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PHYTOCHEMICAL SCREENING AND XRD ANALYSIS OF BLACK PEPPER PIPER NIGRUM

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ARTICLE INFO ABSTRACT

Article History:

Received 17th April, 2017 Received in revised form 10th May, 2017 Accepted 25th June, 2017 Published online 28th July, 2017 The aim of the study was to investigate the phytochemical compounds present in fruit part of *Piper nigrum* extract with petroleum ether, ethanol and chloroform as solvents. The phytochemical screening of plant extracts revealed the presence of steroids, terpenoids, alkaloids, flavonoids, anthraquinone, quinine, phenolic compounds, carbohydrates and proteins.

Key words:

Phytochemicals, XRD, P.nigrum.

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INTRODUCTION

Plants constitute a source of novel chemical compounds which are of potential use in medicine and other applications. Plants contain many active compounds such as alkaloids, steroids, tannins, glycosides, volatile oils, fixed oils, resins, phenols and flavonoids which are deposited in their specific parts such as leaves, flowers, bark, seeds, fruits, roots, etc. The beneficial medicinal effects of plant materials typically result from the combination of these secondary products (Tonthubthimthong et al., 2001). Phytochemicals are known to possess antioxidant (Wong et al., 2009), antibacterial (Nair et al., 2005), antifungal (Khan and Wassilew, 1987), antidiabetic (Singh and Gupta, 2007; Kumar et al., 2008a), antiinflammatory (Kumar et al., 2008b), and radioprotective activity (Jagetia et al., 2005) and due to these properties they are largely used for medicinal purpose. The main purpose of the present study was collection and identification of plant and screening for presence of various materials phytochemicals in these medicinal plants. Herbs and spices are very important and useful as therapeutic agent against many pathological infections (Gull et al., 2012).

MATERIALS AND METHODS

Collection and preparation of extracts

The medicinal plant *Piper nigrum* was collected from surrounding areas of Coimbatore district. Fruit part of *P.nigrum* of medicinal plant were separated, washed with distilled water, shade dried. They were ground in to powder and stored in room temperature.

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Preliminary phytochemical screening

The different qualitative chemical tests were performed for establishing the profile of given extracts to detect various phytoconstituents present in them.

Qualitative analysis

The different qualitative chemical tests were performed for establishing the profile of given extracts to detect various phytoconstituents present in them. Preliminary screening of the extracts and identification was done by colour tests adapting standard methods by Raman (2006). Tests for alkaloids, flavonoids, steroids, terpenoids, anthraquinones, proteins and carbohydrates were carried out.

Particle characterization

The X- ray diffraction (XRD) patterns of the samples were recorded on a PANalytical X'Pert PRO X-ray diffractometer using Cu K α radiation ($\lambda = 0.15406$ [']). The crystallite size of nanocrystalline samples was measured from the line broadening analyses using Debey- Schherer formula after accounting for instrumental broadening (Equation 1):

$$XRD = 0.89 \lambda / \beta \cos \theta$$

.....(1)

Where λ – wavelength of X-ray radiation used in ['], θ is the diffraction angle, β is the full width at half maximum (FWHM) in radians in the 2 θ scale, D XRD is the crystallite size in nm.21.

RESULTS

D

The phytochemical analysis of ethanol, chloroform and petroleum ether extracts of *P.nigrum*, *M.charantia* and *Z.officinale*, revealed the presence of phytochemicals in varying proportions (Table1). The presence of alkaloid has

seen in chloroform and ethanolic extracts but absent in petroleum ether extract. Alkaloids are one of the diverse groups of secondary metabolites found to have antimicrobial activity by inhibiting DNA topoisomerase. Carbohydrate which constitutes the major edible part of the plant is present in all the above three medicinal plant extracts. A considerable amount of flavanoids, anthraquinone, protein and phenol were present in all three extracts and steroids were present in ethanol and petroleum ether. The compound which possess large amount of flavonoids has found to have inherent ability to modify the body reactions to allergens, viruses and carcinogens. XRD pattern were analyzed and all experimental peaks were matched with theoretically generated one and indexed (Fig. 1).

DISCUSSIONS

Secondary metabolite studies of above medicinal plant have shown that the presence of carbohydrates, flavonoids, alkaloid, quinine, phenol, protein, terpenoids which are of great importance in the field of drug research. These classes' alkaloids, flavonoids are known to have activity against pathogens and therefore aid the antimicrobial activities of

S.No	Qualitative Tests		Petroleum Ether	Ethanol	Chloroform
		i) Mayers	-	-	+
1.	Alkaloids	ii) Wagners	-	+	+
		iii) Hagers	-	-	+
2.	Flavanoids	i) Sod.Hydroxide test	-	-	+
		ii) Sulphuric acid test	+	-	-
3.	Steroids	i) Libermann-Burchard	+	-	-
4.	Terpenoids	i) Libermann-Burchard	-	+	-
5.	Anthraquinone	i) Borntragers	+	+	+
	Protein	i) Ninhydrin (Aq)	-	-	-
6.		ii) Ninhydrin (Acetone)	-	-	-
		iii) Biuret	+	-	-
7.	Phenols	i) Ferric Chloride	-	+	-
		ii) Libermann	-	-	+
8.	Quinone	i) Conc HCl test	-	-	+
0	Carbohydrate	i) Molish	+	+	-
9.		ii) Fehlings A &B	-	-	+

Table 1 Phytochemical constituents of Piper nigrum

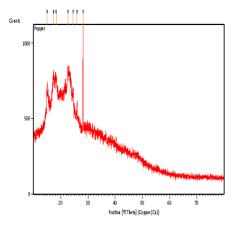


Figure 1 XRD Analysis of Piper nigrum

Table 2 XRD Analysis of Piper nigrum

Pos.[°2Th.] d-	Height [cts]	FWHMLe ft[°2Th.]	spac[Å]	Ret.Int. [%]
14.9906	219.92	0.2676	5.91005	53.24
17.3196	209.66	0.4684	5.12021	50.76
18.2742	163.11	0.4015	4.85484	39.49
22.6332	188.73	0.6691	3.92874	45.69
24.5391	141.09	0.2007	3.62775	34.16
26.0016	94.96	0.2007	3.42692	22.99
28.2043	413.08	0.1004	3.16410	100.00

XRD can be used to characterize the crystallinity of nanoparticles and it gives the average diameters of all the nanoparticles. The fine particles were characterized by XRD for structural determination and estimation of crystallite size.

medicinal plants (Ghosh *et al.*, 2010). In any research in phytotherapy, it is necessary to choose solvent according to biological activity required and not that which gives a high amount of bioactive compounds. The higher concentrations of more bioactive flavonoids compounds were detected with 70% ethanol due to its higher polarity than pure ethanol (Chantal *et al.*, 2005).

CONCLUSION

From the present study it is found that crude extract express good biological capacity which indicates that the substance with powerful biological effect exists in this extract and must be isolated and purified to confirm its pharmacological and medical use.

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