Antiobesity Potential and Complex Phytochemistry of 
Momordica charantia Linn. with Promising Molecular Targets

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Karale et al.: Antiobesity potential of bitter melon

Momordica charantia L. grows in many tropical and subtropical regions, the fruits of which are gradually becoming popular for treating diabetes and associated diseases. Momordica charantia L. appears to be an inimitable species that synthesizes a diverse range of chemical constituents in the fruits, leaves, stem and roots. Over 248 compounds belonging to the lipids, phenolics and terpenoids classes have been reported in various studies. The cucurbitane types of triterpenes exist in various parts of this plant as aglycones as well as glycosylated forms. Momordica charantia L. has been comprehensively studied worldwide for therapeutic properties to treat a number of diseases like diabetes, dyslipidaemia, obesity and certain cancers. The bitter melon seems to exert lipid lowering and antiobesity effects via several mechanisms like PPARs, LXRs, SREBPs, and Sirts-mediated fat metabolism in various tissues, prevent adipocyte hypertrophy and visceral fat accumulation. This article presents an extensive literature review on the vast potential of bitter melon as antiobesity agent with supporting data on complex phytochemistry.

Key words: Obesity, phytochemicals, bitter melon, cucurbitane-type terpenoids

Momordica charantia L. is a tropical or subtropical creeper belonging to family Cucurbitaceae and widely used as a medicinal herb from ancient time (fig. 1). M. Charantia L. or bitter melon, also known as balsam pear or Karela, is a vegetable and common food in Indian cuisine and has been used comprehensively in folk medicine[1]. The Latin name Momordica means ‘to bite’ referring to the serrated edges of the leaf, which appear as if they have been bitten[2]. The major regions of M. charantia L. cultivation are Asia including China, India, Sri Lanka and Thailand, Central and South America and North America[3]. In Ayurveda, the fruit is considered as tonic, stomachic, stimulant, emetic, antibilious, laxative and alterative[4]. Bitter melon has been used in various Asian traditional medicine systems for a long time. It is well recognized, the plant is extensively in use in the Chinese, Ayurvedic and Indonesian systems of medicines as well as in Japan[5].

The therapeutic significance of the plant is symbolized by the fruits which contain about half a dozen seeds per gram of the fresh fruit. As the name implies, the fruits are bitter and bitterness enhance with the level of maturity and hence earlier harvesting required to battle bitter taste. The leaves and young shoots of bitter melon recognized to be used in traditional medicine as a herbal tea. The range of pharmacological activities reported for bitter melon is rapidly increasing in recent years and its claimed uses and potential applications for cancer and other diseases have been extensively reviewed[3]. Likewise, the range of medicinal claims range from diabetes, hypertension, obesity, cancer as well as AIDS.

During the past 2 decades, it has been reported that M. charantia L. possessed various beneficial effects, including antiinflammatory[6], antioxidant[7], cytotoxic[8], anticancer[9-11], antiobesity[12], antiviral[13], antiHIV[14], antiulcer[15], cholesterol lowering[16], inhibition of protein tyrosine phosphatase 1B[17] and

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This review aims to highlight the complex phytochemistry and extensive review on antiobesity potential of bitter melon with possible targets.

**Botanical description:**

*M. charantia* L. was known previously by several synonyms including the *Cucumis argyi* H.Lev., *Cucumis intermedius* M. Roem., *Momordica anthelmintica* Schumach. & Thonn., *M. charantia* subsp. *abbreviata* (Ser.) Greb., *M. charantia* var. *abbreviata* Ser., *M. charantia* f. *muricata* (Willd.), *M. charantia* var. *muricata* (Willd.) Chakrav., *M. charantia* var. *psuedobalsamina* Griseb., *M. charantia* var. *zeylanica* Hitche., *M. elegans* Salisb., *M. indica* L., *M. jagorana* K.Koch., *M. muricata* Willd., *M. papillosa* Peckolt ex Rosenthal., *M. roxburghiana* G. Don., *M. senegalensis* Lam., *M. thollonii* Cogn., *M. zeylanica* Mill. The botanical description of different parts of the plant is presented in Table 1[19].

Cucurbitaceae family is known to comprise some 101 accepted genera and the genus *Momordica* L. itself comprises of some 50 accepted species within the family[20]. The taxonomic hierarchy of bitter melon within the plant kingdom is as follows: Kingdom: Plantae; Subkingdom: Viridiplantae; Superdivision: Embryophyta; Division: Tracheophyta; Subdivision: Spermatophytina; Class: Magnoliopsida; Super order: Rosanae; Order: Cucurbit ales; Family: Cucurbitaceae; Genus: Momordica Linn.; Species: *Momordica charantia* L.-balsampear.

The existence of many varieties of bitter melon has been noted[21], for example, it was reported that the fruits of the wild variety of *M. charantia* L. growing in Trinidad and Tobago were of one-fifth in size as compared to the cultivated Indian ones.

**Phytochemistry:**

The main constituents of bitter melon are triterpenoids, saponins, protein, polysaccharides, steroids, alkaloids, lipids, and phenolic compounds[22]. Several bioactive compounds of *M. charantia* L. have been recorded and literature reported that these are responsible for various pharmacological effects as depicted in Table 2[23-49].

**Triterpenoids:**

The most abundant phytochemical components of bitter melon fruits are the triterpenoid class of secondary metabolites and are well-known for their bitterness and toxicity. These are divided in 2 types primarily the cucurbitane-type and to a less extent oleane-type, which may occur either in their glycosylated or aglycone forms (fig. 2). The fruits are predominant source of terpenoids with great deal of structural diversity but the leaves, stems and roots have also been shown to be good sources of these compounds[23]. An extremely large number (193) of cucurbitane-type triterpenes were isolated from bitter melon that exhibited various pharmacological effects (Table 3). The sugar monomers as β-D-glucopyranosyl, β-D-allopyranosyl, β-D-xylpyranosyl occur in cucurbitane-type triterpenes either by their own as O-linked glycosides, or in different combinations as disaccharides or polysaccharides. The rare glycoside in these compounds is the 3-keto-glucoside[50].

Momordicosides A and B were isolated firstly from the seeds of bitter melon fruits[51]; while, momordicosides C, D and E were isolated as minor components of

![Fig. 1: Plant parts of *M. charantia* L.](a) Leaves; (b) unripe fruit; and (c) ripe fruit and seeds of plant
the seeds. The study on the fruits of *M. charantia* L. further added a momordicosides F1, F2, G, I, K and L as novel compounds in previously reported ones. The momordicosides I and momordicosides M along with other compounds have also been isolated from the fruits of bitter melon. Further additions of momordicosides M-O as new compounds were confirmed by chemical examination of the fresh fruits. The Vietnamese origin dried fruits of *M. charantia* L. had shown existence of an auxiliary three pioneering momordicosides U, V and W.

The cucurbitane-type triterpenes, which are known to be responsible for the bitterness of the leaves and vines of the *M. charantia* L. are momordicines. The momordicine IV and the malonyl derivative of momordicine II, momordicine V are readily available in the leaves of bitter melon. The isolation of momordicines VI, VII, and VIII were first confirmed from the stems and leaves of bitter melon. The goyaglycosides-a, -b, -c, -d, -e, -f, -g, and -h were isolated from the fresh fruits of Japanese *M. charantia* L. A novel compound goyaglycoside I was an

| Plant parts | Description |
|-------------|-------------|
| Leaves      | Broadly ovate to orbicular in outline, cordate, narrowly decurrent on to petiole, sparsely pubescent to densely villous on veins beneath, deeply palmately 3-7-lobed, lobes variously sinuate-dentate or lobulate. Leaf lamina 10×12.5 cm. Flowers are monoeocious and solitary. Male flowers: peduncle 0.3-5 cm long; bract 2-17 mm long, broadly ovate or reniform, sessile, cordate, amplexicaul, pedicel 2-9.5 cm long. Receptacle-tube 1-5 mm long; lobes 3-7 mm long, ovate-lanceolate. Petals 1.0-2.5 cm long, pale to deep yellow, ovate to obovate. |
| Flowers     | Fruits: peduncle 0.2-5 cm long; bract 1-12 mm long; pedicel 1-10 cm long; ovary 8-11×2-4 mm, ovoid-rostrate to fusi-form, ridged, pilose on ridges, tuberculate; receptacle-tube 1-3 mm long, lobes 2-5 mm long, lanceolate; petals 0.7-1.2 cm long. Female flowers: peduncle 0.2-5 cm long; bract 1-12 mm long; pedicel 1-10 cm long; ovary 8-11×2-4 mm, ovoid-rostrate to fusi-form, ridged, pilose on ridges, tuberculate, bright orange-red, dehiscent into 3 valves; fruit-stalk 3.4-15 cm long. |
| Fruits      | Fruit 2.5-4.8×1.5-2.3 cm, ovoid-rostrate or ellipsoid, longitudinally ribbed and tuberculate, bright orange-red, dehiscent into 3 valves; fruit-stalk 3.4-15 cm long. |
| Seeds       | Seeds 8-11, 4.5-8×2-3.5 mm, enveloped in sticky red pulp, ovate-elliptic to oblong in outline; faces flattened, sculptured, with sinuate edges; margins grooved. Petiole Petiole 0.5-7 cm long. |

### TABLE 1: BOTANICAL DESCRIPTION OF MOMORDICA CHARANTIA LINN.

| Bioactive compounds       | Distribution                          | Pharmacological effects                                      | References     |
|---------------------------|---------------------------------------|--------------------------------------------------------------|----------------|
| Triterpenoids             | Leaves, stem, fruits                  | Cancer chemo protective, anticancer antioxidant, antidiabetic | [23-26]        |
| Peptides and proteins     | Seed                                  | Antiviral, anti-tumour, immune suppressant, antimicrobial     | [27-30]        |
| Phenolics                 | Fruit and seed                        | Antioxidant, anti-inflammatory, immunostimulant              | [31-34]        |
| Saponin                   | Fruit, root, seed                     | Antihyperglycemic, hypolipidemic, antiviral, bacteriostatic  | [35-39]        |
| Polysaccharide            | All parts of plant                    | Antioxidant, antidiabetic, immune enhancement, neuroprotective, antitumor | [40-44] |
| Lipid                     | Seed                                  | Anti-tumor, antioxidant                                      | [45,46]        |
| Steroids                  | Fruit and pericarp                    | Antimicrobial                                               | [47-49]        |

**Fig. 2:** The cucurbitane and oleane type triterpene skeleton of *M. charantia* L.
### TABLE 3: CUCURBITANE TYPE OF TRITERPENOIDS OF *MOMORDICA CHARANTIA* L.

| Phytochemical Name | Plant parts | Chemical structure | Pharmacological effects |
|--------------------|-------------|--------------------|------------------------|
| Momordicoside A & B | Fruits and seeds | ![Chemical structure](image1.png) | Antidiabetic, antiobesity, anticancer |
| Momordicoside K | Leaves, fruits and roots | ![Chemical structure](image2.png) | Antiproliferative, hypoglycemic, antiobesity, antioxidant |
| Momordicoside I & F1 | Fruits | ![Chemical structure](image3.png) | Antiproliferative, hypoglycemic, antiobesity, disaccharidase |
| Momordicoside G & F2 | Leaves and fruits | ![Chemical structure](image4.png) | Antiproliferative, hypoglycemic, antiobesity |
| Momordicine I & II | Leaves, vines and fruits | ![Chemical structure](image5.png) | Cytotoxic, antiinflammatory, antiviral, immunomodulatory, antiobesity |
| Goyaglycoside-a & b | Leaves and vines | ![Chemical structure](image6.png) | Antiproliferative, hypoglycemic, antiobesity, anticancer |
| Goyaglycoside-c & d | Fruits | ![Chemical structure](image7.png) | Antiproliferative, hypoglycemic, antiobesity, anticancer |
| Goyaglycoside-e | Fruits | ![Chemical structure](image8.png) | Cytotoxic, antiobesity, antidiabetic |
| Goyaglycoside-f | Fruits | ![Chemical structure](image9.png) | Antiobesity, antidiabetic |
| Goyaglycoside-g | Fruits | ![Chemical structure](image10.png) | Hypoglycemic, antiobesity |
Goyaglycoside-h  Fruits  Hypoglycemic, antiobesity

Kuguacin B  Roots  Anticancer, antidiabetic

Kuguacin C & D  Roots  AntiHIV-1, antidiabetic

Kuguacin E  Roots  AntiHIV-1, hypoglycemic

Kuguacin F  Leaves and vines  Hypoglycemic, lipid lowering

Kuguacin G  Leaves and vines  Antidiabetic, antiobesity, anticancer

Kuguacin K  Leaves and vines  Anticancer, antiproliferative, hypoglycemic

Kuguacin H  Leaves and vines  Hypoglycemic, antiproliferative

Kuguacin I  Leaves and vines  Anticancer, lipid lowering

Kuguacin J  Leaves and vines  Anticancer, hypoglycemic
Kuguacin L  Leaves and vines  Antiproliferative, hypoglycemic

Kuguacin M  Leaves and vines  Hypoglycemic, anticancer

Kuguacin P & Q  Leaves and vines  Hypoglycemic, antiproliferative

Kuguacin R  Leaves, stems and fruits  Antioxidant, hypoglycemic, lipid lowering

Kuguacin S  Leaves and vines  Lipid lowering, hypoglycemic

Charantoside I  Fruits  Hypoglycemic, Antiobesity

Charantoside II  Fruits  Hypoglycemic, antiobesity

Charantoside III  Fruits  Hypoglycemic, Antiobesity

Charantoside IV  Fruits  Hypoglycemic, antiobesity

Charantoside V  Fruits  Hypoglycemic, antiobesity
Charantoside VII Fruits Hypoglycemic, antiobesity

Charantoside VIII Fruits Hypoglycemic, antiobesity

Goyasaponin I Fruits Hypoglycemic, antiobesity

Goyasaponin II Fruits Hypoglycemic, antiobesity

Goyasaponin III Fruits Hypoglycemic, antiobesity

Karavilagenin C Fruits Antidiabetic

Karavilose I, II & III Fruits Antiproliferative, antidiabetic, hypolipidemic

Kuguaglycoside G Roots Cytotoxic, antiproliferative, hypoglycemic, anticancer

Momordicinin Fruits Antidiabetic, antiobesity, lipid lowering
additional triterpene isolated from the immature fruit of bitter melon\(^6\). The novel cucurbitane-type triterpene called karavilagenins A-C and 5 new triterpene glycosides called karavilosides I-V were isolated from the dried fruit of \(M.\) charantia \(\text{L.}\)\(^6\).

The methanol extracts of the fruits of Japanese \(M.\) charantia \(\text{L.}\) showed charantosides I, II, III-VI, VII and VIII, however charantoside IX and X were the other novel compounds reported\(^6\). The 3 novel compounds, charantosides A, B and C were obtained from fruits of bitter melon\(^\text{[22]}\). Another group of interesting triterpenoids are those known by their trivial names kuguacins. kuguacins A-E and kuguacins F-S were isolated from the roots and the leaves of bitter melon plant, respectively\(^6\). Kuguacins II-VI were novel compounds isolated together with various other known compounds from the fruit of \(M.\) charantia \(\text{L.}\)\(^6\). From the aqueous ethanol extracts of fresh fruits 8 novel cucurbitane-type glycosides were isolated, which were named kuguasaponins A-H\(^8\). The ethanol extract of fruits of \(M.\) charantia \(\text{L.}\) identified 15 cucurbitane-type triterpene glycosides including 4 new compounds, kuguasides A-D\(^3\).

Flavonoids and phenolic compounds:

A number of phenolic compounds with many biological activities have been isolated from bitter melon including, coumaric, caffeic, and ferulic acids as well as the caffeic acid ester, chlorogenic acid, benzoic, gallic and gentisic acid\(^6\). The major flavonoids and phenolic acids in the dried leaves of bitter melon were also analyzed and found to be rutin, gentisic acid and coumaric acid\(^6\). While the phenolic acids and flavonoids as well as their glycosides could be readily extracted with water, their non-polar derivatives may be present in the oil components of the plant\(^6\).

Other components:

Other than the bioactive compounds, unsaturated fatty acids, alkaloids, amino acids minerals and vitamins are also present in bitter melon\(^\text{[69-71]}\). The extracts of bitter melon showed the presence of 9 kinds of unsaturated fatty acids\(^7\). It has also been demonstrated that 12, 13 and 12 fatty acids are found in the young, mature and senescent leaves of \(M.\) charantia \(\text{L.}\) representing 87.3, 95.25, and 83.11 % of the total fatty acids\(^7\). The contents of total amino acids and the free amino acids of \(M.\) charantia \(\text{L.}\) were 11.99 and 2.36 % as determined by acid hydrolysis and amino acid analysis\(^\text{[74]}\). In addition, bitter melon is a natural source for vitamins; ascorbic acid was detected in the range of 440-780 mg in the fruit fraction\(^\text{[75]}\). Vicine, an alkaloidal agent was isolated from the seeds of bitter melon, which is responsible for hypoglycemic activity\(^\text{[76]}\).

Antiobesity and lipid lowering effects:

The action of bitter melon in lowering fat has been supported by plentiful studies, its effect on the level
of serum FFAs have been contradictory with some authors showing reduction, some shown same level, and others reported an increased levels. For example, the serum FFAs concentration increased in obese rats treated with bitter melon as reported by Chen et al. [77]. An increased level of TG and LDL-C in the serum that is representative of either overproduction by the liver or defective removal from the circulation or the overall dyslipidaemia in diabetes. It is not clear why bitter melon increases the serum level of FFAs. It may facilitate fat mobilization due to suppression of lipogenesis or lipid deposition. While other studies revealed that M. charantia L. could lower the serum and liver TG levels [78]. Various experimental studies reported a decrease in serum TC, TG and LDL-C concentrations and an increase in serum high density lipoprotein-cholesterol (HDL-C) by bitter melon [79,80].

The findings of experimental research concluded that M. charantia L. might reduce fasting insulin, TG, cholesterol and epididymal fat, which were increased by HFD. The dwindling of insulin resistance, improved glucose tolerance, and increased insulin signaling under HFD-induced insulin-resistance and elevated serum lipids may also been shown by bitter melon. Administration of an aqueous extract of unripe fruits of bitter melon improved glucose and insulin tolerance together with inhibition of plasma apoB-100 and apoB-48. An animal study had shown evidence for a potent inhibitor of apoB secretion and TG synthesis as well as plasma lipid and VLDL levels in bitter melon juice [81]. Overall many studies on the fruits, seeds and aerial parts of M. charantia Linn have shown reduced adiposity, lowered serum insulin and normalized glucose tolerance in rats fed with a HFD. The body weight and visceral fat mass of bitter melon-treated obese rats were shown to be lowered [77,82].

While another study revealed that bitter melon supplementation into HFD notably suppressed the levels of fatty acid synthase (FAS), acetyl-CoA carboxylase-1 (ACC-1), lipoprotein lipase (LPL) and adipocyte fatty acid binding protein [83]. Water extract of M. charantia L. fruits at a dose of 1 g/kg revealed to be effective in improving the obesity-induced hyperglycaemia and hyperleptinemia [84], indicating that bitter melon can reduce insulin resistance, visceral fat accumulation and adipocyte hypertrophy probably by down-regulating the expression of key lipogenic genes or proteins in adipose tissues. Aqueous fruit extract of M. charantia L. significantly reduced levels of serum TG, TC, LDL and VLDL at a dose of 350 mg/kg in experimental rats [85]. Numerous animal studies have reported efficacy of bitter melon in ameliorating the weight gain and regulation of lipid metabolism [86,87].

The methanol extract of fruits of bitter melon showed antidiabetic and antihyperlipidemic action during different seasons of the year, this suggested that antidiabetic and hypolipidemic activity of M. charantia L. may fluctuate according to the quantity and quality of active constituents during different seasons of the year and reach the peak during spring [88]. Bitter melon seed oil had shown significantly decreased in body weight, Lee’s index, fat index and adipose size in the HFD mice. Meanwhile, serum FFAs levels returned to normal at a dose of 10 g/kg [89]. M. charantia L. extracts have antiobesity effects and the ability to modulate lipid profile of mice fed a HFD by suppressing body weight gain, visceral tissue weight, plasma and hepatic lipid concentrations, and lipid peroxidation along with increasing lipid metabolism. The plasma TG, TC, and LDL-C levels along with hepatic TG and TC concentrations were considerably lowered in HFD-fed mice by M. charantia L. extracts. Also elevated plasma HDL-C levels and fecal TG concentration were shown to be elevated in animals treated with these extracts. The extracts exhibited antiobesity effects in HFD-fed mice by inhibiting lipid peroxidation while increasing lipid metabolism [90]. Bitter melon extract showed useful benefit on body weight gain and fat deposition, . 3 shows the probable molecular targets for improving obesity. Mechanism of action of M. charantia L. extracts for antiobesity effect appeared to be, increasing fatty acid oxidation, which could be responsible for reducing weight; decrease in weight of epididymal white adipose tissue (WAT) and adipose leptin; enhanced hepatic and muscle mitochondrial carnitine palmitoyl transferase-I (CPT-I) and acetyl-CoA dehydrogenase enzyme; prevent adipocyte hypertrophy; reduce visceral fat accumulation through dropping mRNA level of fatty acid synthase, lipoprotein lipase, adipocyte fatty acid binding protein and decreasing the lipogenic genes in adipose tissue; downregulation of PPARγ, SREPP and perilipin mRNA gene expression.

Moreover, bitter melon extracts reduced lipid accumulation during differentiation from a pre-adipocyte to adipocyte and down-regulated PPARγ [91]. PPARγ is considered the master regulator of adipogenesis during differentiation of pre-adipocyte to adipocyte [92]. Bitter melon juice inhibited adipocyte differentiation by reducing PPARγ, SREBP, and perilipin mRNA gene expression and by increasing
lipolysis in primary human adipocyte\textsuperscript{[93]}. Another study showed that bitter melon reduced TG and LDL levels and increased HDL levels in high sucrose fed rats\textsuperscript{[94]}. Female Zucker rats supplemented with ground bitter melon seeds showed decreased TC and LDL-C and increased HDL-C\textsuperscript{[95]}. Several transcriptional regulatory factors like AMPK, PPAR-\(\gamma\), and PGC-1\(\alpha\) regulate the mitochondrial biogenesis, which would be a possible way of increasing lipid metabolism and utilization in energy demanding cells and tissues\textsuperscript{[96,97]}. AMPK synchronized PPAR-\(\gamma\) and PGC-1\(\alpha\) activation encouraged most of the transcriptional signal to augment fatty acid oxidation and mitochondrial function\textsuperscript{[98-100]} . PGC-1\(\alpha\) stimulates mitochondrial biogenesis and respiration in multiple cell types and modulates biological programs normally associated with increased oxidative metabolism. Also decreased plasma level of TGs, cholesterol, and FFA was found in plasma of rats fed a HF diet revealed by bitter melon supplementation due to up regulation and activation of PGC-1\textsuperscript{[101]} . A recent investigation also reported that increased hepatic AMPK p, AMPK \(\alpha1\), AMPK \(\alpha2\), and Sirt1 content in HFD-fed mice with supplementation of 1.2 % bitter melon extract\textsuperscript{[102]}.

LXRs were first recognized as orphan members of the nuclear receptor plays an important role in lipid and cholesterol metabolism and oxidized derivatives of cholesterol act as ligands for the LXRs. A high cholesterol diet fed mice develop enlarged fatty livers, degeneration of liver cells, high cholesterol levels in liver, and impaired liver function by LXR\(\alpha\)knockout\textsuperscript{[103]} . The \textit{M. charantia} L. extract supplementation decreased hepatic LXR\(\alpha\), which was responsible for decreased serum TC and LDL-C, HDL-C in high cholesterol diet Wistar rats\textsuperscript{[104]} . Bitter melon extract was a potent inhibitor of lipogenesis and stimulator of lipolysis in 3T3-L1 pre-adipocyte shown by researcher. The study shown that increasing the ethanol extract concentration led to a decrease in lipid accumulation in adipocyte both during and after differentiation in adipogenesis assay with a near 50 \% reduction\textsuperscript{[105]}.

**Toxic effects:**

Severe adverse reactions were not reported during short term studies while extensive data on the potential toxic effect of bitter melon are not available. Bitter melon fruits are edible and assumed to be well tolerated, at the same time toxicological evidences were reported to discover its therapeutic potential for diabetes. Two cases of acute intoxication reported after taking bitter melon tea\textsuperscript{[106]} . Abdominal pain as a side effect has also been reported in some studies\textsuperscript{[107]} . The antifertility and abortifacient effects of the \textit{M. charantia} L. reported in animals also suggests further investigation\textsuperscript{[108-111]} . An acute disease favism characterized by hemolytic anemia, in individuals with a hereditary loss of the enzyme glucose-6-phosphatase has been shown by vicine found in fava bean\textsuperscript{[112]} . Consequently, the presence of vicine in bitter melon seeds was also suggested to put patients with glucose-6-phosphatase deficiency at risk\textsuperscript{[113]} . Although there have been no reports on favism induced by bitter melon, individuals susceptible to the disease should avoid eating the fruit.

Current research is directed to reduce the bitterness of \textit{M. charantia} L. preparations attributes to the triterpene compounds and increasing tolerability by the general public through various formulation approaches. Some recent studies used \(\beta\)-cyclodextrin at 0.25-2 \% concentrations to improve sensory quality, total phenolic content, antioxidant activity and antidiabetic potential of \textit{M. charantia} L. juice\textsuperscript{[114]} . Various encapsulation methods of bitter melon extracts along with optimized spray-drying techniques were also scrutinized to obtain the powder\textsuperscript{[115]} .

To date, \textit{M. charantia} L. has been broadly studied globally for its medicinal properties and to treat a number of diseases like diabetes, dyslipidemia, obesity, and certain cancers. Isolated compounds from this plant like cucurbitane type terpenoids, flavonoids and phenolic acid and extracts possess antiobesity potential similar to its crude extracts. The extracts of fruits and different compounds seem to exert their beneficial effects via several mechanisms like PPARs, LXRs and SREBPs mediated fat metabolism in various tissues, reduces visceral fat accumulation and adipose hypertrophy. These mechanisms will be directly related to controlling and treating diabetes mellitus, dyslipidemia, obesity and related cardiovascular complications. Thus, further studies are required to conduct more double blind randomized trials with bitter melon extracts in obese population. Numerous \textit{in vitro} studies have also been employed to demonstrate these effects and further establish. In this review, we summarized phytoconstituents of bitter melon and its antiobesity potential with mechanism of action. This compilation of phytochemicals and antiobesity activity of \textit{M. charantia} L. would help the researchers in dereplication and designing new untried strategies.

**REFERENCES**

1. Habicht SD, Kind V, Rudloff S, Borsch C, Mueller AS, Pallauf
1. J, et al. Quantification of anti-diabetic extracts and compounds in bitter gourd varieties. Food Chem 2011;126:172-76.

2. Subratty AH, Gurib-Fakim A, Mahomoodally F. Bitter melon: An exotic vegetable with medicinal values. Nutr Food Sci 2003;35:143-47.

3. Grover JK, Yadav SP. Pharmacological actions and potential uses of Momordica charantia: A review. J Ethnopharmacol 2004;93:123-32.

4. Bailey CJ, Day C, Turner SL, Leatherdale BA. Cerasee: a traditional treatment for diabetes. Studies in normal and streptozotocin diabetic mice. Diabetes Res 1985;2:81-4.

5. Raman A, Lau C. Anti-diabetic properties and phytochemistry of Momordica charantia L. (Cucurbitaceae). Phytomedicine 1996;2:349-62.

6. Manabe M, Takenaka R, Nakasa T, Okinaka O. Induction of anti-inflammatory responses by dietary Momordica charantia L. (bitter gourd), Biosci. Biotechnol Biochem 2003;67(12):2512-17.

7. Shan B, Xie JH, Zhu JH, Peng Y. Ethanol modified supercritical carbon dioxide extraction of flavonoids from Momordica charantia L. and its antioxidant activity. Food Bioprod Process 2012;90(3):579-87.

8. Zeng K, He YN, Yang D, Cao JQ, Xia XC, Zhang SJ, et al. Cucurbite-type glycosides from the fruits of Momordica charantia and their hypoglycaemic and cytotoxic activities. J Funct Foods 2014;6:564-74.

9. Ganguly C, De S, Das S. Prevention of carcinogen-induced mouse skin papilloma by whole fruit aqueous extract of Momordica charantia, Eur J Cancer Prev 2000;9(4):283-88.

10. Li CJ, Tsang SF, Tsai CH, Tsai HY, Chyuan JH, Hsu HY. Momordica charantia extract induces apoptosis in human cancer cells through caspase-and mitochondria-dependent pathways. Evid Based Complement Altern Med 2012;26:1971.

11. Nerurkar P, Ray RB. Bitter melon: antagonist to cancer. Pharm Res 2010;27(6):1049-53.

12. Chan LL, Chen QX, Go AG, Lam EKY, Li ET. Reduced adiposity in bitter melon (Momordica charantia)-fed rats is associated with increased lipid oxidative enzyme activities and uncoupling protein expression. J Nutr 2005;135(11):2517-23.

13. Puri M, Kaur I, Kanwar R, Gupta R, Chauhan A, Kanwar J. Ribosome inactivating proteins (RIPs) from Momordica charantia for anti-viral therapy. Curr Mol Med 2009;9(9):1080-94.

14. Fang EF, Ng TB. Bitter gourd (Momordica charantia) is a cornucopia of health: a review of its credited anti-diabetic, anti-HIV, and anti-tumor properties. Curr Mol Med 2011;11(5):417-36.

15. Alam S, Asad M, Asdaq SMB, Prasad VS. Antiulcer activity of methanolic extract of Momordica charantia L. in rats. J Ethnopharmacol 2009;123(3):464-9.

16. Jayasooriya AP, Sakono M, Yukizaki C, Kawano M, Yamamoto K, Fukuda N. Effects of Momordica charantia powder on serum glucose levels and various lipid parameters in rats fed with cholesterol-free and cholesterol-enriched diets. J Ethnopharmacol 2000;72(1):331-6.

17. Zeng K, He YN, Yang D, Cao JQ, Xia XC, Zhang SJ, et al. New compounds from acid hydrolyzed products of the fruits of Momordica charantia L. and their inhibitory activity against protein tyrosine phosphatases1B. Eur J Med Chem 2014;81:176-80.

18. Yang ZG, Shi WZ, Shen ZG, Zhang YP, Yang J. Preventive Effect of Momordica charantia L. saponins on osteoporosis in ovariectomized rats. Food Sci 2010;31(7):272-5.

19. Momordica charantia L. Plants of the World online. [cited 30 Dec 2018] Available from: http://powo.science.kew.org/taxon urn:lsid:ipni.org:names:293413-1.

20. Joseph B, Jini D. Antidiabetic effects of Momordica charantia (bitter melon) and its medicinal potency. Asian Pac J Trop Dis 2013;3(2):93-102.

21. Svorobodova BB, Calhelha L, Hellen RC, Alves S, Walcott MJ. Bioactive properties and phenolic profile of Momordica charantia L. medicinal plant growing wild in Trinidad and Tobago. Ind Crop Prod 2017;95:365-73.

22. Shuo J, Mingyue S, Zhang F, Xie J. Recent Advances in Momordica charantia: Functional Components and Biological Activities. Int J Mol Sci 2017;18:2555.

23. Akihisa T, Higo N, Tokuda H, Ukiya M, Akazawa H, Tochigi Y, et al. Cucurbite-type triterpenoids from the fruits of Momordica charantia and their cancer chemopreventive effects. J Nat Prod 2007;70:1233-9.

24. Agrawal RC, Beohar T. Chemo preventive and anticarcinogenic effects of Momordica charantia extract. Asian Pac J Cancer Prev 2010;11:371-5.

25. Liu CH, Yen MH, Tsang SF, Gan KH, Hsu HY, Lin CN. Antioxidant triterpenoids from the stems of Momordica charantia. Food Chem 2010;118:751-6.

26. Chou CH, Liao MH, Chen TM, Cheng CH, Anggriani R, Tsai CP et al. Bitter melon triterpenes work as insulin sensitizers and insulin substitutes in insulin-resistant cells. J Funct Foods 2015;13:214-24.

27. Upadhyay A, Agrahari P, Singh DK. A review on salient pharmacological features of Momordica charantia. Int J Pharmacol 2015;11:405-13.

28. Meng Y, Liu S, Li J, Meng Y, Zhao X. Preparation of an antitumor and antivirus agent: Chemical modification of a-MMC and MAP30 from Momordica charantia L. with covalent conjugation of polyethylene glycol. Int J Nanomed 2012;7:3133.

29. Leung SO, Yeung HW, Leung KN. The immunosuppressive activities of two abortifacient proteins isolated from the seeds of bitter melon (Momordica charantia). Immunopharmacology 1987;13:159-71.

30. Jabeen U, Khanum A. Isolation and characterization of potential food preservative peptide from Momordica charantia L. Arabian J Chem 2017;10:S3982-9.

31. Anila L, Vijayalakshmi NR. Beneficial effects of flavonoids from Sesamum indicum, Emblica officinalis and Momordica charantia. Phytother Res 2000;14:592-5.

32. Qader SW, Abdulla MA, Chua LS, Najim N, Zain MM, Hamdan S. Antioxidant, total phenolic content and cytotoxicity evaluation of selected Malaysian plants. Molecules 2011;16:3433-43.

33. Bajpai M, Pande A, Tewari SK, Prakash D. Phenolic contents in selected fruits and vegetables, as well as their stimulatory effects on mouse splenocyte proliferation. Food Chem 2007;101:140-7.

34. Lin JY, Tang CY. Determination of total phenolic and flavonoid contents in selected fruits and vegetables, as well as their stimulatory effects on mouse splenocyte proliferation. Food Chem 2007;101:140-7.

35. Hsiao PC, Liaw CC, Hwang SY, Cheng HL, Zhang LJ, Shen CC, et al. Anti-proliferative and hypoglycemic cucurbitane-type glycosides from the fruits of Momordica charantia. J Agric Food Chem 2013;61:2979-86.

36. Keller AC, Ma J, Kavalier A, He K, Brillantes AMB, Kennelly EJ. Saponins from the traditional medicinal plant Momordica
52. Chang CI, Chen CR, Liao YW, Cheng YW, Chen YC.
51. Begum S, Ahmed M, Siddiqui BS, Khan A, Saify ZS, Arif M.
50. Saeed S, Tariq P. Antibacterial activities of
49. Dhar P, Chattopadhyay K, Bhattacharyya D, Roychoudhury
48. Miyahara Y, Okasbe H, Yamauchi T. Studies on the constituents
47. Momordica charantia
46. Isolation and characterization of momordicosides A ang B,
45. Chen JC, Lau CBS, Chan JYW, Fung KP, Leung PC, Liu JQ
44. Kuguacins F-S, cucurbitane triterpenoids from
43. Chen JC, Liu WQ, Lu L, Qiu MH, Zheng YT, Yang LM.
42. Kashiwagi T, Mekuria DB, Dekebo A, Sato K, Tebayashi
41. Yue J, Sun Y, Xu J, Cao J, Zhao Y. Cucurbitane triterpenoids
40. Tsuzuki T, Tokuyama Y, Igarashi M, Miyazawa T. Tumor
growth suppression by eleostearic acid, a linolenic acid
isomer with a conjugated triene system, via lipid peroxidation.
Carcinogenesis 2004;25:1417-25.
39. Saced S, Tariq P. Antibacterial activities of Mentha piperita,
Pisum sativum and Momordica charantia. Pak J Bot 2005;37:997.
38. Begum S, Ahmed M, Siddiqui BS, Khan A, Saify ZS, Arif M.
Triterpenes, a sterol and a monocyclic alcohol from Momordica charantia. Phytochemistry 1997;44:1313-20.
37. Horax R, Hettiarachchy N, Chen P. Extraction, quantification,
and antioxidant activities of phenolics from pericarp and seeds of bitter melons (Momordica charantia) harvested at three maturity stages (immature, mature, and ripe). J Agric Food Chem 2010;58:4428-33.
36. Zhang M, Hettiarachchy NS, Horax R, Chen P, Over KF. Effect of maturity stages and drying methods on the retention of selected nutrients and phytochemicals in bitter melon (Momordica charantia) leaf. J Food Sci 2009;74(6):C441-8.
35. Minh NP. Extraction of polyphenol in bitter melon (Momordica charantia). JIMRD 2014;1(4):115-25.
34. Chuang CY, Hsu C, Chao CY, Wein YS, Kuo YH, Huang CJ. Fractionation and identification of 9c, 11t, 13t-conjugated linolenic acid as an activator of PPAR in bitter gourd (Momordica charantia L.). J Biomed Sci 2006;13:763-72.
33. Krawinkel MB, Keding GB. Bitter gourd (Momordica charantia); A dietary approach to hyperglycemia. Nutr Rev 2006;64:331-7.
32. Chang CI, Chen CR, Liao YW, Cheng YW, Chen YC.
31. Okabe H, Miyahara Y, Yamauchi T. Cucurbitane-type triterpenoids from Momordica charantia. J Nat Prod 2006;69(8):1168-71.
30. Okabe H, Miyahara Y, Yamauchi T. Studies on the constituents of Momordica charantia L. I. Isolation and characterization of momordicosides A ang B, glycosides of a pentahydroxy cucurbitane triterpene. Chem Pharm Bull 1980;28:2753-62.
29. Miyahara Y, Okasbe H, Yamauchi T. Studies on the constituents of Momordica charantia L. II. Isolation and characterization of minor seed glycosides, momordicosides C, D and E. Chem Pharm Bull 1981;29:1561-6.
28. Okabe H, Miyahara Y, Yamauchi T. Structures of momordicosides F1, F2, G, I, K and L, novel cucurbitacin in the fruits of Momordica charantia L. Tetrahedron Lett 1982a; 23(1):77–80.
71. Lucas EA, Dumancas GG, Smith BJ, Clarke SL, Arjmandi BH. Health benefits of bitter melon (Momordica charantia). Bioact Foods Promot Health 2010;35:525-49.
72. Kenny O, Smyth TJ, Hewage CM, Brunton NP. Antioxidant properties and quantitative UPLC-MS analysis of phenolic compounds from extracts of fenugreek (Trigonella foenum-graecum) seeds and bitter melon (Momordica charantia) fruit. Food Chem 2013;141:4295-302.
73. Sarkar N, Mukherjee A, Barik A. Olfactory responses of Epilachna dodecastigma (Coleoptera: Coccinellidae) to long-chain fatty acids from Momordica charantia leaves. Arthropod Plant Interact 2013;7:339-48.
74. Ke LJ, Lu W, Chang JL, Yuan FY, Rao PF, Zhou JW. Effects of heat drying process on amino acid content of Momordica charantia L. Amino Acids Biol Resour 2010;32:14-6.
75. Han C, Zuo J, Wang Q, Xu L, Wang Z, Dong H, Gao L. Effects of 1-MCP on post harvest physiology and quality of bitter melon (Momordica charantia L.). Sci Hortic Amsterdam 2015;182:86-91.
76. Handa G, Singh J, Sharma ML, Kaul A, Zafar R. Hypoglycaemic principle of Momordica charantia seeds. Nat Prod 1990;6(1):16-9.
77. Chen Q, Chan LL, Li ET. Hypoglycemic effect of semi purified peptides from Momordica charantia L. Var. abbreviata Ser. in alloxan-induced diabetic mice. J Food Biochem 2008;32:107-21.
78. Mohammad I, Elattar S, Mohammed S, Ewais M. An evaluation of anti-diabetic and anti-lipidemic properties of Momordica charantia (bitter melon) fruit extract in experimentally induced diabetes. Life Sci J 2012;9(2):363-74.
79. Nerurkar PV, Lee YK, Motosue M, Adeli K, Nerurkar VP. Momordica charantia (bitter melon) inhibits primary human adipocyte differentiation by modulating adipogenic genes. BMC Complement Altern Med 2010;10:34.
80. Chaturvedi P, George S. Momordica charantia maintains normal glucose levels and lipid profiles and prevents oxidative stress in diabetic rats subjected to chronic sucrose load. J Med Food 2010;13(3):520-7.
81. Popovich DG, Li L, Zhang W. Bitter melon (Momordica charantia) triterpenoid extract reduces predipocyte viability, lipid accumulation and adiponectin expression in 3T3-L1 cells. Food Chem Toxicol 2010;48(6):1619-26.
82. Yu Y, Zhang XH, Ebersole B, Ribnicky D, Wang ZQ. Bitter melon extract attenuating hepatic steatosis may be mediated by FGF21 and AMPK/Sirt1 signaling in mice. Sci Rep 2015;5:1206-14.
metabolism are impaired in mice lacking the nuclear oxysterol receptor LXRα. Cell 1998;93(5):693-704.

104. Matsui S, Yamane T, Takita T, Oishi Y, Kobayashi-Hattori K. The hypocholesterolemic activity of Momordica charantia fruit is mediated by the altered cholesterol- and bile acid-regulating gene expression in rat liver. Nutr Res 2013;33(7):580-5.

105. Chikkavadaragudi RS, Vishwanath P, Prashant A, Rangaswamy C, Maduvanahalli NS, Hattur B. Fifty percent ethanolic extract of Momordica charantia inhibits adipogenesis and promotes adipolysis in 3T3-L1 pre-adipocyte cells. Rep Biochem Mol Biol 2017;6(1):23-32.

106. Hulin A. Intoxication aigue par Momordica charantia (sorossi). A propos de deux cas (acute intoxication due to Momordica charantia (sorossi). Study of two cases). Sem Hop 1988;64:2847-8.

107. Dans AM, Villarruz MV, Jimeno CA, Javelosa MA, Chua J, Bautista R. The effect of Momordica charantia capsule preparation on glycemic control in type 2 diabetes mellitus needs further studies. J Clin Epidemiol 2007;60:554-9.

108. Dixit VP, Khanna P, Bhargava SK. Effects of Momordica charantia L. fruit extract on the testicular function of dog. Planta Med 1978;34:280-6.

109. Saksena SK. Study of antifertility activity of the leaves of Momordica Linn (Karela). Indian J Physiol Pharmacol 1971;15:79-80.

110. Stepka W. Antifertility investigations on Momordica. Lloydia 1974;37:645.

111. Chan WY, Tam PP, Yeung HW. The termination of early pregnancy in the mouse by β-momorcharin. Contraception 1984;29:91-100.

112. Basch WE, Gabardi S, Ulbricht C. Bitter melon (M. charantia): a review of efficacy and safety. Am J Health Syst Pharm 2003;60:356-9.

113. Dutta PK, Chakravarty AK, Chowdhury US, Pakrashi SC. Vicine, a Favism-inducing toxin from Momordica charantia Linn. seeds. Indian J Chem 1981;20:669-71.

114. Deshaware S, Gupta S, Singhal RS, Joshi M, Varyar PS. Debittering of bitter gourd juice using β-cyclodextrin: Mechanism and effect on antidiabetic potential. Food Chem 2018;262:78-85.

115. Tan SP, Kha TC, Parks SE, Stathopoulos CE, Roach PD. Effects of the spray-drying temperatures on the physiochemical properties of an encapsulated bitter melon aqueous extract powder. Powder Technol 2015;281:65-75.