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Review Article

ANTIDIABETIC PROPERTIES OF CHANDRAPRABHA VATI Sakshi Bhardwaj^{1*}, Shailza Bhatnagar², Kalpana Yadav³, Pankaj Yogi¹, Pooja Yadav¹, Deepak Jangidh¹, Ambika¹

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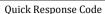
ABSTRACT

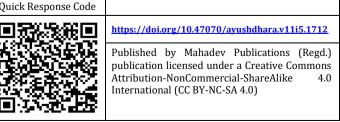
Diabetes mellitus is a group of metabolic disorders characterized by elevated blood glucose levels and symptoms like polyuria, polydipsia and polyphagia. It arises from insufficient insulin secretion, insulin resistance, or both. Ayurveda mentions this disease as Prameha. Chandraprabha vati is an effective formulation widely used in its management. It is an Avurvedic formulation available in classical Vati form. The indications of Chandraprabha vati include Prameha and its complications like Vibandha (constipation), Anaha (distension of abdomen due to obstruction to passage of urine and stools), Kustha (diseases of skin), Netra roga (eve disorder) etc. It contains 37 herbomineral ingredients which exhibit various actions. The Rasayana, Vrishya, Shothhara, Deepana paachana and Vatanulomana karma are present in this formulation which makes it a drug of choice in chronic cases of Prameha as add on therapy to regular treatment. All the ingredients possess anti-diabetic activity. The article discusses about its role in the management of Prameha in detail.

INTRODUCTION

Diabetes mellitus has become a primary health concern these days. It is a metabolic disease characterized by hyperglycemia which occurs due to impairment in insulin secretion, functioning or both. In Ayurveda it is correlated with Prameha. Prameha is primarily viewed as a lifestyle related disorder. Key contributing factors in the Prameha are unhealthy eating pattern and sedentary lifestyle. Prameha is associated with the depletion of *Dhatu* (bodily tissues) ultimately leading to depletion of *Ojas* (vital energy and immunity)^[1]. Rasayana therapy is essential for rejuvenating tissues, replenishing *Ojas*, balancing Doshas. enhancing metabolism, preventing complications, and improving mental health. By incorporating Rasayana into the treatment plan of Prameha, patients can achieve better quality of life. Ayurveda and other traditional systems of medicines describe number of formulations for treatment of

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diabetes. Many herbal formulations are being evaluated for their effectiveness in controlling diabetes. *Chandraprabha vati* is one of such formulations which can be used as a rejuvenating agent in Prameha.

Objective

To review anti-diabetic properties of each content of Chandraprabha vati.

Scientific review of Chandraparabha vati

A. Review as per Ayurveda

Chandraprabha vati is mentioned in the Madhyam Khand of Sharangdhar Samhita in the context of Vati. Different Acharyas have mentioned different number of contents for the formation of Chandraprabha Vati. Acharya Sharangdhara has described 36 Dravya in Chandraprabha Vati. The contents are listed in table no 1. Its indications are Vibandha (constipation), Anaha (distension of abdomen due to obstruction to passage of urine and stools), Shula (colicky Pain), Granthi (anemia), (cyst), Pandu Kamala (jaundice), Mutrakricchra (dysuria), Ashmari (calculus), Arsha (hemorrhoids), Arbuda (tumor), Mutraghata (urinary obstruction), Antra-Vriddhi (hernia), Kati Shula (lower backache), Kustha (diseases of skin), Kandu (itching), Plihodara (disorder of spleen, ascites associated with splenomegaly), Bhagandara (fistula in ano), Dantoroga

(dental disease), *Netra Roga* (eye disorder), *Aruchi* (tastelessness), *Mandagni* (impaired digestive fire), *Striroga* (gynaecological disorders), *Artava Ruja* **Properties and actions of contents of Chandraprabhe**

(dysmenorrhoea), *Shukra Dosha* (vitiation of semen), *Daurbalya* (weakness) and *Prameha*^[2].

Properties and actions of contents of *Chandraprabha vati* in the perview of Ayurveda Table 1: *Chandraprabha vati* ingredients, their properties^[3] and actions^[4]

S.No.	Drug name	Usable Part	Rasa	Guna	Virya	Vipak	Action
1.	Chandraprabha (Cinnamomum camphora)	Sublimated extract	Tiktaa, Katu, Madhur	Laghu, Tikshna	Shita	Katu	Vata Kaphahara, Deepan, Vrishya, Lekhana, Dahaprashman, Ruchya, Uttejaka
2.	Vacha (Acorus calamus)	Rhizome	Katu, Tiktaa	Laghu, Tikshan	Ushna	Katu	Kapha nissaraka, vaatanulomaka, deepana, pachana, Vibandhahara, Adhmanhara, Shulhara, Mala- Mootravishodhaka, Lekhana
3.	Musta (Cyperus scariosus)	Rhizome	Tikta, Katu, Kashya	Laghu, Ruksha	Shita	Katu	Deepan, Pachan, Vaatanulomaka, Kaphaghna, Trishnanigrahan, Kandughna, Balya
4.	Kirattikta (Swertiya chirayita)	Plant	Tikta	Laghu, Ruksha	Ushna	Katu	Deepan, Pachan, Virechan, Dahaghna, Pittavirechaka, Shothaghna
5.	Guduchi (Tinospora cardifolia)	Stem	Tikta, Kashaya	Guru, Snigdha	Ushna	madhura	Tridoshaghna, Rasayana, Balya, Jwarhara, Deepan, Mutrajanan, Twakroghara, Pramehahara
6.	Devdaru (Cedrus deodaru)	Heart Wood	Tikta	Laghu, Snigdha	Ushna	Katu	Swedjanan, Mutrajanan, Vaatanulomaka, Vata- kaphahara Twagdoshahara, Vibandhhara, Adhmanhara, Tandranigraha, Pramehaghna, Kaphaghna
7.	Haridra (Curcuma longa)	Rhizome	Tikta, Katu	Ruksha, Laghu	Ushna	katu	Twakdoshhara, Deepan, Kaphaghna, Vaathara, Prameha Nashaka
8.	Ativisha (Aconitum hetrophyllum)	Root Tuber	Tikta, Katu	Laghu, Ruksha	Ushna	Katu	Deepan-Pachan, Balya, Pittashamak, Lekhan, Shothaghna
9.	Daruharidra (Berberis aristats)	Stem	Tikta, Kashaya	Laghu, Ruksha	Ushna	Katu	Balya, Deepan, Pachan, Shleshmaghna, Rasayana, Twak doshhara
10.	Pippalimula (Piper longum)	Root	Katu	Laghu, Snigdha, Tikshna	Anush nshita	Madhura	Rasayana, Deepan- Pachan, Vaathara, Kaphaghna, Agnimandyahara, Uttejaka, Bhedaka, Anulomaka
11.	Chitraka (Plumbago zeylanica)	Root	Katu	Laghu, Ruksha, Tikshna	Ushna	Katu	Agideepak, Rasayana, Vata- Kaphahara, Lekhan
12.	Dhanyaka (Coriandrum sativum)	Fruit	Kashaya, Tikta, Madhura, Katu	Laghu, Snigdha	Ushna	Madhura	Deepan-Pachan, Vatanulomaka, Pipasaghna, Virechaka, useful in Netra Roga, Rochak, Dahaprashman

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13.	Haritaki (Terminalia chebula)	Pulp	Kashaya prdhan panch rasa lavanvarjit	Laghu, Ruksha	Ushna	Madhura	Excellent Mridu Virechaka, useful in Netraroga and Ajeerna, Vranropaka, Rasayana, Balya, Deepan-Pachan
14.	Bibhitaka (Terminaliya bellirica)	Pulp	Kashaya	Laghu, Ruksha	Ushna	madhura	Virechaka, Vednasthapaka, Shothaghna, Kandughna, Dhatuvardhak, Deepan-Pachan, Anuloman
15.	Amalaki (Embelika officinalis)	Pulp	Amal prdhan panch rasa lavan varjit	Guru, Ruksha, Shita	Shita	madhura	Rasayana, Vrishya, Mridu- Virechak, Deepan, Pramehaghna, Deepan, Dahaprashman
16.	Cavya (Piper retrofractum)	Stem	Katu	Laghu, Ruksha	Ushna	katu	Deepan-Pachan
17.	Vidanga (Embelia ribes)	Fruit	Katu, Kashaya	Laghu, Ruksha, Tikshna	Ushna	Katu	Shoolaghna, Aadhmanhara, Balya, Deepan-Pachan, Anuloman, Rasayana, Vaat- kaphahara
18.	Gajapippali (Scindapsus officinalis),	Fruit	Katu	Laghu, Snigdha, Tikshna	Anushn ashita	Madhura	
19.	Shunthi (Zingiber officinale)	Rhizome	Katu	Laghu, Snigdha	Ushna	Madhura	Bhedan, Agnideepan, Ruchi, Dahaprashman, Shothahara, Vednasthapan, Vaatanuloman, Kaphaghna, Vrishya, Balya
20.	Marica (Piper nigrum)	Fruit	Katu	Laghu, Tikshan	Ushna	Katu	Agnideepan, Vaatshamaka, Chedan, Ruchya, Lekhana, Medohara, Uttejaka
21.	Pippali (Piper longum	Fruit	Katu	Laghu, Snigdha, Tikshna	Anusha nshita	Madhura	Agnideepan, Vrishya, Rasayana, Rechaka, Pachaka, Ajeernahara
22.	Makshika Dhatu bhasma	Calx	Amla, Kashaya, Lavan	Laghu	Shita	Katu	Bastiruk prashaman, Mehnashan, Shothhara, Tridoshhara, Vrishya, Rasayana
23.	Yava kshara	-	Katu	Laghu, Snigdha, Sukshma	Shita	Katu	Agnideepan, Shoolhara, BVaathara, Shleshmahara, Anahhara
24.	Sarji kshara		Katu	Laghu, Snigdha, Sukshma	Shita	Katu	Shoolhara
25.	Saindhava Iavana	-	Lavana	Laghu, Snigdha, Sukshma	Shita	Madhura	Deepan-Pachan, Ruchya, Vrishya
26.	Sauvarcala lavana	-	Lavana	Laghu, Ruksha, Tikshna, Vyavayi	Ushna	Madhura	Ruchya, Bhedana, Deepan- Pachan, Vibandhahara, Sulaghna, Vaatanulomaka
27.	Vida lavana	-	Lavana	Laghu, Sukshma	Ushna	Madhura	Deepan, Ruchya, Vyavaayi, Vibandhanashaka, Vaatanulomaka, Shoolaghna
28.	Trivrit (Operculina	Root	Tikta, katu	Laghu, Ruksh,	Ushna	Katu	Bhedan, Virechaka, Shothaghna, Vaatshamaka

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	turpethum),			Tikshan			
29.	Danti (Baliospermum montanum),	Root	Katu	Guru, Tikshan	Ushna	Katu	Sara, agnideepan, dahahara, rechan, Adhmanhara, Shothaghna
30.	Patraka (Cinnamomom tamala)	Leaf	Katu, Tikta, madhura	Laghu, Ruksha, Tikshna	Ushna	Katu	Deepan, Ruchya, Kapha- Vaatahara
31.	Tvak (Cinnamomum verum)	Stem Bark	Katu, Tikta, madhura	Laghu, Ruksha, Tikshna	Ushna	Katu	Deepan-Pachan, Uttejaka, Anuloman, Trishnanigrahan, Balya
32.	Ela (Elettaria cArdamom)	Stem	Katu, Madhura	Laghu, Ruksha	Shita	Madhura	Trishnanigraha, Deepan- Pachan, Ruchya, Vaatanulomaka, Uttejaka
33.	Vaṃśarocana (Bambusa arundinance)	Silicacious Concretion	Madhura, Kashaya	Ruksha, Laghu, Tikshna	Shita	Madhura	Chedana, Bastishodhaka, Shothaghna
34.	Hata loha	Calx	Tikta	Guru, Ruksha, Sara	Shita	Madhura	Medohara, Mehahara, Shothhara, Kaphapittaghna,
35.	Sita	Crytaline Sugar	Madhura	Guru, Snigdha	Shita	Madhura	Ruchya, Vaata-Pitta hara, Dahahara
36.	Silajatu,	Sublimate d extract	Tikta, Lavana	Laghu, Tikshna	Shita	Katu	Rasayana, Chedana, Kapha- Medshamaka, Shothhara, Udardoshhara
37.	Guggulu (Commiphora mukula)	Exudate	Tikta, katu	Laghu, Ruksha, Tikshna, Vishada, SukshmaS ara, Sugandhit	Ushna	Katu	Tridoshaghna, Rasayana, Balya, Vrishya, Deepan, Shothhara, Vaatanuloman, Vednasthapan

Method of preparation^[2]

The *Churna Dravyas* are powdered and mixed well. In a vessel filled with warm water *Shudha Shilajatu, Shudha Guggulu* are added and melted. When it gets cooled above powdered drugs are added and fine bolus is formed. Afterwards rest of *Bhasma Dravya* along with *Sharkara* is added. Adding little warm water a fine mixture is made and the pills are rolled and kept in shade for drying.

Anti-diabetic properties actions of contents of *Chandraprabha vati*

Chandraprabha (Karpura) (Cinnamomum Camphora)- Camphor is a natural product of the *Cinnamomum camphora*, used in general to control excessive sweating and burning sensation of skin. It has scraping property, useful in balancing *Kapha*^[5]. A study on Diabetic rats shows suggests that its antihyperglycemic properties are likely attributed to its antioxidant effects^[6]. It exhibited inhibitory activities for α -amylase and α -glucosidase assays which are helpful for starch breakdown into glucose. It helps in pancreatic β -cell restoration and insulin secretion also^[7].

Vaca (*Acorus Calamus*)- It increase levels of plasma insulin, tissue glycogen, and G6PD. Ethyl acetate extracted from *Vacha* increases Secretion of GLP-1 and lowers blood glucose levels. It also enhanced insulin secretion in HIT-T15 cells and blocked the αglucosidase in vitro activity^[8]. Treatment with βasarone in obese rats resulted in weight loss and also inhibited metabolic changes, glucose intolerance, elevated cholesterol levels, and variations in adipokine levels^[9]. The in vitro antioxidant activity of acetone, acetonitrile, alcoholic, and aqueous extracts exhibited free radical scavenging activity.^[10]

Nagarmotha (Cyperus Scariosus)- Antihyperglycemic activities are reported in the leaves of this plant^[11]. The active constituents of root extracts and volatile oil include sesquiterpenes, cyperone, selinene, and cyperene which exhibit various pharmacological properties such as antioxidant, and antihyperglycemic activities^[12]. Gallic acid, quercetin, and 4-hydroxycinnamic acid has significant antidiabetic and radical scavenging activities.^[13,14]

Bhūnimba (**Kiratatikta**)- Ethanolic root extract showed a significant anti-diabetic effect, with a notable decrease in blood sugar levels. It significantly reduced cholesterol and lipid levels. Mangiferin, in particular, is recognized for its blood glucose-lowering properties^[15]. The anti-diabetic activity was evaluated by assessing the inhibitory effect on the enzyme α amylase, which is involved in the breakdown of starch to produce glucose^[16].

Guduchi (*Tinospora Cardifolia*)(*Amritaa*)- It has been reported to mediate its anti-diabetic potential through various biologically active phytoconstituents isolated from different parts of plant, including alkaloids, tannins, cardiac glycosides, flavanoids, saponins and steroids. The isoquinoline alkaloid rich fraction from stem, includes palmatine, jatrorrhizine, and magnoflorine which have been reported for insulin mimicking and insulin releasing effect both *in vitro* and *in vivo*. By adenosine monophosphate-activated protein kinase activation, it decreases the blood sugar and cholesterol level and maintains the blood pressure.^[17].

Daruka (Devadaru) (Cedrus Deodaru)- *C.deodara* possesses strong antioxidant properties^[18]. The ethanolic extract has an anti-hyperglycemic effect in rats with streptozotocin-induced diabetes along with reduction SGPT, SGOT, cholesterol, and triglycerides. Histopathological studies revealed that it enhanced islet regeneration in the pancreas and restored normal cellular size of the islets with hyperplasia^[19]. In in-vivo experiments, capsules made from the plant's wood extract have shown significant anti-diabetic properties^[20].

Haridra (Curcuma Longa)- It can delay the development of type 2 diabetes, enhance β-cell function, prevent β-cell death, and reduce insulin resistance in animal studies^[21]. A 9-month treatment with curcumin extract can effectively and safely prevent the progression from prediabetes to type 2 diabetes^[22]. The main constituents Curcumin, Dimethoxy Curcumin, Bis-demethoxy Curcumin, and Arturmerone exhibit PPAR-γ ligand binding activity, enhancing the transcription of several insulin-responsive genes and improving insulin resistance in type 2 diabetes mellitus^[23].

Ativisha (*Aconitum Hetrophyllum*)- Oral administration of A. heterophyllum extract in dietinduced obese rats significantly reduced serum total cholesterol, triglycerides, and LDL cholesterol levels, and inhibited intestinal fat absorption. Its extract possesses potential hypolipidemic^[24] and anti-obesity activities^[25]. **Darvi** (Berberis Aristats)- A systematic study conducted on the roots of *Berberis aristata* indicates that both aqueous and methanolic root extracts exhibit a hypoglycemic effect when administered to normal and alloxan-induced diabetic albino rabbits and Wistar rats by stimulating insulin production^[26].

Pipplimūla (Pippali) (Piper Longum)- It was found that the oral administration of ethanolic extract restores the blood glucose level which ultimately stimulates the liver activities to maintain the normal homeostasis of blood glucose level^[27]. Aqueous extract also showed significant antidiabetic properties^[28]. Another study showed that the oral administration of the dried fruits showed antioxidant, anti-lipid peroxidative and antihyperglycemic activities in diabetic rats when compared with standard reference drug glibenclamide^[29].

Chitraka (Plumbago Zeylanica)- Plumbagin increased the activity of hexokinase and decreased the activities glucose-6-phosphatase and fructose-1. 6of bisphosphatase significantly in treated diabetic rats. Enhanced GLUT4 mRNA and protein expression were observed in diabetic rats after treatment with plumbagin^[30]. Extracts of Plumbago zeylanica show a significant in-vitro inhibitory effect on α glucosidase^[31].

Dhanyaka (Coriandrum sativum)- The ethanolic extract of coriander leaves had the antidiabetic activity by improving and regenerating the β cell in pancreas and inhibiting the α -glucosidase enzyme in small intestine^[32]. The anti-diabetic activity of *Coriandrum* sativum seeds was studied by incorporating the seeds into the diet of rats fed a high-fat diet with added cholesterol. The administration of coriander seeds had a significant hypolipidemic effect. The levels of lowdensity lipoprotein (LDL) and very low-density lipoprotein (VLDL) cholesterol decreased, while highdensity lipoprotein (HDL) cholesterol levels increased in the experimental group compared to the control group. The increased plasma LCAT activity and enhanced degradation of cholesterol to fecal bile acids and neutral sterols were likely responsible for the hvpocholesterolemic effect^[33].

Haritaki (Terminalia Chebula)- Alcoholic extract of *Terminalia chebula* induced efficient anti-hyperlipidemic property^[34]. Oral administration of the water extract of *T. chebula* improved glucose utilization during a glucose tolerance test (GTT), Similarly, the extract reduced fasting blood glucose (FBG) to near-normal levels within two weeks^[35].

Bibhitaka (Terminaliya bellirica)- Continuous administration of a dried 75% methanolic extract significantly prevented hyperglycemia in alloxan-induced hyperglycemia. Similarly there was significant

increase in the activity of catalase and glutathione in blood and liver^[36].

Amalaki(Embelika officinalis)- It have some important constituents (including gallic acid, gallotanin, ellagic acid and corilagin), possess antidiabetic effects through their antioxidant and free radical scavenging properties. It has also been reported to prevent/reduce hyperglycemia, diabetic nephropathy and neuropathy^[37].

Vidanga(Embelia Ribes)- The ethanolic extract reduces serum leptin, adiponectin, lipase, hepatic glycogen content, and glucose-6-phosphatase levels towards normal in diabetic rats^[38]. These findings suggest that ER extract has the potential to prevent lipid abnormalities and pancreatic β-cell dysfunction in type-2 diabetes mellitus (DM). *Embelin* was found to induce PPARγ (a key molecular target for treating type-2 DM, is down-regulated during tissue insulin resistance) expression in the epididymal adipose tissue of diabetic rats^[39]. A case study stated on the efficacy of *Vidangadi Kwatha* on patients of type 2 diabetes mellitus^[40].

Gajapippali (Scindapsus Officinalis)

The water extract demonstrated strong antidiabetic activities, significantly enhancing glucose consumption in insulin-resistant HepG2 cells compared to the model group^[41].

Shunthi (Zingiber Officinale)- Treatment with *Zingiber officinale* significantly increased insulin levels and decreased fasting glucose levels in diabetic rats. Additionally, it lowered serum cholesterol, serum triglycerides, in diabetic rats. It potentially involves 5-47 HT receptors^[42].

Marica (Piper nigrum)- Research indicates that *Piper nigrum* exerts a protective effect in alloxan-induced diabetic rats, helping to reduce glucose levels.^[43] It was observed in one study that nine out of twenty novel piperine analogues (5b, 6a-h) exhibited significantly higher antidiabetic activity compared to the standard drug, rosiglitazone^[44].

Pippali (Piper Longum)- thanolic extract of dried Piper longum fruits possesses potent antihyperglycemic and antilipidperoxidative activities in alloxan-induced diabetes^[45]. Aqueous extract on STZ-induced diabetic rats resulted in a significant decrease in fasting blood glucose (FBG) levels and correction of diabetic dyslipidemia.

Trivrit (Operculina Turpethum)- Its flavoured fraction primarily functions as an alpha-amylase inhibitor and an agonist for PPAR gamma, underscoring its potential role in the management of type 2 diabetes mellitus as an antihyperglycaemic^[46].

Danti (Baliospermum Montanum)- Its extract demonstrated significant effects in reducing blood glucose levels and lipoproteins in diabetic animals. Traditionally, Baliospermum montanum is used as a diaphoretic, promoting the elimination of body fluids through the skin. Its active constituents include flavonoids, both of which contribute to its efficacy in treating diabetic nephropathy^[47].

Patraka (*Cinnamomom Tamala*)- In a study evaluating the ethanolic extract of *C. tamala* leaves, exhibited both hypoglycemic and antihyperglycemic properties in normoglycemic (and STZ-hyperglycemic rats.^[48,49] Its ethanolic extract prevented the rise in total cholesterol (TC), triglycerides (TG), low-density lipoprotein (LDL), and very-low-density lipoprotein (VLDL) cholesterol levels, indicating its beneficial effects on STZ-induced hyperlipidemia.^[50]

Ela (Sukshmaila)- In a clinical trial, carried out on obese or overweight pre-diabetic women (3 gm cardamom for 2 months) disclosed that Total Cholestrol and LDL-C significantly reduced. It also showed a protective effect on HDL-C amount in prediabetic subjects.^[51] Cardamom has been shown to control certain inflammatory and oxidative stress parameters in pre-diabetic people.^[52] The administration of cardamom caused an increase in SIRT1 (Sirtuin 1) in the non-alcoholic fatty liver patient^[53]. It was also found that administration of cardamom powder prevented obesity in high-fat dietinduced obese rats^[54]. Cardamom can ameliorate high blood glucose, insulin resistance, and glucose metabolic disorders. E. cardamomum and its active constituents can control insulin secretion, insulin resistance through increasing the amount of SIRT1, PPAR- γ coactivator-1 alpha (PGC-1 α), and attenuating the function of nuclear factor kappa-light-chainenhancer of activated B cells (NF- κ B) as well as controlling glucose metabolism by inhibiting α glucosidase and α -amylase^[55].

Vamsharocana (Bambusa Arundina)- Aqueous ethanolic seed extract has shown maximum reduction in blood glucose level. The significant antidiabetic activity of these seeds may be attributed to their ability to inhibit free radical generation and prevent subsequent tissue damage caused by alloxan. Additionally, these seeds may enhance the effect of plasma insulin by either stimulating the pancreatic secretion of insulin from existing beta cells or promoting its release. This is evidenced by the significant improvement in glucose and protein levels, as insulin inhibits gluconeogenesis from proteins^[56].

Guggulu (Commiphora Mukula)- In an experimental model of diabetes, *C. mukul* and *P. emblica* dramatically increased DPP-IV levels. The medicinal herbs' DPP-IV

inhibitory activity also translated into considerable antidiabetic efficacy, as seen by a considerable reduction in HBA1C levels^[57]. Docking experiments against DPP-IV revealed that the active compounds in *C. mukul* (Gluggusterone E, Gluggusterone Z) have considerable inhibitory activity against the enzyme (DPP-IV)^[58]. The hypolipidemic activity was shown in animals as well as in patients of obesity and hypercholesterolemia^[59]. It contains Z-guggulsterones and E-guggulsterones which are purported to be the compounds responsible for the hypoglycaemic and hypolipidemic activity of the *Guggulu*.^[60]

DISCUSSION

Chandraprabha Vati is an excellent formulation whose ingredients have undergone various in vivo and clinical studies, demonstrating significant anti-diabetic effects. The formulation contains a wide array of potent herbs and substances, each contributing unique therapeutic properties. For analgesic effects, Vacha, Vidanga, Sarjika kshara, Sauvarcala lavana, Vida lavana, and Yava kshara are included. Rasavana agents like *Guduchi*, *Haritaki* and others offer vital support in promoting overall health. Deepana and Pachan actions are enhanced by herbs such as Musta Kirattikta and Haritaki. Carminative effects are provided by Vacha, Musta, and Pippalimula, while Kaphaghna drugs like Haridra and Devdaru reduce excess Kapha. Shothaghna properties are ensured by Kirattikta and Ativisha among others. For skin disorders Haridra and Daruharidra are included. Jwarahara benefits are provided by Guduchi, while Shothhara effects are supported by *Devdaru* and *Guggulu*. Anti-diabetic (Pramehaghna) properties come from Guduchi, Haridra, and Amalaki. The formulation also includes anti-obesity Medohara agents like Marica and Shilajatu, as well as Chedana drugs like Shilajatu and Vamsarochana. Tridoshaghna herbs like Guduchi and *Guggulu* balance all three doshas, while *Lekhana* and Vyavaayi actions are supported by ingredients like Vacha, Ativisha, and Vida lavana. This comprehensive blend ensures a holistic approach to managing various health conditions, particularly Prameha. Despite its long history of use and the known anti-diabetic and lipid-lowering properties of its components, systematic scientific studies are still needed to fully validate its therapeutic utility in controlling diabetes. *Chandraprabha vati's* ingredients offer various benefits.

CONCLUSION

Diabetes mellitus, a chronic metabolic disease leads to damage in various body systems, causing complications that worsen the diabetic state and impact the quality of life. The ingredients of this formulation have demonstrated significant antidiabetic effects in various studies. The properties mentioned above play a crucial role in managing and controlling *Prameha*.

REFERENCES

- Drdhabala C. Caraka Samhita by Agnivesha with Ayurveda Dipika commentary of Cakrapanidatta. Vaidya Jadavji Trikamji Acharya, editor. Varanasi: Chowkhamba Krishnadas Academy, 2015; 738 p. (66).
- Śārngadhara, Brahmānanda Tripāţhī, Agniveśa. Śārngadhara samhitā. Vārāņasī: Caukhambā Surabhāratī Prakāśana; 2004. P. 201.
- 3. Bhāvamiśra. Bhāvaprakāśa of Bhāvamiśra, commentary by Dr. Bulusu Sitaram. Chaukhambha Orientalia. 2000.
- D Shanthkumar Lucas, Jyotirmitra. An introduction to nighaņţus of Āyurveda. Varanasi: Chaukhambha Sanskrit Bhawan; 2006.
- 5. Therapeutic and Medicinal Uses of Karpura-A Review. 2015; 6(4).
- Drikvandi P, Bahramikia S, Alirezaei M. Modulation of the antioxidant defense system in liver, kidney, and pancreas tissues of alloxan-induced diabetic rats by camphor. J Food Biochem. 2020 Dec; 44(12): e13527. doi: 10.1111/jfbc.13527. Epub 2020 Oct 20. PMID: 33084110.
- 7. Bharti SK, Krishnan S, Kumar A, Kumar A. Antidiabetic phytoconstituents and their mode of action on metabolic pathways. Ther Adv Endocrinol Metab. 2018 Mar; 9(3): 81-100. doi: 10.1177/2042018818755019. Epub 2018 Feb 12. PMID: 29492244; PMCID: PMC5813859.
 - Prisilla D.H., Balamurugan R., Shah H.R. Antidiabetic activity of methanol extract of Acorus calamus in STZ induced diabetic rats. Asian Pac. J. Trop. Biomed. 2012; 2: S941–S946. doi: 10.1016/S2221-1691(12)60341-9.
 - Liu Y.X., Si M.M., Lu W., Zhang L.X., Zhou C.X., Deng S.L., Wu H.S. Effects and molecular mechanisms of the antidiabetic fraction of Acorus calamus L. on GLP-1 expression and secretion in-vivo and Invitro. J. Ethnopharmacol. 2015; 166: 168–175. doi: 10.1016/j.jep.2015.03.014.
 - Si M.M., Lou J.S., Zhou C.X., Shen J.N., Wu H.H., Yang B., Wu H.S. Insulin releasing and alpha-glucosidase inhibitory activity of ethyl acetate fraction of Acorus calamus In-vitro and in-vivo. J. Ethnopharmacol. 2010; 128: 154–159. doi: 10.1016/j.jep.2009.12.044.
 - 11. Utreja D, Sharma P, Ekta X. Chemistry and biology of Cyperus scariosus: An overview. Curr. Chem. Bio. 2015; 9: 2–9. doi: 10.2174/2212796809999150 630115456.

- 12. Utreja D, Sharma P, Ekta X. Chemistry and biology of Cyperus scariosus: An overview. Curr. Chem. Bio. 2015; 9: 2–9. doi: 10.2174/2212796809999150 630115456.
- Naaz J, Roqaiya M, Fatima KMH, Ahad A, Wahid MA. Review on Cyperusscariosus: A Potential Medicinal Plant. Research & Reviews: Journal of Crop Science and Technology [Internet]. 2021 May 8 [cited 2024 May 28]; 10(1): 28–32.
- 14. Kakarla L, Katragadda SB, Tiwari AK, Kotamraju KS, Madhusudana K, Kumar DA, Botlagunta M. Free radical scavenging, α-glucosidase inhibitory and anti-inflammatory constituents from Indian sedges, Cyperus scariosus R.Br and Cyperus rotundus L. Pharmacogn Mag. 2016 Jul; 12
- 15. Sekar BC, Mukherjee B, Chakravarti RB, Mukherjee SK. Effect of different fractions of Swertia chirayita on the blood sugar level of albino rats. J Ethnopharmacol. 1987; 21(2): 175-81.
- Rafatullah S, Tariq M, Mossa JS, al-Yahya MA, al-Said MS, Ageel AM. Protective effect of Swertia chirata against indomethacin and other ulcerogenic agentinduced gastric ulcers. Drugs Exp Clin Res. 1993; 19(2): 69-73.
- 17. Sharma R, Amin H, Galib, Prajapati PK. Antidiabetic claims of *Tinospora cordifolia* (Willd.) Miers: critical appraisal and role in therapy. Asian Pac J Trop Biomed. 2015 Jan 1; 5(1): 68–78.
- 18. Gupta S, Walia A, Malan R. Phytochemistry and pharmacology of Cedrus deodera: an overview. I J Pharm Sci Res 2011; 2: 2010.
- 19. Singh, Pradeep & Khosa, R & Mishra, Garima. (2013). Evaluation of antidiabetic activity of ethanolic extract of Cedrus deodara (Pinaceae) stem bark in streptozotocin induced diabetes in mice. Nigerian Journal of Experimental and Clinical Biosciences | January-December 2013 | Vol 1 | Issue 1 and 2. Nigerian Journal of Experimental and Clinical Biosciences. 33-38. 10.4103/2348-0149.123961.
- 20. Ahmad R, Srivastava SP, Maurya R et al. . Mild antihyperglycaemic activity in Eclipta alba, Berberis aristata, Betula utilis, Cedrus deodara, Myristica fragrans and Terminalia chebula. Indian J Sci Technol 2008; 1: 1–6.
- Jang EM, Choi MS, Jung UJ, et al. Beneficial effects of curcumin on hyperlipidemia and insulin resistance in high-fat-fed hamsters. Metabolism 2008; 57: 1576–1583
- 22. Chuengsamarn S, Rattanamongkolgul S, Luechapudiporn R, Phisalaphong C, Jirawatnotai S. Curcumin extract for prevention of type 2 diabetes. Diabetes Care. 2012 Nov; 35(11): 2121-7. doi:

10.2337/dc12-0116. Epub 2012 Jul 6. PMID: 22773702; PMCID: PMC3476912.

- 23. Chattopadhyay I, Biswas K, Bandyopadhyay U et al. Turmeric and curcumin: Biological actions and medicinal applications. Current Science, Vol. 87, No. 1, 10 July 2004: Page 44-53.
- 24. Subash AK, Augustine A. Hypolipidemic effect of methanol fraction of Aconitum heterophyllum wall ex Royle and the mechanism of action in dietinduced obese rats. Journal of Advanced Pharmaceutical Technology and Research. 2012; 3(4): 224–8.
- 25. Jain SK, Jain R, Jain N. Antiobesity potential of Aconitum Heterophyllum roots. International Journal of Pharma and Bio Sciences. 2019; 10(3): 65–74.
- 26. Semwal BC, Gupta J, Singh S, Kumar Y, Giri M. Antihyperglycemic activity of root of Berberis aristata DC in alloxan-induced diabetic rats. Int J Green Pharm (IJGP). 2009; 3(3). doi: 10.22377/ijgp.v3i3.97.
- 27. Ali MA, Alam NM, Yeasmin MS, Khan AM, Sayeed MA, Rao VB. Antimicrobial screening of different extracts of Piper longum Linn. Res J AgriBiol Sci., 2007; 3(6): 852-7.
- 28. Reddy KR. Remedial merits of Piper longum Linn with astonishing antidiabetic potential. International Journal of Green Pharmacy (IJGP), 2018 Nov 4; 11(04).
- 29. Hua S, Wang B, Chen R, Zhang Y, Zhang Y, Li T, Dong L, Fu X. Neuroprotective effect of dichloromethane extraction from piper nigrum L. and piper longum L. on permanent focal cerebral ischemia injury in rats. Journal of Stroke and Cerebrovascular Diseases, 2019 Mar 1; 28(3): 751- 60. -
- 30. Christudas, Sunil & Duraipandiyan, Veeramuthu & Agastian, Paul & Ignacimuthu, Savarimuthu. (2012). Antidiabetic effect of plumbagin isolated from Plumbago zeylanica L. root and its effect on GLUT4 translocation in streptozotocin-induced diabetic rats. Food and chemical toxicology: an international journal published for the British Industrial Biological Research Association. 50. 4356-4363. 10.1016/j.fct.2012.08.046.
- 31. Aels I, Env. Bulletin of Environment, Pharmacology and Life Sciences Screening for Antimicrobial, Antioxidant and Antidiabetic activity of Plumbago zeylanica Plant Extracts. Pharmacol Life Sci [Internet]. 2022 [cited 2024 Jun 4]; 11(9): 115–21.
- 32. Widhya Aligita, Elis Susilawati, Harini Septiani, Raihana Atsil(2018). "Antidiabetic activity of coriander (Coriandrum Sativum L) leaves' ethanolic extract", International Journal of

PharmaceuticalandPhytopharmacologicalResearch, 8(2), pp.59-63

- 33. Dhanapakiam, P & Joseph, J & Ramaswamy, V & Moorthi, Mahaly & Kumar, A. (2008). The cholesterol lowering property of coriander seeds (Coriandrum sativum): Mechanism of action. Journal of environmental biology/Academy of Environmental Biology, India. 29. 53-6.
- 34. Eltimamy M, Elshamarka M, Aboelsaad M, Sayed M, Moawad H. Effects of alcoholic extract of Terminalia Chebula dried fruit on blood biochemical profile in diabetic rats. J Diabetes Metab Disord. 2022 Jan 22; 21(1): 159-170. doi: 10.1007/s40200-021-00951-8. PMID: 35673508; PMCID: PMC9167356.
- 35. Murali YK, Chandra R, Murthy PS. Antihyperglycemic effect of water extract of dry fruits of Terminalia chebula in experimental diabetes mellitus. Indian J Clin Biochem. 2004 Jul; 19(2): 202-4. doi: 10.1007/BF02894285. PMID: 23105484; PMCID: PMC3454220.
- 36. Sabu MC, Kuttan R. Antidiabetic and antioxidant activity of Terminalia belerica. Roxb. Indian J Exp Biol. 2009 Apr; 47(4): 270-5. PMID: 19382723.
- 37. D'souza JJ, D'souza PP, Fazal F, Kumar A, Bhat HP, Baliga MS. Anti-diabetic effects of the Indian indigenous fruit Emblica officinalis Gaertn: active constituents and modes of action. Food Funct. 2014 Apr; 5(4): 635-44. doi: 10.1039/c3fo60366k. PMID: 24577384.
- D.N. Guhabakshi, P. Sensarma, D.C. Pal, A. Lexicon Medicinal Plants of India Naya Prakashan, Calcutta, India (2001), p. 135
- 39. G.R. Gandhi, A. Stalin, K. Balakrishna, S. Ignacimuthu, M.G. Paulraj, R. Vishal Insulin sensitization via partial agonism of PPARγ and glucose uptake through translocation and activation of GLUT4 in PI3 K/p-Akt signaling pathway by embelin in type 2 diabetic rats Biochim. Biophys. Acta, 1830 (2013), pp.
- 40. Premlata, Ram B, Vimlesh, Sachin Choudhary, Jitendra Khachariya. Role of Vidangadi Kwatha in Madhumeha (Diabetes Mellitus Type II). Archives of Clinical and Medical Case Reports 5 (2021): 629-633.
- Zhao L, Wang Y, Wang Z, Niu T, Yu J, Yue T. Secondary metabolites from Scindapsus officinalis (Roxb.) Schott. with in vitro antidiabetic activities. Fitoterapia. 2024 Mar; 173: 105822. doi: 10.1016/j.fitote.2024.105822. Epub 2024 Jan 13. PMID: 38224899.
- 42. Akhani SP, Vishwakarma SL, Goyal RK. Anti-diabetic activity of Zingiber officinale in streptozotocininduced type I diabetic rats. J Pharm Pharmacol.

2004 Jan; 56(1): 101-5. doi: 10.1211/ 0022357 022403. PMID: 14980006.

- 43. Kaleem M, Sheema, Sarmad H, Bano B. Protective Effects of Piper nigrum andVinca rosea in Alloxan induced Diabetic rats. Indian J Physiol Pharmacol 2005; 49 (1): 65–71
- 44. Kharbanda C, Alam MS, Hamid H, Javed K, Bano S, Ali Y, Dhulap A, Alam P, Pasha MA. Novel Piperine Derivatives with Antidiabetic Effect as PPAR-γ Agonists. Chem Biol Drug Des. 2016 Sep; 88(3): 354-62. doi: 10.1111/cbdd.12760. Epub 2016 May 27. PMID: 27037532.
- 45. Manoharan S, Silvan S, Vasudevan K, Balakrishnan S. Antihyperglycemic and Antilipidperoxidative effects of Piper longum (Linn.) Dried Fruits in Alloxan Induced Diabetic Rats. J Biol Sci. 2007; 6(1): 161–168.
- 46. Choudhary N, Khatik GL, Sharma R, Khurana N, Lobo R, Bhatt S, Tewari D, Suttee A. Ameliorative potential of Operculina turpethum against streptozotocin-induced diabetes in rats: biochemical and histopathological studies. 3 Biotech. 2021 Jun; 11(6): 309. doi: 10.1007/s13205-021-02811-x. Epub 2021 Jun 2. PMID: 34194901; PMCID: PMC8172823.
- 47. Siddha, Mohanraghupathy. (2013). Effect of Hydroalcoholic extract of Baliospermum montanum roots against diabetic nephropathy on rats. International journal of Pharmacology & Pharmacotherapeutics. 2. 263.
- 48. Chakrobarty U, Das H. Antidiabetic and antioxidant activities of Cinnamomum tamala leaf extract in Stz-treated diabetic rat. Glob J Biotech Biochem. 2010; 5: 12–8.
 - 49. Bisht S, Sisodia SS. Assessment of antidiabetic potential of Cinnamomum tamala leaves extract in streptozotocin induced diabetic rats. Indian J Pharmacol. 2011 Sep; 43(5): 582-5. doi: 10.4103/0253-7613.84977. PMID: 22022005; PMCID: PMC3195132.
 - Bisht S, Sisodia SS. Assessment of antidiabetic potential of Cinnamomum tamala leaves extract in streptozotocin induced diabetic rats. Indian J Pharmacol. 2011 Sep;43(5):582-5. doi: 10.4103/0253-7613.84977. PMID: 22022005; PMCID: PMC3195132.
 - 51. Fatemeh Y, Siassi F, Rahimi A, Koohdani F, Doostan F, Qorbani M, et al. The effect of cardamom supplementation on serum lipids, glycemic indices and blood pressure in overweight and obese prediabetic women: a randomized controlled trial. J Diabetes Metab Disord. 2017; 16: 1–9.
 - 52. Kazemi S, Yaghooblou F, Siassi F, Rahimi Foroushani A, Ghavipour M, Koohdani F, et al.

Cardamom supplementation improves inflammatory and oxidative stress biomarkers in hyperlipidemic, overweight, and obese pre-diabetic women: A randomized double-blind clinical trial. J Sci Food Agric. 2017; 97: 5296–5301.

- 53. Daneshi-Maskooni M, Keshavarz SA, Qorbani M, Mansouri S, Alavian SM, Badri-Fariman M, et al. Green cardamom increases Sirtuin-1 and reduces inflammation in overweight or obese patients with non-alcoholic fatty liver disease: a double-blind randomized placebo-controlled clinical trial. Nutr Metab. 2018; 15: 1–12.
- 54. Rahman MM, Alam MN, Ulla A, Sumi FA, Subhan N, Khan T, et al. Cardamom powder supplementation prevents obesity, improves glucose intolerance, inflammation and oxidative stress in liver of high carbohydrate high fat diet induced obese rats. Lipids Health Dis. 2017; 16: 1–12.
- 55. Yahyazadeh R, Ghasemzadeh Rahbardar M, Razavi BM, Karimi G, Hosseinzadeh H. The effect of Elettaria cardamomum (cardamom) on the metabolic syndrome: Narrative review. Iran J Basic Med Sci. 2021 Nov; 24(11): 1462-1469. doi: 10.22038/IJBMS.2021.54417.12228. PMID: 35317114; PMCID: PMC8917848.
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- 56. Anisetti, Ravinder N & Macherla, Saivivek & Goli, Venkateshwarlu & Dasarapu, Santhosha. (2012). Antidiabetic activity of bambusa arundinaceae stem extracts on alloxan induced diabetic rats. Journal of Chemical, Biological and Physical Sciences. 2. 2012.
- 57. Borde MK, Mohanty IR, Maheshwari U. DPP-4 inhibitory activity and myocardial salvaging effects of Commiphora mukul in experimental diabetes. Int J Basic Clin Pharmacol. 2019; 8: 575–83.
- 58. Mohanty IR, Kumar CS, Borde M. Antidiabetic activity of Commiphora mukul and Phyllanthus emblica and Computational analysis for the identification of active principles with dipeptidyl peptidase IV inhibitory activity. Indian J Pharmacol. 2021 Sep-Oct; 53(5): 384-387. doi: 10.4103/ijp.IJP _69_19. PMID: 34854407; PMCID: PMC8641738.
- 59. Satyavati G. V. Gum guggul (Commiphora mukul)the success story of an ancient insight leading to a modern discovery. Indian Journal of Medical Research. 1988; 87(4): 327–335.
- 60. Sharma B., Salunke R., Srivastava S., Majumder C., Roy P. Effects of guggulsterone isolated from Commiphora mukul in high fat diet induced diabetic rats. Food and Chemical Toxicology. 2009; 47(10):2631–2639. doi:10.1016/j.fct.2009.07.021.

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