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RESEARCH ARTICLE

Standardization and Detailed Aspects of Chopchinyadi Churna: A Potent Anti-Arthritic Medicine

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Abstract:

Background:

Chopchinyadi Churna is a powdered Ayurvedic churna, commonly used for treating insect bite, rheumatoid arthritis, gout arthritis.

Objective:

The current research is oriented for the evaluation of ingredients and other aspects of Churna.

Materials and Methods:

The Churna was standardized as per the parameters of Ayurvedic Formulary of India for the organoleptic characters, microscopy, physicochemical, chromatographic, rheological properties and phytochemical screening for the detection of major phytoconstituents.

Results:

The parameters were found to be significant and offered future benefits for the advanced evaluation of Churna.

Conclusion:

Herbal based anti-arthritic medicine Chopchinyadi Churna has been evaluated on the basis of various parameters, which can serve as references for developing the pharmacopoeial standards.

Keywords: Chopchinyadi, Churna, Anti-arthritic, Medicine, Standardization, Phytoconstituents.

Article History

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1. INTRODUCTION

Plants are the richest source of pharmaceutical lead molecules and their contribution to the drug discovery process is remarkable. Many herbs and herbal medicines have been used since time immemorial to cure many disorders/diseases, including arthritis [1]. In the traditional system of Indian medicine, combined extracts of individual plants are preferred rather than individual ones to achieve maximum therapeutic efficacy [2]. Nature has gifted us enormous wealth in the form of herbal plants, distributed all over the world as medicinal agents for the treatment of various diseases [3]. According to the WHO, 80% of the world's population depends on herbal medicines for their primary health care requirements. Herbal medicines are irreplaceable and an important part of human so-

ciety to combat diseases from a civilization [4]. The medicinal properties of these herbal plants owe to the presence of chemical constituents that produce a desired physiological action on the body [5]. The preparation of a drug according to the ancient methods has minimized due to the commercialization of Ayurvedic pharmacy, therefore quality control is essential [6]. Chopchinyadi Churna is an Ayurvedic medicine, used in treating insect bite, rheumatoid arthritis, and gout, used commonly in North India [7]. Standardization of a herbal formulation is essential in order to ensure the quality, purity, safety and efficacy of the constituent drugs. Chopchinyadi-churna is a polyherbal formulation comprising 11 ingredients *i.e.* madhusnuhi (smilax china), vidanga (embeliaribes), triphala, sugar, cinnamon, tvakpippali (piper longum), lavanga (clove), akarakarabha (Anacyclus pyrethrum), kokilaksha (Asteracanthalongifolia), marica (piper nigrum), and sunthi (Zinziberofficinale). Smilax china L. is known as chobchini in

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Hindi, madhusnuhi in Sanskrit and china root in English. It shows medicinal properties as being anti-inflammatory, diuretic, anti-diabetic, antipsoriatic, and digestive [8 - 11]. *Embelia ribes* Burm F., is also known as false black pepper or vidanga. It is a valuable medicinal plant exhibiting anthelmintic, carminative, antibacterial, antibiotic, hypoglycemic, and antifertility actions [12, 13]. All the other constituents of the formulation also have anti-arthritis potential. Though the contents of the formulation are rich enough to act against arthritis as most of them are potent anti-inflammatory agents. The present research focuses on the detailed aspects and complete standardization of Chopchinyadi churna composed of the constituents, which are used in the treatment of arthritic pains.

2. MATERIALS AND METHODS

Chopchinyadi churna was procured as shown in Fig. (1) from the local market of Kanpur. It contains eleven ingredients, which are mentioned in Table 1. All of the constituents present act as potent anti-arthritis agents.



Fig. (1). Chopchinyadi churna.

2.1. Role of Individual Constituents as an Anti-Arthritic Agent

2.1.1. Chopchini

Smilax chinensis, belonging to the family Liliaceae, actively acts as an anti-diabetic, anti-inflammatory, antioxidant, anticancer, analgesic, inflammation agent [14, 15]. The aqueous extract of *Smilax china* L. shows anti-inflammatory and anti-nociceptive activities [16].

2.1.2. Triphala

An Indian traditional Ayurvedic herbal formulation Triphala, which is composed of amla, bahera and harde has already been evaluated for the antiarthritic effect on adjuvant-induced arthritis in mice [17]. Triphala, containing *Terminalia chebula* Retz., *Terminalia bellerica* Roxb. and *Embelia officinalis*, is used for treating bowel-related complications, inflammations and gastritis [18]. Ayurveda doctors recommend

this formulation to relieve symptoms of inflammatory joint problems like rheumatoid arthritis, psoriatic arthritis and gout [19].

Table 1. Constituents of chopchinyadi Churna.

S.No	Herb	Botanical Name
1.	Chop chini	<i>Smilax china</i>
2.	Sarkara	Sugar
3.	Triphala	<i>Embelia officinalis</i> , <i>Terminalia bellerica</i> , <i>Terminalia chebula</i>
4.	Pippalimoola	<i>Piper longum</i>
5.	Maricha	<i>Piper nigrum</i>
6.	Lavanga	<i>Syzygium aromaticum</i>
7.	Akarakarabha	<i>Anacyclus pyrethrum</i>
8.	Kokilaksha	<i>Asteracantha longifolia</i>
9.	Shunti	<i>Zingiber officinalis</i>
10.	Vidanga	<i>Embelia ribes</i>
11.	Cinnamon	<i>Cinnamomum zeylanicum</i>

2.1.3. Pippali

The root of this plant is known as Pippali Mula in Ayurveda and its fruits found in the form of the spike are mainly used for Rasayana purpose. Several pharmacological activities like anti-ulcer, anti-amoebic, anti-oxidant and anti-inflammatory activities have been reported for the fruit of this plant [20, 21].

2.1.4. Maricha

Maricha is also botanically known as *piper nigrum*. The main constituent of it is piperine, which shows anti-inflammatory, antinociceptive and antiarthritic effects. Black pepper or *Piper nigrum* is commonly used as a spice, but it also has multiple uses as a medicine, a preservative, and in perfumery. Its extract from the plant is rich in active phenolic component, piperine [22]. Its anti-inflammatory activities have been exhibited in rat models of carrageenan-induced paw edema [23]. Constituents of the piper species are responsible for in vitro inhibitory activity against the enzymes responsible for leukotriene and prostaglandin biosynthesis [24].

2.1.5. Clove

Clove contains essential oil, which possesses anti-inflammatory and anti-arthritis properties [25]. At a lower concentration of 0.03% v/v, the anti-inflammatory activity of eugenol, which is the important constituent of its oil has been demonstrated in human gingival fibroblast and pulp cells [26, 27].

2.1.6. Akarakarabha

Anacyclus pyrethrum is widely used in Moroccan traditional medicine. It is active against inflammatory and painful diseases. Antinociceptive and anti-inflammatory activities have been exhibited by the aqueous and methanol extracts of roots of *Anacyclus pyrethrum* [28]. N-isobutyldienedynamide and polysaccharides are present in its roots [29 - 32]. It treats rheumatic arthritis by increasing circulation [33].

2.1.7. *Asteracantha Longifolia*

also known as askokilasha, helps in alleviating the symptoms of rheumatoid arthritis [34]. It is rich in phytoconstituents as alkaloids, flavonoids, terpenoids, essential oil. It has been used in the traditional systems of medicine. Many pharmacological studies have been performed on the plant signifying its antioxidant and analgesic potential [35].

2.1.8. *Zingiber Officinale*

Ginger is primarily known as curative [36]. It has also been used medicinally as an anti-inflammatory agent since antiquity [37 - 39]. Gingerols, the phenolic compounds are responsible for its pungent taste and usage in treating inflammatory disorders as arthritis [40]. The potent anti-arthritis effects of the extracts of ginger have been proven in an experimental model of rheumatoid arthritis. However, crude extracts containing both gingerols as well as essential oils have been found to be even more potent in inhibiting joint swelling as compared to gingerols alone [41].

2.1.9. *Embelia Ribes*

It is commonly known as Vidanga. It is used as a powerful anthelmintic in Ayurveda [42]. It has also been evaluated to have antifertility action [43]. Analgesic property has been reported for embelin and its derivatives [44]. It is used as an anti-inflammatory drug in the treatment of rheumatism and fever [45].

2.1.10. *Cinnamomum Zeylanicum*

Cinnamon is used for its aroma in the essence industries. Due to its fragrance, it is applicable for use in different varieties of foodstuffs, perfumes, and medicinal products [46]. Cinnamon bark is rich in ingredients as procyanidins and catechins [47]. The components of procyanidins include both procyanidin A and B [48 - 50]. Therefore it has been used as an anti-inflammatory agent [51 - 53]. The major pharmacological uses of cinnamon bark are due to its anti-inflammatory and anti-microbial properties [54 - 57].

2.2. Phytochemical Screening

Plant extracts when tested for preliminary phytochemical screening, the presence of phytoconstituents as alkaloids, carbohydrates and reducing sugars, glycosides, proteins and amino acids, steroids and triterpenoids, tannins, flavonoids, fixed oils, volatile oils were detected. The methanolic extracts of the formulation were concentrated for further use.

2.3. Tests for Volatile Oils

The non-permanent staining of filter papers, along with the characteristic odour indicated the presence of volatile oils.

2.4. Tests for Alkaloids

Methanolic extracts of all the drugs were acidified and then filtered to obtain the filtrate to carry out further tests.

2.5. Hager's Test

A yellow precipitate was formed with it.

2.6. Tests for Carbohydrates and Reducing Sugars:

2.6.1. Molisch's Test

To drug extracts, a few drops of alcoholic α -naphthol solution, followed by concentrated sulphuric acid were added. A purple ring at the junction of two liquids indicated the presence of carbohydrates.

2.6.2. Fehling's Test

Fehling A and Fehling B solutions were mixed with equal volumes of the filtrates and heated for 5 minutes. First yellow, followed by brick red precipitate indicated the presence of reducing sugars.

2.7. Tests for Glycosides

2.7.1. Foam Test

Drugs in small quantities were shaken vigorously with water. The formation of persistent foam indicated the presence of saponin glycosides.

2.8. Tests for Phenolic Compounds and Tannins

With 5% Ferric chloride solution, they gave a deep blue-black color [58, 59].

2.9. Organoleptic Evaluation

Dried powdered churna was taken in order to examine the macroscopic features of the formulation (Fig. 2). The morphological features were determined with the use of a simple microscope [60].



Fig. (2). Churna sample.

2.10. Microscopic Evaluation

Coarsely powdered, air-dried samples were used for the purpose of microscopic evaluation. These were then treated with reagents like phloroglucinol and hydrochloric acid, followed by treatment with clearing agents, such as lactophenol and chloral hydrate in order to remove the impurities adhered. Then finally, they were mounted in glycerine. Glycerine, being

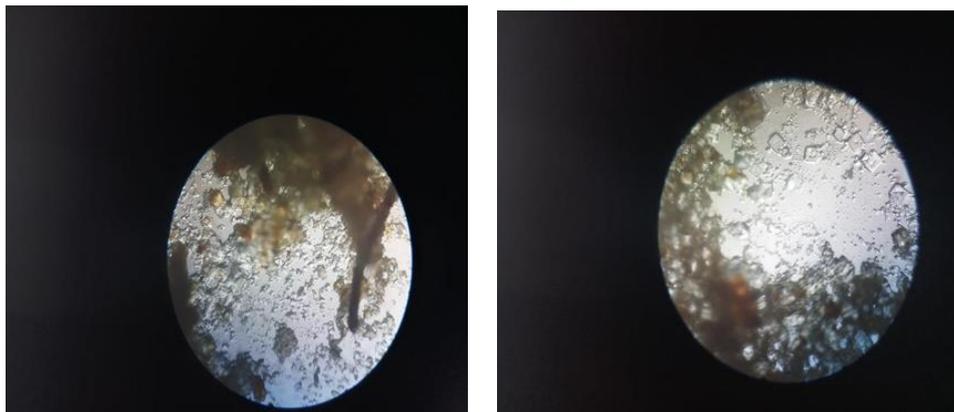


Fig. (3). Microscopy of Churna.

an emollient, saves the sample from drying. Different cell components were studied as in Figs. (3a and 3b) per the standard methods with the help of different freshly prepared reagents [61 - 64].

2.11. Physico-Chemical Evaluation

Physicochemical standards as ash values, moisture content and extractive values were determined as per official methods and procedures.

2.12. Thin Layer Chromatography

A solvent system was used i.e toluene:ethyl acetate (9:3). Plates with the spotted sample were kept in the TLC chamber. Then, the chamber was closed with a lid. After spot development, the plates were removed and dried. Then sample spots were visualized in a UV light chamber (Fig. 4).

2.13. Rheological Evaluation

Coarsely powdered crude drug was screened for different parameters. Bulk density, angle of repose, compressibility, and Hausner's ratio were calculated according to standard methods.

2.13.1. Bulk Density

10g of the powdered drug was taken in a graduated measuring cylinder and tapped on a wooden surface. Bulk density = weight taken / Bulk volume Tapped density = weight of drug taken / volume (tapped).

2.13.2. Angle of Repose

The angle of repose was determined by using the funnel method. The powder was made to flow through a funnel fixed on a stand to form a heap. The height and the radius gave the angle of repose. The angle of repose $\tan\theta = h/r$ $\theta = \tan^{-1} (h/r)$. Where, h = height of heap, r = radius of the heap.

2.13.3. Compressibility / Carr's Index

It is: Bulk density (Tapped) – Bulk density (Untapped)/ Bulk density (Tapped) Hausner's Ratio. The formula used to determine Hausner's ratio is Bulk density (Tapped) x 100 /Bulk density (untapped) [65, 66]. In spectral analysis, the

churna was subjected to spectral techniques as Infra –red and ultraviolet spectroscopy [67 - 69].

3. RESULTS

Phytochemical analysis revealed the presence of alkaloid, tannins (due to the presence of amla, bahera, myrobalans in the form of triphala) reducing sugars, volatile oil (due to the presence of ginger, clove and piper species) and the absence of carbohydrate, saponins. It has been concluded that Churna is free from heavy metals. From the all above values, efficacy of the Churna is revealed (Table 2).

Table 2. Phytochemical screening of Churna.

S. No.	Chemical Tests	Reagents	Result
1.	Test for Alkaloid	Hager's Reagent	+ve
2.	Test for Carbohydrate	Molisch Reagent	-ve
3.	Test for Volatile oil	Filter paper	+ve
4.	Test for Reducing Sugar	Fehling Sol. (A&B)	+ve
5.	Test for Tannins	FeCl ₃	+ve
6.	Test for Saponin	Foam Test	-ve

Botanical parameters revealed that Churna was light brown, moderately fine powder, odour- characteristic, taste- slightly sweet (Table 3).

Table 3. Organoleptic evaluation of Churna.

S.no	Parameter	Result
1	Colour	Light Brown
2	Odour	Characteristic
3	Taste	Slightly Sweet

Results of quantitative analysis for loss on drying at 105 C, total ash, acid insoluble ash, water soluble ash, alcohol soluble extractives and water soluble extractive were calculated and results are shown (Table 4). Ash value helps to assure the authenticity and purity of the drug. The loss on drying indicates the amount of moisture content present in the drug. Moisture content was found to be 8% w/w. The less value of moisture content prevents bacterial, fungal or yeast growth. The obtained values as water soluble and alcohol soluble indicate the

amount of active constituent in the given amount of drug when extracted with the respective solvent. The Rf value was found to be 0.63.



Fig. (4). TLC of Churna.

Table 4. Physico-chemical evaluation of Churna.

S.no	Parameter	Result
1	Moisture content	8%
2	Total ash	9.0%
3	Acid insoluble ash	3.2%
4	Water soluble ash	5.2%
5	Water soluble extractive	41.5%
6	Alcohol soluble extractive	11.5%
7	Rf Value	0.63

Rheological properties of churnas shown in Table 5 like bulk density, tapped density, Hausner's ratio and Carr's index indicate very poor flowability and the angle of repose indicates Poor-must agitate, with vibrate flow property.

3.1. Spectral Analysis of Churna

UV spectrophotometry revealed the absorbance 0.173 at 260 nm. Infrared spectroscopy revealed the presence of peaks at 1700 cm^{-1} and 3400 cm^{-1} , indicating the presence of keto and aldehyde groups containing phytoconstituents in the formulation (Fig. 5).

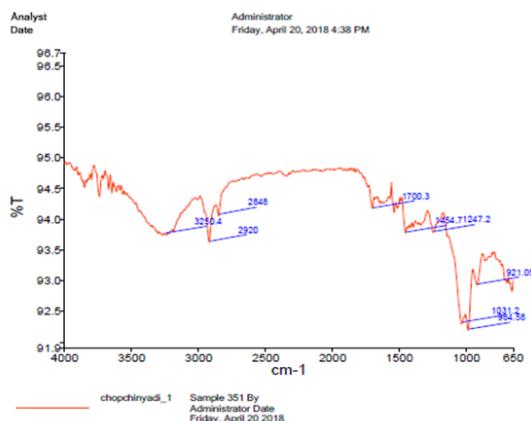


Fig. (5). IR Spectra of Churna.

4. DISCUSSION

Phytochemical screening of churna showed the absence of carbohydrates and saponins. It revealed the presence of tannins, reducing sugars, volatile oil and alkaloids. Morphology discussed the light brown, slightly sweet taste of it. The moisture content was found to be 8% w/w. The less value of moisture content prevents bacterial, fungal or yeast growth. The extractive values as water soluble and alcohol soluble indicate the amount of active constituent in the given amount of drug when extracted with the respective solvent. Powder on its flow study showed that it has a very poor flowability and the angle of repose indicates poor must agitate, with vibrate flow property. Spectral analysis indicated the presence of keto and aldehyde groups containing phytoconstituents in the formulation [70, 71].

CONCLUSION

Various evaluation parameters like physicochemical standards as total ash, acid insoluble ash, water and alcohol-soluble extractive values, loss on drying, microscopy, spectroscopy, phytochemical analysis, flow properties, and morphological features were evaluated. From their results, it was revealed that the formulation of Churna is effective with uniform characteristics of an ideal Churna. It is beneficial, effective and economic as well. The efficacy of the Churna can be proved by the pharmacology which suggests the future scope of R & D. Though the contents of the formulation are rich enough to act against arthritis as most of them are potent anti-inflammatory agents. The study shows that the ingredients of churna present are in accord with the prescribed guidelines of the WHO. All these observations are not mentioned in the standard literature, which could be helpful in authentication

Table 5. Rheological evaluation of Churna.

S.no.	Parameter	Result
1	Tapped density	40.2
2	Untapped density	33.4
3	Angle of repose	33.5
4	Hausner ratio	1.2

and assurance of Churnas. The result of the present study will help to set standards as a reference monograph in the process of drug formulation. Ayurvedic Churna has been standardized by the intervention of modern scientific quality control measures in the traditional preparation described in classical texts. Pharmacognostic characters established for the raw materials could be employed as Q.C. standards for evaluating their identity and can be used for routine analysis.

ETHICAL APPROVAL AND CONSENT TO PARTICIPATE

Not applicable.

HUMAN AND ANIMAL RIGHTS

No animals/humans were used for studies that are the basis of this research.

CONSENT FOR PUBLICATION

Not applicable.

AVAILABILITY OF DATA AND MATERIALS

The authors confirm that data supporting the findings of this research are available within the article.

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CONFLICT OF INTEREST

The authors declare no conflict of interest, financial or otherwise.

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REFERENCES

- Parasuraman S, Thing GS, Dhanaraj SA. Polyherbal formulation: Concept of ayurveda. *Pharmacogn Rev* 2014; 8(16): 73-80. [<http://dx.doi.org/10.4103/0973-7847.134229>] [PMID: 25125878]
- Jayakumar RV. Herbal medicine for type-2 diabetes. *Int J Diabetes Dev Ctries* 2010; 30: 111-2. [<http://dx.doi.org/10.4103/0973-3930.66501>]
- Kalaria P, Gheewala P, Chakraborty M, Kamath J. A phytopharmacological review of *Alstoniascholaris*: A panoramic herbal medicine. *IJRAP* 2012; 3: 367-1.
- Sudha K, Mathanghi SK. Traditional underutilized green leafy vegetables and its curative properties *Int J Pharm* 2012; 2: 786-93.
- Singh V, Patel H, Suvagiya V, Singh K. Some traditionally used anti-arthritis herbs a review. *Int Res J Pharm* 2011; 2: 43-5.
- New Delhi: Govt of India, Ministry of Health and Family Welfare. Anonymous. *The Ayurvedic Formulary of India* 1976.
- Chopchinyadichurna Benefits, Dosage, Ingredients, Side Effects. *Ayurvedic Medicine Information ByDr JV Hebbar* cited from: <https://ayur.medinfo.com/2012/05/29/chopchinyadichurna-benefits-dosage-ingredients-side-effects/>
- Jayprakasham R, Ravit K, Athira K, AshaJyothi B. Quantitative determination of Diosgenin in polyherbal formulation and various extracts of *Smilax china* Linn using standard marker by validated analytical technique. *Universal Journal of Pharmacy* 2013; 02(04): 83-90.
- Khare CP. *Indian Herbal Remedies rational western therapy, ayurvedic and other traditional uses, botany*. Springer 2004; p. 428.
- Saravanakumar S, Christilda F, Sundarapandian S. Phytochemical screening of the methanol extract of root tuber of *Smilax china*. *International Journal of Pharmacognosy and Phytochemical Research* 2014-15; 6(4): 963-6.
- Rajesh B, Anupama S, Vikas A, Veerma R, Anil B. Pharmacognostical standardization, extraction and anti-diabetic activity of *Smilax china* L. rhizome. *Asian J Tradit Med* 2011; 6(5): 218-22.
- Syed A, Ramangdang R. Pharmacognosy of *Embelia ribes* Burm F IJ-RPC 1(4): 1236-50.2011;
- Sudani RJ, Akbari BV, Vidyasagar G, Sharma P. Quantitative and chromatographic fingerprint analysis of *Embeliaribes* churna formulations by HPLC method. *Int J Pharm Biol Arch* 2011; 2(2): 611-8.
- Huang ZQ, Huang YR. Clinical application of single *Smilax china* L. on prolong life of the patients suffer from liver neoplasms *Chinese National Folk Medicine Journal* 2000; 45: 212-13.
- Khare C P. *Indian medicinal plants An Illustrated dictionary* Springer science 2007; 79.
- Shu XS, Gao ZH, Yang XL. Anti-inflammatory and anti-nociceptive activities of *Smilax china* L. aqueous extract. *J Ethnopharmacol* 2006; 103(3): 327-32. [<http://dx.doi.org/10.1016/j.jep.2005.08.004>] [PMID: 16387460]
- Rasool M, Sabina EP. Antiinflammatory effect of the Indian Ayurvedic herbal formulation *Triphala* on adjuvant-induced arthritis in mice. *Phytother Res* 2007; 21(9): 889-94. [<http://dx.doi.org/10.1002/ptr.2183>] [PMID: 17533629]
- Kalaiselvan S, Rasool MK. The anti-inflammatory effect of *Triphala* in arthritic-induced rats. *Pharm Biol* 2015; 53(1): 51-60. [<http://dx.doi.org/10.3109/13880209.2014.910237>] [PMID: 25289531]
- Psoriasis management. *Triphala for Psoriasis and Psoriatic Arthritis*. by Ashish Agarwal November 6, 2017. cited from <http://www.psoriasisselfmanagement.com/detoxification-ideas/triphala-leaky-gut-psoriasis-psoriatic-arthritis/>
- Warrier PK, Nambiar VP, Raman KC. *Madras, India: Orient Longman Ltd: Piper longum, Indian medicinal Plants* 1995; 4: p. 290.
- Dahanukar SA, Karandikar SM. Evaluation of anti-allergic activity of *Piper longum*. *Indian Drugs* 1984; 21: 377-83.
- Srinivasan K. Black pepper and its pungent principle-piperine: A review of diverse physiological effects. *Crit Rev Food Sci Nutr* 2007; 47(8): 735-48. [<http://dx.doi.org/10.1080/10408390601062054>] [PMID: 17987447]
- Mujumdar AM, Dhuley JN, Deshmukh VK, Raman PH, Naik SR. Anti-inflammatory activity of piperine. *Jpn J Med Sci Biol* 1990; 43(3): 95-100. [<http://dx.doi.org/10.7883/yoken1952.43.95>] [PMID: 2283727]
- Stöhr JR, Xiao PG, Bauer R. Constituents of Chinese *Piper* species and their inhibitory activity on prostaglandin and leukotriene biosynthesis *in vitro*. *J Ethnopharmacol* 2001; 75(2-3): 133-9. [[http://dx.doi.org/10.1016/S0378-8741\(00\)00397-4](http://dx.doi.org/10.1016/S0378-8741(00)00397-4)] [PMID: 11297843]
- Han X, Parker TL. Anti-inflammatory activity of clove (*Eugenia caryophyllata*) essential oil in human dermal fibroblasts. *Pharm Biol* 2017; 55(1): 1619-22. [<http://dx.doi.org/10.1080/13880209.2017.1314513>] [PMID: 28407719]
- Prashar A, Locke IC, Evans CS. Cytotoxicity of clove (*Syzygium aromaticum*) oil and its major components to human skin cells. *Cell Prolif* 2006; 39(4): 241-8. [<http://dx.doi.org/10.1111/j.1365-2184.2006.00384.x>] [PMID: 16872360]
- Koh T, Murakami Y, Tanaka S, Machino M, Sakagami H. Re-evaluation of anti-inflammatory potential of eugenol in IL-1 β -stimulated gingival fibroblast and pulp cells. *In Vivo* 2013; 27(2): 269-73. [PMID: 23422489]
- Manouze H, Bouchatta O, Gadhi AC, Bennis M, Sokar Z, Bamm'amed S. Anti-inflammatory, antinociceptive, and antioxidant activities of methanol and aqueous Extracts of *Anacyclus pyrethrum* Roots. *Front Pharmacol* 2017; 8: 598. [<http://dx.doi.org/10.3389/fphar.2017.00598>] [PMID: 28928658]
- Crombie L. Isolation and structure of an n-isobutyldienedynamide from pellitory (*Anacyclus pyrethrum* dc.). *Nature* 1954; 174: 832-3. [<http://dx.doi.org/10.1038/174832a0>]
- Bendjeddou D, Lalaoui K, Satta D. Immunostimulating activity of the hot water-soluble polysaccharide extracts of *Anacyclus pyrethrum*, *Alpinia galanga* and *Citrullus colocynthis*. *J Ethnopharmacol* 2003; 88(2-3): 155-60. [[http://dx.doi.org/10.1016/S0378-8741\(03\)00226-5](http://dx.doi.org/10.1016/S0378-8741(03)00226-5)] [PMID: 12963136]
- De Spiegeleer B, Boonen J, Sharma V, Dixit V. 2011. *New N-Alkylamides from Anacyclus pyrethrum*. Ph.D. thesis, Faculteit Farmaceutische Wetenschappen, Ghent
- Boonen J, Sharma V, Dixit VK, Burvenich C, De Spiegeleer B. LC-MS N-alkylamide profiling of an ethanolic *Anacyclus pyrethrum* root

- extract. *Planta Med* 2012; 78(16): 1787-95.
[http://dx.doi.org/10.1055/s-0032-1315371] [PMID: 23047251]
- [33] Usmani A. Pharmacognostic and phytopharmacology study of *Anacyclus pyrethrum*: An insight. *J Appl Pharm Sci* 2016; 6(03): 144-50.
[http://dx.doi.org/10.7324/JAPS.2016.60325]
- [34] Thankamma A. Rheumatoid arthritis and *astercantha longifolia*. *Anc Sci Life* 1999; 18(3-4): 247-9.
[PMID: 22556897]
- [35] Imtiyaz S, Rahman K. *Asteracanthalongifolia* Linn: An Overview *American Journal of PharmTech Research* 2013; 3(1): 15-26.
- [36] Barnes PM, Powell-Griner E, McFann K, Nahin RL. Complementary and alternative medicine use among adults: United States, 2002. *Adv Data* 2004; 343(343): 1-19.
[http://dx.doi.org/10.1016/j.sigam.2004.07.003] [PMID: 15188733]
- [37] Palatty PL, Haniadka R, Valder B, Arora R, Baliga MS. Ginger in the prevention of nausea and vomiting: A review. *Crit Rev Food Sci Nutr* 2013; 53(7): 659-69.
[http://dx.doi.org/10.1080/10408398.2011.553751] [PMID: 23638927]
- [38] Ali BH, Blunden G, Tanira MO, Nemmar A. Some phytochemical, pharmacological and toxicological properties of ginger (*Zingiber officinale* Roscoe): A review of recent research. *Food Chem Toxicol* 2008; 46(2): 409-20.
[http://dx.doi.org/10.1016/j.fct.2007.09.085] [PMID: 17950516]
- [39] Grzanna R, Lindmark L, Frondoza CG. Ginger an herbal medicinal product with broad anti-inflammatory actions. *J Med Food* 2005; 8(2): 125-32.
[http://dx.doi.org/10.1089/jmf.2005.8.125] [PMID: 16117603]
- [40] Chrubasik S, Pittler MH, Roufogalis BD. *Zingiberis rhizoma*: A comprehensive review on the ginger effect and efficacy profiles. *Phytomedicine* 2005; 12(9): 684-701.
[http://dx.doi.org/10.1016/j.phymed.2004.07.009] [PMID: 16194058]
- [41] Funk JL, Frye JB, Oyarzo JN, Timmermann BN. Comparative effects of two gingerol-containing *Zingiber officinale* extracts on experimental rheumatoid arthritis. *J Nat Prod* 2009; 72(3): 403-7.
[http://dx.doi.org/10.1021/np8006183] [PMID: 19216559]
- [42] Hördegen P, Cabaret J, Hertzberg H, Langhans W, Maurer V. *in vitro* screening of six antihelminthic plant products against larval *Haemonchus contortus* with a modified methyl-thiazolyl-tetrazolium reduction assay. *J Ethnopharmacol* 2006; 108(1): 85-9.
[http://dx.doi.org/10.1016/j.jep.2006.04.013] [PMID: 16725288]
- [43] Gupta S, Sanyal SN, Kanwar U. Antispermatic effect of embelin, a plant benzoquinone, on male albino rats *in vivo* and *in vitro*. *Contraception* 1989; 39(3): 307-20.
[http://dx.doi.org/10.1016/0010-7824(89)90063-2] [PMID: 2714091]
- [44] Zutshi U, Johri RK, Atal CK. Possible interaction of potassium embelate, a putative analgesic agent, with opiate receptors. *Indian J Exp Biol* 1989; 27(7): 656-7.
[PMID: 2561116]
- [45] Kapoor VK, Chawla AS, Kumar M, Kumar P. Anti-inflammatory agent in Indian Laboratories. *Indian Drugs* 1983; 30: 481-8.
- [46] Huang T-C, Fu H-Y, Ho C-T, Tan D, Huang Y-T, Pan M-H. Induction of apoptosis by cinnamaldehyde from indigenous cinnamon *Cinnamomum osmophloeum* Kaneh through reactive oxygen species production, glutathione depletion, and caspase activation in human leukemia K562 cells. *Food Chem* 2007; 103(2): 434-43.
[http://dx.doi.org/10.1016/j.foodchem.2006.08.018]
- [47] Nonaka G-I, Morimoto S, Nishioka I. Tannins and related compounds. Part 13. Isolation and structures of trimeric, tetrameric, and pentamericproanthocyanidins from cinnamon. *J Chem Soc, Perkin Trans 1* 1983; 2139-45.
[http://dx.doi.org/10.1039/p19830002139]
- [48] Anderson RA, Broadhurst CL, Polansky MM, *et al*. Isolation and characterization of polyphenol type-A polymers from cinnamon with insulin-like biological activity. *J Agric Food Chem* 2004; 52(1): 65-70.
[http://dx.doi.org/10.1021/jf034916b] [PMID: 14709014]
- [49] Peng X, Cheng K-W, Ma J, *et al*. Cinnamon bark proanthocyanidins as reactive carbonyl scavengers to prevent the formation of advanced glycation endproducts. *J Agric Food Chem* 2008; 56(6): 1907-11.
[http://dx.doi.org/10.1021/jf073065v] [PMID: 18284204]
- [50] Tanaka T, Matsuo Y, Yamada Y, Kouno I. Structure of polymeric polyphenols of cinnamon bark deduced from condensation products of cinnamaldehyde with catechin and procyanidins. *J Agric Food Chem* 2008; 56(14): 5864-70.
[http://dx.doi.org/10.1021/jf800921r] [PMID: 18558701]
- [51] Chao LK, Hua K-F, Hsu H-Y, Cheng S-S, Liu J-Y, Chang S-T. Study on the antiinflammatory activity of essential oil from leaves of *Cinnamomum osmophloeum*. *J Agric Food Chem* 2005; 53(18): 7274-8.
[http://dx.doi.org/10.1021/jf051151u] [PMID: 16131142]
- [52] Tung Y-T, Chua M-T, Wang S-Y, Chang S-T. Anti-inflammation activities of essential oil and its constituents from indigenous cinnamon (*Cinnamomum osmophloeum*) twigs. *Bioresour Technol* 2008; 99(9): 3908-13.
[http://dx.doi.org/10.1016/j.biortech.2007.07.050] [PMID: 17826984]
- [53] Tung Y-T, Yen P-L, Lin C-Y, Chang S-T. Anti-inflammatory activities of essential oils and their constituents from different provenances of indigenous cinnamon (*Cinnamomum osmophloeum*) leaves. *Pharm Biol* 2010; 48(10): 1130-6.
[http://dx.doi.org/10.3109/13880200903527728] [PMID: 20815702]
- [54] Oussalah M, Caillet S, Lacroix M. Mechanism of action of Spanish oregano, Chinese cinnamon, and savory essential oils against cell membranes and walls of *Escherichia coli* O157: H7 and *Listeria monocytogenes*. *J Food Prot* 2006; 69(5): 1046-55.
[http://dx.doi.org/10.4315/0362-028X-69.5.1046] [PMID: 16715803]
- [55] Lee HS, Kim BS, Kim MK. Suppression effect of *Cinnamomum cassia* bark-derived component on nitric oxide synthase. *J Agric Food Chem* 2002; 50(26): 7700-3.
[http://dx.doi.org/10.1021/jf020751f] [PMID: 12475291]
- [56] Hayashi K, Imanishi N, Kashiwayama Y, *et al*. Inhibitory effect of cinnamaldehyde, derived from *Cinnamomi* cortex, on the growth of influenza A/PR/8 virus *in vitro* and *in vivo*. *Antiviral Res* 2007; 74(1): 1-8.
[http://dx.doi.org/10.1016/j.antiviral.2007.01.003] [PMID: 17303260]
- [57] Cabello CM, Bair WB III, Lamore SD, *et al*. The cinnamon-derived Michael acceptor cinnamic aldehyde impairs melanoma cell proliferation, invasiveness, and tumor growth. *Free Radic Biol Med* 2009; 46(2): 220-31.
[http://dx.doi.org/10.1016/j.freeradbiomed.2008.10.025] [PMID: 19000754]
- [58] Sane RT. Standardization, quality control and GMP for herbal drug. *Indian Drugs* 2002; 39(3): 184-90.
- [59] Khandelwal KR. *Practical Pharmacognosy, Techniques and Experiments*. 12th ed. Pune: Nirali Prakashans 1996.
- [60] Trease and Evans. *Pharmacognosy In: Singapore: Harcourt Brace and company Asia Pvt* 1997.
- [61] Sass JE. *Elements of Botanical Micro Technique* 1940; 222.
- [62] Johanson DA. *Plant Micro Technique* 1940.
- [63] KR. *The practical evaluation of phytopharmaceuticals* 1975.812
- [64] Pal RS, Pal Y, Wal P, Wal A. *Pharmacognostic Evaluation of Roots of Benincasa Hispida (Thunb.) Cogn.(Cucurbitaceae) The Open Plant Science Journal* 2018; 11: 1-6.
- [65] Lachman L, Lieberman HA, Kanig JL. *The theory and practice of industrial pharmacy*. 3rd ed. 1987; pp. 183-316.
- [66] Aulton ME. *Pharmaceutics, the science of dosage forms designs*. 2nd ed. 2002; pp. 205-21.
- [67] Mukherjee PK. *Quality control of Herbal drugs an approach to evaluation of botanicals*. 1st ed. New Delhi: Business Horizons Pharmaceutical Publications 2002.
- [68] Harbone JB. *Phytochemical method a guide to modern technique of plant analysis* 2nded. New York: Champman and Hall 1984.
- [69] *The Ayurvedic Pharmacopoeia of India*, Government of India, Ministry of Health and family welfare, Department of AYUSH In: 2001; 48-9: pp. 173-4.
- [70] Haslam E. Natural polyphenols (vegetable tannins) as drugs: Possible modes of action. *J Nat Prod* 1996; 59(2): 205-15.
[http://dx.doi.org/10.1021/np960040+] [PMID: 8991956]
- [71] Jones GA, Mc Allister T, Muir AD, Chang KJ. *Appl Environ Microbiol Lett* 1994; 46: 223-7.