

Pharmacognostic aspect of *Acalypha indica*, *Vitex negundo* and *Coriandrum sativum*

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ABSTRACT

A study was carried out at Avinashilingam University, Coimbatore to observe the pharmacognostic, phytochemical and biochemical activities of *A. indica*, *V. negundo* and *C. sativum*. The pharmacognostic studies involved organoleptic study, fluorescence analysis, preliminary phytochemical studies and biochemical analysis. Different observation revealed the presence of biologically active compounds. The results of organoleptic study offer a scientific basis for the traditional use. The variation in Colour and taste were observed. Fluorescence analysis gave different colours of powders and treatment with chemicals exhibited changes in reactivity of powders. The biochemical activity also was found to vary significantly among tested plant powders.

Key words: Medicinal plants, *A. indica*, *V. negundo*, *C. Sativum* and Pharmacognostic property, phytochemical characters and biochemical activities.

INTRODUCTION

Now-a-days using plants for medical purposes become essential feature of the culture and the tradition. A Major part of the population depends directly on the traditional medicine for the primary health care. The use of higher plants and their extracts to treat infection is the usual practices. The biochemistry of herbs and the pharmacology of their various constituents is an aspect of phytotherapy that initiates many herbalists to do many researches. An investigation has been carried out to evaluate the pharmacognostic, phytochemical and biochemical aspect of *Acalypha indica*, *Vitex negundo* and *Coriandrum sativum*.

MATERIAL AND METHODS

To assess the activity of selected medicinal plants, pharmacognostic studies like organoleptic study, fluorescences analysis, preliminary phytochemical studies and biochemical analysis

were carried out. Organoleptic studies were based on method of Jackson and Snowdown (1968) and fluorescences properties following Kokoski *et al* (1958) and Chore and Pratt (1949). The biochemical parameters like protein (Lowry *et al.*, 1951), carbohydrates (Hedge and Hofreiter, 1962), reducing sugar (Miller, 1972), total soluble sugar (Mahadevan and Sridhar, 1986) phenol (Malick and Singh, 1980) and chlorophylls (Arnon, 1949) were analysed.

RESULTS AND DISCUSSION

Pharmacognostic study

By screening the various leaf extracts of three test plants, the pharmacognostic features were examined through the following parameters.

Organoleptic study

The investigation of organoleptic study of leaf power of *A. indica*, *V. negundo* and *C. sativum* indicated the characters like colour, odour and taste.

The colour of the leaf powder and the taste were observed and the results are shown in Table 1. The colour varied from dark green to light green in all the three plants with bitter taste and pungent odour. Pandey *et al.* (1984) and Gupta (1986) treated the plant powders with different chemical reagents and observed their behaviors. Here also the above three plant powders were treated with different chemical reagents to find out their diagnostic features.

Fluorescence analysis

All the plant powders treated with various chemicals exhibited various colours in the visible and UV light when compared to control, the three leaf powders revealed colour changes from green to brown in 3rd and 4th treatments. When the powders were treated with aqueous 1N NaOH and ethanol

all the plants exhibited varied green colours in visible and UV light and the result are depicted in Table 2.

With the help of fluorescence analysis, we can identify and discriminate *Aloe vera*, *Acorus calamus* and *Symplocos racemosa* from other species. This coincides with the study of Mary *et al.* (1980) who discriminated two species of *Valerina* L. on the basis of morphology, fluorescent analysis and microscopic characters.

Behavior of the leaf powders

It is evident from the results that the leaf powders of three test plants treated with chemicals like FeCl₃, NaOH, H₂O, I₂, HCl, +H₂O, KOH, ethanol, HNO₃ and H₂SO₄, various shades of green, red and brown colours were obtained. The powder as such

Table 1: Organoleptic study of the powders of *A. indica*, *V. negundo* and *C. sativum*

S. No	Name of the plant	Colour	Odour	Taste
1.	<i>A. indica</i>	Green	Pungent	Little bitter
2.	<i>V. negundo</i>	Light green	Pungent	Bitter
3.	<i>C. Sativum</i>	Dark green	Pungent	Little bitter

Table 2: Fluorescence analysis of the powders of *A. indica*, *V. negundo* and *C. sativum*

S. No	Leaf powder	Treatment with chemical reagent	UV light	Visible light
1.	<i>A. indica</i>	Powder as such	Dark green	Greenish yellow
	<i>V. negundo</i>		Pale green	Reddish brown
	<i>C. sativum</i>		Light green	Reddish yellow
2.	<i>A. indica</i>	Powder + aqueous 1 N NaOH	Pale green	Greenish yellow
	<i>V. negundo</i>		Light green	Yellow green
	<i>C. sativum</i>		Dark green	Reddish green
3.	<i>A. indica</i>	Powder + 1N HCl	Yellowish green	Greenish brown
	<i>V. negundo</i>		Pale green	Green
	<i>C. sativum</i>		Pale green	Greenish brown
4.	<i>A. indica</i>	Powder + 50 percent H ₂ O	Green	Green
	<i>V. negundo</i>		Pale green	Greenish brown
	<i>C. sativum</i>		Red	Red
5.	<i>A. indica</i>	Powder + ethanol	Pale green	Green
	<i>V. negundo</i>		Greenish brown	Yellowish green
	<i>C. sativum</i>		Light brown	Greenish brown

expressed varied green colours and when it was dissolved in water it showed no change in its colour. Various behaviors of powders with different chemical reagents are depicted in Table 3.

During a pharmacognostic study carried out on the flower of *Pterospermum cicerifolium* (L.) by Shome and Mehrotra (1990) greenish purple colour was noted on treatment with 1N HCl and nitro-cellulose.

Table 3: The behaviour of the leaf powders of *A. indica*, *V. negundo* and *C. sativum* when treated with different chemical reagents

S.NO.	Name of the plant	Treatment with chemical	Observation
1.	<i>A. indica</i>	Power as such	Green
	<i>V. negundo</i>		Dark green
	<i>C. sativum</i>		Yellowish green
2.	<i>A. indica</i>	Power + 2 % FeCl ₃	Dark green
	<i>V. negundo</i>		Yellowish green
	<i>C. sativum</i>		Dark green
3.	<i>A. indica</i>	Powder + 10 % NaOH	Green
	<i>V. negundo</i>		Yellowish green
	<i>C. sativum</i>		Dark green
4.	<i>A. indica</i>	Powder + 5 % KOH	Green
	<i>V. negundo</i>		Orange
	<i>C. sativum</i>		Dark green
5.	<i>A. indica</i>	Powder + water shake	No change
	<i>V. negundo</i>		No change
	<i>C. sativum</i>		No change
6.	<i>A. indica</i>	Powder + Iodine	Green
	<i>V. negundo</i>		Brown
	<i>C. sativum</i>		Dark green
7.	<i>A. indica</i>	Powder + HCl	Dark green
	<i>V. negundo</i>		Light green
	<i>C. sativum</i>		Dark green
8.	<i>A. indica</i>	Powder + NaOH + H ₂ O	Light green
	<i>V. negundo</i>		Brown
	<i>C. sativum</i>		Bright green
9.	<i>A. indica</i>	Powder + Ethanol	Green
	<i>V. negundo</i>		Bright green
	<i>C. sativum</i>		Green
10.	<i>A. indica</i>	Powder + Nitric acid	Brown
	<i>V. negundo</i>		Orange
	<i>C. sativum</i>		Brown
11.	<i>A. indica</i>	Powder + H ₂ SO ₄	Bright green
	<i>V. negundo</i>		Deep yellowish green
	<i>C. sativum</i>		Deep green

Phytochemical screening

Phytochemical analysis intends to serve as a major resource for information on analytical

and instrumental methodology in plant science (Table 4). A preliminary study was undertaken to assess the active constituents of *A. indica*,

Table 4: Analysis of phytochemicals present in *A. indica*, *V. negundo* and *C. sativum*

S. No.	Reagent	Nature of Colour change	Phytochemical changes
1.	Substance + FeCl ₃ <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i>	Blackish brown band is formed	Presence of phenol
2.	The powder is put into the test tube and covered with methanol and conc. HCl (4:1) and stoppered. The tube is allowed to stand with occasional shaking for 4- 5 hours <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Green Colour Greenish yellow No change	Presence of Tannin
3.	Powder + Sudan111 <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Shining orange Colour	Presence of fixed oil and fat
4.	Substance + 10 % NaOH <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Green dark green	Presence of flavonoid
5.	Substance shaken in water <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Frothing absent	Presence of saponin
6.	Substance + Chloroform + drop of acetic acid and then heated + conc. H ₂ SO ₄ <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Green Dark green Deep green	Presence of steroid
7.	Powder + conc. HCl <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Deep green Green Deep green	Presence of quinone
8.	Substance + FeCl ₃ <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Green Brown Black band	Presence of phenols
9.	Powder + Iodine followed by H ₂ SO ₄ <i>A. indica</i> <i>V. negundo</i> <i>C. sativum</i> .	Black Dark black Dark black	Presence of cellulose

V. negundo and *C. sativum*. All the phytochemical tests showed positive results in all the test plants. The phytochemicals screened were cellulose, protein, fat and oil, flavonoids, saponin, steroid, phenol, quinone and tannin. No starch was rated from any of the species of *A. indica*, *V. negundo* and *C. sativum*.

The preliminary phytochemical investigation of selected ethno-medicinal plants of Dindigul district showed the presence of phenolics, flavonoids, terpenoids and alkaloids respectively in 60, 58, 50, and 37 plant species (Karuppusamy *et al.*, 2005).

The phytochemical screening of the polyherbal powder showed the presence of alkaloids, carbohydrates, phytosterol and flavonoids. Saponin was absent in all cases. Phytochemicals like alkaloids, carbohydrates, phytosterol, sterols, tannins, proteins, amino acids, saponins, fixed oils, fats and flavonoids were analysed in *Solanum xanthocarpum* by Udayakumar *et al.* (2003).

pH

The pH of the leaf extracts of *A. indica*, *V. negundo* and *C. sativum* was estimated and found

Maximum as 6.5 in *V. negundo*. The pH 6 showed the acidic nature of the extracts of *A. indica* and *C. sativum* (Table 5).

Table 5: pH of plant extracts

S. No.	Name of the plant	pH
1.	<i>A. indica</i>	6
2.	<i>V. negundo</i>	6.5
3.	<i>C. sativum</i>	6

Biochemical parameters of test plants

Total protein content (table 6)

Biochemical studies on the leaf powders of three plants (*A. indica*, *V. negundo* and *C. sativum*) revealed that the protein content in *V. negundo* has been found to be maximum of 6.08 g per 100 g powder while in *A. indica*, it has been 3.91 g. In the case of *C. sativum* the protein content was minimum, 5.12 mg per 100 g. Only slight variation was observed in the values of protein in water and powder extract. Udayakumar *et al.* (2003) studied the amount of protein present in *Solanum xanthocarpum*.

Table 6: Estimation of total protein, total carbohydrates and phenol

S. No.	Plant name	Protein (mg/100g)	Carbohydrates (mg/100g)	Phenol (mg/100g)
1.	<i>A. indica</i>	3.91	5.02	3.9
2.	<i>V. negundo</i>	6.08	8.32	4.22
3.	<i>C. sativum</i>	5.12	7.13	5.61

Table 7: Estimation of chlorophyll

S. No.	Plant Name	Chlorophyll 'a' (mg/100g)	Chlorophyll 'b' (mg/100g)	Total Chlorophyll (mg/100g)
1.	<i>A. indica</i>	0.09	0.07	0.16
2.	<i>V. negundo</i>	0.12	0.10	0.22
3.	<i>C. sativum</i>	0.08	0.11	0.19

Total carbohydrates content

The values obtained for the carbohydrates content of three test plants ranged between 5.02 mg/100 as the minimum in *A. indica*, 8.32 mg/100g as the maximum in *V. negundo* and 7.13 mg/100g in *C. sativum* (Table 6).

Naseer Banu *et al.* (2003) estimated carbohydrates contents in *Amaranthus viridis* and *Spinacea oleracea* in which, *A. viridis* showed higher carbohydrate content (3.562 mg/g) than *Spinacea oleracea*. Amount of carbohydrates in polyherbal powder and various extracts was found to be less when compared with *A. viridis*.

Phenol content

There were differences in the values obtained for phenol content of three leaf extracts. A maximum of 5.61mg/100 g phenol was estimated from *C. sativum* and a minimum of 3.9mg/100 g from *A. indica* (Table 6).

Amudhan *et al.* (1999) estimated the total phenol profile in some rice varieties in relation to infestation by Asian rice gall or *Seolia oryzae*. The amount of phenol in the polyherbal powder (6.09) was greater than the other extracts.

Chlorophyll content

Estimation of chlorophyll in the leaf powder of *A. indica* showed 0.09 mg of chlorophyll 'a', 0.07 mg of chlorophyll 'b' and 0.16 mg of 'total' chlorophyll per 100 g. While that of *V. negundo* leaf powder

contained 0.12 mg of chlorophyll 'a', 0.10 mg of chlorophyll 'b' and 0.22 mg of 'total' chlorophyll content per 100 g. *C. Sativum* leaf powder contained 0.08 mg, 0.11 mg and 0.19 mg chlorophyll 'a', chlorophyll 'b' and 'total' chlorophyll respectively. (Table 7)

Sims and Gamon (1999) used spectral reflectance for estimation of chlorophyll, anthocyanin, carotenoid concentration, in which the amount of chlorophyll 'a', chlorophyll 'b' and total chlorophyll were calculated. Results of the chlorophyll contents of the above mentioned plants were in accordance with the above findings.

Reducing sugar and total soluble sugar content

The amount of reducing sugar present in all the three plants are shown in Table 8. 100 g of leaf powder of *C. sativum* showed 4.12 g/100 g reducing sugar, while in *V. negundo* the presence of reducing sugar has been 3.14 g/100 g. The content of reducing sugar in *A. indica* has been 1.02 g/100 g. Among the three plants *C. sativum* leaf powder showed higher content of reducing sugar.

The amount of total soluble sugar present in 100 g of *V. negundo* has been 6.9 g while in *A. indica* the presence of total soluble sugar has been 4.2 g/100 g. The content of total soluble sugar in *C. sativum* has been 3.2 g/100 g. Among the three plants *V. negundo* showed higher content of total soluble sugar.

Table 8: Estimation of reducing sugar and total soluble sugar

S. No.	Plant name	Reducing sugar (g/100g)	Total soluble sugar (g/100)
1.	<i>A. indica</i>	1.02	4.2
2.	<i>V. negundo</i>	3.14	6.9
3.	<i>C. sativum</i>	4.02	3.2

The presence of reducing sugar, resins, *etc.* was reported by Wahi *et al.* (1984) in *Aganosma dichotoma* (Roth). Amount of total soluble sugar

present sugar present in extracts and dried powders of *A. vera*, *A. calamus* and *S. racemosa* was observed (Habib *et al.*, 2003).

CONCLUSION

The results organoleptic study offer a scientific basis for the traditional use of *A. indica*, *V. negundo* and *C. sativum* which possess characters like varied green, light green, dark green and pungent odour and bitter taste.

The fluorescence studies revealed that the powders of three test plants showed varied degrees of green and brown colours, when it is exposed to visible and UV rays. As the powders were treated with chemicals like FeCl₃, NaOH, KOH, I, HCl, NaOH

+ H₂O, ethanol, HNO₃ and H₂SO₄, the colour changes were noted in the treated powders and colour varied from green, orange and brown.

The phytochemical screening of the test plants may be attributed to the nature of biological active components. The pH of the test plants were found to be acidic and neutral in nature. Of the powders analyzed, *V. negundo* leaf powder showed the maximum protein content, carbohydrates, chlorophylls and total soluble sugars. *C. sativum* exhibited highest phenol and reducing sugar.

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