

## Comparative Study of antioxidant activity on aqueous, ethanolic & petroleum ether extract of passiflora incarnate linn. Flower.

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Submitted: 20-06-2023

Accepted: 29-06-2023

### ABSTRACT

To study and compare the antioxidant activity of aq. ethanol & petroleum ether extract of passifloraincarnate linn. flower powder. Plants belonging to the Genus Passiflora have been commonly used intraditional medicine for a variety of health conditions. The major chemical constituents present in these plants are identified as carbohydrates, alkaloids, glycosides and tannin compounds. Ferric Reducing Antioxidant Power (FRAP) has been used to determine the antioxidant activity of passiflora incarnate linn.

This plant was used widely in traditional medicine in West India, Mexico, the Netherlands, South America, Italy and Argentina. One of the species of this genus named as Passiflora incarnate is more popular than its other species. Passiflora contains several compounds including alkaloids, phenols, glycosyl flavonoids and cyanogenic compounds. In some experiments, it has potential effects for treatment of some diseases like anxiety, opiate withdrawal, insomnia, attention-deficit hyperactivity disorder and cancer. Passion flower is also known as maypop, apricot vine, passion Vine, and granadilla. It grows as much as 30ft(10m) tall, with a thick, woody stem.

### TAXONOMY

Kingdom: Plantae – Plants

Division: Magnoliophyta –  
Flowering plants

Class: Magnoliopsida – Dicotyledons

Family: Passifloraceae – Passion-  
flower family

Genus: Passiflora L. – Passion flower

Species : P. incarnate L. – Purple Passion  
flower

Flower extract against gram  
positive and negative bacteria.

### I. INTRODUCTION

#### Passionflower:

The genus Passiflora consists of 500 species that are mostly found in warm and tropical regions. Passiflora comes from the Latin word “Passio” that was first time discovered by Spanish discoverers in 1529 and was described as a symbol for “Passion of Christ”

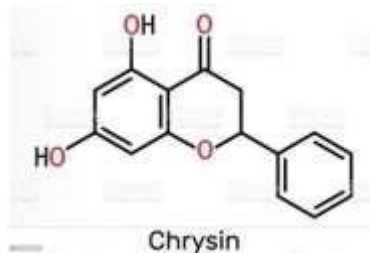


Sr.no	Morphological Characters	Observation
1	Stem	herbaceous or woody, generally climbing, very rarely arborescent.
2	Leaves	Alternate, sometimes simple, entire, lobed or palmate, sometimes compound, imparipinnate; stipules germinate at the base of petioles, rarely absent; tendrils axillary, arising from sterile pedicels.
3	Flowers	bisexual or unisexual, regular
4	Stamens	inserted either at the bottom of the perianth, or at the base or top of gynophore.
5	Seeds	Numerous; funicle dilated into a pulpy cupuliform or saccate aril; testa crustaceous, foveolate,
		easily separable from the membranous endopleura, which bears a longitudinal raphe
6	Ovary	Superior, more or less stipitate, very rarely sessile, unilocular, of 3–5 united carpels Containing several or many anatropous ovules on parietal placentas

**Chemical constituents:-**

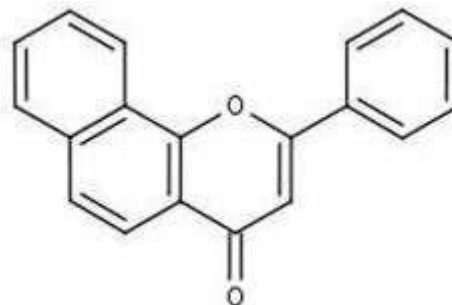
The main chemical constituents of the Passionflower are the flavonoids (0.25%) such as asvitexin, isovitexin, orientin, isorientin, apigenin, kaempferol and quercetin. The indole alkaloids (0.1%) based on the beta-carboline ring system such as harman, harmin, harmalin, harmol and harmalol. Some other isolated plant constituents have been identified such as glycosides, carbohydrates, amino acids, benzopyrones, cyanogenic glycosides such as cyanocardin, pyrone derivatives such as maltol and ethyl maltol. Two important constituents like chrysin and tri-substituted benzoflavone moiety (BZF) have been isolated.

**Chrysin:** C<sub>15</sub>H<sub>10</sub>O<sub>4</sub> (5, 7-dihydroxy-2-phenyl-9CI)



Chrysin is a naturally occurring flavone chemically extracted from the blue passionflower (*Passiflora caerulea*). Chrysin acts as an aromatase inhibitor supplement to bodybuilders and athletes. It has been shown to induce an anti-inflammatory effect, most likely by inhibition of COX-2. In rodent in vivo studies, chrysin was found to be anxiolytic. In herbal medicine, it is recommended as a remedy for anxiety. Chrysin exhibited an anxiolytic effect, which was shown by an increase in locomotor activity. This effect was linked to GABA benzodiazepine receptors in the brain because the anxiolytic effect was blocked by an injection of Flumazenil, which is a benzodiazepine antagonist. Chrysin and apigenin have been shown to inhibit the growth of breast carcinoma cells, human thyroid cancer cells and human prostate tumors. Apigenin is considered antimutagenic because it reduces the effects of mutagens.

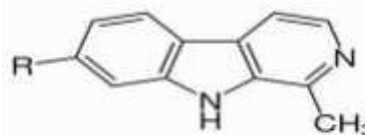
**Benzoflavone**



The -Naphthoflavone, also known as 5,6-benzoflavone, is a potent agonist of the aryl hydrocarbon receptor and an inducer of detoxification enzymes such as cytochromes P450 (CYPs) and uridine 5'-diphosphoglucuronosyltransferases (UGTs) (Chlouchi et al., 2007).

Naphthoflavone is a putative chemopreventive agent. Harmala alkaloids: C<sub>13</sub>H<sub>12</sub>N<sub>2</sub>O (7-Methoxy-1-methyl-9H-pyrido[3,4-b]indole)

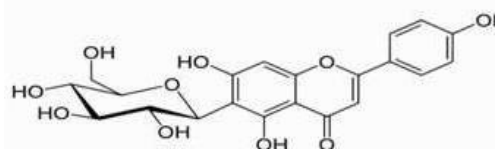
The Passiflora family contains small amounts of harmala alkaloids, harmaline (passiflorine), and possibly harmine (telepathine), harmaline, harmol, and harmalol. The presence of the last four in *P. incarnata* is disputed because they are contained in only very small amounts (0.01% or less).



**Isoorientin (Luteolin-8-C-glucoside)**

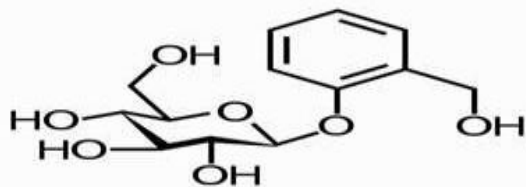
Orientin is a flavone, a chemical flavonoid-like compound found in the passion flower, the Açai palm and *Anadenanthera peregrina*.

Orientin is also reported to be in millets. Isoorientin (or homoorientin) is the luteolin-6-C-glucoside. It can be isolated from the passion flower, *Vitex negundo*, the Açai palm and *Swertia japonica*.



### Glycosides

Glycosides are molecules in which a sugar is bound to a non-carbohydrate moiety, usually a small organic molecule. Glycosides play numerous important roles in living organisms. Many plants store chemicals in the form of inactive glycosides which can be activated by enzyme hydrolysis (Brito-Arias, 2007). Leaf and stem material of *P. edulis* contain the new cyanogenic glycosides (2R)-allopyranosyloxy-2-phenylacetonitrile and (2S)-allopyranosyloxy-2-phenylacetonitrile, along with smaller amounts of (2R)-prunasin, (2S)-sambunigrin. Many different types of glycosides are present in passion flower such as apigenin, homoorientin, 7-isoorientin, isoshaftoside, isovitexin, kaempferol, luteolin, norientin, passiflorine (named after the genus), quercetin, rutin, saponarin, saponarin, shaftoside, vicenin and vitexin.



### Other organic compounds

Passion flower contains many alkaloids, flavonoids as well as many organic compounds such as organic acids. This genus is rich in formic, butyric, linoleic, linolenic, malic, myristic, oleic and palmitic acids as well as phenolic compounds, and the amino acid -alanine. Some species contain ester such as ethyl butyrate, ethyl caproate, n-hexylbutyrate and n-hexylcaproate which give the fruits their flavor and appetizing smell. Sugars, contained mainly in the fruit, are mostly d-fructose, d-glucose and raffinose. Among enzymes, *Passiflora* was found to be rich in catalase, pectin methyl esterase and phenolase. Apart from glycosides, phenols and alkaloids, various miscellaneous phyto-constituents which were also reported to be in *P. edulis* include, Edulans I and II and pectin.

### PHARMACOLOGICAL ASPECTS:

#### Cannabinoids:

**Reversal** The newly reported benzoflavone (BZF) moiety from the plant *P. incarnate* (Linn) has been evaluated in light of traditional reports on the use of this plant in

breaking down cannabis addiction. In the modern or allopathic system of therapeutics, there has been no suitable remedy to combat these severe withdrawal effects of various cannabis products, including marijuana, hashish, ganja, etc., the world-wide consumption of which has attained alarming proportions especially among the younger generation. It has been reported that the BZF of *P. incarnate*, when administered concurrently with cannabinoids, prevented the development of tolerance and dependence of cannabinoids in mice. In this study the mice were given a 10 mg/kg twice-daily dose of delta9-tetrahydrocannabinol (delta9-THC) by oral route for six days to make them dependent upon cannabinoids.

Concurrently, other groups of mice were administered delta9-THC along with a 10 or 20 mg/kg twice-daily dose of the BZF moiety from *P. incarnate* orally for 6 days.

Upon measuring locomotor activity during the treatment regimen, it was noticed that the mice receiving the

*P. incarnate* extract and delta9-THC together developed significantly less tolerance and dependence, relative to the mice receiving delta9-THC alone. Even an acute administration of the BZF significantly blocked the expression of withdrawal effects in cannabinoid dependence. So these studies suggested that the BZF may have a beneficial role in cannabinoid reversal.

#### Nicotine

**Reversal** In light of various reports mentioning the usefulness of *P. incarnate* in tobacco addiction, studies have been performed using four doses (1, 5, 10 and 20 mg/kg) of the bioactive BZF moiety isolated from the aerial parts of *P. incarnate*. In a 7-day experimental regimen, mice were given nicotine hydrogentartrate (2 mg/kg) and combinations of nicotine with four doses of BZF by the subcutaneous route. At the end of the 7 days of treatment, naloxone was given to the mice from all groups to induce a nicotine withdrawal syndrome. The mice that had been treated with 10 and 20 mg/kg dose of BZF concurrently with nicotine showed a significantly fewer number of withdrawal jumps relative to the group treated with nicotine alone. Separately, in a 14-day treatment regimen, mice were administered nicotine (2 mg/kg) and combinations of nicotine with four doses of BZF by the subcutaneous route.

Spontaneous physical and behavioural

signs of nicotine dependence were observed 3 hours after cessation of treatments on the 14th day. Mice administered with combinations of nicotine and 5, 10 and 20 mg/kg doses of BZF, exhibited less intensity and severity of withdrawal effects compared to the mice treated with nicotine alone. Those mice treated with the two highest doses of BZF, in combination with nicotine, showed significantly fewer

**nicotine**-abstinence withdrawal jumps and normal ambulatory behaviour. BZF treatment prevented weight loss and resulted in normal performance in the swimming endurance test, which may be a measure of stress and/or depression. Similarly, acute administration of a single 20 mg/kg dose of BZF prevented some of the nicotine-withdrawal effects; lower doses were almost inert. These studies, although preliminary, suggest that the BZF may have value in treating nicotine addiction. **Alcohol Withdrawal**

A BZF moiety has been reported recently to be responsible for the multifarious CNS effects of *P. incarnate*. In the light of the established usefulness of the BZF moiety in counteracting the withdrawal effects of substances like cannabinoids and nicotine by the authors, the bioactive BZF moiety has been tested in mice treated with an addictive dose of ethyl alcohol, in order to evaluate its effectiveness in countering alcohol dependence. The chronic administration of *P. incarnate* with alcohol had better preventive effects than the single acute treatment with *P. incarnate* in alcohol-dependent mice. These results suggested that the treatment of *P. incarnate* extract could be used as safe and alternative drug for alcohol withdrawal.

#### Anticonvulsant

The current therapeutic treatment of epilepsy with modern antiepileptic drugs (AEDs) is associated with side-effects, dose-related and chronic toxicity, and teratogenic effects, and approximately 30% of the patients continue to have seizures with current AEDs therapy.

Natural products from folk remedies have contributed significantly in the discovery of modern drugs and can be an alternative source for the discovery of AEDs with novel structures and better safety and efficacy profiles. Evidence for anticonvulsant activity of *P. incarnate* in the clonic seizure of pentylenetetrazole model has been tested in mice. As the protective effects of *P. incarnate* in clonic seizure, it suggests that it could

be useful for treatment of absence seizure.

Furthermore, the important role of benzodiazepine receptor in the effects of *P. incarnate* should be considered.

#### Antianxiety

Herbal medicines are popularly used worldwide and could be a natural option for treating anxiety if shown to be effective and safe. Passion flower extract is one of these compounds (35). *P. incarnate* has been used to cure anxiety and insomnia since time immemorial. A fraction derived from the methanol extract of *P. incarnate* has been observed to exhibit significant anxiolytic activity in mice using elevated plus-maze (EPM) model of anxiety. The possibility of a phytoconstituent having BZF nucleus as the basic moiety being responsible for the bioactivity of *P. incarnate* is highly anticipated.

The potential anxiolytic effects of chrysin, a *Passiflora* extract, and the purported modulation of the benzodiazepine receptor on the GABA (A) receptor in laboratory rats has been tested. It has been hypothesized that chrysin decreases anxiety via interaction with the GABA (A) receptor in laboratory rats as measured by elevated plus-maze, corticosterone, and catecholamine assays.

In this study, each group of animals received an intraperitoneal injection of vehicle (DMSO 4%), chrysin, 2 mg/kg, midazolam, 1.5 mg/kg, or flumazenil, 3 mg/kg and chrysin, 2 mg/kg. The EPM was used to evaluate the behavioral component of anxiolysis, and catecholamine and corticosterone assays were examined to measure the neurohormonal effects of anxiety. No statistical difference was found among groups in catecholamine and corticosterone levels. The data suggested that chrysin may have anxiolytic properties similar to midazolam but to a lesser magnitude at the 2 mg/kg dose used in this study (37).

**Aphrodisiac** The isolation of a tri-substituted BZF moiety as the main bioactive phyto-constituent of *P. incarnate* has been an encouraging breakthrough in elucidating the mode of action of this plant, which finds mention in the ancient ayurvedic medical writings as a promising cure for male-impotence, post-menopausal decline in libido in females, menstrual irregularity, morphinism, alcoholism and tobacco addiction.

BZF speeds up the restoration of sexuality in rats upon cessation of the administration of substances like alcohol, nicotine and alcohol-nicotine combinations, which have

severe detrimental effects upon male sexuality, fertility and vigour. BZF, the strongest inhibitor of aromatase enzyme (a member of cytochrome P-450 enzyme family, i.e., CYP3A4) prevents the metabolic conversion of androgens (testosterone) to its metabolites, thereby, increasing the testosterone levels in the gonadal tissue, thus, increasing the free testosterone and decreasing free estrogen.

**HORMONAL**

The testosterone levels in the plasma have an effect upon the gonadotropins (luteinizing hormone LH and follicle-stimulating hormone FSH) which regulate spermatogenesis and maturation of sperms. BZF, when administered concurrently with substances like alcohol and nicotine restores sexual virility, libido and vigour in male rats by maintaining the blood-testosterone level high (40). The aphrodisiac properties of the methanol extract of leaves of *P. incarnate* has been evaluated in mice by observing the mounting behaviour. So this study suggested that the *P. incarnate*, may cause sexual desire in human beings as well.

**Antiasthmatic**

The methanol extract of the leaves of *P. incarnate* was evaluated for its antiasthmatic effects against acetylcholine chloride-induced bronchospasm in guinea-pigs. This may be due to defective alpha-adrenoceptor function reported after excessive or continuous administration of an alpha-receptor agonist. **Antitussive**

The methanolic extract of leaves of *P. incarnate* (100 and 200 mg/kg, p.o.) exhibited significant antitussive activity on sulfur dioxide-induced cough in mice, the cough inhibition being comparable to that of codeine phosphate. These results corroborate the folklore claim on the effectiveness of the plant in managing, "tough" cough conditions. Moreover, *P. incarnate*, that has not been reported anywhere to possess addiction-liabilities, could present advantages over available cough-suppressants (opiates, antihistaminic) which, though acting fast, have several adverse effects including CNS depression, dryness of mouth, blurred vision, severe gastrointestinal effects, and burning micturition.

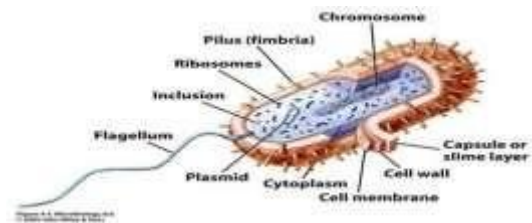
Further studies are, therefore, necessary to evaluate better the potential of *P. incarnate* as an effective cough suppressant. Anticancer The phytochemical composition of passionfruit juice

was hypothesized to have valuable anti-cancer activity. Chrysin, a passion flower extract, may be beneficial because of its potential to attenuate surgical suppression of natural killer (NK) cell activity, thereby minimizing metastatic spread of cancer.

**Hypertension**

Despite improved pharmacotherapies and mechanical treatments, cardiovascular disease remains a principal cause of morbidity and mortality worldwide, with every chance that this burden will increase (46). *P. incarnate* which is an allied species of *Passiflora nepalensis* has already been reported to possess antihypertensive effects. The antihypertensive effect of *P. incarnate* is contributed due to presence of water soluble substance isolated as a mercury salt ( $C_{10}H_{22}O_8NHgCl_2$ ) (3) and flavonoids. *P. nepalensis* is used in folklore medicine for treating hypertension.

**BACTERIA**



**Bacteria structure:-**

The structure of bacteria is known for its simple body design. Bacteria are single celled microorganisms with the absence of the nucleus and other cell organelles; hence, they are classified as prokaryotic organisms. They are also very versatile organisms, surviving in extremely inhospitable conditions.

**Bacteria Used**

1. *S. aureus*:

*Staphylococcus aureus* is a Gram-positive bacterium that typically resides asymptotically in the anterior nares and the skin of mammals. Since its discovery in the 1880s, it has been recognized as a major opportunistic pathogen in humans, responsible for various diseases, ranging from minor skin infections to severe bacteremia and necrotizing pneumonia. Before the era of antibiotics, the mortality rate of patients infected with *S. aureus* exceeded 80%.

1. S.aureus:



2. P.aeruginosa:



*Pseudomonas aeruginosa* is a Gram-negative, rod-shaped, asporogenous, and monoflagellated bacterium. It has a pearlescent appearance and grape-like or tortilla-like odor. *P. aeruginosa* grows well at 25°C to 37°C, and its ability to grow at 42°C helps distinguish it from many other *Pseudomonas* species. *P. aeruginosa* is a ubiquitous microorganism which has the ability to survive under a variety of environmental conditions.

4. S.epidermis:

*S. epidermidis* is a Gram-positive bacterium. Its cell wall teichoic acid is formed by polymerized glycerol, glucose, and N-acetylglucosamine. *Staphylococcus epidermidis*, normally found on human skin, is capable of biofilm formation when it expresses polysaccharide intracellular adhesin (PIA). Production of PIA is a virulence factor that is associated with *S. epidermidis* strains found in opportunistic infections. Phase variation of PIA can occur by transposition of IS256 into biosynthetic genes for PIA, *icaA*, and *icaC*.



3. K.pneumoniae:

*Klebsiella pneumoniae* (Friedlander's bacillus) is a rare cause of community-acquired pneumonia but accounts for a higher proportion of pneumonia acquired in hospital, where patients are more likely to be treated with antibiotics that permit this bacterium to dominate the pharyngeal flora. *K. pneumoniae* is also a particularly common inhabitant of the oral cavity in those with poor dental hygiene and such persons are accordingly at increased risk of *Klebsiella pneumoniae*.

**II. MATERIALS & METHODS (EXPERIMENTAL WORK)**

1. PLANT COLLECTION & EXTRACTION METHOD

**Collection of flower:-**  
 The fresh flowers of *Passiflora incarnata* were collected from cultivated farms and the open fields of Thane district. Fresh flowers were identified and authenticated prior to photochemical analysis. The flowers were separately cut into small bits, and air dried on shadow for two weeks. After dry they were grinded into powder with 1 mm size by using a Grinder machine before being subjected to photochemical screening. The powder was stored in an air tight container and placed in a cool, dry and dark place.

**Powder preparation:-**

The flower samples were rinsed to discard undesirable particles and the samples

were dried inside the room by using an electric fan at ambient for five days. Later on, the dried samples were crushed into coarse powder individually by using a massive duty blender machine. The coarse flowers powder samples (300 mg) were poured into the Soxhlet apparatus.

**Extraction method:-**

**1. Aqueous extraction:-**

Materials: - Powdered flowers, Water distilled, magnetic stirrer. Magnetic beads. Conical flask.

Procedure:- A properly washed conical flask was filled with 250ml of water and accurately weighed with 30gm of flowers powder was mixed in the flask properly. It was placed on a magnetic stirrer and kept for 48 hours. Now the whole aqueous extract was separated using vacuum filtration (Buchner funnel). The extract was concentrated by heating directly and then was finally dried on Hot plate. A brown colored powder was obtained. The powdered extract was collected using a scraper and was collected and stored in a self-sealable pouch.

**2. Ethanol & Petroleum extraction:-** Materials: - Powdered flower of passiflora incarnate, Soxhlet apparatus, heating mantle, Cotton Plug, Round bottom flask.

Procedure: - Properly washed and dried the Soxhlet apparatus and filled accurately weighed 60 gm. of powdered flower. Then the RBF was filled with 150ml of both solvents and fitted with the Soxhlet apparatus. 100ml of solvents was added to the thimble directly.

The Soxhlet extraction was placed on a heating mantle at 45-50°C for about 7 hours.

The extract was then filtered using sintered glass filter and stored in a conical flask. The extract was concentrated using direct heat with constant stirring.



**Chemical Test**

Chemical Test	Observation		
	Aqueous	Ethanolic	Petroleum ether
Protein	Present	Present	Absent
Amino acid	Present	Absent	Absent
Alkaloids	Present	Absent	Absent
Glycosides	Present	Present	Present
Flavonoids	Absent	Present	Absent
Tannins	Absent	Present	Absent
Steroids	Absent	Absent	Absent
Cyanogenic glycoside	Absent	Absent	Absent
carbohydrate	Present	Absent	Present

**AIM:**

Using the FRAP test, the scavenging activity of passiflora flower extract was evaluated.

Chemical: Phosphate buffer (monobasic sodium phosphate and dibasic sodium phosphate, 6.6 pH), plant extract concentration 0.1 percent ferric chloride, 10% trichloroacetic acid, and 0.2 M 1% potassium ferrocyanide.

**• CHEMICAL PROCESS OF MAKING**

1. A mixed phosphate buffer with a pH of 6.8

dissolve to make 1000 ml, mix 13.872 g of potassium

dihydrogen phosphate and 35.084 g of disodium hydrogen phosphate with water.

.Store in a cool area.

2. Potassium Ferrocyanate, also known as potassium hexacyanoferrate(III)  $K_3Fe(CN)_6$ , is a grade of commercial analytical reagent. Crystal in ruby red.

3. Trichloroacetic acid ( $Cl_3COOH=163.40$ ) is a colourless, very lustrous crystal or crystalline mass with a distinctive, mild to pungent aroma. Its melting point (MP) is about 56 degrees.

Store with light protection.

Produce a trichloroacetic acid solution by dissolving 10g of the acid in enough water.

4.  $FeCl_3$  solution at 0.1%

Solution I: Mix 100 ml of hydrochloric acid with the weight of  $Ng$  hexahydrate.

Second option: dissolve Potassium ferricyanide in water, 3.5g per 100ml.

**Procedure:**

1. Take plant extract in different concentrations, 10, 20, 40, and 80. After adding 2.5 ml of phosphate buffer, add 2.5 ml of ferro cyanide at 1%.

2. Combine thoroughly, cover with aluminium foil, and incubate. After 20 minutes of boiling in a water



bath (at a temperature of 50 degrees Celsius), chill it.

2.5 cc of trichloroacetic acid is added after shaking. After centrifuging, take the upper layer and transfer around 2.5 ml to a new test tube. Add 2.5 ml of distilled water and ferric chloride, and the blue colour will absorb.

Assay for ferrous-reducing antioxidant capacity: Calculating the antioxidant impact  
 Antioxidant impact (%) = (control absorbance) - (sample absorbance) x100/(control absorbance).

**Antioxidant activity:**

Using ascorbic acid as a reference, the FRAP method was used to assess the antioxidant activity of the aqueous, ethanolic, and petroleum ether

extracts of the Passifloraincarnata flower.

The aqueous extract's highest antioxidant activity was 96.22 at 80 ug/ml.

The ethanol extract's maximal level of antioxidant activity was 82.83 at 80 ug/ml.

Petroleum ether extract demonstrated the highest percentage of antioxidant activity, which was 97.59 at 80ug/ml.

When comparing the antioxidant activity of the passiflora incarnate flower's aqueous, ethanol, and petroleum ether extracts, it is confirmed that the petroleum ether extract exhibits more powerful antioxidant activity.

**1. ResultforAntioxidant:**

RecordedunderUVspectrophotometeratawavelengthof 700nm.

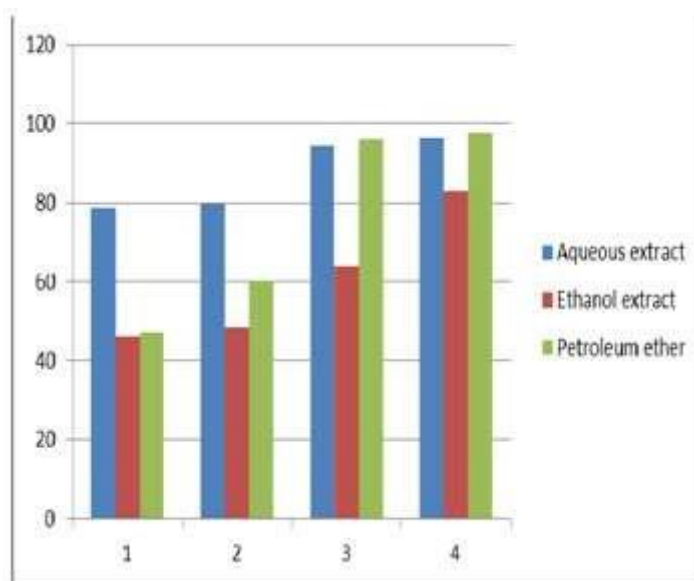
**%ANTIOXIDANTACTIVITY:**

Resultofantioxidant:

Graphicalrepresentationof%antioxidantactivity.

Concentration(ug/ml)	Antioxidanteffect		
	Aqueousextract	Ethanolextract	Petroleume ther
10	0.186	0.470	0.463
20	0.178	0.318	0.348
40	0.049	0.316	0.034
80	0.033	0.15	0.021

Concentration(ug/ml)	%Antioxidant effect		
	Aqueous extract	Ethanol extract	Petroleum ether
10	78.71	46.21	47.01
20	79.63	48.58	60.17
40	94.39	63.84	96.16
80	96.22	82.83	97.59



### III. DISCUSSION:

#### 1. Chemical test:

Preliminary phytochemical analysis revealed that the presence of secondary metabolites like carbohydrates, protein, amino acid, alkaloids, tannin, glycosides in flower of passiflora incarnate.

#### 2. Antioxidant activity:

The scavenging activity of aqueous, ethanol and petroleum ether extract of passiflora incarnate flower extract was determined by FRAP method using ascorbic acid as a standard.

The maximum antioxidant activity was shown by aqueous extract was

96.22 at 80 ug/ml. The maximum antioxidant activity was shown by ethanol extract was 82.83 at 80 ug/ml. The maximum antioxidant activity was shown by petroleum ether extract was 97.59 at 80 ug/ml. While comparing the antioxidant activity between aqueous, ethanol and petroleum ether extract of passiflora incarnate flower confirms that the petroleum ether extract shows more potent antioxidant activity as it shows

#### IV. CONCLUSION:

The research mentioned above demonstrates antioxidant properties, while *P. incarnata*'s petroleum ether extract is the only one to demonstrate antibacterial properties. Modern testing and evaluation (pre-clinical and clinical trials) in various medical conditions have proven the medicinal efficacy of *P. incarnata*, a plant widely employed in Indian system of medicine.

According to these investigations, this natural remedy is a cutting-edge option for drug development and bioprospecting for the treatment of conditions like anxiety, sleeplessness, convulsions, sexual dysfunction, cough, cancer, and postmenopausal syndrome. There are endless opportunities for research into this plant's medical uses and more recent aspects of its function.

Therefore, these plants' phytochemicals and minerals will make it possible to utilise them for therapeutic purposes.

#### V. REFERENCE:

- [1]. G.R. Kinghorn. Passion, stigma and STI. *Sex Transm Inf.* 77:370–75(2001).
- [2]. K. Dhawan, S. Dhawan and A. Sharma. *Passiflora: a review update.* *J Ethnopharmacol.* (2004).
- [3]. The Wealth of India. A Dictionary of Indian Raw Materials and Industrial Products. (2001).
- [4]. S. Akhondzadeh, H.R. Naghavi, M. Vazirian, A. Shayeganpour, H. Rashidi and M. Khani. Passionflower in the treatment of generalized anxiety: a pilot double-blind randomized controlled trial with oxazepam. *J Clin Pharm Ther.* (2001).
- [5]. F.H. Reginatto, F. De-Paris, R.D. Petry, J. Quevedo, G.G. Ortega, G. Gosmann and E.P. Schenkel. Evaluation of anxiolytic activity of spray dried powders of two South Brazilian *Passiflora* species. *Phytother Res.* (2006).
- [6]. D. Wheatley. Medicinal plants for insomnia: a review of their pharmacology, efficacy and tolerability. *J Psychopharmacol.* (2005)
- [7]. Dhawan K, Sharma A. Prevention of chronic alcohol and nicotine-induced azospermia, sterility and decreased libido, by a novel tri-substituted benzoflavone moiety from *Passiflora incarnata* Linn. in healthy male rats. *Life Sci* 2002;71:3059- 69.
- [8]. Dhawan K, Kumar S, Sharma A. Aphrodisiac activity of methanol extract of leaves of *Passiflora incarnata* Linn in mice. *Phytother Res* 2003;17:401-3.
- [9]. Dhawan K, Kumar S, Sharma A. Antiasthmatic activity of the methanol extract of leaves of *Passiflora incarnata*. *Phytother Res* 2003;17:821-2
- [10]. Dhawan K, Kumar S, Sharma A. Antitussive activity of the methanol extract of *Passiflora incarnata* leaves. *Fitoterapia* 2002; 73:397-9.
- [11]. Beaumont DM. The effects of chrysin: a *Passiflora incarnata* extract, on natural killer cell activity in male Sprague-Dawley rats undergoing abdominal surgery. *AANA J* 2008;76:113-7.
- [12]. Benson VL, Khachigian LM, Lowe HC. DNAzymes and cardiovascular disease. *Br J Pharmacol* 2008;154:741-8.
- [13]. Strohmalm H, Dregus M, Wahl A, Engel KH. Enantioselective analysis of secondary alcohols and their esters in purple and yellow passion fruits. *J Agric Food Chem* 2007;55:10339-44.
- [14]. Dhawan K, Kumar S, Sharma A. Nicotine reversal effects of the benzoflavone moiety from *Passiflora incarnata* Linn in mice. *Addict Biol* 2002;7:435-41
- [15]. Dhawan K, Kumar S, Sharma A. Suppression of alcohol-cessation-oriented hyper-anxiety by the benzoflavone moiety of *Passiflora incarnata* Linn. in mice. *J Ethnopharmacol* 2002;81:239-44.
- [16]. Miyasaka LS, Atallah AN, Soares BG. *Passiflora* for anxiety disorder. *Cochrane Database Syst Rev* 2007;24:CD004518.
- [17]. Dhawan K, Kumar S, Sharma A. Anti-anxiety studies on extracts of *Passiflora incarnata* Linnaeus. *J Ethnopharmacol* 2001;78:165- 70.



- [18]. Brown E, Hurd NS, McCall S, Ceremuga TE. Evaluation of the anxiolytic effects of chrysin: A *Passiflora incarnata* extract, in the laboratory rat. *AANA J* 2007;75:333-37.
- [19]. Chen S, Kao YC, Laughton CA. Binding characteristics of aromatase inhibitors and phytoestrogens to human aromatase. *J Steroid Biochem Mole Biol* 1997;61:107-15.
- [20]. Dhawan K, Sharma A. Prevention of chronic alcohol and nicotine-induced azoospermia, sterility and

- decreased libido, by a novel tri-substituted benzoflavone moiety from *Passiflora incarnata* Linn. in healthy male rats. *Life Sci* 2002;71:3059-69.
- [21]. Dhawan K, Kumar S, Sharma A. Antiasthmatic activity of the methanol extract of leaves of *Passiflora incarnata*. *Phytother Res* 2003;17:821-2.
- [22]. Dhawan K, Kumar S, Sharma A. Antitussive activity of the methanol extract of *Passiflora incarnata* leaves. *Fitoterapia* 2002;73:397-9.
- [23]. Beaumont DM. The effects of chrysin: A *Passiflora incarnata* extract, on natural killer cell activity in male Sprague-Dawley rats undergoing abdominal surgery. *AANA J* 2008;76:113-7