Phytochemical analysis of the flowers of Chrysanthemum indicum L. and Calendula officinalis

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**Article History**
- Received on: 09-02-2021
- Revised On: 04-03-2021
- Accepted on: 11-03-2021

**Abstract**
Plant extracts have been used as a source of medicines for a wide variety of human ailments. The phytochemical analysis of flowers of ornamental plants of Chrysanthemum indicum L. and Calendula officinalis are very useful in identifying new sources of therapeutically important compounds like saponins, terpenoids, flavonoids and steroids are present in the flower extract of Chrysanthemum indicum L and carbohydrates, flavonoids, terpenoids are present in the flower extract of Calendula officinalis. Chrysanthemum indicum L : Hindi :Guldaudi, Sanskrit : Bahupatrika, Odiya: Shevathi Calendula officinalis: English : Marigold, Hindi : Genda.

**Keywords**: Phytochemical Analysis, flowers, Chrysanthemum indicum L, Calendula officinalis.

**DOI**: https://doi.org/10.46796/ijpc.vi.148

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**Introduction**
Phytochemical screening refers to the extraction, screening and identification of the medicinally active substances found in plants. Some of the bio-active substances that can be derived from plants are flavonoids, alkaloids, carotenoids, tannin, anti-oxidants and phenolic compounds. Phytochemists study phytochemicals by first extracting and isolating compounds from the origin plant, followed by defining their structure or testing in laboratory model systems, such as cell cultures, in vitro experiments, or in vivo studies using laboratory animals. Challenge in that field include isolating specific compounds and determining their structures, which are often complex, and identifying what specific phytochemical is primarily responsible for any given biological activity. Phytochemicals are chemicals produced by plants. These are the chemicals that plants use to defend themselves against disease [1,2].

Extraction methods involved in this project are Maceration, Percolation, Soxhlet Extraction, Supercritical fluid extraction, Microwave assisted extraction, Ultrasound assisted extraction, Accelerated solvent extraction. The formation of yellow colour indicated the presence of flavonoids while the brown colour formation indicated the presence of alkaloids and terpenoids. The phenol content was maximum in roots (82.13 mg/gdw) followed by seed leave, stem and fruit.
The sugar content was highest in leaves (8.27 mg/gdw) followed by fruits, stem, root, and seed. The protein content was maximum in fruits (55.59 mg/gdw) followed by seed leaves, stem, and root [3].

**Chrysanthemum indicum L.**

**Scientific classification**
Kingdom: Plantae  
Phylum: Tracheophyta  
Class: Angiosperms  
Clade: Asterids  
Order: Asterales  
Family: Astreaceae  
Genus: Chrysanthemum  
Species: C. Indicum L.

**Uses**
- The whole plant is antiphlogistic, blood tonic, depurative, febrifuge, and vulnerary, dissipating heat, detoxifying, and dissipating blood stasis. In conjunction with black pepper it is used in the treatment of gonorrhea and Hypertension.
- The flowers are aperient, bitter, hypotensive, stomachic, and vasodilator.
- They contain the glycoside chrysanthemin that yields glucose and cyanidin on hydrolysis, together with stachydrine and an essential oil.
- They have an antibacterial action, inhibiting the growth of Staphylococcus, E. Coli, streptococcus, C. Diphtheriae, Bacillus dysenteria.
- The flowers are used in the treatment of furuncle, scrofula, deep-rooted boils, inflammation of the throat, eyes, and cervix, eczema, itchiness of the skin.
- They have a rejuvenating effect when used over a long period of time.
- An essential oil obtained from the plant contains chrysanthene, this is active on the brain center affected by Parkinson’s disease.

**Phyto chemistry**
Phytochemicals are biologically active chemical compounds which are derived from plants. They have many health benefits for humans further than those attributed to macronutrients and Chrysanthemum spp. Flower contains octa-cosyl alcohol, β-sitosterol, lupeol, α-amyrin, daucosterol, ineupatorolide B, syringin, chlorogenic acid, petasphenol, physcion, acacetin, eupatilin, quercetin, diosmetin, luteolin, apigenin, apigenin-7-O-β-D-glucopyranoside, quercetin-3-O-β-D-glucopyranoside, luteolin-7-O-β-D-glucopyranoside, apigenin-7-O-β-D-neosperoside, and acacetin-7-O-β-D-glucoside. Most of the Chrysanthemum flowers contain anthocyanins, cyanidin-3-glucoside and cyanidin-3-(3′-malonoyl) glucoside and carotenoids: lutein, zeaxanthin, β-cryptoxanthin, 13-cis-β-carotene, α-carotene, trans-β-carotene, and 9-cis-β-carotene. The major volatile compounds present in the plants are camphor, α-pinene, chrysanthene, safanal, myrcene, eucalyptol.

Chlorogenic acid

**2,4,5,6,7ab-hexahydro-1H-indene, verbenone, β-phellandrene and camphene [3,4].**

**Quercetin**
Calendula officinalis

**Scientific classification**

- **Kingdom**: Plantae
- **Phylum**: Tracheophyta
- **Class**: Angiosperms
- **Clade**: Asterids
- **Order**: Asterales
- **Family**: Astreaceae
- **Genus**: Calendula
- **Species**: C. Officinalis

**Synonyms**

- Calendula aurantiaca kotschy ex Boiss.
- Calendula eriocarpa DC.
- Calendula hydruntina (Fiori) Lanza
- Calendula prolifera Hort. ex Steud.
- Calendula x santamariae Font Quer
- Caltha officinalis (L.) Moench

**Uses**

Calendula is an active ingredient in several diaper creams for babies. It is also used to make natural hand lotions and first-aid creams. It is useful in making a lotion for skin problems like eczema, psoriasis, acne, sunburn, wounds, athletes foot and ringworm. It can add to homemade DIY shampoo and foot soak a yellow dye obtained from the boiled flowers can also be used to add colour to hair.

**Phytochemistry**

Identification of primary and secondary constituents has become the utmost important tool for the presence of active moiety. The phytochemical screening of Petroleum ether, Methanol and Ethanol extracts of calendula officinalis flowers. Petroleum extracts show the presence of fatty acids, ethanol extracts show the presence of triterpenes and sterols. Methanol extract shows saponins, phenolic substances. Various flavonoids have been isolated from the ethanol extract of the flowers of C. officinalis. They include quercetin, isorhamnetin, isoquercetin, isorhamnetin-3-O-β-D-glycoside, narcissin, calendoflaside, calendoflavoside, calendoflavobioside, rutin, isoquercitrin, neohesperidoside, isorhamnetin-3-Oneohesperidoside, isorhamnetin-3-O-2G - rhamnosyl rutinoside, isorhamnetin-3-Orutinoside, quercetin-3-O-glucoside and quercetin-3-O-rutinoside. Coumarins The ethanol extract of the inflorescence of the C. officinalis reported to contain coumarins - scopoletin, umbelliferone and esculetin. Quinones reported from C. officinalis were plastoquinone, phylloquinone, α-tocopherol in the chloroplast, ubiquinone, phylloquinone, α-tocopherol in mitochondria, and phylloquinone in the leaves . Volatile oil C. officinalis flowers contain maximum volatile oil at full flowering stage (0.97 %) and minimum during the preflowering stage (0.13 %) . The composition also showed different patterns at different phases of vegetative cycles. Various monoterpenes and sesquiterpenes have been reported in the volatile oil : α-thujene, sabinene, β-piene, limonene, 1,8-cineol, p-cymene, trans-β-ocimene, γ-terpene, δ-3-carene, nonanal, terpine-4-ol, 3-cylohexene-1-ol, aphellandrene, α-terpeneol, geraniol, carvacrol, bornyl acetate, sabinyl acetate, acubebe, α-copeane, α-bourbonene, βcubebe, α-gurjunene, aromadendrene, βcaryophyllene, α-ylangene, α-humulene, epibicyclo-sequiphellandrene, germacrene D, alloaromadendrene, β-saliene, calarene, muurolene, δ-cadinene, cadina 1,4-diene, αcadinene, nerolidol, palustron, endobourbonene, oplopenone, α-cadinol, Tmuurolol
The essential oil was found to be rich in α-cadinene, α-cadinol, t-muurolol, limonene, and 1,8-cineol with p-cymene at lower levels at the post-flowering period.

**Methodology**

**Methodology of Chrysanthemum indicum L.**

**Collection of plant material**
- The flowers of Chrysanthemum indicum L. were collected from Tagarupuvalasa market, Vizianagaram district, Andhra Pradesh, India in December 2019.

**Preparation of extract**
The collected flowers were washed with distilled water and dried in shade for 4-6 days. They were then coarsely powdered and extracted completely by cold maceration. The solvents used for maceration were petroleum ether, methanol and Ethanol (95%).

**Procedure for Maceration**
By soaking 77.4 g of each samples in 250ml of solvents (petroleum ether, methanol and ethanol) by cold maceration and allow them to stand with intermittent shaking for 3 days. The extracts were filtered using Whatman no 41 filter paper. The filters were collected and made up to known volume and stored in refrigerator at 4°C.

**Qualitative Analysis of Phytochemical constituents**
The flower extracts (ethanol, methanol, petroleum ether) were subjected to preliminary phytochemical studies to qualitatively analyze the active components present in them.

**Phytochemical evaluation**
1. **Test for Alkaloids**
The extracts of flower were warmed separately with 2% H₂SO₄ for 2 minutes. It was filtered and few drops of following reagents were added, which indicated the presence of alkaloids.
   - Dragendroff’s reagent: A red precipitation indicated the positive test.
   - Mayer’s reagent: A creamy white color indicated the positive test.
   - Picric acid: A yellow precipitation indicates the positive test.

2. **Test for Flavonoids**
A small quantity of the extract was heated with 10ml of ethyl acetate in boiling water for 3mins. The mixture was filtered and the filtrates were used for the following tests. The filtrate was shaken with 1ml of dil. Ammonia solution(1%). The layers were allowed to separate. A yellow coloration was observed in ammonia layer indicating the presence of flavonoid. The filtrate was shaken with 1ml of 1% Ammonium Chloride solution, where light yellow colour
was observed. It indicated the presence of flavonoid.

3. **Test for carbohydrates**
The extracts were shaken vigorously with water and filtered. A few drops of molisch’s reagent was added to the aqueous filter, followed by vigorous shaking again. Concentrated H₂SO₄ (1ml) was carefully added to form a layer below the aqueous solution. A brown ring at the interface indicated the positive test.

4. **Test for Saponins**
A small quantity of different extracts was diluted with 4ml of distilled water. The mixture was shaken vigorously and then observed on standing for stable brake, which indicated positive test.

5. **Test for steroids**
2 ml of Acetic anhydride and 2ml H₂SO₄ were added to the extracts. The color changed from violet to blue or green, which indicated the presence of steroids.

6. **Test for Anthraquinone glycosides** *(Borntrager’s test)*
Dil.H₂SO₄ was added to the extracts and boiled. Then filtered and cooled. To the cold filtrate, 3 ml of benzene was added and mixed. The benzene layer was separated and ammonia (2ml) solution was added to it. A rose pink to red color in ammonical layer was observed, which indicated positive test.

7. **Test for cardiac glycosides (Legal’s test)**
To each extract, 1ml of pyridine and 1ml of sodium nitroprusside solution were added and observed. A deep red color was observed indicating the positive test.

8. **Test for terpenoids (salkowski test)**
Each extract was mixed with 2ml of chloroform and then concentrated H₂SO₄ (3ml) was carefully added to form a layer. A reddish brown coloration at the interface indicated positive result for the presence of terpenoids.

9. **Test for gum and mucilages**:
Each extract was dissolved in 10ml of distilled water and 25 ml of absolute alcohol was added to it with constant stirring. White or cloudy precipitate indicated the presence of gum and mucilages.

10. **Test for Amino acids and proteins**
Each extract was dissolved in 10 ml of distilled water and the filtrate was subjected to test the presence of proteins and amino acids.

a) **Biuret Test**
2 ml filtrate was treated with one drop of 2% Copper Sulphate solution and then 1ml of ethanol (95%) was added to it followed by excess potassium hydroxide pellets. Pink color in the Ethanolic layer indicated the presence of proteins.

b) **Ninhydrin Test**
Two drops of ninhydrin solution (10 mg of ninhydrin in 200 ml of acetone) were added to 2ml of aqueous filtrate. A characteristic purple color indicated the presence of amino acids.

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**Results**

*Chrysanthenum indicum L.*

a) Initial weight of dried flowers = 231g
b) Weight of the powdered flowers = 130g
c) Weight of the powder used for the experiment = 77.
d) Weight of the petroleum ether extract = 1.1g
e) Weight of the methanol extract = 3.6g
f) Weight of ethanol extract 2.9
Phytochemical screening results

<table>
<thead>
<tr>
<th>S.no</th>
<th>Phytochemical constituents</th>
<th>Petroleum ether</th>
<th>Methanol</th>
<th>Ethanol</th>
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<tbody>
<tr>
<td>1</td>
<td>Carbohydrates</td>
<td>-</td>
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<tr>
<td>2</td>
<td>Saponins</td>
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<td>+</td>
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<tr>
<td>3</td>
<td>Alkaloids</td>
<td>-</td>
<td>+</td>
<td>+</td>
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<tr>
<td>4</td>
<td>Cardiac glycosides</td>
<td>-</td>
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<td>+</td>
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<tr>
<td>5</td>
<td>Amino acids and proteins</td>
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<tr>
<td>6</td>
<td>Gums and mucilage</td>
<td>-</td>
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<tr>
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<td>8</td>
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<tr>
<td>9</td>
<td>Steroids</td>
<td>+</td>
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<td>+</td>
</tr>
</tbody>
</table>

‘+’ indicates presence ‘−’ indicates absence

Methodology of calendula officinalis
Collection of plant material
The flowers of Calendula officinalis were collected from Brahmapur, Ganjam district, Odisha, India in November 2020.

Preparation of extract
The collected flowers are washed with distilled water and dried in shade for 4-6 days. They were then coarsely powdered and extracted completely by cold maceration. The solvents used for maceration were petroleum ether, methanol and Ethanol(95%).

Procedure for Maceration of Candula Officinalis
By soaking 133.3 g of each samples in 400ml of solvents (petroleum ether, methanol and ethanol) by cold maceration and allow them to stand with intermittent shaking for 3 days. The extracts were filtered using Whatman no 41 filter paper. The filters were collected and made up to known volume and stored in refrigerator at 4°C. Qualitative Analysis of Phytochemical constituents : The flower extracts (petroleum ether, methanol and ethanol ) were subjected to preliminary phytochemical studies to qualitatively analyze the active components present in them.

Results
Calendula officinalis
- Initial weight of dried flowers
- Weight of the powdered flowers
- Weight of the powder used for the experiment = 133.3g
- Weight of the petroleum ether extract = 0.9g
- Weight of the methanol extract
- Weight of ethanol extract

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Discussion
The medicinal value of these plants lies in some chemical substances that have a definite physiological action on body [1,2]. The most important of these bioactive constituents of the plants are alkaloids, terpenoids flavonoids, steroids, cardiac glycosides and protein compounds. Results showed that the compounds like saponins, terpenoids, flavonoids and steroids are present in the flower extract of Chrysanthemum indicum L. Results showed that the compounds like carbohydrates, flavonoids, terpenoids
are present in the flower extract of Calendula officinalis [4-6].

**Summary**
The thesis consists of phytochemical investigation of some potent indigenous medicinal plants of traditional use namely Chrysanthemum indicum L. and Calendula officinalis (flowers).

**Conclusion**
The plants studied here can be seen as a potential source of useful drugs. It also justifies the folklore medicinal uses and claims about the therapeutic values of these plants as curative agent. We therefore, suggests further the isolation, purification and characterization of the bioactive compounds of the flower part of Chrysanthemum indicum L. and Calendula officinalis with a view to obtain some useful chemotherapeutic agents.

**Author Contribution**
All authors Contributed Equally.

**Conflict of Interest**
The authors declares no conflict of Interest.

**Funding**
No Funding

**References**
7. Suma Tagadur Suresh Chandra, Vijay Barve vernacular names of chrysanthemum, FRLHT’s ENVIS centre on medicinal plants, Bangalore 2016.
8. Jhingnam Girija- vernacular names of calendula officinalis, botany, the botanical dictionary.
10. National Chrysanthemum Society, USA.
15. Arana, Lide; Salado, Clarisa; Vera, Sandra; Tatiana; Usobiaga, Aresatz; Arrondo, Jose Luis; Alonso, Alicia (2015-11-01).