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1. Recombination in the alsh2 region.

Backcross data on recombination rate in three different genetical backgrounds, having in common the "Texas" male sterile cytoplasm, are reported for \underline{a}_1 \underline{sh}_2 markers in coupling in the following table:

enetical	No. of	Total no.	A ₁ sh ₂ seeds	a ₁ Sh ₂ seeds	% of recombination	
ackground	GSTD		9	14	0.107	
1158	3 6	21386	•	·	0.072	
122	33	9632	3	4	0.114	
	31	12215	. 8	6	Q. 114	
NF9	_	1.0000	20	21,	0.101	
Totals	100	43233	CU CU		Pet of Pet 19 at 18 at 18 at 19 at 18 at 1	

The crossover value is clearly lower than that previously reported (0.27%). However, it is undecided whether this is due to different environmental conditions (including the T type cytoplasm) or to a difference in nuclear genotypes, although the latter interpretation appears less likely because of the uniform behavior of the different appears of the inbred lines used. It is interesting that this backgrounds of the inbred lines used. It is interesting that this value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intracistron value is of the same order of magnitude as at least one intraci

A. Bianchi G. Lapietra

2. Effect of storage on x-rayed pollen.

By applying Everett's technique it has been possible to prolong the pollen life-span and to study the effect of storage on x-rayed pollen (1750 roengten in about 7!).

The data appearing in the following table are roughly in agreement with those obtained with storage of dry irradiated seeds:

Temperature and relative humidity		Storage periods (in hours)	Chromosome and markers	Kernels scored	% of germless kernels	% of endosperm mutations
3°±1°C	70-90%	1 4 60	3,5 a ₁ sh ₂ ,pr	988 1130 980 498	9.6 16.6 9.1 13.2	14.6 19.6 17.6 13.6
20-30°C	60-70%	121 1 4 22	idem	663 587 153	11.7 19.2 17.9	10.2 19.0 22.2 18.2
3° <u>+</u> 1°C	70 - 90%	30 1 48 78	idem*	136 716 193 340 68	22.0 15.5 18.6 19.1 19.1	13.7 19.8 11.9 14.0
3° <u>+</u> 1°C	70 - 90%	96 1/2 148 1 1/2 52 148	9 C sh bz wx	217 493 558 212	23.9 4.2 6.6 6.6	16.4 2.6 4.3 3.3

The data marked with an asterisk refer to a hybrid type, homozygous nevertheless for the appropriate markers involved: the radio-sensitivity of the hybrid appears somewhat greater as compared with that obtained from an inbred line.

The recovery following the initial increase of damage could partially be ascribed, either to the active metabolism of pollen and/or to aplontic selection. The types of mutations detected are of the kind common in maize, and interpretable as chromosomal deletions.

G. Gavazzi

3. Mutagenic activity of nebularine and ethyl methane sulphonate.

Mutagenic activity of such chemicals has been studied on the basis of endosperm mutations, following treatment of mature pollen. Solutions are administered with a vial applied to tassels newly shedding pollen 24 hours before a first pollination, which was, then, repeated for 2 more successive days.