To determine whether a component of the instability was due to the different growth factors in the media (2,4-D; 2,4,5-T; or 2,4,5-TP), all counts were pooled for this assessment according to media type (Table 2). Both diploid and haploid cultures did not appear to have been affected by the different growth factors in the media.

Whether complete chromosome stability would be realized upon continued growth of these cultures cannot be as yet determined. However, finding such a high proportion of cells with the stable karyotype after nine months of culturing suggests that they serve as the primary progenitors in future callus growth. Otherwise their relative frequency should decline with time.

The data reported here show that studies of somatic mutability would be profitably undertaken under such a technical regime of tissue culture. In the case of haploid callus it would seem at this time to be workable material for a study of induced mutation rates. Since a completely defined media is available, mutations affecting basic metabolic pathways could be selected for. It does not now seem out of place to suggest that this technology of tissue culturing could yield genetic information of the order found in microbial genetics of today with the exception of not having a high resolution recombination system.

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1. Linkage relationship between Y-y and Rf2-rf2 on chromosome 6.

From tests involving $\underline{Rf_2}$ and a series of chromosomal translocations, it appears that $\underline{Rf_2}$ is located on the short arm of chromosome 6 (Beckett, Maize Genetics Coop. News Letter 36:31, 1962).

The data which follow were obtained from crosses involving $\underline{Y}-\underline{y}$ and $\underline{Rf}_2-\underline{rf}_2$. Since $\underline{Y}-\underline{y}$ is reported to be located approximately 13 crossover units from the distal end of the short arm of chromosome 6, it was expected that $\underline{Y}-\underline{y}$ would be closely linked with $\underline{Rf}_2-\underline{rf}_2$.

From the following linkage data it appears that these two genes are not linked and therefore Rf2-rf2 is not on the short arm of chromosome 6.

Cross #1	Parent <u>RfjRfj</u>	A rforf	2 <u>YY</u> T	x :		foriovy
			Expec F2 Ratio	ted	Actual F ₂ Ratio	
X ² for <u>Y</u> -y Segregati	on :	3.43	(9)			Fertile
X ² for <u>Rf-rf</u> Segrega	tion	3.43	(3)			Sterile
	•	1.37	(3)	35		Fertile
X ² for Linkage X ² for Total		8.24	(1)	<u>14</u> 153	White	Sterile
P = .5020 for L	inkage	<u>x</u> 2				
Cross #2	Dorei		<u>rf2YY</u>	x		rf2rf2YY
			Expe F ₂ Rati	cted o	Actu F ₂ Rati	o
X ² for Y-Y Segregate X ² for Rf-rf Segre X ² for Linkage	tion	0.04) 16		w Fertile
	gation	2.97			•	W Sterile
		1.63		٠.		e Fertile e Sterile
x ² for Total		4.64	+ (7) 4		5 5002
P = 0.20 for Links	ige X2	······································				

Backcross #1

Parent C Parent D rf1rf1Rf2Rf2YY x Rf1Rf1rf2rf2yy Parent D

		Expected BC Ratio	Actual Backcross Ratio
X ² for Y-y Segregation	4.32	(1)	160 Yellow Fertile
X ² for Rf-rf Segregation	կ. 32	(1)	150 Yellow Sterile
X ² for Linkage	3.11	(1)	151 White Fertile
K ² for Total	11.75	(1)	213 White Sterile
P = .1005 for Linkage	χ2		

Cross #3

Parent C Parent D rf_rf_Rf_Rf_YYY x Rf_Rf_rf_2rf_2yy

(F₁) Rf1rf1Rf2rf2Yy x Rf1Rf1Rf2rf2yy Parent E

(3)	141 Yellow Fertile
(1)	38 Yellow Sterile
(3)	127 White Fertile
(1)	لِوع White Sterile
	(3)

Cross #4	Parent A <u>Rf_lRf_lrf₂r</u>	f ₂ YY	x	Parent F <u>rf₁rf₁Rf₂Rf₂yy</u>
		Expec F ₂ Ratio	ted	Actual F ₂ Ratio
X ² for <u>Y-y</u> Segregati	n 1.04	(27)	125	Yellow Fertile
x ² for <u>Rf-rf</u> Segregat	_ •	(21)	114	Yellow Sterile
X ² for Linkage	0.50	(9)	50	White Fertile
X ² for Total	1.78	(7)	<u>40</u> 329	White Sterile
P = .50 for Linkage	^x 2			
Cross #5	Parent A Rf1Rf1rf2	rf2YY.	x	Parent G rflrflRf2Rf2YY
		Expec F ₂ Ratio		Actual F ₂ Ratio
x ² for <u>Y-y</u> Segregati	on .06			Yellow Fertile
X ² for <u>Rf-rf</u> Segrega		(21)	46	Yellow Sterile
X ² for Linkage	.92	(9)	18	White Fertile
x ² for Total	0.98		22 154	White Sterile
P = .5020 for Li	nkage X ²			

The parents involved in each cross were test crossed to lines of know Rf genotypes in order to verify parental Rf genotypes.

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