Barratt, R. W. and P. St. Lawrence. Anti-

metabolite inhibition of mod-5.

tation can be rationalized as consequences of a change in permeability that facilitates the entry of a number of metabolites into the organism. They observed that <u>mod-5</u> strains were completely inhibited by concentrations of the antimetabolites p-fluorophenvlalanine and 4-methyltryptophane which had little or no effect on unmodified cultures.

The data in Table 1 (P. St. Lawrence) and Table 2 (R. W. Barratt) support the above observations and indicate that the use of these antimetabolites is a gwd method for scoring for the presence of the <u>mod-5</u> mutation. The results are expressed as <u>mycelial</u> dry weight in In 1964 St. Lawrence, Maling, Altwerger and Rachmeler (Genetics 50: 1384) reported the genetics and physiology of a gene designated as <u>mod-5</u> (modifier of permeability) induced in a <u>tryp-3</u> (td16) stock and concluded that all of the phenotypic manifestations of the mod-5 mu-

Table ]. Inhibition of mod-5 by antimetabolites in cultures grown at 34°C.

Strain	p-fluorophenylalanine (conc.in ४∕ml)		4-methyltryptophan (conc.in ∛ml)	
	0. 1	1.0	1.1	11.0
wild type (isolate 2.3)	94.9	53.7	59.0	48.1
mod-5 (FGSC <sup>#</sup> 1603)	80.3	0.0	64.2	0.0
wild type (isolate 6. 1)	90.8	86.9	w.2	71.1
<u>mod-5 (</u> isolate 6.3)	59.	0.5	13.2	0.0

milligrams from 72-hour stationary cultures (except where noted) grown in 20 ml of Vogel's minim | N containing 2% sucrose plus the indicated antimetabolite (added after autoclaving). The inoculum was approximately 10<sup>8</sup> conidia per flask.

Strain	Temperature	p-fluorophenylalanine (conc.in /ml)		4-methyltryptophan (conc.in ∛/ml)	
		0.0	1.0	0.0	11.0
wild type (FGSC <sup>#</sup> 987)	25°C*	55.3	1.2	35.0	38.0
	34°C	48.6	34.0	70.5	43.9
mod-5 (FGSC#1603)	25°C*	46.6	0.0	51.4,	0.0
	34°C	102.2	0.8	39.4	2.3
Yield measured in mg dr	y weight. *Har	vested at 9	6 hrs.		

Table 2. Inhibition of mod-5 by antimetabolites in cultures grown at 25°C and 35°C.

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