Comparative study of High-performance thin-layer chromatography and Antioxidant potential of Hydrocotyle asiatica mother tincture used in Homoeopathy

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Abstract

Background: Hydrocotyle asiatica has a therapeutic significance in the Indian system of medicine due to its rich antioxidant activity. In Homoeopathy, Hydrocotyle asiatica is used for the treatment of jaundice, skin diseases, dropsy, elephantiasis, leprosy, gonorrhea, leucorrhea, and nervous debility. It contains the abundant triterpene glycoside Asiatic acid which shows cytotoxic activity on cancer cells. Its homoeopathic mother tincture is a major source of antioxidant compounds, which is responsible for its overall pharmacological activity.

Objectives: This study was done to evaluate antioxidant activity and High-Performance Thin-Layer Chromatography study of Hydrocotyle asiatica in-house homoeopathic mother tincture and market samples.

Materials and Methods: Antioxidant activity of in-house homoeopathic mother tincture (A) and three market samples (B, C, and D) were determined by 2, 2-diphenyl-1-picryl-hydrazyl-hydrate (DPPH) free-radical scavenging activity, total phenol, and 2,2'-azino-bis 3-ethylbenzothiazoline-6-sulfonic acid (ABTS) assay methods. High-performance thin-layer chromatography (HPTLC) study performed on precoated silica gel 60 F254 TLC plate, mobile phase used was toluene: ethyl acetate: formic acid (5.5:4.5:1, v/v/v) and UV detection were performed at 254 and 366 nm. For derivatization, an anisaldehyde sulfuric acid reagent was used.

Results: The homoeopathic mother tincture of Hydrocotyle asiatica had prominent antioxidant activity. HPTLC study indicated the presence of triterpene glycoside compound Asiatic acid in chloroform extract of Hydrocotyle asiatica.

Conclusion: The mother tinctures prepared by authenticated plant samples showed maximum active constituents and prominent antioxidant activity as compared to the mother tinctures procured from the market. The present study justifies the homoeopathic usage of Hydrocotyle asiatica and highlights its healing properties.

Acknowledgments and Source of Funding

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Comparative study of high-performance thin-layer chromatography and antioxidant potential of Hydrocotyle asiatica mother tincture used in homoeopathy

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Abstract

Background: Hydrocotyle asiatica has a therapeutic significance in the Indian system of medicine due to its rich antioxidant activity. In Homoeopathy, Hydrocotyle asiatica is used for the treatment of jaundice, skin diseases, dropsy, elephantiasis, leprosy, gonorrhea, leucorrhea, and nervous debility. It contains the abundant triterpene glycoside Asiatic acid which shows cytotoxic activity on cancer cells. Its homoeopathic mother tincture is a major source of antioxidant compounds, which is responsible for its overall pharmacological activity. Objectives: This study was done to evaluate antioxidant activity and High-Performance Thin-Layer Chromatography study of Hydrocotyle asiatica in-house homoeopathic mother tincture and market samples. Materials and Methods: Antioxidant activity of in-house homoeopathic mother tincture (A) and three market samples (B, C, and D) were determined by 2, 2-diphenyl-1-picryl-hydrayl-hydrate (DPPH) free-radical scavenging activity, total phenol, and 2,2’-azino-bis 3-ethylbenzothiazoline-6-sulfonic acid (ABTS) assay methods. High-performance thin-layer chromatography (HPTLC) study performed on precoated silica gel F 254 TLC plate, mobile phase used was toluene: ethyl acetate: formic acid (5:5:4.5:1, v/v/v) and UV detection were performed at 254 and 366 nm. For derivatization, an anisaldehyde sulfuric acid reagent was used. Results: The homoeopathic mother tincture of Hydrocotyle asiatica had prominent antioxidant activity. HPTLC study indicated the presence of triterpene glycoside compound Asiatic acid in chloroform extract of Hydrocotyle asiatica. Conclusion: The mother tinctures prepared by authenticated plant samples showed maximum active constituents and prominent antioxidant activity as compared to the mother tinctures procured from the market. The present study justifies the homoeopathic usage of Hydrocotyle asiatica and highlights its healing properties.

Keywords: 2,2’-Azino-bis 3-ethylbenzothiazoline-6-sulfonic, 2,2-Diphenyl-1-picryl-hydrayl-hydrate, Antioxidant activity, Homoeopathic mother tincture, High-performance thin-layer chromatography, Hydrocotyle asiatica

Introduction

Hydrocotyle asiatica (Centella asiatica) belongs to the family of perennial plants Umbelliferae (Apiaceae).[1] It is a tasteless, odorless plant that thrives in and around water [Figure 1].[2] Hydrocotyle asiatica is a very important medicinal herb used in different orient and is becoming a popular medicine in the west.[3,4] Hydrocotyle asiatica is commonly known as brahmi in Hindi, mundukparni, kodavanin Ayurveda, Gotu kola in China and Sri Lanka, buakbok in Thailand, kaki kuda in Indonesia, and yuhong-yuhong in the Philippines.[5-7] In Chinese medicine, Hydrocotyle asiatica is known as one of the ‘miracle elixirs of life’ known over 2000 years ago.[8] It is native to wetlands in Asia such as India, Sri Lanka, China, Indonesia, Malaysia, South Africa, and Madagascar.[9] Its active constituents include pentacyclic triterpene derivatives.[10] Hydrocotyle asiatica is an important medicinal plant that is widely used as a homoeopathic medicine due to its bioactive compounds such as Asiatic acid, rutin, kaempferol, quercetin, gallic acid, luteolin, and catechin.[11,12] It has been used in folk herbal medicine for centuries for memory enhancement, depression, wound healing, psoriasis, and the treatment of related chronic diseases. This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

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Percentage loss on drying was 30.5. In view of that, we designed the study to investigate the presence of Asiatic acid in the in-house mother tincture sample (A) and three market samples (B, C, and D) of Hydrocotyle asiatica by HPTLC method. Furthermore, the whole plant of Hydrocotyle asiatica is a rich source of antioxidant compounds. Antioxidant components are micro constituents that inhibit lipid oxidation by inhibiting the initiation or propagation of oxidizing chain reactions and are involved in the scavenging of free radicals. In view of that, we designed the study to evaluate the antioxidant potential of Hydrocotyle asiatica by various assay methods. The present study is helpful for the determination and quantification of antioxidant compounds which are useful for producing Hydrocotyle asiatica-based drugs for the treatment of various ailments of human beings.

**Methods**

**Collection of plant materials**

The whole plant of the specimen Hydrocotyle asiatica was collected and authenticated by staff at the Center of Medicinal Plants Research in Homoeopathy, Tamil Nadu. The voucher specimen was deposited in the herbarium and in the laboratory of Dr. D. P. Rastogi Central Research Institute (Homoeopathy) Noida, Uttar Pradesh, India, for future reference with collection number 9647. The whole plant was shade-dried and powdered mechanically, and the fine powder was used for the preparation of the mother tincture. Asiatica acid (C_{39}H_{48}O_{5} M. P. 325–330°C) with purity by high performance liquid chromatography (HPLC) >98% w/w purchased from Sigma Aldrich, USA. Solvents used were ethanol, methanol, HPLC water, and chloroform of analytical grade purity (Merck Ltd., India).

**Physicochemical studies for raw drug standardization**

**Loss on drying**

Loss on drying method was used for determination of moisture content as per methods recommended in Homoeopathic Pharmacopoeia of India. Percentage loss on drying was calculated.

**Foreign matter determination**

For foreign matter determination, 100 g of plant raw material was taken and outspread in the form of a thin layer. The sample was examined by a 6× lens or with an unaided eye, the foreign organic matter was picked manually. The ratio of total foreign matter weighed, and the weight of drug taken gave the % of foreign matter.

**Total ash value determination**

In the drug, the impurity present in the form of organic matter was determined with the help of the total ash value. For its determination, 2 g of the dried raw drug was weighed in powdered form in a pre-weighed silica crucible. The sample was incinerated in a silica crucible by gradually increasing the temperature up to 450°C for 4 h or until it became carbon-free. The crucible was cooled and weighed until a constant weight was obtained. Percent of total ash value was then calculated by taking the ratio of loss in weight to the weight of the sample taken.

**Acid-insoluble ash value determination**

After total ash value determination, 25 mL of 5 M hydrochloric acid was added to the dried ash and boiled in a water bath for 10 min. The solution was concentrated till its color changed to yellow. Acid insoluble matter was filtered using ashless Whatman paper number 1 followed by washing with distilled water. The paper was again ignited in a crucible at a temperature not more than 450°C for 4 h, after which the crucible was kept in a desiccator, cooled, and weighed. With reference to the originally taken air-dried powdered drug, the percentage of acid-insoluble ash value was calculated.

**Water-soluble extractive value determination**

For determination of water extractive value, 2 g of sample was accurately weighed, and air-dried powdered drug was put in a conical flask with 100 mL water added to it. The
solution was allowed to stand for 24 h with intermittent shaking of the flask after every 4 h. The water-soluble extractive was filtered using Whatman filter paper. 25 mL of this filtrate was completely dried on a pre-weighed Petri plate at 105°C. The increase in weight of the petri dish was noted to calculate the water-soluble extractive value determination. With reference to the originally taken air-dried powdered drug, the percentage of water-soluble extractive value was calculated.

**Alcohol-soluble extractive value determination**

For determination of alcohol soluble extractive value, accurately weighed 2 g air-dried powdered drug was put in a conical flask, and 100 mL absolute alcohol was added to it. The whole solution was left for 24 h for complete extraction at room temperature. The solution was shaken vigorously for a few minutes after every 6 h. The extract was filtered with the help of Whatman filter paper taking precautions to avoid evaporation loss of alcohol from the extract weighed the empty flat-bottomed Petri dish. The Petri dish with 25 mL of filtrate was heated at 105°C in an electric oven then cooled in a desiccator and weighed. With reference to originally taken air-dried powdered, the percentage of alcohol-soluble extractive value was calculated and is shown in Table 1.

**Preparation of crude extract/in-house mother tinctures**

100 g of coarsely dried powdered *Hydrocotyle asiatica* whole plant was taken in which 300 mL distilled water and 730 mL strong alcohol (95%) were added to make one thousand milliliters of the mother tincture using the percolation method[27] (as per Homoeopathic Pharmacopoeia of India). This tincture was transferred to a tightly packed amber glass container and stored for further study.

**Qualitative phytochemical screening**

Phytochemical tests were performed on crude extract for qualitative estimation of phytochemicals present in in-house mother tincture of *Hydrocotyle asiatica* with all respective testing procedures as described in the textbook by Harborne [Table 2].[28]

**Standardization of mother tincture**

Standardization of mother tincture was conducted to identify the organoleptic and physicochemical properties of mother tincture. Organoleptic properties measurement was done for color, odor, and clarity of solution. The samples were tested for various physicochemical properties such as sediments, pH, total solids, weight/mL, and total alcohol content.

**Preparation of standard Asiatic acid**

Dissolved 5 mg of Asiatic acid in 5 mL ethanol in a volumetric flask and sonicated for 10 min to prepare a working standard of asiatic acid with a concentration of 1 mg/mL.

**Preparation of chloroform extract**

25 mL of mother tincture was taken in a 50 mL beaker. The solution was evaporated on a water bath to remove the ethanol and extracted three times with 20 mL chloroform. Combined and concentrated chloroform was extracted up to 2 mL volume. TLC of chloroform extract of mother tincture was carried out and standard asiatic acid on silica gel 60 F254 pre-coated plate was referenced.

**HPTLC fingerprinting profile study**

For HPTLC fingerprinting study a densitometric HPTLC Camag Linomat 5 (Switzerland) system was used.[31] As a sample applicator, Camag Linomat 5 was used for spotting TLC plate. Spots were made on silica gel 60 F254 pre-coated plate (Merek) 20 × 10 cm plate with an aid of a sampling machine and the solvent front was run up to 70 mm height. For the development of the mobile phase, a saturating chamber Camag Twin Trough glass chamber was used. Camag TLC Scanner and software vision CATS were used for scanning purposes. HPLC grade solvents were used for all the extracts solutions. Volume applied for standard 2–6 µL and for sample 2–6 µL. Asiatic acid was used as a reference standard. For detection of triterpene glycoside Asiatic acid various mobile phase was used toluene: chloroform:ethanol (4:4:1, v/v/v), chloroform: methanol:formic acid (7:3:0.5, v/v/v) and chloroform: methanol (9:1, v/v), toluene: ethyl acetate: formic

### Table 1: Results of test of physicochemical properties of raw drug material

<table>
<thead>
<tr>
<th>Name of test</th>
<th>Result (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Foreign matter</td>
<td>2.00</td>
</tr>
<tr>
<td>Loss on drying</td>
<td>4.98</td>
</tr>
<tr>
<td>Total ash value</td>
<td>6.80</td>
</tr>
<tr>
<td>Acid-insoluble ash value</td>
<td>1.20</td>
</tr>
<tr>
<td>Water-soluble extractive value</td>
<td>26.82</td>
</tr>
<tr>
<td>Alcohol-soluble Extractive value</td>
<td>10.40</td>
</tr>
</tbody>
</table>

### Table 2: Results of Phytochemical tests for screening of various phytochemicals present in mother tincture of *Hydrocotyle asiatica*

<table>
<thead>
<tr>
<th>Name of phytochemical tests</th>
<th>Procedure</th>
<th>Observation</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flavonoid (Lead acetate)</td>
<td>Mother tincture with a few drops of 10% lead acetate solution</td>
<td>Yellow precipitate</td>
<td>Positive</td>
</tr>
<tr>
<td>Alkaloids (Dragendorff’s test)</td>
<td>In chloroform solution add dilute hydrochloric acid with a few drops of Dragendorff’s reagent</td>
<td>Red-brown precipitate</td>
<td>Positive</td>
</tr>
<tr>
<td>Glycosides (Sodium hydroxide reagent test)</td>
<td>Mother tincture with 1 mL water and sodium hydroxide solution</td>
<td>Yellow color</td>
<td>Positive</td>
</tr>
<tr>
<td>Tannin (Ferric chloride test)</td>
<td>Mother tincture mixed with few drops of ferric chloride solution</td>
<td>Green color</td>
<td>Positive</td>
</tr>
<tr>
<td>Saponins (Foam test)</td>
<td>Mother tincture shaken with little quantity of water</td>
<td>Foam produced</td>
<td>Positive</td>
</tr>
</tbody>
</table>
acid (5:5:1, v/v/v). TLC spots were visualized after illumination at 254 nm, 366 nm, and after derivatization.

**Study of antioxidant potential**

**Determination of total phenolic content (TPC)**

The TPC of the extracts was determined by Folin-Ciocalteu’s reagent procedure reported by Singleton.[30] The total phenol content was estimated in *Hydrocotyle asiatica* in-house mother tincture sample A and its market samples (B, C and D). Ascorbic acid was used as the reference standard. Different concentrations (0.2661-8.517 mM) of ascorbic acid were prepared and analyzed at 736 nm and a calibration curve was plotted as absorbance versus concentration. TPC was estimated by using Ascorbic acid as standard approximately 50 μL of the mother tincture was mixed with 5 mL of 10% Folin-Ciocalteu’s (phenol reagent) and 4 mL of sodium carbonate. The mixture was allowed to stand for 1 h in dark. After 1 h the color changed from yellow to blue. The absorbance of the solutions was measured at $\lambda_{max}$ 736 nm using a UV-VIS spectrophotometer (U.V. Spectrophotometer SPECORD 200 plus Analytik Jena, Germany). The TPC was calculated from the calibration curve and in a result TPC of the *Hydrocotyle asiatica* sample A (in-house mother tincture) and market sample B, C, and D were calculated as the ascorbic acid equivalents (AAEs) using ascorbic acid as standard ($Y=0.0753x$ $+0.0256, R^2=0.9995$). TPC was expressed in mM concentration of ascorbic acid equivalent.

**2.2-Diphenyl-1-picryl-hydrazyl-hydrate (DPPH) radical scavenging assay**

The free radical scavenging activity of *Hydrocotyle asiatica* sample in-house mother tincture (A) and market sample (B, C, and D) was measured by DPPH radical scavenging assay. The standard solution of DPPH was prepared by dissolving 0.025 g in 25 mL methanol and different concentrations of standards/mother tincture sample (100 μL) were mixed with 4 mL methanol and 1 mL of DPPH standard. The mixture was allowed to stand for 1 h in dark, after which the absorbance was measured at 517 nm using a UV-VIS spectrophotometer (UV Spectrophotometer SPECORD 200 plus Analytik Jena, Germany). The percentage inhibition was determined by comparing the result of the test and the control (methanol used as solvent blank).[31] Percentage degradation was calculated by the formula:

$$DPPH \text{ radical scavenging } (\%) = \left( \frac{B - A}{B} \right) \times 100 + B$$

Where,

A=Absorbance of the sample

B=Absorbance of control

The inhibiting effects of the mother tincture showed varied levels of DPPH radical scavenging activity, expressed as percentage degradation.

**Determination of 2,2'-azino-bis 3-ethylbenzothiazoline-6-sulfonic (ABTS) assay**

Free radical scavenging activity of in-house mother tincture sample (A) and market samples (B, C, and D) were determined by ABTS radical cation decolorization assay. ABTS$^+$ cation radical was produced by the reaction between 7 mM ABTS in water and 2.45 mM potassium persulfate (1:1) stored in the dark at room temperature for 16 h before use. ABTS$^+$ solution was then diluted with methanol to obtain an absorbance of 0.700 at 746 nm. After the addition of 10 μL of mother tincture/standard in 2 mL of diluted ABTS$^+$ solution, the absorbance was measured at 5 min after the initial mixing. An appropriate solvent blank (methanol) was run in each assay.

Percent inhibition of absorbance at 746 nm was calculated using the formula:

$$ABTS \text{ ion scavenging effect } (\%) = \left[ \frac{(AB - AA) \times 100}{AB} \right] + AB$$

Where,

AB is the absorbance of ABTS radical + methanol

AA is the absorbance of the ABTS radical+ sample/standard.

Trolox was used as a standard substance.

**Results**

**Results of physicochemical and phytochemical studies**

The physicochemical properties of the tinctures of in-house drug sample (A) for parameters like sediments, pH, total solids, alcohol content, and weight per mL were analyzed and tabulated in Table 3. The results obtained for various physicochemical studies of raw drug are tabulated in Table 1. Phytochemical tests performed on the crude extract of the whole plant of *Hydrocotyle asiatica* showed positive results for various tests as mentioned in Table 2. Organoleptic observations of the prepared in-house mother tincture indicated the formation of a clear green solution with characteristic tincture odors.

**Result of HPTLC study**

Based on extensive literature reviews, various combinations of solvent systems were studied with an aim to have an appropriate mobile phase composition for the best and most efficient HPTLC chromatographic separation of Asiatic acid in *Hydrocotyle asiatica* chloroform extract. In the mobile phase toluene: chloroform: methanol:formic acid (7:3:0.5, v/v/v), and chloroform: methanol (9:1, v/v) no appropriate resolution of the band was observed, whereas in mobile phase toluene: ethyl acetate: formic acid (5:5:1, v/v/v) efficient band resolution

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Parameter</th>
<th>In-house sample A</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Sediments</td>
<td>Nil</td>
</tr>
<tr>
<td>2.</td>
<td>pH</td>
<td>5.90</td>
</tr>
<tr>
<td>3.</td>
<td>Total solid</td>
<td>1.05% w/v</td>
</tr>
<tr>
<td>4.</td>
<td>wt/mL</td>
<td>0.90 g</td>
</tr>
<tr>
<td>5.</td>
<td>Alcohol content</td>
<td>64.0% v/v</td>
</tr>
<tr>
<td>6.</td>
<td>$\lambda_{max}$</td>
<td>331 nm</td>
</tr>
</tbody>
</table>
of Asiatic acid was observed with improved $R_f$ value of 0.49. Among all the mobile phase combinations studied, toluene: ethyl acetate: formic acid (5:5:1, v/v/v) was finalized to be the ideal one for the evaluation of compound Asiatic acid in *Hydrocotyle asiatica*. Thus, it was finalized the best appropriate mobile phase composition for the entire HPTLC method development study. Table 4 recorded various mobile phase combinations used for the preliminary screening study for the best possible separation of bands.

**Qualitative HPTLC study of in-house mother tincture and market samples**

HPTLC study of *Hydrocotyle asiatica* chloroform extract of the in-house sample (A), three market samples (B, C, and D), and standard Asiatic acid was carried out using selected mobile phase toluene: ethyl acetate: formic acid in the ratio of volume (5:5:1, v/v/v). At UV light 254 nm and 366 nm, no spots of Asiatic acid were observed for any of the samples [Figures 2 and 3]. Therefore, for better resolution, an anisaldehyde-sulfuric acid reagent was used as derivatizing agent. After derivatizing the plate with anisaldehyde-sulfuric acid reagent, a blue spot of Asiatic acid was observed at $R_f$ 0.49 [Figure 4] in in-house sample (A) and in the market sample (B, C, and D). 3D diagram of HPTLC dendrogram displayed the presence of standard Asiatic acid in mother tincture of in-house sample A and market samples B, C, and D displayed in [Figure 5], respectively.

**Result of antioxidant activity**

In the present study, the TPC of *Hydrocotyle asiatica* in-house sample A and its market sample B, C, and D were determined by Folin–Ciocalteu method and reported as AAE. The study reveals TPC found in *Hydrocotyle asiatica* in-house sample A, market samples B, C, and D (75μL) was 12.20, 2.80, 10.68, and 3.59 AAE [Table 5].

In the present study, the DPPH assay of *Hydrocotyle asiatica* in-house sample A and market samples B, C, and D were determined by DPPH radical scavenging assay method and reported as AAE. The study reveals *Hydrocotyle asiatica* in-house sample A and market samples B, C, and D were able to decolorize DPPH free radical, the DPPH scavenging increased with the concentration of the extract. The result showed a greater rate of DPPH scavenging activity found in the in-house sample as compared to the market samples. The percentage of inhibition found in 100 μL volume of

| Table 4: Comparison of various mobile phase combinations used for preliminary screening study for best possible chromatographic separations of Asiatic acid |
|-----------------|-----------------|-----------------|
| Used mobile phase combinations for evaluation of asiatic acid | $R_f$ value | Observations |
| Toluene: chloroform: ethanol (4:4:1, v/v/v) | 0.40 | Poor resolution of band |
| Chloroform: methanol: formic acid (7:3:0.5, v/v/v) | 0.10 | No appropriate resolution of band |
| Chloroform: methanol (9:1, v/v/v) | 0.23 | No appropriate resolution of band |
| Toluene: ethyl acetate: formic acid (5:5:1, v/v/v) | 0.49 for Asiatic acid | Efficient band resolution with improved $R_f$ |

| Table 5: Result of total phenolic content in mother tincture of *Hydrocotyle asiatica* in-house sample A and market samples B, C, and D |
|-----------------|-----------------|-----------------|
| S. No. | Sample | Concentration in (mM) of AAE | Absorbance |
| 1. | *Hydrocotyle asiatica* in-house sample A | 12.20 | 0.9439 |
| 2. | *Hydrocotyle asiatica* market sample B | 2.80 | 0.2366 |
| 3. | *Hydrocotyle asiatica* market sample C | 10.68 | 0.8298 |
| 4. | *Hydrocotyle asiatica* market sample D | 3.59 | 0.2957 |
| 5. | Control | 0.11 | 0.0172 |

AAE: Ascorbic acid equivalents

*Hydrocotyle asiatica* in-house sample A and market samples B, C, and D were 88.55%, 38.74%, 40.43%, and 24.10%, respectively [Table 6]. The order of DPPH scavenging against *Hydrocotyle asiatica* in-house sample A and market samples B, C, and D was found to be sample A > sample C > sample B > sample D.

In the DPPH assay, a significant correlation coefficient ($R$, 0.9955) was found between the antioxidant activity of alcoholic extracts (mother tinctures) of *Hydrocotyle asiatica* in-house sample A and market samples B, C, and D. The hydrogen radical scavenging action is known to be one of the important mechanisms for measuring antioxidant activity. This assay determines the scavenging of stable radical species DPPH by antioxidants compounds present in the mother tincture. The results showed a greater rate of DPPH scavenging activity in in-house sample A as compared to market samples B, C, and D probably due to the presence of high content of phenolic compound. Our study clearly indicated that the mother tincture of in-house sample A of *Hydrocotyle asiatica* exhibited high content of phenolic compound, that is, 12.20 mM which was significantly correlated with DPPH radical scavenging activity %, that is, 88.55% [Tables 5 and 6].

In the ABTS’ assay of *Hydrocotyle asiatica*, in-house sample (A) and market sample (B, C, and D) were determined by ABTS’ assay method and reported in terms of Trolox equivalents. A significant correlation coefficient ($R$, 0.9901) was found between the antioxidant activity of alcoholic extracts (mother tinctures) of *Hydrocotyle asiatica* in-house sample
Figure 2: High-performance thin layer chromatography fingerprints of *Hydrocotyle asiatica* at ultraviolet 254 nm. Standard Asiatic acid Track (1-3), Track (4-6) In-house sample A color rendering index (h), Track (7-9) commercial market sample B, Track (10-12) market sample C, Track (13-15) market sample D.

Figure 3: High-performance thin-layer chromatography fingerprints of *Hydrocotyle asiatica* at ultraviolet 366 nm. Standard Asiatic acid Track (1-3), Track (4-6) In-house sample A color rendering index (h), Track (7-9) commercial market sample B, Track (10-12) market sample C, Track (13-15) market sample D.

Figure 4: High-performance thin layer chromatography fingerprints of *Hydrocotyle asiatica* after derivatization with anisaldehyde sulfuric acid reagent viewed in white light. Standard Asiatic acid Track (1-3), Track (4-6) In-house sample A color rendering index (h), Track (7-9) commercial market sample B, Track (10-12) market sample C, Track (13-15) market sample D.
(A) and market sample (B, C, and D). The study reveals Hydrocotyle asiatica in-house sample (A) and market sample (B, C, and D) were able to decolorize ABTS⁺ free radical, the ABTS radical cation scavenging activity increased with the concentration of the extract. The result showed the greater rate of ABTS cation scavenging activity found in Hydrocotyle asiatica in-house sample A as compared to the market sample (B, C, and D). The percentage of inhibition found in 10 µL volume of Hydrocotyle asiatica in-house sample A and market sample (B, C, and D) were 99.89%, 99.40%, 99.82%, and 98.27%, respectively [Table 7]. The order of ABTS radical cation scavenging activity against Hydrocotyle asiatica was found to be in the order sample A > sample C > sample B > sample D.

**Discussion**

Several factors relating to climate, altitude, rainfall, and other conditions are responsible for the growth of the plant Hydrocotyle asiatica which affects the concentration of bioactive constituents grown in the same country. These conditions may produce significant variations in the bioactive constituents and thus cause a variation in the therapeutic efficacy. The variations in the different sample results may change the efficacy as the concentration of bioactive compounds is different in different samples analysed by us through HPTLC and UV spectrophotometer. The whole plant of Hydrocotyle asiatica is a natural source of important antioxidant substances. The results of various physicochemical studies of raw drugs tabulated in Table 1 indicate higher water-soluble extractive values as compared to alcohol-soluble extractive values. Since water is more polar than alcohol; therefore, all the polar compounds present in the plant are soluble in water. As a result, we found a greater extractive value in water as compared to alcohol. Whereas in the antioxidant study, high contents of phenolic compounds and a significant linear correlation between the values of the concentration of phenolic compounds and antioxidant activity indicate that these compounds contribute to the strong antioxidant activity of Hydrocotyle asiatica. Antioxidants are used to prevent aging, diabetes, heart diseases, cancer, and many other illnesses. The strong potential of tested Hydrocotyle asiatica mother tinctures as antioxidants in the present study suggests that the effect of Hydrocotyle asiatica mother tincture in the treatment of various diseases may be due to this whole plant antioxidant activity.

The present study revealed that as part of the preformulation study, the alcoholic extracts i.e., mother tincture of the whole plant of Hydrocotyle asiatica showed promising physicochemical characteristics. The result of the HPTLC fingerprinting profile study confirms the presence of triterpene glycoside Asiatic acid in chloroform extract of Hydrocotyle asiatica homoeopathic in-house mother tincture (A) as well as in commercial market samples (B, C, and D) at R_f = 0.49.
The antioxidant activity of *Hydrocotyle asiatica* in-house mother tincture and the commercial market sample was investigated by different antioxidant assay methods such as total phenol, DPPH, and ABTS assay. Results show that the antioxidant potential of *Hydrocotyle asiatica* mother tincture demonstrated the highest antioxidant activity found in in-house mother tincture sample as compared to the market samples. The present study demonstrates that the studied homeopathic mother tincture of *Hydrocotyle asiatica* has a significant concentration of polyphenols. The high polyphenol content correlated with the significant antioxidant activity and can be the explanation for the beneficial effect of *Hydrocotyle asiatica* mother tincture in homeopathic treatments.

**CONCLUSION**

The present study may push forth further research work to increase the usefulness of the plant *Hydrocotyle asiatica* in alternative systems of medicine. Quantitative estimation of other compounds present in this plant may also be evaluated in future studies responsible for its other pharmacological activities. Further, it is highlighted that the mother tinctures prepared by authenticated plant samples bear better quantity and quality of active constituents and thus displayed more prominent antioxidant activity, in this case.

**REFERENCES**

19. Kumar A, Prakash A, Dogra S. *Centella asiatica* attenuates D-galactose-induced...
Das Ergebnis zeigte, dass der antioxidative Test der homöopathischen Urtinktur der antioxidative Aktivität der hauseigenen homöopathischen Urtinktur zuständig ist.

Zielsetzungen: Ziel der Studie war es, die antioxidative Aktivität und die Hochleistungs-Dünnschichtchromatographie (HPTLC) der teinture mère homöopathique interne d’Hydrocotyle asiatica und des échantillons des marchés.


Vergleichende Studien zur Hochleistungs-Dünnschichtchromatographie und zum antioxidativen Potenzial der in der Homöopathie verwendeten Urtinktur von Hydrocotyle asiatica

Estudios comparativos de cromatografía en capa fina de alto rendimiento y potencial antioxidante de la tintura madre de Hydrocotyle asiatica utilizada en homeopatía

Antecedentes: Hydrocotyle asiatica tiene un gran significado terapéutico en el sistema indio de la medicina debido a su rica actividad antioxidante. En homeopatía, Hydrocotyle asiatica se utiliza para el tratamiento de la ictericia, enfermedades de la piel, hidropesía, elefantiasis, lepra, gonorrhea, leucorrea y debilidad nerviosa. Contiene el glucósido triterpeno más abundante ácido asiático que muestra actividad citotóxica en las células cancerosas. La tintura madre homeopática de este fármaco es una fuente importante de compuestos antioxidantes, que es responsable de su actividad farmacológica general. Objetivos: El objetivo del estudio fue evaluar la actividad antioxidante y el estudio de cromatografía en capa fina de alto rendimiento (HPTLC) de tintura madre homeopática y muestras de mercado de Hydrocotyle asiatica. Materiales y Métodos: La actividad antioxidante de la tintura madre homeopática interna (A) y de tres muestras de mercado (B, C y D) se determinó mediante métodos de ensayo de 2,2-difenil-1-picril-hidrazil-actividad de depuración de radicales libres, fenol total y 2,2-azino-bis-(3-etilbenzotiazolina-6-sulfonato. Estudio realizado con HPTLC en placa de TLC de gel de sílice 60 F254 prerecubierta, la fase móvil utilizada fue tolueno:acetato de etilo:ácido fórmico (5,5:4,5:1, v/v/v) y detección UV a 254 y 366 nm. Para la derivación se utilizó un reactivo de ácido sulfúrico anisaldehído. Resultados: El resultado reveló que el análisis antioxidante de la tintura madre homeopática de Hydrocotyle asiatica tenía una actividad antioxidante prominente. HPTLC研究表明三萜苷化合物亚洲酸在Hydrocotyle asiatica的氯仿提取物中存在. Conclusión: Las tinturas madre preparadas por muestras vegetales autenticadas mostraron constituyentes activos máximos y actividad antioxidante prominente en comparación con las tinturas madre obtenidas del mercado. El presente estudio justifica el uso homeopático de Hydrocotyle asiatica y destaca sus propiedades curativas.

[医]亚洲水样母町用于顺势疗法的高效薄层色谱与抗氧化潜力的比较研究

背景资料: [医]亚洲水样由于其丰富的抗氧化活性，在印度医学体系中具有重要的治疗意义。在顺势疗法中，[医]亚洲水样用于治疗黄疸，皮肤病，水肿，象皮病，麻风病，淋病，白带和神经衰弱。它含有最丰富的三萜糖苷亚洲酸，对癌细胞显示细胞毒活性。这种药物的顺势疗法母町是抗氧化化合物的主要来源，其对其整体药理活性负责。目标: 本研究的目的是评价抗氧化活性和高效薄层色谱（HPTLC）研究的[医]亚洲水样内部的顺势疗法; 顺势疗法母町和市场样品。材料和方法: 采用2,2-二苯基-1-吡啶基-肼基-水合二氢自由基清除活性、总苯酚和2,2-叠氮基-双-(3-乙基苯并噻唑啉-6硫代硫酸钠。研究在HPTLC板上进行HPTLC研究，使用的流动相为甲苯：乙酸乙酯：甲酸（5.5：4.5：1），并用254和366nm处进行UV检测。为了衍生，使用了茴香醛硫酸试剂。结果: 研究结果表明，亚洲水样顺势疗法母町的抗氧化试验具有突出的抗氧化活性。HPTLC研究表明三萜苷化合物亚洲酸在[医]亚洲水样的氯仿提取物中存在。结论: 与从市场上采购的母町相比，经认证的植物样品制备的母町显示出最大的活性成分和突出的抗氧化活性。本研究证明了亚洲氢钾的顺势疗法的使用，并强调了其治疗作用。