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Note

Rapid separation and detection of some systemic fungicides by paper chromatography

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As fungicides are now being used extensively in controlling plant diseases, it is necessary to determine that the plant parts used for food are free from harmful quantities of the residues of the fungicides used. The analysis of fungicide residue from plants involves their extraction in a suitable solvent, followed by separation and detection by several methods. Of the many methods available, paper and thin-layer

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STEMIC FUNGICIDES WITH THEIR COINED, LABEL AND CHEMICAL NAMES, AND STRURAL FORMULA AND MANUFACTURERS

ined name	Label name	Chemical name	Structural formula	Manufacturer
rboxin	Vitavax	5,6-Dihydro-2-methyl- 1,4-oxathiin-3-carbox- anilide		Uniroyal Chemical Co. Bethany, Conn., U.S.A
ycarboxin	Plantvax	2,3-Dihydro-5-carbox- anilide-6-methyl-1,4- oxathiin-4,4-dioxide		Uniroyal Chemical Co.
iabendazole	Mertect	2-(4-Thiazolyl)benzimid- azole		Merck Chemical Divisie Rahway, N.J., U.S.A.
idemorph	Calixin	N-Tridecy1-2,6-dimethyl- morpholine	$CH_3 \rightarrow C_{13}H_{27}$	Badische Anilin & Soda Fabrik (India), Bombay India
ia W 524	Triforine	N,N'-Bis(1-formamide- 2,2,2-trichloroethyl)- piperazine.	CI ₃ C CH NH CHO	Cela Landwirtschaftlich Chemikalien, Ingelheim Rhein, G.F.R.

chromatography have recently proved to be the simplest and most sensitive for separation and identification. Thin-layer chromatography has been used for the analysis of fungicides such as thiram, ziram, zineb^{1,2}, Botran³, carboxin, oxycarboxin^{4,5} and thiabendazole (Mertect)⁶. In this paper, a simple and quick method for separation and identification of five systemic fungicides, viz., Vitavax, Plantvax, Mertect, Calixin and Triforine, by ascending paper chromatography is described.

MATERIALS AND METHODS

The fungicides used were: carboxin, oxycarboxin, thiabendazole, tridemorph and Cela W 524 (Table I).

Standard solutions of the fungicides and their mixtures were prepared in acetone. Samples were spotted on 25×13 cm paper with the help of a micropipette. After allowing each spot to dry for a few minutes, the paper was placed in a chromatographic chamber which was previously saturated with the solvent system, acetonewater (1:6). Keeping the spotted edge downwards, the paper was gently immersed in a trough containing the solvent system. The upper end of the paper was held by clips. The chamber was covered immediately, and the solvent system was allowed to ascend by a distance of 20 cm. The chromatogram was removed from the chamber and dried at 110° for 5 min in an oven.

The fungicides were located on the chromatogram by allowing the paper to react with the iodine vapour for 1 min. The R_F values were then recorded (Table II).

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R _F	VALUES	OF	SOME	SYSTEMIC	FUNGICIDES	AND	THEIR	MIXTURES

Fungicides and their mixtures	R_F values in acetone-water (1:6)				
Vitavax	0.81				
Plantvax	0.89				
Mertect	0.49				
Calixin	0.00				
Triforine	0.71				
Vitavax - Plantvax	0.81				
	0.89				
Vitavax -+- Mertect	0.83				
	0.49				
Vitavax + Calixin	0.81				
	0,00				
Vitavax Triforine	0.82				
	0.69				
Plantvax Calixin	0,87				
	0.00				
Plantvax + Triforine	0.87				
	0.71				
Mertect + Calixin	0.49				
	0.00				
Mertect + Triforine	0.49				
	0.74				

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RESULTS AND DISCUSSION

Table II gives the R_F values of five fungicides and eight of their binary mixtures. The R_F values of individual fungicides in a mixture are more or less similar to the R_F values of the pure fungicides. Thus it is possible to separate and identify systemic fungicides singly or in mixtures. The technique is being employed in this laboratory in the determination of the persistence of fungicides in plants. The separation can be accomplished in about 3 h.

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