



Passiflora edulis: A Bioactive Bounty – A Comprehensive Review

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Abstract

Passiflora edulis, commonly known as passion fruit, transcends its delightful taste by harbouring many medicinal properties. Rich in antioxidants like vitamin C and polyphenols, passion fruit displays robust free-radical-scavenging effects, suggesting potential anti-inflammatory and cardiovascular benefits. Studies underscore its anti-cancer potential, necessitating further exploration. Passion fruit, especially its leaves, emerges as a potent source of antioxidants, as demonstrated through in vitro and in vivo assessments. Compounds like vitexin and isoorientin contribute to its antioxidative effects. Passion fruit exhibits analgesic and anti-inflammatory properties, with butanoic fractions and C-glucosyl flavones showing promise in alleviating pain and suppressing inflammation. The fruit peel demonstrates anti-inflammatory effects in inflammatory bowel disease models, reinforcing its potential therapeutic role. Studies unveil passion fruit's anti-hypertensive prowess, attributed to compounds like luteolin and γ-aminobutyric acid. Yellow passion fruit pulp and peel extract effectively reduce blood pressure, showcasing its vasodilatory effects. Passion fruit emerges as a hypolipidemic agent, countering hyperlipidemiarelated risks. Juice, peel flour, and seed extracts exhibit lipid-lowering effects, attributed to compounds like pectin and linoleic acid. In the realm of antidiabetic activities, passion fruit demonstrates glucose-lowering effects in diabetic models. Aqueous leaf extracts mitigate diabetes-related complications, emphasising their potential as a preventive measure. Passion fruit peel flour improves insulin sensitivity and protects against insulin resistance. Remarkably, passion fruit unveils antitumor potential, hindering tumour growth and inducing apoptosis in various cancer cells. Polysaccharide fractions and ethanolic extracts exhibit significant anticancer effects. Finally, passion fruit manifests antidepressant properties, with cyclobutane triterpenoids identified as potential bioactive components. Ethanol extracts reduce immobility time, suggesting a role in modulating dopaminergic and serotonergic pathways. In essence, passion fruit, with its diverse bioactive compounds, holds promise as a multifaceted therapeutic agent, warranting further exploration for its extensive medicinal applications.

Keywords: Antidepressant, Anti-hypertensive, Anti-inflammatory, Lipid-lowering Effects, Passiflora edulis

1. Introduction

The *Passiflora* genus, which includes perennial, evergreen climbing vines with woody stems, belongs to the Passifloraceae family and is found extensively in the tropical areas of America, Asia, and Africa^{1,2}. The *Passiflora* genus comprises more than 500 species, mostly grown for gardening and decorative uses, with around sixty of these species producing fruits that are edible^{3,4}. The leading species, *P. edulis*, commonly known as "passion fruit," has seen a surge in popularity for its economic, medicinal, and culinary value. Noted for its

unique taste and fragrant properties, *P. edulis* extends its usefulness beyond just being a tasty fruit. As such, it has become an important focus of research in horticulture, agriculture, and nutritional science. Among tropical fruits, passion fruit ranks third in importance, following pineapple and mango³. With its complex aroma that echoes the scents of over 130 different fruits, including apple, guava, banana, strawberry, mango, and pineapple, passion fruit has been dubbed the "king of fruit juice," as well as "*maracujá*," "love fruit," "fruit lover," and "spice fruit". This diverse fragrance profile has inspired a wide range of passion fruit-derived products, such as jelly, ice

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cream, yoghurt, cake, tea, wine, vinegar, jam, soup bases, condiment sauces, and more. Additionally, passion fruit is used in traditional medicine across various cultures and has found its way into the cosmetics industry in numerous countries as a moisturizing agent⁵. Significant strides have been taken in recent decades to conduct pharmacological research on P. edulis. Various extracts from passion fruit, as well as its fruit juices and isolated compounds, have demonstrated a spectrum of biological effects, including hepatoprotective⁶, neuroprotective⁷, anti-hypertensive⁸, antitumour⁹, antimicrobial¹⁰. sedative¹¹, antidepressant¹², anxiolytic-like properties¹³, lung-protective activities¹⁴, free radical scavenger activity¹⁵, anti-diabetic¹⁶, and anti-inflammatory properties¹⁷. The myriad health benefits of passion fruit are closely tied to its rich content of bioactive compounds, including polyphenols, triterpenes, flavonoids, and polysaccharides. This fruit is also packed with essential nutrients like minerals, dietary fibre, and vitamins, making it a prime candidate for classification as a "functional food." However, despite its nutrientdense profile and significant potential for promoting health, passion fruit remains underappreciated in several areas, highlighting a clear need for greater awareness and exploration. The primary goal of this concise review is to provide a comprehensive summary of recent progress in the understanding of the chemical and biological properties of various parts of P. edulis. This includes an investigation into the unique chemical compositions and biological effects of the fruit, stems, leaves, and peel of P. edulis, aiming to consolidate and present the existing knowledge on these aspects.

2. Medicinal Properties of Passiflora edulis

Passiflora edulis is celebrated not just for its delicious flavour but also for its medicinal qualities, thanks to its high antioxidant content, including vitamin C and polyphenols. These compounds help combat free radicals, aiding in health maintenance. The fruit is known for its anti-inflammatory effects, which could play a role in treating inflammation-related disorders. Moreover, it has been suggested that *P. edulis* can enhance heart health by lowering cholesterol and maintaining blood pressure levels. Although preliminary studies suggest its antioxidants might

offer anti-cancer benefits, more research is needed to confirm these findings.

2.1 Antioxidant Activity

Research has highlighted the importance of passion fruit as a potent source of natural antioxidants. These antioxidants are essential for moderating the body's oxidative balance, either by neutralizing free radicals or inhibiting their actions. Various parts of P. edulis, such as the seeds, fruit, peel, leaves, and bark, have been examined for their antioxidant capabilities and their ability to scavenge radicals, utilizing a range of in vitro methods (including ABTS, ORAC, DPPH, AAPH, FRAP, HOCl scavengers, and ferrous ion assays). Moreover, numerous in vivo studies have been undertaken to assess the efficacy of these extracts in real biological systems^{18,19}. In this study, both *in vitro* and *in* vivo methods were used to investigate the antioxidant properties of passion fruit leaves. Through HPLC-PDA and ESI-MS/MS analyses, researchers probed for bioactive compounds, with a focus on polyphenols, present in the water extract of P. edulis leaves. In animal tests, the leaf extract demonstrated antioxidant activity by decreasing TBARS levels in the liver, increasing GSH levels in the kidneys, and affecting the activities of enzymes like SOD, GR, and GPx. The extract was found to contain significant compounds such as vitexin, isovitexin, and isoorientin. Moreover, its consumption resulted in alterations in the gut microbiota and an increase in short-chain fatty acid production. The study concludes that P. edulis leaf extract may act as a valuable antioxidant source, offering a defence against oxidative stress²⁰. This review examined the antioxidant and anti-inflammatory properties of a water-based extract derived from P. edulis leaves, containing isoorientin, vitexin, and isovitexin. The study found that oral administration of P. edulis extracts at a concentration of 1100 µg/mL significantly increased intrinsic antioxidant levels and reduced lipid peroxidation in the bloodstream, liver, and colon in a colitis model induced by 2,4,6-trinitrobenzene sulphonic acid. Additionally, the extract demonstrated anti-inflammatory effects in the colon, notably reducing levels of cytokines such as TNF- α and IL-1 β . These results suggest that the *P. edulis* leaf extract has therapeutic potential for managing oxidative stress and inflammation, observed in both healthy rats and a colitis-induced model²¹.

2.2 Analgesic and Anti-inflammatory Activity

Comparative studies in thermal stimulation pain models revealed that the analgesic effectiveness of n-butanol extracts obtained from P. edulis leaves is dependent on the dosage administered²². Moreover, the analgesic effects were exhibited by the polysaccharide extracted from the dried fruit of P. edulis in models involving acetic acid-induced writhing and formalininduced paw licking. Interestingly, results from the hot plate test did not show significant increases in response times, suggesting that the analgesic mechanism of the polysaccharide may be associated with peripheral pathways¹⁷. The study delved into the anti-inflammatory properties of individual compounds and sub-fractions isolated from the butanol fraction of P. edulis var. flavicarpa leaves. The butanol fraction demonstrated significant anti-inflammatory effects, particularly in its ability to impede the infiltration of leukocytes and neutrophils. Notably, sub-fraction C exhibited remarkable efficacy among the sub-fractions tested. Compounds isolated from sub-fraction C, including Isoorientin, Vicenin-2, and Spinosin, displayed antiinflammatory activity by suppressing both neutrophils and leukocytes. Intriguingly, both isoorientin and the butanol fraction also showed a decrease in myeloperoxidase activity. These findings suggest that C-glucosylflavones extracted from P. edulis leaves may play a role in the observed anti-inflammatory effects in a mouse model of pleurisy²³. This study examined the impact of P. edulis peel, known for its high dietary fibre and polyphenol content, on inflammatory bowel disease. In an animal model, the consumption of P. edulis peels demonstrated anti-inflammatory effects in the intestines, effectively reducing colonic damage induced by DSS. This was evidenced by a decrease in disease activity index values and improvements in histological assessments. Biochemical and molecular analyses indicated a strengthened intestinal protective barrier and a decrease in the expression of proinflammatory cytokines. Additionally, there was an increase in the production of short-chain fatty acids, which could have prebiotic effects. Mice treated with the polysaccharide fraction obtained from the dried fruit of P. edulis at a dose of 3 mg/kg exhibited a reduction in paw oedema caused by various inflammatory agents

such as 48/80, carrageenan, histamine, 5-HT, and prostaglandin E2. Furthermore, it significantly reduced vascular permeability and lowered the levels of TNF- α and IL-1 β^{17} .

2.3 Anti-hypertensive Activity

The study explored the potential antihypertensive properties of luteolin at a dose of 50 mg/kg and the methanol extract obtained from the rind of P. edulis at concentrations of 10 mg/kg and 50 mg/kg. When administered orally to Spontaneous Hypertensive Rats (SHRs), both the extract and luteolin exhibited a significant decrease in systolic blood pressure. Quantitative analysis using liquid chromatographytandem mass spectrometry (LC-MS/MS) revealed that the extract contained luteolin (20 µg/g dry weight) and luteolin-6-C-glucoside (41 µg/g dry weight). Additionally, y-aminobutyric acid (GABA), recognized for its blood pressure-lowering effects, was detected at levels of 2.4 mg/g dry weight (determined by LC-MS/MS) or 4.4 mg/g dry weight (based on amino acid analysis). The findings suggest that the main antihypertensive impact of the extract in SHRs is associated with its substantial GABA content, complemented by the vasodilatory properties of luteolin and other polyphenols²⁵. In this investigation, five consecutive days were spent giving yellow passion fruit pulp orally via gavage at varying doses (5, 6, or 8 g/kg b.w.). Tandem mass spectrometry and highperformance liquid chromatography with photodiode array were used to establish the presence of phenolic chemicals, ascorbic acid, carotenoids, and flavonoids in vellow passion fruit pulp. The highest dose of passion fruit pulp showed a noteworthy drop in Thiobarbituric Acid Reactive Substances (TBARS), an increase in glutathione (GSH), and a considerable reduction in systolic blood pressure. Remarkably, there were no alterations observed in renal function parameters or the frequency of micronuclei in bone marrow cells. The research concludes that the antihypertensive effects of yellow passion fruit pulp can be attributed, at least in part, to its elevated antioxidant content. However, for a comprehensive understanding of the mechanisms underlying this observed effect, further investigation is deemed necessary⁸. The study demonstrated the antihypertensive efficacy of compounds derived from

1206

both yellow and purple passion fruit in spontaneously hypertensive rats. Administration of *P. edulis* peel extract orally to these rats resulted in reductions in blood pressure, serum nitric oxide levels, and hemodynamic parameters according to studies^{26,27}.

2.4 Hypolipidemic Activity

The study underscores the considerable health risks associated hyperlipidemia, with encompassing conditions like atherosclerosis, inflammation of the pancreas and coronary artery disease. Passion fruit emerges as a promising preventive measure against hyperlipidemia. The offspring of diabetic Wistar rats exhibited a significant reduction in Total Cholesterol (TC), Triglycerides (TG), and Low-Density Lipoprotein Cholesterol (LDL-C) levels after receiving 580 mg/ kg of passion fruit juice daily for 30 days. There was also a discernible rise in High-Density Lipoprotein Cholesterol (HDL-C) levels at the same time²⁸. Additionally, in the context of diet-induced obesity in rats, peel flour derived from P. edulis exhibited positive effects by reversing cumulative body weight gain, reducing adiponectin levels and adiposity, and increasing leptin levels²⁹. The research validated the effectiveness of orally administering pectin derived from the peel of P. edulis fruit at doses between 0.5 and 25 mg/kg for 5 days, leading to a significant reduction in triglyceride levels in diabetic rats³⁰. Moreover, the nonsoluble fibre extracted from P. edulis seeds demonstrated a cholesterol-lowering effect by increasing the levels of bile acids, total lipids, and cholesterol in the faeces of golden Syrian hamsters.

Concurrently, it led to a reduction in serum TG, TC, and liver cholesterol levels³¹. In this research, the hypolipidemic potential of *P. edulis* variety seed oil (PE) from west Cameroon was explored in both females and males. Rats that were administered PE showed a notable reduction in TG, TC, and LDL-C levels. In contrast to untreated male rats, the groups receiving PE demonstrated a significant elevation in HDL-C levels. A similar pattern was observed in female rats concerning TG, with the lowest values observed with 1 ml of olive oil. The study suggests that the elevated linoleic acid content in PE may contribute to its hypolipidemic effects, resembling those of olive oil³². The group administered *P. edulis* juice showed reduced levels of free fatty acids and low-density lipoprotein-cholesterol,

along with increased levels of high-density lipoproteincholesterol compared to the control group. There were no statistically significant differences observed in SOD activity, GSH concentration, TG levels, and LDL levels between the two groups. However, the concentration of thiobarbituric acid-reactive chemicals significantly decreased in the treated group, indicating a reduction in lipid peroxidation. These results suggest that *P. edulis* juice contributed to a decrease in lipid peroxidation and had positive effects on the lipid profile in Wistar rats under the specified experimental conditions³³.

2.5 Antidiabetic Activity

Several studies suggest that P. edulis peel flour, juice, and seeds may exert anti-diabetic properties by dropping glucose tolerance in diabetic rats and mice. In a specific study, streptozotocin-induced diabetic rat pups exhibited a significant decrease in glucose levels when orally administered passion fruit juice (580 mg/ kg) once daily for thirty days²⁸. The study persuaded diabetes in Wistar albino rats using alloxan and explored the potential anti-diabetic properties of the water leaf extract of P. edulis Sims. After the aqueous extract was given continuously for 30 days at a dose of 200 mg/kg, lipid parameters, glycosylated haemoglobin, and blood glucose levels all significantly decreased. Additionally, the extract markedly elevated haemoglobin, HDL, and plasma insulin levels. P. edulis also demonstrated a significant increase in glycogen and liver protein levels, accompanied by the normalization of enzymes involved in carbohydrate metabolism. Based on these findings, the study concluded that P. edulis significantly mitigates diabetes risk in rats induced with alloxan³⁴. Furthermore, db/db mice and rats induced with diabetes via streptozotocin exhibited reduced blood glucose levels following the administration of extracts derived from passion fruit seeds or leaves^{35,36}. Administering diabetic rats, induced with alloxan, a daily dosage of pectin extracted from P. edulis fruit peel (ranging from 0.5 to 25 mg/kg) for five consecutive days led to a notable decrease in blood glucose levels. This indicates a potential novel treatment strategy for type 2 diabetes³⁰. In a study conducted by Lima et al., rats fed a high-fat diet to induce obesity exhibited improved insulin sensitivity when supplemented with P. edulis peel flour. This improvement was evidenced by elevated levels of glucagon-like peptide-1 (GLP-1) and Glucose-Dependent Insulinotropic Polypeptide (GIP). Moreover, *P. edulis* peel flour was found to shield rats from insulin resistance induced by a low-fructose diet. Furthermore, the leaf extract of *P. edulis* enriched with flavonoids demonstrated health benefits in diabetic conditions by delaying the onset of associated complications^{37,38}.

2.6 Antitumor Activity

The study focused on a polysaccharide fraction (PFCM) extracted from P. edulis using hot aqueous extraction. When administered to Sarcoma 180 tumours, PFCM demonstrated a growth inhibition ratio ranging from 40.59% to 48.73%, depending on the dosage and method of administration (either orally or intraperitoneally). Toxicological evaluations revealed no significant changes in biochemical markers or organs such as the kidneys and liver, but rather an increase in cell types associated with essential defense mechanisms. Intriguingly, compared to treatment with 5-fluorouracil, PFCM treatment increased spleen weight⁹. The study aimed to investigate the potential inhibitory effects of the ethanolic extract from P. edulis leaves on colorectal adenocarcinoma cells, specifically Caco-2 and SW480. The extract exhibited a significant impact on cell viability and antiproliferative activity, assessed through clonogenic and 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide tests. Flow cytometry analysis revealed that the extract could control cell cycle progression and trigger apoptosis. After 48 hours of treatment, a notable inhibition of cellular proliferation occurred in both cell lines, with determined half maximum inhibitory concentration (IC50) values for the extract. Moreover, the extract increased the proportion of necrotic and apoptotic cells, upsetting the cell cycle phase distribution by elevating SubG1 and G2/M populations and initiating pathways that result in cell death. All of these findings suggest that P. edulis could be a valuable source of phytochemical substances with notable anticancer effects, especially when it comes to the treatment of colorectal cancer³⁹. The administration of yellow passion fruit ethanol extract in vivo showed a significant 48.5% inhibition of tumour development. Moreover, the extract increased the lifetime of male Balb/c mice carrying Ehrlich cancer cells by nearly 42 per cent. With particular emphasis on lauric acid,

the presence of medium and long-chain fatty acids is probably responsible for this advantageous result⁴⁰.

2.7 Antidepressant Activity

The antidepressant capabilities of extracts from P. edulis stems and leaves have been verified through in vivo studies. Over seven days, mice given oral ethanol extracts from P. edulis aerial parts at doses equal to 10 and 2 g/kg of the plant material showed signs of an antidepressant-like effect by showing a reduction in immobility time in forced swim and tail suspension tests. Furthermore, at a dosage of 50 mg/kg, cycloartane triterpenoids, namely cyclopassiflosides IX and XI, showed an antidepressant-like effect, indicating that these compounds may be the main bioactive ingredients accountable for P. edulis's antidepressant properties⁴¹. Oral dosages of *P. edulis* Sims extracts (300 mg/kg), ethyl acetate (50 mg/kg), and butanol (50 mg/ kg) exhibited a reduction in the length of immobility time in mice in the forced swimming test; these effects were comparable to those observed with fluoxetine and nortriptyline. Notably, the ethyl acetate and butanol extracts enriched with flavonoids exhibited a preference for displaying antidepressant qualities. The antagonistic actions of sulpiride, a-methyl-DL-tyrosine chloride, and p-chlorophenylalanine indicated a specific relationship with the modulation of dopaminergic and serotonergic transmission through D2 receptor, 5-HT, and catecholamine pathways⁴².

3. Conclusion

In conclusion, the multifaceted properties of *P. edulis*, commonly known as passion fruit, present a captivating narrative of its potential contributions to human health. Beyond its delightful taste, passion fruit emerges as a powerhouse of medicinal benefits. Rich in antioxidants, such as vitamin C and polyphenols, it demonstrates formidable free-radical-scavenging effects, contributing to overall well-being. The anti-inflammatory attributes of passion fruit suggest promising prospects in managing inflammatory conditions, while its recognized natural sedative and anxiolytic effects offer a calming influence, potentially aiding in anxiety and insomnia. Moreover, *P. edulis* may play a role in cardiovascular health by reducing cholesterol levels and supporting optimal blood

pressure. The fruit's fibre content not only promotes digestive health but may also hint at anti-cancer potential, although further research is warranted for conclusive evidence. Laden with essential vitamins and minerals, passion fruit proves to be not just a culinary delight but a nutritional powerhouse contributing to holistic well-being. The extensive research on its antioxidant, anti-inflammatory, analgesic, antihypertensive, hypolipidemic, antidiabetic, antitumor, and antidepressant activities underscores its potential as a versatile and valuable natural resource in the realm of preventive and therapeutic health interventions.

4. References

- Zhang J, Koike R, Yamamoto A, Ukiya M, Fukatsu M, Banno N, *et al.* Glycosidic inhibitors of melanogenesis from leaves of *P. edulis*. Chem Biodivers. 2013; 10(10):1851-65. https://doi.org/10.1002/cbdv.201300181 PMid:24130028
- Hu Y, Jiao L, Jiang MH, Yin S, Dong P, Zhao ZM, *et al.* A new C-glycosyl flavone and a new neolignan glycoside from *P. edulis* Sims peel. Nat Prod Res. 2018; 32(19):2312-8. https:// doi.org/10.1080/14786419.2017.1410809 PMid:29199463
- Anderson JD, Vidal RF, Brym M, Stafne ET, Resende Jr MF, Viana AP, Chambers AH. Genotyping-by-sequencing of passion fruit (*Passiflora* spp.) generates genomic resources for breeding and systematics. Genet Resour Crop Evol. 2022; 69(8):2769-86. https://doi.org/10.1007/s10722-022-01397-4
- Dhawan K, Dhawan S, Sharma A. Passiflora: A review update. J Ethnopharmacol. 2004; 94(1):1-23. https://doi. org/10.1016/j.jep.2004.02.023 PMid:15261959
- Xu FQ, Wang N, Fan WW, Zi CT, Zhao HS, Hu JM, *et al.* Protective effects of cyclobutane triterpenoids from *P. edulis* Sims against glutamate-induced neurotoxicity in PC12 cell. Fitoterapia. 2016; 115:122-7. https://doi.org/10.1016/j. fitote.2016.09.013 PMid:27693740
- Zhang YJ, Zhou T, Wang F, Zhou Y, Li Y, Zhang JJ, et al. The effects of Syzygium samarangense, P. edulis and Solanum muricatum on alcohol-induced liver injury. Int J Mol Sci. 2016; 17(10):1616. https://doi.org/10.3390/ijms17101616 PMid:27681723 PMCid: PMC5085649
- Tal Y, Anavi S, Reisman M, Samach A, Tirosh O, Troen AM. The neuroprotective properties of a novel variety of passion fruit. J Funct Foods. 2016; 23:359-69. https://doi. org/10.1016/j.jff.2016.02.039
- 8. Konta EM, Almeida MR, Amaral CL, Darin JD, de Rosso VV, Mercadante AZ, *et al.* Evaluation of the antihypertensive properties of yellow passion fruit pulp (*P. edulis* Sims f. *flavicarpa* Deg.) in

spontaneously hypertensive rats. Phytother Res. 2014; 28(1):28-32. https://doi.org/10.1002/ptr.4949 PMid:23436457

- Silva DC, Freitas AL, Barros FC, Lins KO, Alves AP, Alencar NM, *et al.* Polysaccharide isolated from *P. edulis*: Characterization and antitumor properties. Carbohydr Polym. 2012; 87(1):139-45. https://doi.org/10.1016/j. carbpol.2011.07.029 PMid:34662942
- Dzotam JK, Touani FK, Kuete V. Antibacterial and antibiotic-modifying activities of three food plants (*Xanthosoma mafaffa* Lam., *Moringa oleifera* (L.) Schott and *P. edulis* Sims) against Multidrug-Resistant (MDR) Gram-negative bacteria. BMC Complement Altern Med. 2015; 16:1-8. https://doi.org/10.1186/s12906-016-0990-7 PMid:26753836 PMCid: PMC4709887
- Klein N, Gazola AC, de Lima TC, Schenkel E, Nieber K, Butterweck V. Assessment of sedative effects of *P. edulis* f. *flavicarpa* and *Passiflora alata* extracts in mice, measured by telemetry. Phytother Res. 2014; 28(5):706-13. https://doi. org/10.1002/ptr.5043 PMid:23893399
- Alves JS, Marques JI, Demarque DP, Costa LR, Amaral JG, Lopes NP, *et al.* Involvement of isoorientin in the antidepressant bioactivity of a flavonoid-rich extract from *P. edulis* f. *flavicarpa* leaves. Rev Bras Farmacogn. 2020; 30(2):240-50. https://doi.org/10.1007/s43450-020-00003-x
- Li H, Zhou P, Yang Q, Shen Y, Deng J, Li L, et al. Comparative studies on anxiolytic activities and flavonoid compositions of *P. edulis 'edulis*' and *P. edulis 'flavicarpa*'. J Ethnopharmacol. 2011; 133(3):1085-90. https://doi. org/10.1016/j.jep.2010.11.039 PMid:21111038
- Chilakapati SR, Serasanambati M, Manikonda PK, Chilakapati DR, Watson RR. Passion fruit peel extract attenuates bleomycin-induced pulmonary fibrosis in mice. Can J Physiol Pharmacol. 2014; 92(8):631-9. https://doi. org/10.1139/cjpp-2014-0006 PMid:24933624
- Panelli MF, Pierine DT, De Souza SL, Ferron AJ, Garcia JL, Santos KC, *et al.* Bark of *P. edulis* treatment stimulates antioxidant capacity and reduces dyslipidemia and body fat in db/db mice. Antioxidants. 2018; 7(9):120. https:// doi.org/10.3390/antiox7090120 PMid:30205562 PMCid: PMC6162700
- Salles BC, da Silva MA, Taniguthi L, Ferreira JN, da Rocha CQ, Vilegas W, *et al. P. edulis* leaf extract: evidence of antidiabetic and antiplatelet effects in rats. Biol Pharm Bull. 2020; 43(1):169-74. https://doi.org/10.1248/bpb.b18-00952 PMid:31645529
- 17. Silva RO, Damasceno SR, Brito TV, Dias JM, Fontenele AM, Braúna IS, *et al.* Polysaccharide fraction isolated from *P. edulis* inhibits the inflammatory response and the oxidative stress in mice. J Pharm Pharmacol. 2015; 67(7):1017-27. https://doi.org/10.1111/jphp.12399 PMid:25808583
- 18. Phamiwon ZA, John S. Phytochemical investigation and antioxidant activities of *P. edulis* (passion fruit) leaves from

the Ukhrul district, Manipur, India. World J Pharm Res. 2017; 6(14):793-801.

- Rotta EM, Rodrigues CA, Jardim IC, Maldaner L, Visentainer JV. Determination of phenolic compounds and antioxidant activity in passion fruit pulp (*Passiflora spp.*) using a modified QuEChERS method and UHPLC-MS/MS. Lwt. 2019; 100:397-403. https://doi.org/10.1016/j. lwt.2018.10.052
- 20. da Silva JK, Cazarin CB, Colomeu TC, Batista ÂG, Meletti LM, Paschoal JA, *et al.* Antioxidant activity of aqueous extract of passion fruit (*P. edulis*) leaves: *in vitro* and *in vivo* study. Food Res Int. 2013; 53(2):882-90. https://doi. org/10.1016/j.foodres.2012.12.043
- 21. Cazarin CB, da Silva JK, Colomeu TC, Batista ÂG, Meletti LM, Paschoal JA, *et al.* Intake of *P. edulis* leaf extract improves antioxidant and anti-inflammatory status in rats with 2, 4, and 6-trinitrobenzene sulphonic acid-induced colitis. J Funct Foods. 2015; 17:575-86. https://doi.org/10.1016/j. jff.2015.05.034
- 22. Nayak L, Panda SK. Phytochemical investigation and evaluation of analgesic activity of *P. edulis* Linn leaves available in Southeastern Odisha. Int J Pharm Bio Arch. 2012; 3:897-9.
- 23. Zucolotto SM, Goulart S, Montanher AB, Reginatto FH, Schenkel EP, Fröde TS. Bioassay-guided isolation of anti-inflammatory C-glucosylflavones from *P. edulis.* Planta Medica. 2009; 75(11):1221-6. https://doi. org/10.1055/s-0029-1185536 PMid:19353489
- 24. Cazarin CB, Rodriguez-Nogales A, Algieri F, Utrilla MP, Rodríguez-Cabezas ME, Garrido-Mesa J, *et al.* Intestinal anti-inflammatory effects of *P. edulis* peel in the dextran sodium sulphate model of mouse colitis. J Funct Foods. 2016; 26:565-76. https://doi.org/10.1016/j.jff.2016.08.020
- 25. Ichimura T, Yamanaka A, Ichiba T, Toyokawa T, Kamada Y, Tamamura T, *et al.* The antihypertensive effect of an extract of *P. edulis* rind in spontaneously hypertensive rats. Biosci Biotechnol Biochem. 2006; 70(3):718-21. https://doi.org/10.1271/bbb.70.718 PMid:16556991
- 26. Zibadi S, Farid R, Moriguchi S, Lu Y, Foo LY, Tehrani PM, *et al.* Oral administration of purple passion fruit peel extract attenuates blood pressure in female spontaneously hypertensive rats and humans. Nutr Res. 2007; 27(7):408-16. https://doi.org/10.1016/j.nutres.2007.05.004
- Lewis BJ, Herrlinger KA, Craig TA, Mehring-Franklin CE, DeFreitas Z, Hinojosa-Laborde C. Antihypertensive effect of passion fruit peel extract and its major bioactive components following acute supplementation in spontaneously hypertensive rats. J Nutr Biochem. 2013; 24(7):1359-66. https://doi.org/10.1016/j.jnutbio.2012.11.003 PMid:23333089
- 28. Barbalho SM, Damasceno DC, Spada AP, Lima IE, Araújo AC, Guiguer EL, *et al.* Effects of *P. edulis* on the metabolic

 profile of diabetic Wistar rat offspring. J Med Food. 2011;

 14(12):1490-5.
 https://doi.org/10.1089/jmf.2010.0318

 PMid:21663518

- 29. Lima GC, Vuolo MM, Batista AG, Dragano NR, Solon C, Junior MR. *P. edulis* peel intake improves insulin sensitivity, increasing incretins and hypothalamic satietogenic neuropeptide in rats on a high-fat diet. Nutr. 2016; 32(7-8):863-70. https://doi.org/10.1016/j.nut.2016.01.014 PMid:27138107
- Silva DC, Freitas AL, Pessoa CD, Paula RC, Mesquita JX, Leal LK, *et al.* Pectin from *P. edulis* shows anti-inflammatory action as well as hypoglycemic and hypotriglyceridemic properties in diabetic rats. J Med Food. 2011; 14(10):1118-26. https://doi.org/10.1089/jmf.2010.0220 PMid:21554121
- Chau CF, Huang YL. Effects of the insoluble fibre derived from *P. edulis* seed on plasma and hepatic lipids and faecal output. Mol Nutr Food Res. 2005; 49(8):786-90. https://doi. org/10.1002/mnfr.200500060 PMid:15995986
- 32. Ngakou Takam P, Tonfack Djikeng F, Kuate D, Nouemsi Kengne AP, Doungué Tsafack H, Makamwé I, *et al. P. edulis* seed oil from west Cameroon: Chemical characterization and assessment of its hypolipidemic effect in high-fat diet–induced rats. Food Sci Nutr. 2019; 7(11):3751-8. https://doi.org/10.1002/fsn3.1234 PMid:31763024 PMCid: PMC6848813
- 33. de Souza MD, Barbalho SM, Damasceno DC, Rudge MV, de Campos KE, Madi AC, et al. Effects of P. edulis (yellow passion) on serum lipids and oxidative stress status of Wistar rats. J Med Food. 2012; 15(1):78-82. https://doi. org/10.1089/jmf.2011.0056 PMid:21877954
- 34. Kanakasabapathi D, Gopalakrishnan VK. Evaluation of the antidiabetic potential of aqueous extract of *P. edulis* Sims on alloxan-induced diabetes mellitus in Wistar albino rats. Int J Pharm Sci Rev Res. 2015; 34(1):171-7.
- Panchanathan S, Rajendran J. Evidence of the antihyperglycemic and antioxidant effect of *P. edulis flavicarpa* (Sims.) in streptozotocin-induced diabetic rats. Not Sci Biol. 2015; 7(4):383-9. https://doi.org/10.15835/nsb749655
- Uchida-Maruki H, Inagaki H, Ito R, Kurita I, Sai M, Ito T. Piceatannol lowers the blood glucose level in diabetic mice. Biol Pharm Bull. 2015; 38(4):629-33. https://doi. org/10.1248/bpb.b15-00009 PMid:25832644
- Salles BC, da Silva MA, Taniguthi L, Ferreira JN, da Rocha CQ, Vilegas W, *et al. P. edulis* leaf extract: evidence of antidiabetic and antiplatelet effects in rats. Biol Pharm Bull. 2020; 43(1):169-74. https://doi.org/10.1248/bpb.b18-00952 PMid:31645529
- 38. Soares RD, Campos MG, Ribeiro GP, Salles BC, Cardoso NS, Ribeiro JR, *et al.* Development of a chitosan hydrogel containing flavonoids extracted from *P. edulis* leaves and the evaluation of its antioxidant and wound healing properties for the treatment of skin lesions in diabetic mice. J Biomed

1210 Passiflora edulis: A Bioactive Bounty – A Comprehensive Review

Mater Res A. 2020; 108(3):654-62. https://doi.org/10.1002/ jbm.a.36845 PMid:31747098

- 39. Ramirez V, Arango Varela SS, Maldonado Celis ME, Uribe Yunda DF, Aguillón Osma J, Quintero JP, et al. Biological activity of *P. edulis* f. *flavicarpa* ethanolic leaves extract on human colonic adenocarcinoma cells. J Appl Pharm Sci. 2019; 9(2):64–71. https://doi.org/10.7324/ JAPS.2019.90209
- 40. Mota NS, Kviecinski MR, Zeferino RC, de Oliveira DA, Bretanha LC, Ferreira SR, *et al. In vivo* antitumor activity of by-products of *P. edulis* f. *flavicarpa* Deg. Rich in medium and long-chain fatty acids evaluated through oxidative stress markers, cell cycle arrest and apoptosis induction. Food

Chem Toxicol. 2018; 118:557-65. https://doi.org/10.1016/j. fct.2018.06.010 PMid:29886231

- Wang C, Xu FQ, Shang JH, Xiao H, Fan WW, Dong FW, et al. Cycloartane triterpenoid saponins from watersoluble *P. edulis* Sims and their antidepressant-like effects. J Ethnopharmacol. 2013; 148(3):812-7. https://doi. org/10.1016/j.jep.2013.05.010 PMid:23702036
- 42. Ayres AS, Santos WB, Junqueira-Ayres DD, Costa GM, Ramos FA, Castellanos L, *et al.* Monoaminergic neurotransmission mediates the antidepressant-like effects of *P. edulis* Sims fo. *edulis*. Neurosci Lett. 2017; 660:79-85. https://doi.org/10.1016/j.neulet.2017.09.010 PMid:28893593