Table 1 Fractional losses of A Sh following electric-field treatments of ++/a sh males.

Male	1/2	Frac	tiona: 1/8	l cate	gory limit	Total	No. <u>A</u> <u>Sh</u> kernels
		1	4	3	8	16	991
1 (Control) 3 (40 min)		3	3	8	5	19	686
11 (3.5 hr)	5	7	3	15	23	53	1359
Total	5	11	10	26	36	88	3036
TOTAL							

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1. Preferential pairing in trisomic plants containing an irradiated chromosome.

Pollen from plants with normal chromosomes 3 containing the A₁ allele was given 1000 \underline{r} and used to fertilize standard trisome 3 plants which were homozygous for \underline{a}_1 . The gene segregation from the resulting trisome 3 plants ($\underline{A}/\underline{a}/\underline{a}$) when used as the pollen parent is given in Table $\overline{1}$. Corresponding control data are given in Table 2.

The control data in Table 2 indicate that the theoretical ratio of 1 A: 2 a is held to very closely. There is only one progeny out of twenty-five in which the percentage of A gametes is significantly higher than 33.3%, but this may be expected at the .05% level. The interaction chi square of 32.7 with 24 degrees of freedom is not significant. The data are homogeneous.

In the case of the trisomes with an irradiated chromosome, it is an entirely different situation. Six of the 26 plants had transmission frequencies of A gametes significantly lower than 33.3% and six others had rates which were significantly higher than 33.3%. The former was

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<u>.</u>	Table 1									
	No.A gam.	Total gamet.	% <u>A</u>	x ² (1:2)	No.A Total X^2 gam. gamet. $\%$ A (1:2)					
	Ban.		** O O C	31.3**	14 218 650 33.54 0.0					
1	523	1280	40.85	7.6**	15 462 1379 33.50 0.0					
2	207	532	38.91		16 703 2110 33.32 0.0					
3	462	1201	38.47	14.4**	17 248 746 33.24 0.0					
4	572	1493	38.31	16.5**	1/ 2/0 /					
5	178	465	38.28	5.1*	- TO 21 10 1.7					
6	343	931	36.84	5.3*	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7					
7	256	699	36.62	3.4	20 100 100 29 111 4.7*					
8	262	. 728	35.99	2.2	21 101 /01 27 01 10.6*					
9	375	1047	35.82	2.8	22 250 024 275.1*					
10	311	1435	35.61	3.4	23 142 700 2017 67.2°					
11	689	1961	35.14	2.9	24 97 700 27 00 76 43					
		817	34.27	0.4	25 05 722 -2007 /164 2					
12 13	_	410	34.15	_	26 30 1101 2.97 464.2					

Table 2 Control

				003	10101				
	No.A	Total gam.	% <u>A</u>	x2		o. <u>A</u> T	otal gam.	% <u>A</u>	x ²
1 2 3 4 5 6 7 8 9 10	265 463 481 296	468 467 943 484 1485 1581 2765 765 1338 1401 865 911	36.96 36.62 36.48 36.36 35.49 35.17 34.97 34.64 34.60 34.33 34.22 34.14	3.7 2.2 4.3* 2.1 3.1 2.4 3.3 0.6 1.0 0.6 0.3 0.3	15 16 17 18 19 20 21 22 23 24 25	196 202 390 286 167 162 180	831 405 416 673 1641 607 631 1229 932 546 540 602 23377	33.57 33.33 32.93 32.69 32.36 32.29 32.01 31.73 30.69 30.59 30.00 29.90 33.81	0.0 0.0 0.1 0.7 0.3 0.5 1.5 3.0 1.9 2.7 3.3 2.4
1	3 288	851	33.84	0.1	Total	700			

expected, but the latter was not, since there was no indication of "negative preferential pairing" when In 3a chromosomes were used in an earlier experiment. The term, chromosomes were used in an earlier experiment. The term, "negative preferential pairing," may not be a good one. "negative preferential pairing is still preferential and It is possible that the pairing is still preferential and that the tacit assumption that "the greater the structural that the tacit assumption that "the greater is their homology between two chromosomes is, the greater is their pairing affinity at meiosis" needs to be examined critipairing affinity at meiosis" needs to be examined critically. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that have synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally in hybrids in hybrids is cally. It has been observed that synapsis in hybrids is cally. It has been observed that synapsis in hybrids is cally in hybrids in hybrids is cally in hybrids. It has been observed that synapsis in hybrids in hybrids is cally in hybrids. It has been observed that synapsis in hybrids in hybrids is cally in hybrids. It has been observed that synapsis in hybrids in hybrids

Let us consider the spatial orientation of two homologous chromosomes prior to synapsis, i.e., at the leptotene stage of meiosis. To perhaps oversimplify the matter, stage of meiosis. To perhaps oversimplify the matter, they will either be in reverse or non-reverse position in they will either be in reverse or non-reverse position in regard to order of their pairing code units. See figures 1 and 2.

A	В	C	D	E	F	G	Н	I	
Ī	H	G	F	Ε	D	С	В	<u>A</u>	
	Fig. 1								

Reverse Orientation

Fig. 2

Non-reverse Orientation

It may be easily imagined that the initiation of synapsis in the reverse orientation might be very difficult; one of the chromosomes or part of one would have to be rotated 180°. the chromosomes or part of one would have to be rotated 180°. If, however, a small inversion has been induced as is shown in Fig. 3 and Fig. 4, then pairing could take place with in Fig. 3 and Fig. 4, then pairing could take place with ess difficulty when the chromosomes are in the reverse less difficulty when the chromosomes slither past each in their orientation. As the chromosomes slither past each in their random movements, the directionally homologous segments random movements, the directionally homologous segments could make the initial contact and facilitate synapsis along the rest of the chromosome. A small inversion would not the rest of the chromosome. A small inversion would not seriously hamper synapsis when the chromosomes were in the non-reverse orientation.

No cases of "negative preferential pairing" were found when In 3a chromosomes were irradiated. This is probably because an inversion is already present and the chromosomes do not need help in pairing in the reverse orientation.

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Any additional inversions would not help much and would 112 hamper pairing in the non-reverse orientation.

Some derivatives of the irradiated In 3a chromosomes from the original experiment have been reintroduced into trisomic 3 plants and the amount of preferential pairing has been redetermined. This was done by taking plants grown from colored (A) kernels from the testcross, $a/a \times A/a/a$, and crossing them to the standard trisome 3 stock (a/a/a) and crossing the resulting trisomes as the pollen parent to The results are given in Table 3. the a/a tester.

In the first case in Table 3, the \underline{A} locus has probably crossed over onto a normal chromosome. It is significantly tests for a fit to a l A : 2 a ratio. higher than 33.3% A. Perhaps a chromosome aberration which now causes negative preferential pairing has been retained.

Table 3 $(x^2)!$ Total No. of % A Derived gametes Original Trisome Λ gametes . case. # Λ % 5.1* 36.90 905 334 1 17.15 0.0 15.72 369 58 2 8.7** 15.65 13.60 25.6** 463 63 10.90 18.5** 743 18.14 81 10.29 45678 39.3** 447 46 10.73 4.4* 1072 115 15.45 61.3** 16.2** 906 140 11.81 2277 269 15.00 2402 360 14.2** 15.23 7.2** 1057 161 14.58 10 830 0.8 11.54 121 20.49 11 693 142 12 0.2 12.08 886 107 17.0** 13 12.56 14.58 830 0.5 121 11.20 14 375 10.23 42 23.58 15 814 15.8** 7.55 192 16 2000 0.0 151 21.81 17 674 3.7 147 12.43 18 748 1.6 93 12.10 19 347 42 20

See text for explanation of chi square tests

In the cases 2 - 11, 13 - 15, 17, 19, and 20 the chi square tests for a correspondence between the original transmission rate of \underline{A} and the derived one. In cases 2, 13, 15, 19, and 20 the rate has been unaffected. The other cases 3 - 11, 14, and 17 show a shift in the amount of preferential pairing, possibly due to the loss of positively or negatively acting aberrations by crossing over.

In cases 12, 16, and 18, the amount of preferential pairing has reverted to that expected from In 3a alone (22% \underline{A}), again probably by crossing over.

Unfortunately the data are too limited to make any sweeping conclusions. It is apparent that the level of pairing affinity is heritable and that it would be possible to map the location of these "synaptic mutations."

Another method of detecting preferential pairing has been devised and tested. Pollen from wx/wx plants was given 1000 r and used to fertilize trisome 9 plants which were homozygous for $\frac{Wx}{Wx}$. The pollen from the resulting trisomic plants $\frac{(Wx/Wx)}{Wx}$ is stained with iodine and is scored for Wx and wx. The results of this experiment are given in Table 4.

Table 4

Table 4									
						lrradia	ted		
	Cont			No	D. WX	Total gametes	% wx X	(wx = 23.22)	
	No. <u>wx</u> gametes	Total gametes	% <u>wx</u>		metes	gametes 681	21.44	1.2	
1 2 3 4 5	182 182 158 144 129	708 731 684 654 646 3423	25.71 24.90 23.10 22.02 19.97 23.22	6 7 8 9 10 11	146 128 114 119 109 106 88	656 635 676 621 652 605	19.51 17.95 17.60 17.55 16.26 14.54	4.9* 9.6** 12.0* 11.1** 17.4** 31.3**	
	ter. X ²	= 8.23 (not sig	.)12				2.2	

As may be seen in Table 4 the average frequency of wx pollen The interaction chi in the five control trisomes was 23.22%. square was only 8.23 with 4 degrees of freedom, so the data are homogeneous. Testing the frequency of wx pollen in the trisome 9 plants which received an irradiated wx chromosome against the value of 23.22% it was found that six out of seven plants gave an indication of preferential pairing.

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The advantages of this method may be readily apparent. 114 is possible to examine a hundred thousand pollen grains or more if necessary and thus it is possible to detect very small differences in the level of preferential pairing. Also it is feasible to look for rare spontaneous changes in chromosome structure affecting chromosome pairing without having to plant an acre of tester plants. bility, which now can be tested easily, results from nonhomologous pairing of a univalent (the pairing with itself) in a trisome. Crossing over in this non-homologously paired region would lead to the formation of an inversion.

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Japanese local races of maize resistant to the virus disease, corn stunt.

Four virus diseases are known to occur in maize under natural conditions. Stunt disease transmitted by the smaller brown planthopper, Delphaiodes striatella Fallén, is the most harmful one in Japanese maize production. A great deal of damage by virus disease is done to maize cultivation in the southern district of Japan, especially in Kyushu.

Over a period of 2 years many varieties were tested for resistance to stunt disease at the Miyakonozyo Sub-station of the Miyazaki Agricultural Experiment Station, Miyazaki Seventy materials (48 Japanese local races, 17 varieties introduced from foreign countries, and 5 recommended hybrids) were tested in 1963. The results showed that all but 2 Japanese races, Kamigane-l and Suyame-inno-l, had high susceptibility to this disease. Frequency (%) of diseased plants and index of susceptibility* was over 50% and 1.70 respectively. However, Kamigane-1 showed only 13.5% and 0.42, and Suyama-inno-1 showed 24.4% and 0.65 respectively.

In 1964, two hundred eighteen races (151 Japanese local races and 67 races collected from foreign countries, of which 17 were from Thailand), were tested. It is said that most of the Thailand races originated from the progenies of Guatemala Japanese races mentioned above had low resistance to the disease, showing similar values regarding susceptibility as the test in 1963. The values in some Thailand races varied from 9.4% to 45.2% and 0.3 to 1.4. The values of Kamigane-1 were 12.5% and 0.3, and those of Suyama-inno-1 were 6.3% and o I respectively.