen minutes. Similar experiments using other total dosages showed similar results.

Since the chromosome aberrations used in this analysis include only two-hit alterations, the greater frequency of such aberrations at high X-ray intensity must mean that reunion of broken ends occurs within certain limits of time. If the broken ends fuse quickly a low intensity should produce fewer aberrations than a high intensity of radiation be-
cause a break in one chromosome would heal before a second break occurred in an adjacent chromosome, and no aberration would be produced. But if the broken ends remain in an unstable condition for a long time and

![Figure 5](image.png)

**Figure 5.**—The relation between frequency of chromosomal aberrations and intermittent radiation. Four two-minute X-ray doses were given at 50 r/m, with rest periods of 0, 40, 80 and 120 minutes respectively. The base line (B), established by determining the percentage of breaks for a single two-minute exposure and multiplying by 4, coincides with percent of breaks at about the 70 minute rest period.

few fusions occur during the time of raying there should be little effect of the time-intensity factor. The results show that many fusions occur in a few minutes and probably in even less time, but they do not indicate the maximum length of time necessary to complete the fusion of all broken ends.

The time during which a broken end of a chromosome remains in an unstable condition can be determined from intermittent timing experiments. The results of one of these experiments are shown in the form of a curve in figure 5. Five groups of flower buds were irradiated, all at 50 r/m. The first group was given continuous radiation for eight minutes, the second, third, and fourth groups were given four intermittent exposures of two minutes each, with rest periods of 40, 80, and 120 minutes respectively. The fifth group was given only one two-minute exposure. If broken ends of chromosomes can remain unstable and capable of fusion with other broken ends for only a short time, a sufficiently long rest period should prevent fusions between breaks induced at successive exposures. This point
is reached where a further increase in the rest period produces no further decline in frequency of aberrations. The base line can also be determined by multiplying the frequency of aberrations induced by a single exposure by the number of exposures used in the intermittent timing experiment. In the above experiment the base line is at about six percent and is reached at a rest period of somewhat more than an hour. These results show that a break induced at one exposure period may remain open so that a broken end can fuse with another broken end produced by the second exposure an hour later, although most of the fusions occur in a considerably shorter period. Similar experiments with different X-ray intensities show similar results, although at high intensities, where the dose is relatively high for each exposure, there is less effect of the intermittent exposures. These results are based on an analysis of more than 80,000 chromosomes.

The effect of the time-intensity factor is based on the fact that the two-hit aberrations are dependent upon two independent breaks, limited in time of occurrence. According to this interpretation the time-intensity factor should have no effect on one-hit aberrations. This assumption has been tested by comparing the effect of intermittent timing on one-hit and two-hit chromatid aberrations. As expected, the frequency of the one-hit aberrations is independent of the time factor while the frequency of the two-hit aberrations declines with increasing rest periods (Sax 1939).

THE EFFECT OF TEMPERATURE ON X-RAY INDUCED CHROMOSOMAL ABERRATIONS

In an earlier experiment no differences in frequency of chromosomal aberrations were found when the cells were irradiated at different temperatures (Sax 1938). The results were not based on an analysis of individual chromosomes and the frequency of aberrations included all types of alterations obtained at various times after irradiation. More critical experiments show clearly that the temperature during and following irradiation has a marked effect of the frequency of both one-hit and two-hit aberrations (Sax and Enzmann 1939). The results of two of these experiments are shown in graphic form in figure 6.

The inflorescences were immersed in water at the desired temperature for five to ten minutes before raying and kept just under the surface of the water during irradiation. In a few experiments the flower stalks were removed from the water and placed at room temperature soon after raying, but usually they were left in the water for an hour after raying. When pasteboard cartons were used for containers the temperature dropped gradually during this period, but similar temperature effects were obtained when the flowers were put in thermos bottles where the temperatures were maintained at a constant level during the entire period.
The cells which were held at a low temperature during and following irradiation had from two to five times as many chromosome aberrations as those which were subjected to a high temperature. Preliminary work on the frequency of aberrations at various temperatures shows that the frequency does not decrease regularly with increased temperature, but tends to drop sharply at a critical temperature, which varies from less than 20°C to above 30°C, depending on the time of year the work is done. Presumably this seasonal effect is related to the length of the nuclear cycle; in the winter months the nuclear cycle from meiosis to division of

![Figure 6](image)

**Figure 6.**—The effect of varying temperatures during and immediately after X-radiation on the frequency of chromosomal aberrations. The different temperatures at which the greatest decrease in frequency of aberrations occur in the two curves is attributed to seasonal effects (Data from SAX and ENZMANN 1939).

the microspore nucleus requires about 12 days and the critical temperature is about 30°C, while in late spring the nuclear cycle requires about one week and at this season the critical temperature is less than 20°C.

At both mid-prophase and at the resting stage the frequency of X-ray induced aberrations is decreased at high temperatures, but at earliest prophase when the chromosomes are beginning to split into chromatids the higher temperatures cause many more aberrations than are produced by raying at low temperatures. The effect of high (30°C) and low (3°C) temperatures on frequency of X-ray induced aberrations at different prophase stages is shown in table 5. The low frequency of chromatid aberrations induced at early prophase, 48 hours before metaphase, is due to the fact that many of the chromosomes have not split and these produce chromosome aberrations which are not included in this table. The frequency of X-ray induced aberrations, at 24 hours before metaphase, is much higher
in the cold series, but at 48 hours the high temperature produces many more aberrations than the low temperature. This relation holds for both one-hit and two-hit aberrations, although the latter are infrequent in the 48 hour preparations.

INTERPRETATION OF RESULTS

The time of chromosome splitting

The evidence regarding the time of chromosome splitting in the nuclear cycle has been reviewed by Mather (1937). The results obtained from X-ray experiments are considered to be more critical than the interpretations based on direct visual evidence. Certainly the types of aberrations induced by X-rays at the various stages in the nuclear cycle of Tradescantia microspore indicate that the chromosome reacts as a single thread during the resting stage and becomes effectively split into sister chromatids at prophase (cf. Riley 1936). It may be argued that the effect of a single X-ray quantum is not sufficiently localized to be used as a tool for differentiating the number of chromatids or half-chromatids in a chromosome, since a single “hit” can break two adjacent chromatids at late prophase when the sister chromatids are more than a tenth of a micron apart. If the chromosomes were double at the early resting stage a single hit would usually break both of the closely associated chromatids, but such breaks should be followed by lateral fusion of broken ends of adjacent sister chromatids. Aberrations of this type are very seldom found in microspores irradiated in the resting stage, and the rare exceptional cases can be attributed to spontaneous aberrations. The effective split which produces differentiated chromatids, corresponding to those involved in crossing over at meiosis, certainly occurs at the prophase stage in the Tradescantia microspores.

There is some experimental evidence that the effective split may occur

### Table 5

The effect of temperature on frequency of chromatid aberrations at very early prophase (48 hours) and at midprophase (24 hours). Total chromosomes 19,848.

<table>
<thead>
<tr>
<th>Dosage</th>
<th>% One-Hit</th>
<th>% Two-Hit</th>
<th>Total</th>
<th>Dosage</th>
</tr>
</thead>
<tbody>
<tr>
<td>24 hours</td>
<td>Cold</td>
<td>3.6</td>
<td>2.6</td>
<td>6.2</td>
</tr>
<tr>
<td>Hot</td>
<td>1.8</td>
<td>1.6</td>
<td>3.4</td>
<td>160 r</td>
</tr>
<tr>
<td>48 hours</td>
<td>Cold</td>
<td>0.8</td>
<td>0.1</td>
<td>0.9</td>
</tr>
<tr>
<td>Hot</td>
<td>1.1</td>
<td>0.4</td>
<td>1.5</td>
<td>320 r</td>
</tr>
<tr>
<td>48 hours</td>
<td>Cold</td>
<td>1.5</td>
<td>0.2</td>
<td>1.7</td>
</tr>
<tr>
<td>Hot</td>
<td>4.6</td>
<td>1.0</td>
<td>5.6</td>
<td>320 r</td>
</tr>
</tbody>
</table>
at the resting stage preceding nuclear division (Mather 1937), or even at the preceding anaphase. The behavior of the broken ends of inversion bridge chromosomes suggests that the meiotic anaphase chromosomes may be effectively split. In both Tradescantia (Sax 1937) and Zea (McClintock 1938b) the break in an inversion bridge is followed by lateral fusion of sister chromatids to produce a bridge at the following division of the chromosome in the microspore. If the chromosome were single at meiosis a single broken end would have no other broken end with which to fuse, and perhaps such a broken end would remain in an unstable condition until the chromosome is split at the prophase stage of the microspore nucleus, at which time the lateral fusion would occur between the broken ends of sister chromatids. But McClintock has found that adjacent double inversion bridges behave in the same way, even though two broken ends are present in the newly formed microspore nucleus. On the other hand a break in a somatic ring chromosome of Zea is followed by fusion of broken ends of the two arms and not between ends of adjacent sister chromatids (McClintock 1938a). This difference cannot be attributed to differences in behavior at meiosis and mitosis because X-ray induced aberrations in Tradescantia root tips produce double bridges, which, when broken, are followed by lateral fusion of ends of sister chromatids. There is no reason to believe that the X-ray induced and spontaneous breaks differ in their effects on subsequent fusions of broken ends of chromosomes.

The behavior of the induced ring chromosomes is of interest in relation to the nature of splitting. If these ring chromosomes induced at the resting stage are single threads they must split in different planes because the rings may be continuous or interlocked at the following metaphase. This problem has been analyzed in some detail by Husted (1936).

The time of effective chromosome splitting in different microspores is quite variable, ranging from about 23 to 31 hours before metaphase in Tradescantia plants grown in the field during the summer months. The splitting of the chromosomes of a single nucleus is much more limited in time, and the splitting of an individual chromosome must be very rapid since both chromosome and chromatid breaks are rarely found in the same chromosome. There is some indication that effective splitting begins at the proximal end of the chromosome arm and progresses distally (Sax and Mather 1939).

The frequency of types of aberrations in relation to chromosome arrangement

It has been shown that all potential fusions between broken ends of chromosomes are completed within an hour after the breaks are produced. There can be no extensive movement of the chromosomes during this
time, at most periods in the nuclear cycle, so that fusions between broken ends must be limited to those in close proximity. Certainly at the resting stage there could be no extensive movement of the chromosomes, which in the form of relic coils appear to completely fill the nucleus (Belar 1928). With an increase in nuclear size and uncoiling of relic spirals at prophase more movement of the chromosomes could occur, but not sufficient to permit random association of broken ends of chromatids. Accordingly, an analysis of the loci of breaks involved in fusions between different chromosomes or chromosome arms should indicate the arrangement and relations of the chromosomes at various phases in the nuclear cycle.

The chromosomes of Tradescantia are strongly polarized during the resting and early prophase stages so that if little chromosome movement occurs between breakage and fusion, the fusions between different chromosome arms should occur usually at corresponding loci in respect to the distance from the centromeres. Such a relation is found for both chromosome and chromatid aberrations. Unequal exchanges between chromosome arms are rare, and an analysis of the fusions and exchanges between chromatids of different chromosomes shows that the breaks have occurred at approximately the same relative loci of both chromosomes in over 80 percent of the figures examined.

The relative frequencies of dicentric and ring chromosomes also show that proximity facilitates, or conditions, the fusions of broken ends of chromosomes. A broken end of a chromosome arm can fuse with a broken end in the other arm of the same chromosome to form a ring, or it can fuse with a break in any one of the ten arms of the other five chromosomes to form a dicentric union. Random reunion of broken ends should produce a ratio of dicentric to ring chromosomes of 10:1, but the observed ratio is 3.3 to 1 (table 1). It is evident that the distance between breaks, both laterally and longitudinally in respect to the polarized arms of the chromosomes, conditions or determines the occurrence of two-hit aberrations.

The spatial relationships of chromosome arms appear to change as the nuclear cycle progresses from the resting stage into prophase. At the resting stage the ratio of dicentric to rings is 3.3:1, but irradiation at prophase results in a ratio of dicentric to rings of 13:1. This decrease in frequency of ring chromosomes at mid-prophase is attributed to the greater separation of the two arms of a single chromosome at this stage. It is known that chromosomes repel each other, especially when they become split into sister chromatids. The prophase chromosomes also become shorter and the increased size of the nucleus permits greater movement of the chromosomes.

The relative frequencies of one-hit and two-hit aberrations induced at prophase vary with dosage as is shown in figure 3. There is, however, a
striking difference in these frequencies between early and mid-prophase (table 4). The relative frequency of two-hit aberrations induced at early prophase is very much less than at mid-prophase. This difference can hardly be attributed to greater repulsion between arms of chromosomes, and must be related to differences in the nature of the induced breaks. The two-hit breaks—dicentrics, exchanges and rings—induced at mid-prophase may depend upon breaks in only one chromatid of each of the chromosome arms involved in the aberrations. The sister chromatids at this stage are distinctly separated and breaks in only one of the two sister chromatids at a given locus are found at metaphase. At earliest prophase, however, the two sister chromatids must be closely associated so that a single X-ray hit would usually break both chromatids at the same locus. Such breaks would be followed by the fusion of adjacent ends of the broken chromatids in most cases, to produce a terminal deletion which is classed as a one-hit aberration. This interpretation is supported by the fact that two-hit chromosome aberrations, induced before chromosome splitting, but at about the same time, show a normal frequency.

The limitations of fusions imposed by the factor of proximity must mean that most of the breaks induced by X-rays do not lead to the production of aberrations. The two broken ends simply reunite in the original position and no structural alteration of the chromosome is visible at later stages in nuclear development. The chromosome breaks which lead to fusions between chromosome arms must constitute only a very small proportion of the total breaks induced, but simple breaks at the following metaphase are far less frequent than aberrations which involve two breaks in close proximity. The one-hit breaks at prophase are frequent only because two chromatids are broken and lateral fusion of broken ends leads to a permanent aberration. The frequency of breaks which heal in the original position cannot be determined, but such breaks must be far more frequent than those which lead to gross structural changes in the chromosomes.

A comparison of the frequencies of types of aberrations has also been used in the analysis of chromosome arrangements in Drosophila sperm (Catcheside 1938; Bauer, Demerec and Kaufmann 1938). The salivary gland chromosomes permit an analysis of alterations within chromosome arms, as well as between chromosome arms, although the numerous cell divisions between irradiation and chromosome analysis must produce differential elimination of the various types of aberrations. Catcheside finds that exchanges within the same arm are twice as favored as those between different chromosome arms. Exchanges between arms of the same chromosome when compared with the frequency of exchanges between arms of different chromosomes, show no significant deviation from a random dis-
INDUCED ABERRATIONS IN TRADESCANTIA

Induced aberrations in Tradescantia have been obtained by Bauer, Demerec, and Kaufmann. These results must mean that a single chromosome in the Drosophila sperm is folded so that various parts of the same chromosome arm are in close proximity. Presumably, little movement is possible in the closely packed chromosomes of the sperm and induced aberrations must depend upon close proximity of the induced breaks.

The loci of breaks in relation to aberrations

It has been shown that the distribution of breaks involved in chromosome and chromatid aberrations is not at random for the various loci of chromosome arms. The initial breaks must be at random, but those which are involved in chromosome alterations are most frequent at the proximal ends of the chromosome arms. This distribution must be caused by secondary factors, and since the unequal distribution holds for one-hit chromatid breaks the secondary factor must be confined to the individual arms of the chromosomes. The greater frequency of such breaks at the proximal end of the chromosome arm is attributed to stresses imposed by the various coiling mechanisms which would tend to throw the broken ends out of alignment so that reunion in the original position would be inhibited. The breaks induced in the chromosomes of the sperm of Drosophila show a nearly random distribution of breaks with a possible higher frequency at the distal ends (Bauer, Demerec and Kaufmann 1938). Perhaps the more random distribution of effective breaks in Drosophila can be attributed to the close association of the chromosomes in the Drosophila sperm so that proximity of broken ends does not necessitate any appreciable movement of broken ends in order to produce fusions between different chromosomes.

The relation between X-ray dosage and frequency of chromosomal aberrations

In both Drosophila and Tradescantia the frequency of X-ray induced breaks increases approximately as the 1.5 power of the dosage when the various doses are given at the same intensity. But since the time factor must be considered in such experiments the dosage should be varied by changing the intensity and not the time of exposure. When this is done the frequency of two-hit chromosome and chromatid aberrations in Tradescantia varies approximately as the square of the dosage. This relation should be expected since the chance that two or more independent events will happen together is the product of their respective chances of happening.

The frequency of simple deletions is assumed to be induced by single hits and should show a linear relation to dosage, at least for doses which
produce a low frequency of breaks. The data indicate a slight exponential increase of aberrations with increased dosage which may be due to experimental error, or to the occasional production of the deletions by two independent hits in the two chromatids at approximately the same locus.

The number of X-ray hits or quanta necessary to produce an effect has been calculated by comparing the survival curves with those obtained from Poisson's exponential series (Timoféeff-Ressovsky 1937). For one-hit and two-hit chromosomal aberrations the equation can be expressed as: percent B = 1 − e^−x and percent B = 1 − e^−x(1+x), respectively, where x represents the number of effective hits. Using an arbitrary value of x of .00025 per r unit of dosage, we find that the calculated values for the one-hit aberrations are practically identical with the observed values. The calculated values of the two-hit curve also are in close agreement with the observed percentages of breaks at the various doses, but in order to get these theoretical values, the value of x must be increased to .0025 per r unit. The dosage curves, (D/45)\^{1.1} and (D/67)\^{1.9}, do not indicate such different dosages for one- and two-hit aberration frequencies, as is indicated by the Poisson exponential series. It is not clear how these apparent discrepancies can be reconciled. It is evident, however, that the slopes of the two dosage curves are practically identical with the theoretical curves for one-hit and two-hit effects.

In all of the various experiments on the relation between dosage and the mutation rate in Drosophila a linear relation has been found. The frequency of mutation is also independent of the time-intensity factor (Timoféeff-Ressovsky 1937). These facts indicate that most of the induced mutations are produced by single hits and that relatively few of the induced mutations are due to "position effects" resulting from two-hit aberrations such as reciprocal translocations and large inversions.

The frequency of X-ray induced aberrations induced in the Drosophila sperm must also depend on the time factor. With dosage controlled by increasing the time of exposure the relation between frequency of aberrations and dosage is expressed by the equation percent B = (D/K)\^{1.5} in both Drosophila and Tradescantia; but in Tradescantia, with the time of exposure constant, the frequency of aberrations varies as the square of the dosage. Control of the time factor should also increase the exponent from 1.5 to 2.0 for the dosage-aberration curve in Drosophila.

The differential susceptibility of chromosomes to X-rays during the nuclear cycle and in different types of cells

The frequency of gross chromosomal aberrations is much greater at prophase than during the resting stage, when the same X-ray dose is used (figure 3). The equation for the relation between dosage and frequency of
Induced aberrations in Tradescantia is: percent $B = (D/37)^{1.5}$ for all the chromatid aberrations recorded, and percent $B = (D/107)^2$ for the chromosome aberrations. This comparison may not be entirely valid because certain types of aberrations can be detected when induced at prophase, but not when induced at the resting stage. In general, however, the frequency of gross alterations is greatest when the cells are rayed at prophase. This relation is to be expected for the following reasons: there are twice as many threads to be hit at prophase; terminal deletions can be produced at prophase by lateral fusion between broken chromatids while in the resting stage they are produced only when the broken end loses its capacity for fusion; and the secondary factors of chromosome torsion and movement would be expected to be greater at prophase.

A comparison of the frequency of chromosomal aberrations in Tradescantia and Drosophila shows that breaks involving chromosome changes are much more frequent in Tradescantia microspores than in Drosophila sperm. Some of this difference may be attributed to inviable aberrations in Drosophila, since the chromosomes were analyzed many cell generations after raying the sperm. The data of Bauer, Demerec and Kaufmann show that the frequency of breaks, in percent of total chromosomes, may be expressed by the equation, percent breaks $= (D/1150)^{1.5}$. For Tradescantia microspores rayed at the resting stage, with dosage varied by varying the time of exposure, the equation is: percent $B = (D/80)^{1.5}$ (Sax 1938). These data would seem to indicate that the Tradescantia chromosomes are much more sensitive than those of Drosophila. This difference probably can be attributed to differences in chromosome size in the two genera and to differences in chromosome organization in the two types of cells. According to Metz and Boche (1939) both chromosome aberrations and mutations in Sciara are less frequent than in Drosophila when given the same X-ray dosage.

The difference in the frequencies of X-ray induced chromosomal aberrations in different cells of the same organism should provide additional evidence regarding X-ray effects on the chromosome. It is known that both mutations and gross chromosomal alterations occur more frequently in the sperm of Drosophila than in the oocytes for a given X-ray dose (Oliver 1934). In Sciara gross chromosomal alterations are readily secured by raying the sperm, but were not found when the egg cells were irradiated. Metz and Boche (1939) attribute this difference in Sciara to physical differences in the chromosomes of the two types of cells. The close proximity of the chromosomes in the sperm should facilitate illegitimate union of broken ends, while most of the chromosomes of the eggs at the time of treatment were in metaphase or anaphase of the first meiotic division. The diffuse stage at meiotic prophase of meiosis also was found to
be resistant to X-ray effects. Metz (1934) has suggested that the chromosome matrix plays an important part in this differential susceptibility of chromosomes to radiation.

The differential susceptibility to X-rays of different stages of the nuclear cycle, of different cells in the same organism, and of similar cells in different organisms may depend in part on the same secondary factors such as proximity of chromosomes, freedom of chromosome movement, and the physical condition of the chromosomes. There is, however, some evidence that differences in the internal structure of the chromosomes may be responsible for differential X-ray susceptibility. Navashin and his collaborators have found that the aging of seeds increases both the mutation rate and frequency of spontaneous chromosomal aberrations (cf. Giles 1940). Offermann (1938) has shown that X-rays produce more mutations in old than in young sperm of Drosophila. These results suggest that aging of chromosomes induces internal changes, probably of a chemical nature, which makes them more susceptible to both spontaneous and induced alterations.

In any analysis of differential susceptibility of different stages in the nuclear cycle it is essential to differentiate the physiological effect of X-rays from the production of mutations and gross chromosomal aberrations which lead to genetic alterations. The physiological effect of suppressed nuclear activity and clumping of the chromosomes has long been known and this reaction in plant cells has been dealt with in considerable detail by Marquardt (1938). At moderate X-ray doses the inhibition of nuclear development is temporary and the cell recovers. The time of recovery depends on the dosage (Carlson 1938, and unpublished). Heavy X-ray dosage may cause so much clumping of the metaphase chromosomes that normal division is prevented and the effect is lethal. Maximum X-ray effects at metaphase probably can be attributed to physiological factors.

It is probable that the physiological and genetic effects can be differentiated by determining the temperature coefficient of the reaction to X-rays. If chromosome aberrations are responsible for cell injury the temperature coefficient should be one or less than one, while if physiological effects are responsible the temperature coefficient should be appreciably greater than one.

**The behavior of broken ends of chromosomes**

The experimental evidence shows that the induced chromosomal aberrations involving two chromosomes are dependent upon two independent breaks limited in both time and space. The broken ends of chromosomes either reunite in the original position or produce chromosome aberrations within a short time after the breaks are induced by X-rays, except a very small proportion of breaks producing broken ends which appear to have
lost their capacity for fusion. Most of the unions occur within a few minutes, and at the end of about an hour all fusions are completed. This does not necessarily mean, however, that a broken end can remain in an unstable condition for only an hour.

Stadler (1932) has assumed that an acentric fragment may be included in a daughter nucleus by chance and fuse with the original broken chromosome after several cell generations. Such behavior is highly improbable for the following reasons: all potential fusions between broken ends are completed in about an hour in the Tradescantia microspore which has a nuclear cycle of about a week; nearly all the acentric fragments found at metaphase have no broken ends since the two ends of such fragments are the normal ends of chromosomes or chromatids; the few terminal deletions which persist until metaphase appear to have lost their capacity for fusion since chromosome fragments show no lateral fusion of sister chromatids; and very few of the various types of fragments are included in the mitotic daughter nuclei in plant cells. Acentric fragments of meiotic chromosomes often are included in the daughter nuclei in Zea (McClintock 1938b), and in the neuroblasts of the grasshopper such fragments usually are included in the daughter nuclei (Carlson 1938). Even in these cases reunion of acentric fragments is improbable. Moreover, according to McClintock, the acentric fragments occasionally included in the Zea microspores act as normal chromosomes both in their cytological behavior and in their genetic influence. Thus there could be no genetic deficiency by deletion of a chromosome segment, or genetic recovery when the fragment joins the centric segment, as Stadler has postulated.

The temperature effect on induced chromosomal aberrations and mutation

According to Muller (1935), Papalashvili has found that the frequency of chromosomal rearrangements in Drosophila is increased by raying the flies at low temperatures. Mickey (1937) has found a much higher frequency of translocations in flies rayed at 15°C than in those rayed at 28°C–34°C. In Tradescantia microspores both chromosome and chromatid aberrations, induced at either the resting or at mid-prophase, are much more frequent in the cells rayed at low temperatures. The variations in temperature affect both one-hit and two-hit aberrations. The decreased frequency of X-ray induced aberrations at the higher temperatures can be attributed to a more rapid fusion of broken ends or to a more relaxed condition of the chromosomes which inhibits chromosome movements. Either factor would favor the reunion of broken ends in the original position and decrease the frequency of illegitimate unions which produce the
aberrations. It has been shown that the frequency of the initial breaks is relatively independent of the temperature.

The reversed temperature effect on the chromatids at earliest prophase is difficult to explain. The reaction of the chromosomes cannot be due to changes in the nucleus as a whole because the single chromosomes show a decreased frequency of aberrations, and the split chromosomes an increased frequency of aberrations at high temperatures. An increase in the speed of union of broken ends may be involved. At earliest prophase the newly formed chromatids may be so closely associated that a break would separate the broken ends of a single chromatid further than the distance between chromatids. At mid-prophase the chromatids may be separated by a distance greater than the distance between broken ends of a chromatid. Rapid union of broken ends would tend to produce lateral fusions and consequently aberrations at early prophase, but would favor reunion in the original position and consequently fewer aberrations at later prophase stages. This interpretation could hardly be applied to the two-hit chromatid aberrations, which also seem to show the reversed temperature effect at early prophase, although the numbers are very low. An increase in torsional strain induced by high temperatures only at earliest prophase could also be invoked to explain these results, but further work is necessary in order to determine the factors involved in the reversed temperature effect at earliest prophase.

The nature of X-ray induced breaks

The chromosomal aberrations induced by X-rays have been attributed to fusions followed by breaks and to independent breaks followed by fusions between broken ends. The contact hypothesis sponsored by Müller (1932) postulates that the chromosomes in contact or closely associated may be affected by a single X-ray hit. The breakage hypothesis, suggested by Stadler (1931), postulates that independent breaks are followed by illegitimate fusions. The analysis of X-ray induced chromosomal aberrations in Tradescantia confirms Stadler's hypothesis. The dosage curves for one-hit and two-hit aberrations, the effect of the time-intensity factor, and the temperature effect, all prove that aberrations involving two chromosomes are dependent upon two independent hits.

The one-hit chromatid breaks appear to be caused by single X-ray "hits" since the relation between dosage and frequency of such aberrations is approximately linear. Since two sister chromatids may be broken by a single hit the ionization or excitation must cover a relatively large range because these sister chromatids are separated by a distance of at least a tenth of a micron at mid-prophase. The frequency of induced mutations in Drosophila is independent of X-ray wave-length (Timofeeff-Ressov-
The frequencies of X-ray induced chromosomal aberrations in Tradescantia also are independent of wave-length (Rick, unpublished). These results indicate that the X-ray effects are caused by ionization (Goodspeed and Uber 1939).

The broken ends of chromosomes may remain in an unstable condition and capable of fusion for an hour or longer. It is surprising that a molecular complex of such a nature can remain in an unstable condition for so long a period. When fusions between broken ends do occur the union appears to be quite perfect in most cases, and breaks in subsequent chromosome bridges are no more likely to occur at the point of union than at other loci. There must be natural breaking points in the chromosome because breaks and fusions are normal features of practically all chromosomes at the time of meiosis. The reunion of both X-ray induced and crossover breaks appears to have no effect on the physical condition of the chromosome.

About five percent of the visible chromosome alterations induced by irradiation of the resting nucleus consist of terminal deletions. The broken ends of such deletions appear to behave as normal ends. Such a behavior may be caused by breaks at loci other than the natural breaking points.

ACKNOWLEDGMENTS

This work was supported, in part, by a grant from the National Research Council Committee on Radiation, and by research funds of the Biological Laboratories of Harvard University. Most of the preparations were made by Margery Poole, who also scored the types and frequencies of chromosomal aberrations in many of the X-ray experiments.

SUMMARY

An analysis of X-ray induced chromosomal aberrations in the microspores of Tradescantia has led to the following conclusions:

1. The chromosomes react to X-rays as though they were single threads during the resting stage and become effectively split to form sister chromatids at early prophase.

2. The effective splitting of an individual chromosome occurs very rapidly, and there is some evidence that splitting begins at the region of the centromere and progresses distally.

3. The simple terminal deletions are readily induced at prophase because the two sister chromatids may be broken by one hit, followed by lateral fusion of broken ends of sister chromatids.

4. The frequency of these simple deletions shows an approximately linear relation to dosage. The equation for this relation is percent B = (D/K)^1.1, where B is the number of effective breaks in relation to the total number of chromosomes, D the dosage in r units, and K a constant.
5. A simple deletion induced at the resting stage persists until metaphase only when the broken ends lose their capacity for fusion. Such breaks are rare. The broken ends of these chromosomes appear to behave as normal ends.

6. Aberrations involving two chromosomes, or two loci in the same chromosome, are dependent upon two independent breaks, limited in both time and space.

7. The frequency of these two-hit aberrations is dependent upon the radiation intensity. High intensity is more effective than low intensity because the aberrations are dependent upon two adjacent breaks within certain limits of time. The frequency of one-hit aberrations at a given dosage is independent of the time-intensity factor.

8. The frequency of two-hit aberrations increases exponentially with increased dosage. When the dosage is varied by varying the time of exposure the relation between frequency of aberrations and dosage is percent \( B = (D/K)^{1.5} \), but if dosage is varied by varying the intensity the equation is percent \( B = (D/K)^{2.0} \).

9. Intermittent dosage experiments show that broken ends of chromosomes may remain in an unstable condition for as long as an hour before fusion, although most fusions occur in a much shorter time.

10. Only a small proportion of the X-ray induced breaks result in visible chromosome aberrations. Most of the broken ends reunite in the original positions with no evident alteration of the chromosome.

11. The frequency of the initial breaks induced by X-rays is relatively independent of temperature. The production of aberrations, however, is greater when the resting and prophase nuclei are irradiated at low temperatures. High temperatures during or immediately following irradiation increase the reunion of broken ends in the original position thus decreasing the frequency of aberrations. This temperature effect is reversed at very early prophase.

12. The initial breaks induced by X-rays must be distributed at random, but both the one-hit and two-hit aberrations occur more frequently at the proximal ends of the chromosome arms. This distribution of effective breaks is attributed to the effect of secondary factors.

13. Differential X-ray production of chromosomal aberrations in different phases of the nuclear cycle, in different cells of the same organism, and in similar cells of different organisms, can be attributed, in part, to secondary factors, such as proximity of chromosomes, freedom of chromosome movement, and the physical condition of the individual chromosome.

14. The X-ray induced breaks appear to occur usually at natural breaking points. Most of the breaks and fusions following irradiation appear to be comparable to those occurring naturally at the time of crossing over.
ADDENDUM

I have seen, in manuscript, a paper on X-rayed Tradescantia chromosomes submitted to "Genetics" in July 1939 by A. C. Fabergé of University College, London. Fabergé’s conclusions on the time factor and the temperature effect are essentially the same as those reported above. His data were not adapted to an analysis of the dosage curve, and his conclusions in this regard appeared to require reconsideration.*

LITERATURE CITED


Oliver, C. P., 1932 An analysis of the effect of varying the duration of X-ray treatment upon the frequency of mutations. Z.i. A.V. 61: 447–488.


* Dr. Fabergé has in the meantime withdrawn his paper from “Genetics” and has submitted it to the “Journal of Genetics.”


1938 Chromosome aberrations induced by X-rays. Genetics 23: 494-516.


