

preferential segregation with loci in chromosome 3, a number of plants were obtained which were heterozygous for abnormal 10 and carried a single B chromosome. Although our earlier studies had indicated no homology between the extra segment of abnormal 10 and B chromosomes, it was felt that a further examination was called for in view of Ting's (Chromosoma 1958) suggestion that the abnormal 10 which he found arose from a B-10 translocation. If the extra segment of the abnormal 10 found by Longley and subsequently widely studied by others has come from a B chromosome via translocation, pairing between the distal end of abnormal 10 and homologous regions of the B chromosome should occur frequently in plants with a single B and heterozygous for abnormal 10 because of a lack of pairing competition. A large number of pachytene figures were examined in such plants and, except for an occasional adhesion of the knob-like region of abnormal 10 to the distal heterochromatic regions of the B, there was no evidence of homology between the two chromosome segments.

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1. The effect of the independent activator of a-mutable on pale green mutable.

In a previous communication (MCNL 30) it was reported that pale-green-stable (pg^S) does not respond to the activator of a^m . It is now believed that in this case a non-mutable pg^S (similar to a^{dl}) was being tested. Recent experiments with a known mutable pg^S clearly show that pg^{S*} does mutate in the presence of the independent controller of a -mutable. Furthermore, this controller corresponds in every way to Enhancer and so will be designated En.

The test of similarity showing En to cause both a^{mI**} and pg^S to be mutable.

	F ₁ pg-segregating types
+/ pg^{S*} A A x Pg/Pg a sh/a sh ^{***} + mutable factor	-- 1/2 Pg/ pg^S A/a sh
no mutable factor	1/2 Pg/ pg^S A/a sh
+ mutable factor	

* known to respond to En

** a^{mI} is mutable only in the presence of En

*** this a responds to Dt and not to En

	Total No. of ears	In progeny of \bar{E}		In progeny of \bar{E}	
		segregates pg^m & pg^s	segregates only pg^s	segregates pg^m & pg^s	segregates only pg^s
1958					
1370	8	4	0	0	4
1371	4	1	0	0	3
1372	3	2	1****	0	0
1373	12	<u>8</u>	<u>0</u>	<u>0</u>	<u>4</u>
		15	1	0	11

** a^mI is mutable only in the presence of \bar{E}
 *** This a responds to Dt and not to \bar{E}
 **** only exception to correspondence of \bar{E} causing pg^s to be mutable and a^mI to be mutable. In all other cases, the occurrence of pg^m is correlated with a -mutability.

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2. Other factors associated with a-mutable.

Dense factor

Among the tested progeny of one a^m allele, two very distinct pattern types were observed: very dense mutable (almost full color) and fine mutable (like Dotted). The following sample data represents some of the typical segregations of the two phenotypes.

The Cross: $a^m Sh/a sh \times a sh/a sh$
 (Dense) \times colorless shrunken

1957	non-shrunken kernels	
	<u>mutable</u>	
	<u>very dense</u>	<u>fine</u>
410-30 x a sh	93*	89**
31B-1 x a sh	44	52
410-4 \bar{E}	117	49
410-14 \bar{E}	168	42
410-31 x a sh***	76	39

* On further tests, Dense gives rise to 1/2 dense : 1/2 fine.
 ** On further tests, fine mutable gives rise only to fine.
 *** These off-ratios have not as yet been analyzed.