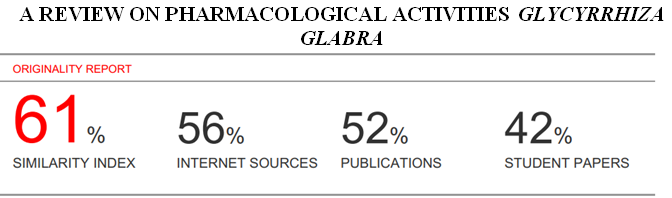
**Reviewer’s Comments**

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**A review on pharmacological activities *glycyrrhiza glabra***

**ABSTRACT:**

Plants have been one of the important sources ofmedicines for human being and animals since the ancient time. At present scenario there is an increasing demand for herbal medicines, health products and pharmaceuticals products. Herbal medicines have attained popularity at global level to replace the synthetic chemicals as they have shown less adverse reactions.

*Glycyrrhiza glabra* Linn is a commonly used herb for different diseases. Present review article deals with chemical constituents present in various parts of *Glycyrrhiza glabra* and pharmacological activities. Present article aim to comply all the updated information on its phytochemical and pharmacological activities,which were performed by widely different methods. *Glycyrrhiza glabra Linn* possesses antibacterial, antioxidant, antimalarial, antispasmodic, anti-inflammatory and anti-hyper glycemic properties. Various other effects like antiulcer, antiviral, antifungal have also been discussed.

This article may be useful for many researchers in discovering potential therapeutic effects and developing new formulations.

**Keywords:** *Glycyrrhiza glabra,* glycyrrhizin, antibacterial,antioxidant, anti malarial.

**INTRODUCTION**

Human beings have used plants for medicinal purposes forcenturies. It has been estimated that such use of medicinal plants possibly go back in time to around 3000 years. Today, a great percentage of the world population, particular in developing countries, uses plants for facing primary needs of medical assistance1. Modern drugs like aspirin, atropine, ephedrine, digoxin, morphine, quinine, reserpine and tubocurarine are examples, which were originally discovered through observations of traditional cure methods of indigenous people2.

*Glycyrrhiza glabra* has been known in pharmacy forthousands of years. In old Chinese pharmacy, it was considered to belong to drugs of the first class and to it was ascribed the rejuvenating property when consumed for long periods3. In ancient Egypt, Greece and Rome licorice was frequently used. It was referred to by Theophrastus. The earliest record of its use in medicine is found in „code Humnubari‟ (2100 BC)4. It was also one of the important plants mentioned in Assyrian herbal (2000BC). Hippocrates (400BC) mentioned its use as a remedy of ulcers and quenching of thirds. The drug was also mentioned by Theophrastus and Dioscorides. In traditional Siddha system of medicine, liquorice is used as a demulcent,expectorant, anti-tussive, laxative and sweetener5.

**CLASSIFICATION6, 7, 8, 9**

|  |  |
| --- | --- |
| Kingdom | Plantae |
| Subkingdom | Tracheobionta-Vascular plants |
| Super division | Spermatophyta-Seed plants |
| Division | Magnoliophyta |
| Class | Magnoliopsida |
| Subclass | Rosidae |
| Order | Fabales |
| Family | Fabaceae-Pea family |
| Genus | *Glycyrrhiza* L*.-* licorice |
| Species | *Glycyrrhiza glabra* L |

**DISTRIBUTION/ HABITAT:**

This plant is cultivated in Russia, UK, USA, Italy, France, Germany, Spain, China and Northern India (Punjab and Sub-Himalayan tracts). It is distributed in Southern Europe, Syria, Iran, Afghanistan, Russia, China, Pakistan and Northern India. Largescale commercial cultivation is seen in Spain, Sicily and England10.

**CULTIVATION11, 12, 13**

It is a perrineal herb/ sub shrub for subtropical and temperate zone. Young pieces of stolons having 2-3 buds of aerial shoot are used for propagation. 3 to 4 feet deep soil having light, loamy, stone-free texture is needed for the plant. It is usually grown continuously on the same land. Licorice enjoys fertile, sandy or clay soil near a river or stream where enough water is available for the plant to flourish in the wild, or under cultivation where it can be irrigated.

The pieces of stolons are planted in March at 2' by 3' distance.The plant attains a maximum height up to 2 m. The underground stem grow horizontally up to 2 m length, highly branched consisting of short taproot with large number of rhizomes. The roots are harvested 3-4 years after planting when they show sufficient growth. Dry conditions at planting time and for the next two months give best chance for a good crop. A yield of two tons of roots per acre for bailing, plus 3-4 cwt of trimmings or offal is considered satisfactory. Rhizomes and roots are dug up in October, from the plants which have not borne the fruits. Drug is washed after removal of buds and rootlets. Some pieces are peeled and divided into small pieces. Thedrug is dried first under sun and then in shades, during which it loses about 50 per cent of its weight.

**MORPHOLOGY14, 15, 16**

**Leaves**- are compound, imparipinnate, alternate, having 4-7 pairs of oblong, elliptical or lanceolate leaflets covered with soft hairs on underside.

**Flowers-** are narrow, typically papilionaceous, borne in axillary spikes, lavender to violet in color. The calyx is short, campanulate, with lanceolate tips and bearing glandular hairs.

**Fruit-** is a compressed legume or pod, upto 1.5 cm long, erect, glabrous, somewhat reticulately pitted, and usually contains, 3-5 brown, reniform seeds.

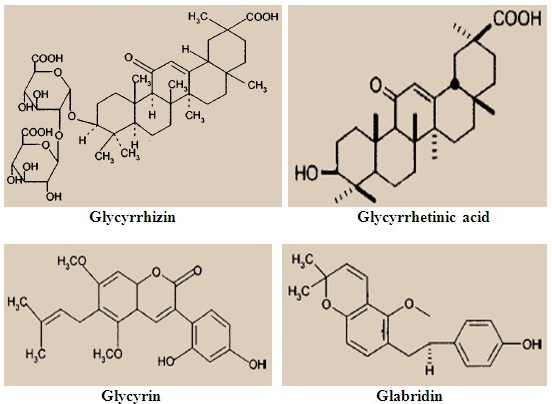
**Root**- is approximately 1.5 cm long and subdivides into subsidiary roots, about 1.25 cm long, from which the horizontal woody stolons arise. They may reach 8 m andwhen dried and cut, together with the root, constitute commercial licorice. It may be found peeled or unpeeled. The pieces of root break with a fibrous fracture, revealing the yellowish interior with a characteristic odor and sweet taste.

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**Figure 1: Leaves, flower, root and powder of *Glycyrrhiza glabra***

**CHEMICAL CONSTITUENTS17, 18, 19**

Flavonoid rich fractions include liquirtin, isoliquertin liquiritigenin and rhamnoliquirilin and five new flavonoids-glucoliquiritin apioside, prenyllicoflavone A, shinflavanone, shinpterocarpin and 1- methoxyphaseolin isolated from dried roots. The roots of *Glycyrrhiza glabra* Linn contain glycyrrhizin, which is a saponin that is 60 times sweeter than cane sugar. Both glycyrrhizin and glycyrrhetic acid can exist in the 18α and 18β stereoisomers. Isolation and structure determination of licopyranocoumarin, licoarylcoumarin, glisoflavone and new coumarin-GU-12 also isolated. Four new isoprenoid-substituted phenolic constituents-semilicoisoflavone B, 1-methoxyficifolinol, isoangustone A, and licoriphenone isolated fromroots. A new prenylated isoflavan derivative, kanzonol R was also isolated.



**Figure 2: Chemical constituents of *Glycyrrhiza glabra***

The presence of many volatile components such as pentanol, hexanol, linalool oxide A and B, tetramethyl pyrazine, terpinen-4-ol, α- terpineol, geraniol and others in the roots is reported. Presence of propionic acid, benzoic acid, ethyl linoleate, methyl ethyl ketine, 2, 3-butanediol, furfuraldehyde, furfuryl formate, 1-methyl-2- formyl pyrrole, trimethyl pyrazie, maltol andany other compounds is also isolated from the essential oil.

**QUANTITATIVE STANDARDS20, 21, 22**

|  |  |
| --- | --- |
| **Total ash** | Not more than 4.5 % |
| **Acid insoluble ash** | Not more than 1.5% |
| **Water soluble ash** | Not more than 1% |
| **Aqueous extractive value** | Not less than 20% |
| **Ethanolic extractive value** | Not less than 15% |
| **Chloroform extractive value** | Not less than 5% |
| **Water soluble extractive** | Not less than 20% |
| **Moisture content** | Not more than 7.45% |

**Pharmacology**

After oral administration of licorice in humans, the main constituent, glycyrrhizic acid, is hydrolyzed to glycyrrhetic acid by intestinal bacteria possessing a specialized ß-glucuronidase. Glycyrrhetic acid is 200-1,000 times more potent an inhibitor of 11-ß-hydroxysteroid dehydrogenase (involved in corticosteroid metabolism) than glycyrrhizic acid23. After oral dosing, glycyrrhetic acid is rapidly absorbed and transported via carrier molecules to the liver. In the liver it is metabolized to glucuronide and sulfate conjugates, which are subsequently rehydrolyzed to glycyrrhetic acid24. Glycyrrhetic acid is thenreabsorbed, resulting in a significant delay in terminal clearance from plasma. In the 24-hour period after oral administration, glycyrrhizin was found in the urine, indicates it’s partly absorption as an intact molecule25.

**TRADITIONAL USES26, 27, 28, 29**

1. A decoction of madhuka or its powder is generally prescribed with honey in anemia.

2. After mixing with cow‟s milk it is used for promoting lactation.

3. 10g powder of it mixed with 10g sugar, pounded with rice water is commonly prescribed in men-metrorrhagia.

4. A confection of rice milk, prepared with Yashtimadhu, is usedtreatment of hoarseness of voice.

5. Charaka prescribed 10 g madhuka powder mixed with honey, followed by intake of milk, as an aphrodisiac and as an intellect-promoting tonic.

6. Charaka prescribed paste of licorice and *Picirrhiza kurroa* with sugar water as a cardio-tonic.

7. Charaka prescribed Yashtimadhu and *Santalum album*, powdered with milk in haematemisis.

8. Sushrata prescribed the paste of Yashti madhu 10g in intrinsic haemorrhage.

9. In oedema, paste of licorice and Sesamum indicum, milk mixed with butter is used.

10. Warm clarified butter mixed with licorice, is used topically on wounds,bruises and burns.

11. A decoction of madhuka is used on erysipelas.

12. A decoction of the root is a good wash for falling and greying of hair.

**Pharmacological activities**

**1. Anti-tussive and Antidemulcent activity**

The liquorice powder and extract was found to be effective in treatment of sore throat, cough and bronchial catarrh. Liquorice has been shown to work as efficiently as codeine in sore throat. It decreases irritation and produces expectorant effects. Carbenoxolone (a semi synthetic compound derived from *Glycyrrhiza*) stimulates gastric mucus secretion. Glycyrrhizin is responsible for demulcent action of liquorice. Liquiritin apioside, an active compoundpresent in the methanolic extract of liquorice which inhibits capsaicin induced cough30.

**2. Antioxidant activity**

*Glycyrrhiza* have a significant free‐radical quenching effect. Liquorice flavonoids have exceptionally strong antioxidant activity. Antioxidant activity of liquorice flavonoids was found to be over 100 times stronger than that of antioxidant activity of vitamin E. Thus, liquorice extract can be efficiently used to formulate cosmetic products for the protectionof skin and hair against oxidative damage31, 32.

**3. Anti-malarial activity**

Licochalcone A (a chalcone) present in liquorice is responsible for antimalarial activity. A previous reported *In and* study against *P. yoelii* in mice with oral doses of 1000 mg kg-1 have shown to eradicate malarialparasite completely33.

**4. Anti-fungal activity**

*Glycyrrhiza glabra* possess good anti fungal activity. In a previous reported study of screening for antifungal compounds from various plant material, licorice extracts with 80% methanol (oil‐based extract of licorice; OEL) was found to have high fungicidal effect against Arthrinium sacchari M001 and Chaetomium funicola M002,and its active compoundwas identified as glabridin34. Thus, liquorice extract has a great potential in formulating cosmetic products with antiseptic activities.

**5. Immunostimulatory activity**

*In and* studies proved that *Glycyrrhiza glabra* at 100μg/ml concentration, showed immunostimulatory effects. It increases production of TCD69 lymphocytes and macrophages from human granulocytes. In a previous reported study, liquoriceroot extract was found to prevent the rise in the amount of immune-complexes related to autoimmune diseases like systemic lupus erythematosus35.

**6. Anti ulcer activity**

Glycyrrhizinic acid, a major component of licorice, has antiulcer effect by raising the local concentration of prostaglandins that promote mucous secretion and cell proliferation in the stomach. In a previous reported study *in and* activity of Extractum liquiritiae (EL), glycyrrhizic acid, glycyrrhetinic acid and a novel lipophilic derivative of glycyrrhetinic acid monoglucuronide (GAMG), acetylated GAMG(aGAMG), against 29 Helicobacter pylori strains. The potent in vitro activity of glycyrrhizic acid against H. pyloriconcludes its beneficial effect on peptic ulcers36.

**7. Anti dysliipidaemic effect**

In a previous reported study ethanolic (95%) extract of root of *Glycyrrhiza glabra* and its fractions were investigated for its antidyslipidaemic activity on dyslipidaemic hamsters.

The reduction in LDL‐cholesterol level by ethanolic extract, ethyl acetate soluble fraction and water soluble fraction were 43.9, 31.0, 33.4 and 24.6%, respectively. The treatment with *Glycyrrhiza glabra* root ethanolic extract and its fractionssignificantly brought down LDL and VLDL in the HFD fed hamsters to various degrees37.

**8. Anti-bacterial Activity**

Secondary metabolites such as; saponins, alkaloids, flavonoids are present in hydro-methanolic root extract of *Glycyrrhiza glabra,* possess potent antibacterial activity. *In and* studies have proved that aqueous and ethanolic extracts of liquorice show inhibitoryactivity on cultures of *Staphylococcus aureus* and *Streptococcus pyogenes38*.

**9. Anti-viral effects**

It is reported that liquorice extract inhibits the growth of viruses, including herpes simplex, Varicella zoster, and of Japanese encephalitis, influenza virus, vesicular stomatitis virus, type A influenza virus. Glycyrrhizin does not allow the virus cellbinding. Thus, it is found to have a prominent antiviral activity39.

**10. Antithrombotic effect**

In a previous reported study the in-vivo effects of extract of *Glycyrrhiza glabra* and also the combined effect with Vitamin K and Heparin was evaluated in Sprague Dawley Rats. Extract of *G.glabra* increased the bleeding time when given in the doses of 180 mg/kg and 360 mg/kg. Blood loss was evaluated 60 minute later as a function of absorbance at 540 nm due to hemoglobin content in water solution. Altogether data indicates that *Glycyrrhiza glabra* is an effective anti thrombotic agent *in and40*.

**11. Hair growth stimulation**

Liquorice has a significant hair growth activity and it can be safely used in herbal formulations in treatment of various types of Alopecia. In a previous reported study hydro-alcoholic extract of liquorice showed good hair growth promoting activity. Comparisonbetween liquorice extract and the standard drug used (Minoxidil 2%) showed that, 2% concentration of liquorice extract showed better hair growth stimulatory activity than 2% Minoxidil41.

**12. Skin lightening activity**

The extract of liquorice is reported to be an effective pigment lightening agent. Glabridin in the hydrophobic fraction of liquorice extract inhibits tyrosinase activity in cultured B16 murine melanoma cells. Some other active compounds in liquorice extract like glabrene, Licochalcone A, Isoliquiritin are also responsible for inhibition of tyrosinaseactivity. Liquiritin present in liquorice extract disperse melanin, thereby inducing skin lightening42.

**13. Anti-inflammatory activity**

It is reported that glycyrrhetinic acid in liquorice extract gives anti-inflammatory effect similar

to glucocorticoids and mineralocorticoids Liquorice root (Glycyrrhiza) extract promotes the healing of ulcers of the stomach and mouth. The fact was known for over 2000 years.

According to *In and* studies, glycyrrhizic acid inhibits all factors responsiblefor inflammation. It inhibits cyclooxygenase activity andprostaglandin formation. It isalso responsible for indirectly inhibiting platelet aggregation43.

**CONCLUSION**

At present scenario there has been an increase in demand for the phytopharmaceuticals all over the world because of the fact that the allopathic drugs have more side effects. This forms a basis for the selection of plant for further phytochemical and pharmacological investigation.

The pharmacological activities reported in the present review confirm the therapeutic value of *Glycyrrhiza glabra.* The plant has beenused since centuries for asthma, bronchitis,ulcers, and an anti-inflammatory. It is reportedto contain essential oil, coumarins, alkaloidsand flavonoids. Extract of root can be found in various herbal preparations that are in market

today.Presence of chemical compounds indicates that the plant couldserve as “lead” for development of novel agents for disorders in the coming years.

In this regard, further studies need to be carried out to explore *Glycyrrhiza glabra* Linn for its potential in preventing and treating diseases. So, the present review gives a direction for future investigators to carry out research on the plant so that they could get some medicinally important drugs or may design new dosage form of active constitutents.

**REFERENCES:**

1. Mazumdar PM., Patnayak SP., Parwani H. Evaluation of immunomodulatory activity of *Glycyrrhiza glabra* roots in combination with zinc. *Asian pac J of Trop Med*. 2012, S15-S20.
2. Sowmya M, Kumar S. Antistress property of *Glycyrrhiza glabra* on stress induced Drosophila melanogaster*. J of Str Phys and Bioch*. 2010, 6: 18-27.
3. Jatav VS, Singh SK, Sharma AK. Recent Pharmacological trends in *Glycyrrhiza glabra*. *Int J of Pharm Fron Res*. 2011, 1: 170-85.
4. Lakshmi T, Geetha RV. *Glycyrrhiza glabra* commonly known as licorice- a therapeutic review. *Int J of Pharm and Pharm Sci*. 2011, 3: 20-25.
5. Kumar A, Dora J. Review on *Glycyrrhiza glabra*: licorice. J of Pharm and Scient Innov. 2012, 1: 1-4.
6. Ramar PS, Peter NP, Ponnampalam G. A compilation of bioactive compounds from Ayurveda; Biomedical Informatics Publishing Group. 2008, 3(3), 100-110.
7. Karahana F, Avsarb C, Ilker IO, Berber I. Antimicrobial and antioxidant activities of medicinal plant *Glycyrrhiza glabra* var. glandulifera from different habitats, Biotechnology and Biotechnological Equipment. 2016:1-8.
8. Tene V, Malago O, Finzi PV, Vidari G. An ethnobotanical survey of medicinal plants used in Loja and Zamora chinchipi*, Equador. J of Ethnopharm*. 2007, 111: 63-81.
9. Kalaiarasi P, Pugalendi KV. Antihyperglycemic effect of 18 beta glycyrrhetinic acid, aglycone of glycyrrhizin on streptozotocin diabetic rats. *Europ J of Pharm*. 2009; 1: 269-73.
10. Mukherjee PK, Wahile A. Integrated approaches towards drug development from Ayurveda and other Indian system of medicines. *J of Ethnopharm*. 2006, 103, 25-35.
11. Al Razzuqii RAM., Al-Hussaini JA. Hepatoprotective effect of *Glycyrrhiza glabra* in CCl4 induced model in acute liver injury. *Journal of Physiology and Pharmacology Advances*. 2012; 2: 259-63.
12. Cooper H, Bhattacharya B, Verma V. Liquorice and Soy sauce, a life-saving concoction in a patient with addisons disease. *Ann Clin Biochem*. 2007, 44: 397-9.
13. Kamei J, Nakamura R, Ichiki H, Kubo M. Antitussive principles of Glycyrrhiza radix, a main component of Kampo preparations Bakumondo-to (Mai-men-dongtang). *E J Pharm*. 2003; 69:159-163.
14. Nitalikar M, Munde KC, Dhore BV. Studies of antibacterial activities of *Glycyrrhiza glabra* root extract. *Int Jof Pharm Tech and Res.* 2010; 2: 899-901.
15. Dhingra D, Parle M, Kulkarni SK. Memory enhancing activity of *Glycyrrhiza glabra* Linn in mice. *J Ethnopharmacology*. 2004; 91(2-3):361-365.
16. Takii H, Kometani T, Nishimura T, NakaeT, Okada S and Fushiki T, Anti-diabetic effect of *Glycyrrhizin* in genetically diabetic KK-Ay mice, *Biol Pharm Bull*. 2000, 24:484-487.
17. Lee CK, Park KK, Lim SS. Effects of licorice extracts against tumor growth and cisplatin induced toxicity in a mouse xenograft model of colon cancer. *Biol Pharm Bulletin*. 2007; 30: 2191-5.
18. Alaaeldin AH. *Curcuma longa*, *Glycyrrhiza glabra Linn* and *Moringa oleifera* Ameliorate Diclofenac-inducedHepatotoxicity in Rats. *American J Pharmacol and Toxicol*. 2007; 2(2):80-88.
19. Jahan Y, Siddique HH. Study of antitussive potential of *Glycyrrhiza glabra* and *Adhatoda vasica* using a cough model induced by SO2 gas in mice. *Int J of Pharm Sci and Res*. 2012, 3: 1668-74.
20. Trivedi R, Sharma K. Hydroalcoholic extract of *Glycyrrhiza glabra* attenuates chronic fatigue stress induced behavioral alterations in mice. *Int J of Pharm and Biolog Sci*. 2011, 2: 996-1001.
21. Kalaigandhi V, Poovendran P. Antimicrobial activity of *Glycyrrhiza glabra* against peptic ulcer produced *Helicobacter pylori*. *Int J of Curr Pharm Res*. 2011; 3: 93-95.
22. Latif M., Iqbal L., Fatima N. Evaluation of antioxidant and urease inhibition activity of roots of *Glycyrrhiza glabra*. *Pakistan J of Pharm Sci*. 2012, 25: 99-102.
23. Visavadiya NP, Soni B, Dalwadi N. Evaluation of antioxidant and anti-atherogenic properties of *Glycyrrhiza glabra* root using In vitro models. Int J of Food Sci and Nut. 2009; 60(2):135-149.
24. Chakravarthi KK., Avadhani R. Effect of *Glycyrrhiza glabra* root extract on learning and memory in Wistar albino rats. *Drug Inv Today*. 2012, 4: 387-90.
25. Gupta N, Belemkar S, Gupta PK, Jain A. Study of *Glycyrrhiza glabra* on glucose uptake mechanism in rats, *Int Jof drug disc and herbal res*. 2011, 1(2): 50-51.
26. Mirmala P, Selvaraj T. Anti-inflammatory and antibacterial activities of *Glycyrrhiza glabra*. J of Agri Techn. 2011, 7: 815-23.
27. Chowdhury B, Bhattamisra SB, Das MC. Anti-convulsant action and amelioration of oxidative stress by *Glycyrrhiza glabra* root extract in pentylenetetrazole- induced seizure in albino rats, *Indian J Pharmacol*. 2013; 45(1): 40–43.
28. Xu-ying W, Ming L, Xiao-dong L and Ping H. Hepatoprotective and anti hepatocarcinogenic effects of glycyrrhizin and matrine. *J Chemico-Biological Interac*. 2009; 181(1):15-19.
29. Yazdi A, Sardari S, Sayyah Md. Evaluation of anticonvulsant activity of leaves of *Glycyrrhiza glabra* grown in Iran as apossible renewable source for anticonvulsant compounds*. Iranian J of Pharm res*. 2011, 10(1), 75-82.
30. Anderson DM, Smith WG. The antitussive activity of glycyrrhetinic acid and its derivatives*. J. Pharm.Pharmacol*. 1961, 13:396‐404.
31. Kiso Y, Tohkin M, Hikino H. Mechanism of antihepatotoxic activity of glycyrhhizin, I: Effect on free radical generation and lipid peroxidation. Planta Medica. 1984; 50:298‐302.
32. Demizu S, Kajiyama K, Takahashi Y, Hiraga S, Yamamoto Y, Tamura K, Kinoshita T. Antioxidant and antimicrobial constituents of *licorice*: isolation and structure elucidation of a new benzofuran derivative. Chem. *Pharm. Bull*. 1988 36:3474-3479.
33. Sianne S, Fanie RVH. Antimalarial activity of plant metabolites. Nat Prod Rep. 2002; 19:675-692.
34. Hojoa H, Satob J. Antifungal Activity of Licorice (*Glycyrrhiza glabra*) and potential applications in beverage foods, *Food Ingredients J*. Japan. 2002, 203.
35. Ju HS, Li XJ, Zhao BL, Han ZW, Xin WJ. Effects of Glycyrrhiza Flavonoids on lipid peroxidation and active oxygen radicals. Acta Pharm Sinicia. 1989; 24(11):807-812.
36. Krausse R, Bielenberg J, Blaschek W, Ullmann U. *In and* anti‐ Helicobacter pylori activity of Extractum liquiritiae, glycyrrhizin and its metabolites. *J Antimicrob Chemother*. 2004, 54(1): 243-246.
37. Kim YW, Kang HE, Lee MG, Hwang SJ, Kim SC, Lee CH. Liquiritigenin, a flavonoid aglycone from licorice, has a choleretic effect and the ability to induce hepatic transporters and phase‐II enzymes. *Am J Physiol Gastrointest Liver Physiol*. 2009; 296(2):372-381.
38. Sharma V, Agrawal RC, Pandey S. Phytochemical screening and determination of anti-bacterial and antioxidant potential of *Glycyrrhiza glabra* root extracts. *J Environ Res Develop*. 2013; 7(4A):1552-1558.
39. Pompei R, Pani A, Flore O, Marcialis MA, Loddo B. Antiviral activity of glycyrrhizic acid. Experientia. 1980, 36:304.
40. Mendes-Silva W, Assafim M, Ruta B, Monteiro RQ, Guimaraes JA, Zingali RB. Antithrombotic effect of Glycyrrhizin, a plant-derived thrombin inhibitor. Thromb Res. 2003; 112:93-98.
41. Roy SD, Karmakar PR, Dash S, Chakraborty J, Das B. Hair growth stimulating effect and phytochemical evaluation of hydro-alcoholic extract of Glycyrrhiza glabra, *Global J res Med plants and indigen med*. 2014; 3(2):40-47.
42. Cronin H, Draelos ZD. Top 10 botanical ingredients in 2010 anti-aging creams. *Journal of Cosmetic Dermatology. 2010; 9(3):218-225.*
43. *Ohuchi K, Tsurufuji A.* A study of the anti-inflammatory mechanism of glycyrrhizin. *Mino med rev*. 1982; 27:188- 193.