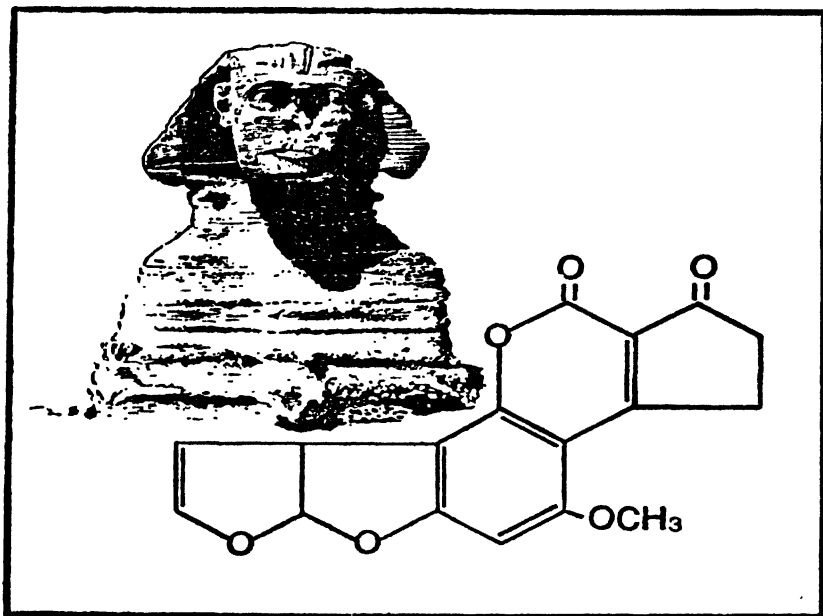


# PROCEEDINGS OF THE INTERNATIONAL SYMPOSIUM ON MYCOTOXINS



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## ***Aflatoxin Production in Groundnut Cultivars Resistant and Susceptible to Seed Invasion by *Aspergillus flavus****

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### **A b s t r a c t**

*The use of groundnut cultivars resistant to seed invasion and colonization by *Aspergillus flavus* is a possible means of preventing or reducing aflatoxin contamination. Such resistance was identified in several cultivars, one of which was the released commercial Indian cultivar J11. No direct relationship was found between resistance to seed colonization by *A. flavus* and the quantity of aflatoxins produced when seeds were colonized by toxigenic strains of the fungus. Some of the cultivars with seed resistance to *A. flavus* colonization have been found resistant to fungal pod rot.*

### **I n t r o d u c t i o n**

Contamination of groundnuts with aflatoxin is a serious problem in many parts of the world. The use of groundnut cultivars resistant to seed invasion and colonization by toxigenic strains of *Aspergillus flavus* has been suggested as an effective means of preventing or at least reducing aflatoxin contamination (1-3). Early research reported varietal differences in aflatoxin production when autoclaved seeds were colonized by toxigenic strains of *A. flavus* (4,5). Although claims of resistance to aflatoxin production were not confirmed by later research (6,7), quantitative varietal differences in aflatoxin production were indicated.

This paper reports on screening of germplasm for resistance to seed invasion and colonization by *A. flavus* and for resistance to aflatoxin production when seeds are colonized by toxigenic strains of the fungus.

## Materials and Methods

### Source of Seed

Seeds of all cultivars were obtained from the 1979 and 1980 rainy season and the 1979-80 and 1980-81 post-rainy season crops grown at ICRISAT Center, Patancheru, A.P., India. The crops were not subjected to drought stress. Cultivars were harvested at optimum maturity and plants were arranged in windrows in the field with pods exposed. After windrow-drying for 2 days in the post-rainy and 3 days in the rainy seasons, the pods were hand-picked and sun-dried on mats until seed moisture contents were reduced to less than 7%. Pods were then stored in cloth bags in the laboratory until seeds were required for testing.

### Seed Colonization Test

The Mixon and Rogers method (1) with some modification (8) was used. For each cultivar, 3 plates of surface-sterilized (soaked in 0.1% aqueous solution of mercuric chloride for 2 min followed by 4 rinses in sterile distilled water) hydrated seeds were inoculated with a spore suspension ( $4.0 \times 10^6$  conidia/ml) of the toxigenic *A. flavus* strain AF8-3-2A. The plates were incubated at 25° C and percentages of seeds colonized were recorded after 8 days.

### Aflatoxin Production Test

Using seed from the 1980 rainy season crop, aflatoxin production tests were carried out on each cultivar by the method described by Mehan and McDonald (8). Surface-sterilized seeds in 3 plates had their testas damaged by scraping them with a sterile needle. The seeds were then inoculated with a spore suspension of *A. flavus* strain AF8-3-2A and incubated as described above for the seed colonization test. After 10 days of incubation, aflatoxins were extracted using Romer's method (9), determined quantitatively by the method of Nabney and Nesbitt (10), and expressed as micrograms per gram of seed.

## Results and Discussion

From 150 cultivars screened, 8 showed resistance to seed colonization. Mean seed colonization percentages are shown in Table 1 for the 8 resistant cultivars, and 2 control cultivars, the susceptible TMV 2 and the highly susceptible OG43-4-1, for seeds from 2 rainy season and 2 post-rainy season crops. The cultivars Ah 7223, Monir 240-30, and UF 71513 showed the highest resistance. Less resistant, but still significantly more resistant than TMV 2, were cultivars J 11, Faizpur, Var. 27, PI 337394 F, and PI 337409. PI 337394 F and PI 337409 have been reported resistant in the United States and in Senegal (2). These 2 cultivars are not agronomically acceptable but J 11 is a released commercial cultivar in India. This cultivar and the other *A. flavus* resistant cultivars Ah 7223, Monir 240-30, and Var. 27 have also been found resistant to a fungal pod rot at ICRISAT Center.

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TABLE 2. Aflatoxin B<sub>1</sub> production in groundnut cultivars resistant and susceptible to seed invasion by toxigenic *A. flavus*

Cultivar	Reaction to seed invasion by <i>A. flavus</i>	Aflatoxin B <sub>1</sub> (µg/g seed)
PI 337394 F. . . . .	Resistant	106.4
PI 337409. . . . .	Resistant	95.5
J 11 . . . . .	Resistant	117.8
Ah 7223 . . . . .	Resistant	115.2
Monir 240-30 . . . . .	Resistant	93.6
Var.27 . . . . .	Resistant	90.3
Faizpur . . . . .	Resistant	113.5
TMV 2 . . . . .	Susceptible	226.2
FESR-11-P11-B2-B1 . . . . .	Highly susceptible	50.0
OG 43-4-1 . . . . .	Highly susceptible	76.3
SD (at 5%)		13.2
CV (%)		7.1

supported production of aflatoxin B<sub>1</sub> and there were significant differences between them in the amount produced. There was no correlation between resistance to seed invasion and colonization by *A. flavus* and the ability of the seed to support aflatoxin production. The cultivar FESR-11-P11-B2-B1, which is known to be highly susceptible to seed invasion and colonization by *A. flavus* (13), had the lowest level of toxin production (Table 2). In another experiment, the rate of accumulation of aflatoxin B<sub>1</sub> was shown to be slower in this cultivar than in cultivars TMV 2, J 11, and PI 337409 (Fig. 1).

It would be most useful if cultivars could be developed which combined seed resistance to invasion and colonization by *A. flavus* with resistance to aflatoxin production in the event of the seeds being colonized by a toxigenic strain of the fungus.







