

## An Overview of Medicinal Importance of SWERTIA CHIRAYITA

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### Abstract

*Herbs orchestrate a resurgence and vegetal awakening is supervened every where in the world. Vegetal commodities currently illustrate assurance as compared to the factitious ones that are contemplate as alarming to humans and environment. Out of 2,50,000 higher plant species on this planet, more than 80,000 types are declared to have in some ways remedial importance and around 5000 species have characteristic analeptic value. Organized storage and commodious plowing of relevant medicinal plant species are thus of ample precedence. An important herb Swertia chirayita, is a medicinal plant aboriginal to clement Himalayas in India, Nepal and Bhutan. Its medicinal usage is declared in American and British pharmacopoeias, Indian Pharmaceutical codex and in different conventional systems of medicines like Ayurvedic, Unani and Sidha. Plants mainly utilize in Ayurveda can contribute organically active compounds and lead structures for the advancement of transformed subordinates with increased activeness and abate virulence. We are well enumerate as the most paramount chirayita producer and vendor based in India. The chief bioactives of Swertia are Xanthones, other active constituents of this genus are the secondary metabolites which played a momentous role in biological activities like being hepatoprotective, digestive, astringent, laxative, anti-inflammatory and anti-malarial. Hence this herb provides potent therapeutic lead compounds, which would be beneficial for mankind.*

**Key Words:** *Swertia Chirayita* , orchestrate, Phyto-constituents, bioactives, momentous .

### Overview

Medicinal plants always played a important role in the health development of mankind. In developing countries, 80% of populations are totally dependent on plants for their primary health care. Over 25% of prescribed medicines in industrialised countries derive directly or indirectly from medicinal plants. A multidisciplinary approach combining botanical, ethnobotanical, phytochemical and biological techniques led to Drug discovery from plant (Newman et.al, 2000). Plants provide us new lead molecules for the development of drugs against various pharmacological targets.

Medicines based on plants were dispensed earlier in the form of crude drugs such as tinctures, teas, powders, and other herbal formulations, which now serve as the basis of novel drug discovery. Discovery of drugs from plants has traditionally been time-consuming ,so faster methods for plant collection, bioassay screening, isolation and development of compound must be adopted (Karan et.al, 1996). Chirayita provides us new lead molecules for the development of drugs against various pharmacological targets. Plants included in this family are annual and persistent herbs or shrubs, indigenous to northern moderate stretch of the world (Daniel and Sabnis, 1978).

*Swertia Chirayita* is also known as Haima, kirata Tikta, Nidrari, Ramasenka, kairata in Sanskrit, in urdu language it is called Chiravata, Chireta in Bengal and in Arabic and Farsi called as Qasabuzzarirah. Chiretta is its market name (Anon, 1978 ; kirtikar and K.R ,1984).

And this herbal drug 'Chiretta' is gathered from dried plants of *Swertia* species. Although full-length plants of *Swertia* are medicinally important but roots are manifold paramount (Anonymous, 1976). Chiretta is available in Indian medical conformity as therapy for different kinds of disorders like diarrhea, never ending fever, anemia, liver function disorders and bronchial asthma. *Swertia chirayita* (Roxb. ex Fleming) H. Karst. is primitive to moderate Himalaya observed at an eminence of 1200–3000 m (4000 to 10,000 ft), from Kashmir to Bhutan, and in the Khasi hills at 1200–1500 m (4000 to 5000 ft) (Kirtikar and K.R., 1984; Clarke and C.B., 1885). *Swertia chirayita* has also been remarked in the biography as *Swertia chirata*, Buch-ham; *Ophelia chirata* Grisebach; *Agathotes chirayita* Don; *Gentiana chirayita* Roxburge (Anon, 1978; Kirtikar et al., 1984; Duke and J.A., 2002; Clarke et al. 1885; Anonymous, 1976). It is distinguished by a parade of names, recommending its extensive applicability. As pinpointed by National Medicinal Plant Board, Government of India, *chirayita* is in the midst of the 32 awful pre eminent medicinal herbs in the affluent biodiversity of Uttarakhand (India).

It is ingathered for drug industry (Bentley and Trimen, 1880). It is called as elixir and immersion in American and British pharmacopoeias (Joshi and Dhawan, 2005). *Chirayita* has an organized vend both domestic in India as well as globally and it is increasing at an estimate of 10% every year. Nepal is affluent in breed variance of *Swertia*, by virtue of enormous assortment of geomorphological aspect and plenty of contrasting environs. *Chirayita* is named in Nepal as tite, chiraito and pothi chiraito and 45% of total *chirayita* in the region of Himalaya gathered from Nepal. Currently in Nepal, more steps have been taken to ethnological studies of the species on the basis of molecular differences.

### Important pharmacological effects

*S. chirayita* is used as antipyretic, anthelmintic, antiperiodic, cathartic and in asthma and leucorrhoea in Ayurveda and as harsh, analeptic, stomachic, mitigate inflammation, relaxing to pregnant uterus and never ending fevers (Kirtikar and Basu, 1984). It is a remedy for ulcers, Gastrointestinal diseases, skin diseases, cough, hiccup, liver and Kidney diseases, Neurological disorders, and urinogenital tract disorders. Also used as purifier of Breast milk, and as a laxative and carminative (Garg 1965 and Sharma 1986). Pharmacological studies on medicinal species belonging to family Gentianaceae were consummate earlier, 1930. Comprehensive work was done on an isoprene alkaloid called Gentianine, enunciate to have divergent pharmacological effects categorizing from anti-inflammatory to diuretic.

### Investigated for Drug reinforcement

Important phytochemicals like Amarogentin and Swerchirin have been investigated for drug reinforcement (Brahamchari et al., 2004).

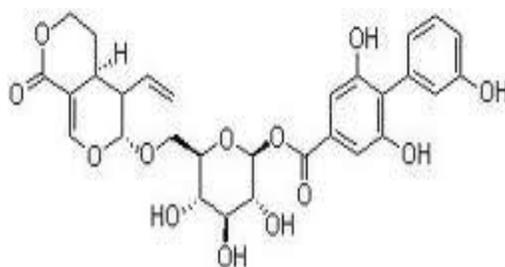
### Medicinal and pharmaceutical importance

*Swertia Chirayita* is known for its medicinal and pharmaceutical importance. It is a filthy provenance of alkaloids and flavanoids, most of them having ample scale exercise. Their roots have considerable antipyretic and analgesic effects and a high rise therapeutic clue. It is having a large number of chemical constituents estimating more than twenty polyhydroxylated xanthenes and some of these are swertinin, swerchirin, mangiferin, decussatin and isobellidifolin; a dimeric xanthone and chiratanin has also been segregated (Bhattacharya et al., 1976). Important photochemicals like Amarogentin and Swerchirin have been investigated for drug reinforcement (Brahamchari et al., 2004). *Chirayita* has an authorized sedentary (India) and global market developing progressively 10% annually.

### Chemical compounds residing in chirayita

#### Amarogentin (chirantin)

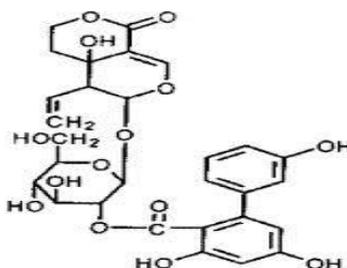
It is secoiridoid glycoside, and is the most acerbic substance found. It tastes bitter even at a dilution of 1:58,000,000 and can be procured from, *Swertia chirayita* (Roxb ex. Flem) Karst (Arino et al., 1997). It acquires Topoisomerase inhibition (Ray et al., 1996), chemo-preventive (Saha and Dass, 2005) and antileishmanial effects (Ray et al., 1996; Medda et al., 1999).



**Fig. 1 : A biphenylcarboxylic acid moiety; biosynthesized by a polyketide-type pathway, with three units of acetyl-CoA and one unit of 3-hydroxybenzoyl- CoA (figure1; chemicalbook.com)**

### Amaroswerin

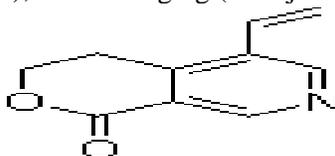
It is a Secoiridoid glycoside collected from *Swertia chirayita* and found to be gastro-sterilizing (Niiho et al., 2005).



**Fig. 2: An iridoidal glycoside having molecular formula  $C_{29}H_{30}O_{14}$  (Figure 2 ; Plant -expert.com)**

### Gentianine

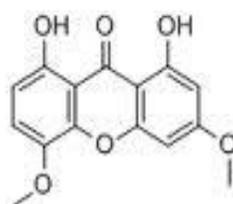
A sullen, translucent monoterpene alkaloid, obtained from several plant species of family Gentianaceae including *Swertia chirayita* (Purushothaman et al., 1973). It possesses anti-inflammatory, anesthetic antihistaminic ,anticonvulsant properties (Song et al., 1958; Tao et al., 1959; Kwak et al., 2005). And also having hypotensive, antipsychotic (Bhattacharya SK et al., 1974) lenitive, diuretic (Mansoor and Malghani, 2005) antimalarial ,antiamoebic and antibacterial properties(Natarajan et al., 1974). It is essential bioactive metabolites of gentiopicroside in rats. Virulency of gentianine is achieved. LD 50 for gentianine: LD50 (mice): 480mg/kg (oral); 300mg/kg (belly injection); 250-300mg/kg (IV injection) (Yang and Song , 2000).



**Figure 3. A pyridine alkaloid having molecular formula  $C_{10}H_9NO_2$**

### Swerchirin

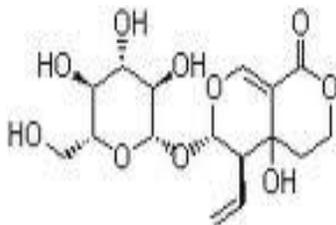
A medicinally foremost xanthone, obtained from several plants of family Gentianaceae including *Swertia chirayita*; having antimalarial, hypoglycemic (Arino A et al., 1997; Bajjpai et al., 1991;Saxena et al.,1996), hepatoprotective, pro-heamatopoitic (Ya et al., 1999) and weak chemo preventive pharmacological effects (Hirkawa et al., 1987).



**Figure4. Methylbellidifolin; 1,8- Dihydroxy- 3,5-dimethoxy-9H-xanthen-9one**

**Swertiamarin**

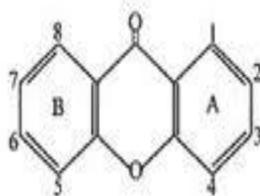
A Secoiridoid glycoside obtained from *Swertia chirayita* (Roxb ex. Flem) Karst; having analgesic property (Lei *et al.*, 1982).



**Figure 5. Swertiamaroside having molecular formula  $C_{16}H_{22}O_{10}$**

**Xanthenes**

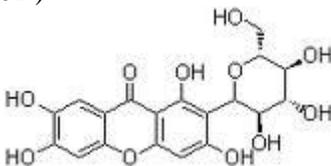
Over all Xanthenes are important bioactive constituent present in the drug which shows CNS down regulation in mice and rats (Bhattacharya *et al.*, 1976)



**Figure 6. An organic compound having molecular formula  $C_{13}H_8O_2$**

**Mangiferin**

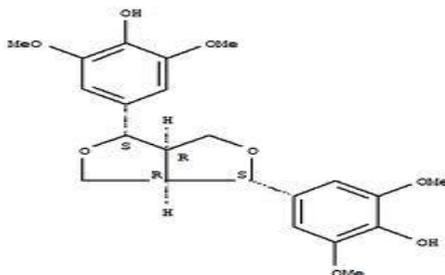
This compound, which is isolated from chirayita species possesses strong anti-inflammatory activity in arthritic mice, and accounted for lowering down TNF-alpha, IL-1beta, IL-6, and IFN-gamma and up regulation of IL-10 in the joint homogenates of mice (Anonymous, 2004; Kumar and Paul, 2003). It is also found to be a strong chemoprotective agent (Yoshimi *et al.*, 2001)



**Figure 7. A natural phenol having molecular formula  $C_{19}H_{18}O_{11}$**

**Lignan**

A lignan (syringaresinol; a negligible fraction of herb) which is hepatoprotective in nature, and the ubiquitous  $\beta$ -sitosterol are also present (Chatterjee and Pakrashi, (eds.) 1995; Rastogi and Mehrotra 1991; Rastogi and Mehrotra 1993; Rastogi and Mehrotra 1995; Rastogi and Mehrotra 1998).



**Figure 8. Linoresinol B having molecular formula  $C_{22}H_{26}O_8$**

### Triterpenoids

Chirayita also contains triterpenoids namely; swertanone, swertenol, episwertinol, gammacer-16-en-3 $\beta$ -ol, 21- $\alpha$ -H-hop-22(29)-en-3 $\beta$ -ol, taraxerol, oleanolic acid, ursolic acid, swerta-7, 9(11)-dien-3 $\beta$ -ol, pichierenol. Among them swertanone has got the anti-inflammatory property. Taraxerol and oleanolic acid are found to be analgesic and emollient respectively. Ursolic acid has anti-inflammatory, chemoprotective and anti-microbial activities. (Chatterjee and Pakrashi 1995; Rastogi and Mehrotra 1991; Rastogi and Mehrotra 1993; Rastogi and Mehrotra 1995; Rastogi and Mehrotra 1998)

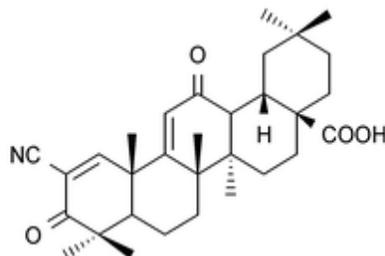


Figure 9. Isoprenoids having chemical structure 2-cyadioxooleana-1,9-dien-28-oate, CDDO,

### Pentacyclic triterpenoids

A class of pentacyclic triterpenoids also belongs to this herb including  $\beta$ -amyrin, friedlin, chiratenol, kairatenol, oleanolic acid, ursolic acid. Among them kairatenol is found to be hypoglycemic in nature. (Chatterjee and Pakrashi 1995; Rastogi and Mehrotra 1991; Rastogi and Mehrotra 1993; Rastogi and Mehrotra 1995; Rastogi and Mehrotra 1998)

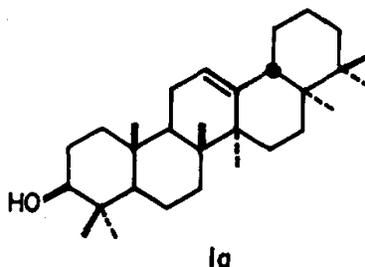


Figure 10. Kairatenol (3S,4S,5R,6R)-1,3,4,5,6,7-hexahydroxyheptan-2-On

Table 1; list of important bioactive constituents isolated from *Swertia chirayita*;

Active constituents	Biological activities	References
<u>Amarogentin (chirantin)</u>	Topoisomerase inhibition, chemo-preventive and antileishmanial effects.	[Ray et al., 1996], [Saha and Dass, 2005], [Ray et al., 1996; Medda et al., 1999].
<u>Amaroswerin</u>	Gastro-shielding [Niiho Y et al., 2005].	[Niiho et al., 2005].
<u>Gentianine</u>	Anti-inflammatory, anesthetic, antihistaminic, anticonvulsant properties, hypotensive, antipsychotic, lenitive, diuretic, antimalarial, antiamoebic and antibacterial properties.	(Song Zhen Yu et al., 1958; Geng Tao et al., 1959; Kwak et al., 2005). (Bhattacharya et al., 1974), (Mansoor and Malghani MAK, 2005), (Natarajan et al., 1974).
<u>Swerchirin</u>	Antimalarial, hypoglycemic, hepatoprotective, pro-heamatopoietic, and weak chemo preventive pharmacological effects.	(Arino et al., 1997), (Bajjpai et al., 1991), (Saxena et al., 1996), (Ya et al., 1999) (Hirkawa et al., 1987).
<u>Swertiamarin</u>	Analgesic property	(Lei et al., 1982).

### **Monetary products**

Some of the Herbaceous medicaments like Ayush-64, Diabecon, Mensturyl syrup and Melicon V ointment restrain chirayita essence in variable expense for its antipyretic, hypoglycemic, antifungal and antibacterial effects (Edwin and Chungath, 1988; Valecha et al., 2000; Mitra et al., 1996).

### **General caliber of *Swertia chirayita***

Pharmaceutical value of chirayita is increasing day by day. On the other hand its counter claim in pharmaceutical industry is increasing in the same speed. There are various factors like less activity and growing percentage of seeds and gentle field handling of plantlets which distress the advancement of agricultural technologies. Due to these problems, crude material for industrial use is being persistently gathered naturally.

### **Role of biotechnology in conservancy**

Biotechnology can play an indispensable role in conservation of *chirayita* species. It can engross different modes of conservation. Farming of *chirayita* at low heights is prohibited because of several environmental factors like fertility and textures of soil, pH, humidity etc. Tissue culture technique has found to be very useful for conservancy of intimate *Swertia chirayita* as little plant material can produce a big number of disease free propagules which can be revived in their innate environment. Systematized cultivation of the plant is important to guarantee continuous supplementation and affirmation of drug. Exercise in quality control of raw material of chirayita is labeling of DNA markers that associate DNA finger printing data with quantity of selected markers. (Joshi and Chavan P, 2004). Research is still going on molecular investigation of chirayita for protection of this wild endangered species, establishing methods for breeding in-vitro and making efforts for assuring continuous supply of its raw material.

### **Conclusion**

Farming of this very essential and endangered medicinal species should be promoted, if we consider its high demand and importance from medicinal and pharmaceutical aspects. Standardized steps for lab growth of *chirayita* should be followed to get maximum yield and extraction of active constituents, this will directly increase foreign exchange, because herb has great demand internationally as well due to its broad-range medicinal effects. Bioactive constituents extracted from *Chirayita* have got a number of beneficial medicinal effects for different kinds of ailments, with no side effects and also they are very cost effective as compared to allopathic medicines.

### **References**

- Anon( 1982) "In The Wealth of India: Raw Materials" Publication and Information Directorate, CSIR: New Delhi, Vol. X, pp. 78-81.
- Anonymous, (1976) "The Wealth of India" CSIR: New Delhi, pp.79-80.
- Anonymous (2004) "Wealth of India (Raw Materials)" Publications and Information Directorate, CSIR: New Delhi, vol.10, pp.77-81
- Ariò A. et al. (1997) "The extraction of yellow gentian root (*Gentiana lutea* L.)" Zeitschrift für Lebensmitteluntersuchung und-Forschung, Vol.205, pp. 295-299 .
- Bajpai MB. et al. (1991) "Hypoglycemic effect of swerchirin from the hexane fraction of *Swertia chirayita*" *Planta Med.* Vol. 57, pp. 102-4.
- Bentley, R. and H. Trimen, (1880) "Medicinal Plant" J and A Churchill :London, pp.183.
- Bhattacharya SK. et al. (1974) "Letter: Chemical constituents of gentianaceae. XI. Antipsychotic activity of gentianine" *J Pharm Sci*, Vol .63, pp.1341-2.
- Bhattacharya SK, Reddy PKSP, Ghosal S, Singh AK, Sharma PV (1976) "Chemical constituents of gentianaceae XIX: CNS-depressant effects of swertiamarin" *J. Pharm. Sci.*, Vol .65, pp.1547-1549.
- Brahmachari G et al (2004) " *Swertia* (Gentianaceae): chemical and pharmacological aspects" *Chem. Biodivers*, Vol .1, pp.1627-51.
- Chatterjee A, and Pakrashi SC, (eds.) (1995) "The Treatise on Indian Medicinal Plants used in Ayurveda" Vol. 4, Publication and Information Directorate, New Delhi: India, Vol. 4, p. 92 .
- Clarke, C. B., ( 1885) "In The Flora of British India" (ed. Hooker, J. D.), L.Reeve and Co:London, , Vol. IV, p. 124.
- Daniel M. and Sabnis SD (1978) "Chemical systematics of family Gentianaceae" *Curr. Sci.*, Vol .47, pp. 109 -111..
- Duke, J.A (2002) "In Handbook of Medicinal Herbs" CRC Press: Washington DC, p. 190.

- Edwin R, and Chungath JI (1988) "Studies in *Swertia chirata*" Indian Drugs, Vol. 25, pp.143–146.
- Garg DS, (editor-in chief) Dhanvantri-Banaushdhi Vishesh Ank.,(1965) , Dhanvantri Karyalaya,Vijaygarh, Aligarh :India, Vol.3, p.94.
- Geng Tao. et al. (1959)Journal of Physiology, Vol. 23, pp.203.
- Hirkawa K. et al (1987) "Chemo preventive action of xanthone derivatives on photosensitized DNA damage" Photochem Photobiol, Vol .81, pp.314-9 .
- Hostettmann-Kaldas M. et al. (1981) " Flavones and secoiridoids of American *Gentiana* species "Phytochemistr, Vol. 20, pp. 443-446.
- Joshi K, Chavan P, Warude D, Patwardhan B (2004)"Molecular markers in herbal drug technology" Current Science, Vol. 87, pp.159-165.
- Joshi, P. and V. Dhawan, (2005) "Swertio, chirriyito',- an overview "Curr. Sci., Vol .89, pp. 635-640.
- Kirtikar,K. R. and Basu, B. D. (eds) (1984) " Indian Medicinal Plants", :Allahabad, vol. III, pp. 1664–1666.
- Kirtikar, KR. and B.D. Basu, (1984) " Indian Medicinal Plants "Latit Mohan Basu, Leader Road, Allahabad: India, pp.1664-1665.
- Karan, M., Vasisht, K. and Handa, S. S., (1996) " In Supplement to Cultivation and Utilisation of Medicinal Plants" (eds Handa, S. S. and Kaul, M. K.). RRL, Jammu-Tawi, pp. 349–354.
- Kumar IV, Paul BN, Asthana R, Saxena A, Mehrotra S, Rajan G (2003) "Nov.,*Swertia chirayita* mediated modulation of interleukin-1beta, interleukin-6, interleukin-10, interferon-gamma, and tumor necrosis factor-alpha in arthritic mice" 25(4), pp. 573-83.
- Kwak WJ. et al. (2005) " Effects of gentianine on the production of proinflammatory cytokines in male Sprague-Dawley rats treated with lipopolysaccharide (LPS) " Biol Pharm Bull, Vol. 28, pp.750-3.
- Lei Wei Ya, et al. (1982) " Swertiamarin's central inhibitory effects" Journal of Chinese Materia Medica, Vol 13, pp. 368 .
- Mansoor A and Malghani MAK. (2005) "Diuretic effect of *Gentiana olivieri* and its alkaloid gentianine "Botany.m plants. August 13-17, Austin, Texas. Oral presentation.
- Mitra SK, Gopumadhavan S and Muralidhar T S (1996) " Effect of D-400, an ayurvedic herbal formulation on experimentally-induced diabetes mellitus" Phytother. Res., Vol. 10, pp.433.
- Natarajan PN, Wan AS, and Zsaman V. (1974) " Antimalarial,antiamobeic and toxicity tests on gentianine "Planta Med, Vol .25,pp.258-260.
- Niiho Y. et al. (2005) "Gastro protective effects of bitter principles isolated from *Gentian* root and *Swertia* herb on experimentally induced gastric lesions in rats" Journal of Natural Medicine, Vol. 60,pp.888.
- Newman DJ, Cragg GM & Snader KM (2000)" The influence of natural products upon drug discovery" Nat. Prod. Rep, Vol .17, pp.215–234.
- Purushothaman . et al. (1973) Chem Abst, Vol. 79, pp.113-21.
- Rastogi RP, and Mehrotra BN "Compendium of Indian medicinal plants" CDRI, Lukhnow and National institute of Science Communication, New Delhi: India Vol. 2(1991), p. 654; Vol. 3(1993), p. 615;Vol. 4(1995), p. 701; Vol. 5(1998), pp. 815.
- Ray S. et al. (1996) "Amarogentin, a naturally occurring secoiridoid glycoside and a newly recognized inhibitor of topoisomerase I from *Leishmania donovani*" J Nat Prod, Vol. 59, pp.27-9 .
- Saha P, and Das S. Bitter(2005) " fraction of *Swertia chirata* prevent carcinogenic risk due to DMBA Exposure "Indian Journal of Medical Research, 22-27 Poster Presentation.
- Saxena AM. et al. (1996) "Mode of action of three structurally different hypoglycemic agents: a comparative study". Indian J Exp Biol, Vol. 34, pp. 351-355 .
- Sharma PV, Dravyaguna-vijnana, (1986) Chaukhambha Bharti Academy, Varanasi:India, Vol. 2.
- Song Zhen Yu. et al. (1958) Journal of Physiology, Vol. 22 , pp.201.
- Suzuki, M.N., Yoshim, R., Ritsuko, F., Fiji & Hiroshi. (1989) Daiko Minami Higoshi Nagya. 461.T.Ph.Skin Valecha N, Devi UC, Joshi H, Shahi VK, Sharma VP, and Lal S.(2000) " Comparative efficacy of ayush-64 vs chloroquine in vivax malaria "Curr. Sci., Vol .78,pp.1120–1122.
- Ya BQ. et al. (1999) "Protective effect of swerchirin on hematopoiesis in 60 Co-irradiated mice" Phytomedicine, Vol. 6, pp. 85-8.
- Yang XF and Song CQ. (2000) "Studies on the metabolism of gentiopicroside by rat intestinal flora " Zhongguo Zhong Yao Za Zhi, Vol. 25,pp.673-6.
- Yoshimi N, Matsunaga K, Katayama M, Yamada Y, Kuno T, Qiao Z, et al.,(2001)"The inhibitory effects of mangiferin a naturally occurring glucosylxanthone , in bowel carcinogenesis of male F344 rats" Cancer Lett., Vol.163, pp.163-170.