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Note

Benzoyl peroxide as a spray reagent for carbazole alkaloids

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There is considerable interest in the chemistry of carbazole alkaloids¹, and thin-layer chromatography has been found to be suitable for their separation and identification. Carbazoles could be detected on the developed chromatogram by their fluorescence or quenching under UV light as well as from the colours developed by spraying with picric acid, DDQ or HCl^{2,3}. However, DDQ has been reported to be toxic³, and acids are not very convenient to use.

We have found that benzoyl peroxide is an effective spray reagent for the detection of carbazoles and has lower detection limits than previously used reagents as found by direct comparison.

Thin-layer glass plates $(20 \times 10 \text{ cm})$ were prepared using silica gel G as the adsorbent and the chromatograms were developed with benzene-chloroform (1:1) in

TABLE I

Compound	Source	R _F	Colour developed
Carbazole	Fluka (Buchs, Switzerland)	0.9	Yellow
Murrayanine	Murraya Koenigii Spreng.	0.35	Yellow
3-Methylcarbazole	Clausena heptaphylla Wt. & Arn.	0.8	Yellow
Mukonal	Murraya koenigii Spreng.	0.31	Yellow
Mukonine	Murraya koenigii Spreng.	0.36	Yellow
Glycozoline	Glycosmis pentaphylla (Retz)	0.69	Yellowish green
Glycozolidine	Glycosmis pentaphylla (Retz) DC	0.51	Yellowish green
Koenimbine	Clausena heptaphylla Wt. & Arn.	0.65	Violet
Koenidine	Clausena heptaphylla Wt. & Arn.	0.25	Violet
Murrayacine	Murraya koenigii Spreng.	0.35	Violet
Heptazolidine	Clausena heptaphylla Wt. & Arn.	0.74	Violet
Girinimbine	Murraya koenigii Spreng.	0.76	Violet
Murravacinine	Murraya koenigii Spreng.	0.35	Yellow
Mahanimbine	Murraya koenigii Spreng.	0.93	Yellow
Murryazoline	Murraya koenigii Spreng.	0.95	Yellow

COLOURS DEVELOPED AFTER SPRAYING WITH BENZOYL PEROXIDE AND RF VALUES

the usual way. After development the chromatograms were dried and sprayed with a 2% solution of benzoyl peroxide in chloroform, and colours formed immediately. The colours and R_F values for various compounds are given in Table I. Carbazoles can be detected at levels down to 0.05 μ g with this reagent, whereas the minimum concentration varies from 10 μ g to 0.1 μ g in the case of previously used reagents.

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