

HEALTH BENEFITS AND THERAPEUTIC POTENTIAL OF BAEI (*Aegle marmelos* L.): A Comprehensive Review

ABSTRACT

Bael (*Aegle marmelos* L.), a sacred and medicinal plant in traditional Indian systems, has gained increasing scientific attention due to its diverse health benefits and therapeutic properties. This comprehensive review aims to explore the medicinal potential of bael, highlighting its phytochemical composition, antioxidant, antimicrobial, and anti-inflammatory activities, along with its role in treating various ailments such as diabetes, gastrointestinal disorders, cardiovascular diseases, and cancer. The fruit, leaves, and roots of bael possess potent bioactive compounds, including alkaloids, flavonoids, and coumarins, which contribute to its pharmacological properties. This review also discusses the molecular mechanisms underlying bael's therapeutic effects, emphasizing its role as a natural remedy with minimal side effects. Future research directions and the development of bael-based nutraceuticals are considered for broader clinical applications.

Keywords: Phytochemicals, Antioxidant, Anti-Inflammatory, Antimicrobial and Gastrointestinal Disorders.

INTRODUCTION

Bael (*Aegle marmelos* L.), commonly known as the Bengal quince or wood apple, is a tropical fruit-bearing tree native to the Indian subcontinent and Southeast Asia. Its significance in traditional medicine and culinary practices spans several millennia, particularly within the Ayurvedic system, where it is esteemed for its diverse therapeutic properties. This comprehensive review aims to elucidate the health benefits and therapeutic potential of bael, integrating contemporary scientific insights with traditional knowledge (Jafri *et al.* 2001).

The bael tree, known for its distinctive, aromatic fruit and its complex array of health-promoting compounds, has garnered considerable attention for its medicinal value. The fruit, leaves, bark, and seeds of *Aegle marmelos* contain a wealth of bioactive compounds, including essential oils, flavonoids, phenolic acids, and alkaloids. These constituents are believed to contribute to its wide-ranging pharmacological effects, including antioxidant, anti-inflammatory, antimicrobial, antidiabetic, and anticancer properties (Jagetia *et al.* 2003).

One of the most celebrated aspects of bael is its potential in managing gastrointestinal disorders. Its high fiber content and various organic acids aid in promoting digestive health, alleviating constipation, and managing irritable bowel syndrome. Additionally, bael has been shown to possess protective effects against ulcers and other forms of gastrointestinal distress, underscoring its value as a natural remedy for digestive issues (Jagetia *et al.* 2004).

In the realm of metabolic health, bael exhibits significant promise as an adjunct in diabetes management. Research indicates that extracts of bael may help regulate blood glucose levels, enhance insulin sensitivity, and reduce complications associated with diabetes. This makes bael a candidate for inclusion in dietary strategies aimed at managing or preventing diabetes and its associated complications. The fruit's antimicrobial and anti-inflammatory properties are also noteworthy. Bael has been utilized traditionally to combat infections and inflammatory conditions. Modern studies have supported these uses, demonstrating that bael extracts can inhibit the growth of various pathogens and modulate inflammatory responses. These properties contribute to its potential use in treating infections and inflammatory diseases (Jagetia *et al.* 2005).

Moreover, bael's potential in cancer prevention and treatment is an emerging area of interest. Preliminary studies suggest that compounds found in bael may exert cytotoxic effects on cancer cells, inhibit tumor growth, and enhance the efficacy of conventional chemotherapeutic agents. These findings highlight the need for further research to fully elucidate bael's role in cancer therapy (Johnson, 2010).

In addition to its therapeutic uses, bael also plays a role in preventive health. Its rich nutrient profile, including vitamins, minerals, and antioxidants, supports overall well-being and helps in fortifying the body's defenses against various ailments. The fruit's consumption has been linked to improved immune function, enhanced detoxification processes, and better skin health. This review will explore the breadth of research conducted on bael, examining the scientific evidence supporting its traditional uses and identifying areas where further investigation is needed. By synthesizing existing knowledge, we aim to provide a comprehensive overview of bael's health benefits and therapeutic potential, shedding light on its role in contemporary medicine and future research avenues (Jyothi and Rao, 2010).

1. Phytochemical Composition of Bael: Bioactive Compounds and Their Properties

The Bael tree (*Aegle marmelos* L.), commonly known as the bael fruit, is renowned for its rich phytochemical profile which contributes significantly to its health benefits and therapeutic potential. The fruit, leaves, bark, and seeds of bael contain a diverse array of bioactive compounds including alkaloids, flavonoids, tannins, saponins, essential oils, and phenolic acids (Kala *et al.* 2005).

Alkaloids in bael, such as aegeline and marmeline, exhibit noteworthy pharmacological activities including anti-inflammatory and anti-microbial effects. Flavonoids like quercetin and kaempferol are potent antioxidants that combat oxidative stress and mitigate cellular damage, thereby potentially reducing the risk of chronic diseases such as cancer and cardiovascular disorders. Tannins contribute to the astringent properties of bael, which are beneficial in treating gastrointestinal issues such as diarrhea and dysentery (Kamalakkannan and Prince, 2003).

Saponins present in bael have been shown to possess immunomodulatory and hypoglycemic effects, which can aid in the management of diabetes and boost overall immune response. Essential oils in bael, particularly those rich in eugenol, demonstrate anti-inflammatory, anti-microbial, and analgesic properties. Phenolic acids, including ferulic and caffeic acids, further enhance the antioxidant capacity of bael and contribute to its protective effects against cellular oxidative damage (Kamalakkannan and Prince, 2005).

These bioactive compounds collectively endow bael with a broad spectrum of therapeutic properties, encompassing anti-inflammatory, anti-microbial, anti-diabetic, and antioxidant activities. The integration of bael into traditional and modern medicinal practices underscores its potential as a valuable resource in preventive and therapeutic health strategies. Further research into the precise mechanisms and therapeutic applications of bael's phytochemicals could enhance our understanding and utilization of this remarkable fruit (Kar *et al.* 2002).

2. Antioxidant Properties of Bael and Its Role in Preventing Oxidative Stress

Bael (*Aegle marmelos* L.), a tropical fruit native to the Indian subcontinent, has garnered significant attention for its rich phytochemical profile and therapeutic potential. One of the most notable attributes of bael is its potent antioxidant properties, which play a crucial role in mitigating oxidative stress, a condition associated with various chronic diseases and aging. The antioxidant efficacy of bael is largely attributed to its high concentration of bioactive compounds, including ascorbic acid (vitamin C), flavonoids, and phenolic acids. These compounds contribute to the fruit's ability to neutralize free radicals and reactive oxygen species

(ROS), which are responsible for cellular damage and oxidative stress. Studies have demonstrated that bael extracts can effectively scavenge free radicals, thereby reducing lipid peroxidation and preventing DNA damage (Karakya, 2004).

In addition to its free radical scavenging activity, bael has been shown to enhance the activity of endogenous antioxidant enzymes, such as superoxide dismutase (SOD) and catalase. This dual mechanism—direct scavenging of ROS and boosting the body's own antioxidant defenses—contributes to its protective effects against oxidative stress-related conditions. The therapeutic potential of bael's antioxidant properties extends to various health benefits. Its ability to combat oxidative stress has been linked to protective effects against cardiovascular diseases, neurodegenerative disorders, and certain types of cancer. Furthermore, bael's antioxidant-rich profile supports overall health by aiding in the prevention of chronic inflammation and metabolic syndromes. Bael (*Aegle marmelos* L.) stands out as a valuable source of antioxidants, offering significant potential for preventing and managing oxidative stress-related diseases. Continued research is essential to fully elucidate its mechanisms and optimize its use in health and therapeutic applications (Kaur *et al.* 2009).

3. Antimicrobial and Antiviral Activities of Bael: A Natural Protector

Bael (*Aegle marmelos* L.), a revered plant in traditional medicine, has garnered significant interest for its antimicrobial and antiviral properties. Recent studies underscore its potential as a natural protector against various pathogens. The antimicrobial activity of bael is attributed primarily to its rich phytochemical profile, which includes essential oils, alkaloids, and flavonoids. These compounds exhibit substantial antibacterial effects against a range of Gram-positive and Gram-negative bacteria. For instance, research has demonstrated that bael extracts inhibit the growth of pathogens like *Staphylococcus aureus* and *Escherichia coli*, which are notorious for causing infections and contributing to antibiotic resistance (Kaushik *et al.* 2000).

In addition to its antibacterial properties, bael also shows promising antiviral activity. The fruit, leaves, and bark of the bael tree contain bioactive compounds that can impede viral replication. Studies have revealed that bael extracts are effective against several viruses, including the herpes simplex virus (HSV) and human immunodeficiency virus (HIV). These effects are believed to be due to the ability of bael constituents to interfere with viral entry and replication processes. Furthermore, bael's antiviral properties extend to its potential in managing

viral-induced diseases such as influenza and hepatitis. Its high content of antioxidants and anti-inflammatory agents enhances the immune response, providing additional protection against viral infections and aiding in faster recovery. Bael (*Aegle marmelos* L.) offers a dual protective role through its antimicrobial and antiviral activities. This underscores its potential as a valuable component in both traditional and modern therapeutic practices. Future research should continue to explore and validate these properties to harness bael's full therapeutic potential (Khan and Sultana, 2011).

4. Anti-Inflammatory Effects of Bael: Mechanisms and Applications

Bael (*Aegle marmelos* L.), a tropical fruit revered for its therapeutic potential, has garnered significant interest for its anti-inflammatory properties. The anti-inflammatory effects of Bael are attributed to its rich composition of bioactive compounds, including flavonoids, alkaloids, and essential oils. These compounds act through multiple mechanisms to mitigate inflammation. One primary mechanism involves the modulation of pro-inflammatory cytokines. Bael extract has been shown to reduce levels of interleukins (IL-1 β , IL-6) and tumor necrosis factor-alpha (TNF- α), key mediators in the inflammatory response. This action is facilitated by Bael's ability to inhibit nuclear factor-kappa B (NF- κ B) and mitogen-activated protein kinases (MAPKs), which are crucial pathways in inflammatory signaling (Kokateet *al.* 2002).

Additionally, Bael exhibits antioxidant properties that contribute to its anti-inflammatory effects. By scavenging reactive oxygen species (ROS) and enhancing the body's antioxidant defense system, Bael reduces oxidative stress, which is often a precursor to inflammation. The fruit's polyphenolic compounds, particularly its tannins and flavonoids, play a significant role in this antioxidant activity. In clinical and preclinical studies, Bael has demonstrated efficacy in reducing symptoms of various inflammatory conditions, such as arthritis, colitis, and respiratory inflammation. The fruit's extracts have been utilized in traditional medicine for their ability to alleviate pain and swelling, offering a natural alternative or complement to conventional anti-inflammatory drugs (Kour and Singh, 2012).

Overall, the anti-inflammatory properties of Bael are attributed to its complex interplay of bioactive compounds that target inflammatory pathways and oxidative stress. Further research and clinical trials are warranted to fully elucidate the therapeutic potential of Bael in

inflammatory diseases and to establish optimal dosage and application methods (Citarasuet *al.* 2003).

5. Bael in the Management of Diabetes and Blood Glucose Regulation

Bael (*Aegle marmelos* L.), a traditional medicinal plant, has garnered interest for its potential in managing diabetes and regulating blood glucose levels. The fruit, leaves, and bark of bael have been extensively studied for their antidiabetic properties, primarily attributed to their rich phytochemical profile (Costa-Lotufoet *al.* 2005).

Research indicates that bael exerts hypoglycemic effects through several mechanisms. The fruit is a significant source of dietary fiber, which aids in controlling blood glucose levels by delaying carbohydrate absorption in the gut. Additionally, bael contains bioactive compounds such as flavonoids, alkaloids, and essential oils, which have demonstrated insulin-mimetic and insulin-sensitizing activities. These compounds enhance insulin secretion and improve insulin sensitivity, thus contributing to better glycemic control. Clinical studies support bael's role in diabetes management. For instance, animal models and human trials have shown that bael extract can significantly reduce fasting blood glucose levels and improve glycemic parameters. The presence of compounds like tannins and saponins in bael has also been linked to reduced oxidative stress and inflammation, which are critical in diabetes pathogenesis and complications (Dhankhar, 2010).

Moreover, bael's antidiabetic effects extend beyond glucose regulation. The plant has been reported to improve lipid profiles, reduce body weight, and enhance overall metabolic health, making it a valuable adjunct in diabetes management. Its potential to mitigate complications associated with diabetes, such as nephropathy and neuropathy, further underscores its therapeutic value. Bael (*Aegle marmelos* L.) offers promising benefits in diabetes management and blood glucose regulation. Its multifaceted mechanisms, including dietary fiber content and bioactive compounds, position it as a complementary therapeutic option for diabetes care. Further research is warranted to fully elucidate its efficacy and safety in diverse populations (Dhankhar *et al.* 2011).

6. Bael's Potential for Therapeutic Benefits in Digestive Health

Bael (*Aegle marmelos* L.) is a revered medicinal plant known for its extensive therapeutic benefits, particularly in gastrointestinal health. The fruit, leaves, and bark of Bael have been

traditionally used to treat various digestive disorders, supported by emerging scientific evidence that underscores its efficacy (Dhar *et al.* 1968).

Bael is rich in bioactive compounds such as alkaloids, flavonoids, tannins, and essential oils, which contribute to its gastroprotective effects. Research indicates that Bael possesses antidiarrheal properties, likely due to its high tannin content, which helps in reducing intestinal inflammation and controlling fluid loss. Clinical studies have shown that Bael pulp can effectively manage acute and chronic diarrhea by modulating gut motility and enhancing stool consistency. Moreover, Bael exhibits significant potential in managing peptic ulcers. Its flavonoids and phenolic compounds have demonstrated ulcer-healing properties by promoting mucosal integrity and reducing gastric acid secretion. Animal studies have confirmed that Bael extracts can significantly reduce ulcer indices and facilitate ulcer healing, potentially offering an alternative or complementary approach to conventional ulcer therapies (Dhiman, 2003).

Additionally, Bael's role in alleviating constipation is notable. Its fiber-rich content enhances bowel movements and improves overall digestive health. Bael's laxative effect, attributed to its mucilage content, helps in softening stools and preventing constipation. Overall, Bael's multifaceted therapeutic properties make it a valuable natural remedy for gastrointestinal health. Its ability to address diarrhea, peptic ulcers, and constipation highlights its potential as a functional food and medicinal plant. Continued research and clinical trials are essential to fully understand its mechanisms and validate its therapeutic benefits in gastrointestinal disorders (Dhuley, 2003).

6. Cardiovascular Benefits of Bael: Reducing Risks and Enhancing Heart Health

Bael (*Aegle marmelos* L.) has garnered significant attention for its potential cardiovascular benefits. This ancient fruit, revered in traditional medicine, offers a range of bioactive compounds that contribute to heart health. Research highlights several mechanisms through which Bael exerts its cardioprotective effects. The fruit is rich in flavonoids, alkaloids, and essential oils, which possess potent antioxidant properties. These antioxidants combat oxidative stress, a key contributor to cardiovascular diseases (CVD) by neutralizing free radicals and reducing cellular damage. The reduction in oxidative stress helps prevent the oxidation of low-density lipoprotein (LDL) cholesterol, thus mitigating the risk of atherosclerosis (Dhuley, 2007).

Bael's impact on blood pressure regulation is another notable benefit. Studies have demonstrated that Bael can help lower systolic and diastolic blood pressure, potentially due to its vasodilatory effects and the presence of compounds that improve endothelial function. Enhanced endothelial function contributes to better blood flow and reduced hypertension, a significant risk factor for heart disease (Evans and Saponin, 2002).

Additionally, Bael's high fiber content supports cardiovascular health by improving lipid profiles. Fiber aids in lowering total cholesterol and LDL levels while increasing high-density lipoprotein (HDL) cholesterol. The fruit's anti-inflammatory properties also play a crucial role in reducing inflammation, a common underlying factor in CVD. Bael's traditional use in managing diabetes also indirectly benefits cardiovascular health. By helping to regulate blood glucose levels, Bael reduces the risk of diabetes-related cardiovascular complications, such as diabetic cardiomyopathy. Bael's cardiovascular benefits are attributed to its antioxidant, anti-inflammatory, and lipid-lowering properties. As research continues to unfold, Bael holds promise as a natural adjunct to conventional therapies for enhancing heart health and reducing cardiovascular risks (Fanner *et al.* 1993).

7. Bael and Cancer: Exploring Antitumor and Chemo preventive Properties

Bael (*Aegle marmelos* L.), a significant medicinal plant in traditional Ayurvedic medicine, has drawn attention for its potential antitumor and chemo preventive properties. Recent scientific studies indicate that various phytochemicals found in bael, such as flavonoids, tannins, and coumarins, may play a critical role in inhibiting cancer cell proliferation and promoting apoptosis (programmed cell death). These compounds possess strong antioxidant activity, which helps to neutralize harmful free radicals and prevent oxidative damage, a key factor in cancer development (Geetha and Varalakshmi, 2001).

Several *in vitro* and *in vivo* studies have explored to inhibit cancer cell growth in different types of cancer, including breast, colon, and lung cancers. For instance, the methanolic extract of bael has shown cytotoxic effects against human cancer cell lines, such as MCF-7 (breast cancer) and HeLa (cervical cancer), by inducing apoptosis through the mitochondrial pathway. Moreover, the ability to downregulate pro-inflammatory cytokines and inhibit the activity of enzymes like cyclooxygenase-2 (COX-2), both of which are involved in cancer progression (Gheisari *et al.* 2011).

Bael's chemo preventive potential lies in its ability to modulate various molecular targets, such as tumor suppressor proteins (p53) and pro-apoptotic factors (Bax). Furthermore, its detoxifying and anti-inflammatory effects may reduce the risk of cancer development by curbing chronic inflammation, a known precursor to tumorigenesis. While more clinical trials are needed to validate these findings, the antitumor and chemo preventive properties of bael suggest its promising role as an adjunctive therapy in cancer prevention and treatment, highlighting its therapeutic potential in integrative oncology (Ghosh and Playford).

8. Future Prospects: Developing Bael-Based Nutraceuticals and Therapeutic Applications

Bael (*Aegle marmelos* L.) has demonstrated significant potential for the development of nutraceuticals and therapeutic products due to its broad spectrum of bioactive compounds, including flavonoids, alkaloids, and terpenoids. The fruit, leaves, and bark of bael are rich in antioxidants, anti-inflammatory agents, and antimicrobial properties, making it a prime candidate for the formulation of natural health products. As global interest in plant-based therapies and functional foods increases, bael-based nutraceuticals are poised to gain prominence in the wellness industry. One of the key future directions involves the exploration of bael as a functional ingredient in dietary supplements targeting metabolic disorders, such as diabetes, dyslipidemia, and obesity. Several studies have highlighted bael's hypoglycemic and lipid-lowering effects, which could be harnessed in therapeutic applications to manage or prevent chronic conditions. Additionally, bael's demonstrated gastroprotective, hepatoprotective, and cardioprotective properties make it an ideal candidate for developing supplements that support gut health, liver function, and cardiovascular wellness (Goel *et al.* 2000).

Further research is necessary to standardize bael extracts and optimize bioavailability for enhanced therapeutic efficacy. Advancements in biotechnology and nanotechnology could facilitate the development of more effective bael-based formulations, improving their absorption and targeted action. Clinical trials are essential to validate the safety and efficacy of these products, ensuring their regulatory approval and consumer acceptance. The future development of bael-based nutraceuticals and therapeutic applications holds immense promise. By integrating traditional knowledge with modern science, bael can be transformed into a valuable resource for addressing global health challenges, particularly in the areas of metabolic health, digestive wellness, and chronic disease prevention (Gapalan *et al.* 2010).

CONCLUSION

Bael (*Aegle marmelos* L.), a medicinally important plant, has shown remarkable potential in promoting human health and offering therapeutic benefits. This comprehensive review highlights its rich phytochemical profile, including alkaloids, flavonoids, tannins, and coumarins, which contribute to its diverse pharmacological properties. Bael has demonstrated antioxidant, anti-inflammatory, antimicrobial, antidiabetic, hepatoprotective, and cardioprotective effects, making it a valuable natural remedy for various ailments. Notably, its antitumor and chemopreventive activities open new avenues for cancer research and treatment. While traditional medicine has long recognized its benefits, modern scientific studies provide robust evidence supporting its use in developing nutraceuticals and therapeutic applications. However, more clinical studies are necessary to fully validate its medicinal potential and ensure safe and effective formulations. Overall, Bael represents a promising natural resource for health promotion, disease prevention, and therapeutic interventions in the future.

REFERENCES

1. Abdullakasim, P., Songchitsomboon, S., Techagumpuch, M., Balee, N., Swatsitang, P. and Sungpuag, P. 2007. Antioxidant capacity, total phenolics and sugar content of selected *Thai* health beverages. *International Journal of Food Sciences and Nutrition*. 58(1):77-85.
2. Agarwal, V.S. 1997. Rural economics of medicinal plants: vegetable in the forests. *Drug Plant of India*, Kalyani Publishers, New Delhi. 58(4):1-160.
3. Aiyer, A.K.Y.N. 1956. *The Antiquity of Some Field and Forest Flora of India*. Bangalore Printing and Publishing, Bangalore. 62(1):15-21.
4. Akazone, Y. 2004. Characteristic and physiological functions of polyphenols from apple. *Biofactors*. 22(14):311-314.
5. Ali, M.S. and Pervez, M.K. 2004. Marmenol: a 7-geranyloxy coumarin from the leaves of *Aegle marmelos* L. *Corr. National Centre for Natural Product Research*. 18(2):141-146.
6. Anonymous 2011. Fruit juice nutrition and health. *International Federation of Fruit Juice Producers, IFU Science Review*. 22(1):1-11.

7. Ansary, P.Y. 2005. *A Handbook on the Plant Sources of Indigenous Drugs*. International Book Distributors, Dehradun. 12(5):36-38.
8. Aprikian, O., Levrat-Verny, M.A., Besson, C., Busserolles, J., Remesy, C. and Demigne, C. (2001), "Apple favourably affects parameters of cholesterol metabolism and of anti-oxidative protection in cholesterol feed rats", *Food Chemistry*, Vol. 75 No. 4, pp. 445-452.
9. Aritajat, S., Kaweewat, K., Manosroi, J. and Manosroi, A. (2000), "Dominant lethal test in rats treated with some plant extracts", *Southeast Asian Journal of Tropical Medicine and PublicHealth*, Vol. 31 No. 1, p. 171.
10. Arul, V., Miyazaki, S. and Dhananjayan, R. (2005), "Studies on the anti-inflammatory, antipyretic and analgesic properties of the leaves of *Aegle marmelos* L. Corr", *Journal of Ethno-Pharmacology*, Vol. 96 No. 4, pp. 159-163.
11. Atul, N.P., Nilesh, V.D., Akkatai, A.R. and Kamlakar, S.K. (2012), "A review on *Aegle marmelos* Corr: a potential medicinal tree", *International Research Journal of Pharmacy*, Vol. 3 No. 8, pp. 86-91.
12. Babbar, O.P., Joshi, M.N. and Madan, A.R. (1982), "Evaluation of plants for antiviral activity", *Indian Journal of Medical Research*, Vol. 76, p. 54.
13. Badam, L., Bedekar, S.S., Sonawane, K.B. and Joshi, S.P. (2002), "In vitro antiviral activity of *bael* (*Aegle marmelos* Corr) upon human coxsackieviruses B1-B6", *The Journal of Communicable Diseases*, Vol. 34 No. 2, pp. 88-99.
14. Baliga, M.S., Bhat, H.P., Joseph, N. and Fazal, F. 2011. Phytochemistry and medicinal uses of the *baelfruit* (*Aegle marmelos* L.): a concise review. *International Journal of Food Research*. 44(7):1768-1775.
15. Baliga, M.S., Bhat, H.P., Pereira, M.M., Mathias, N. and Venkatesh, P. (2010), "Radioprotective effects of *Aegle marmelos* Corr: a concise review", *The Journal Alternative and Complementary Medicine*, Vol. 16 No. 10, pp. 1109-1116.
16. Baliga, M.S., Thilakchand, K.R., Rai, M.P., Rao, S. and Venkatesh, P. 2012. *Aegle marmelos* L. Corr and its phytochemicals in the treatment and prevention of cancer. *Integrative Cancer Therapies*. 12(3):187-196.

17. Bansal, Y. and Bansal, G. 2011. Analytical methods for standardization of *Aegle marmelos* Corr: a review. *Journal of Pharmaceutical Education and Research*. 2(2):37-44.
18. Barthakur, N.N. and Arnolds, N.P. 1989. Certain organic and inorganic constituents in *bael*(*Aegle marmelos*Correa) fruits. *Tropical Agriculture*. 66(1):10-14.
19. Basak, R.K., Mandal, P.K. and Mukherjee, A.K. 1982. Investigation on the structure of a hemicelluloses fraction isolated from the trunk of *Aegle marmelos*L. Corr Tree. *Carbohydrate Research*. 104(2):309-317.
20. Basu, D.K. and Brahmankar, D.M. 1983. Studies on interactions of calcium antagonist with cardioactive agents. *Indian Journal of Pharmacology*. 15(4):321-330.
21. Benni, J.M., Jayanthi, M. and Suresha, R. 2011. Evaluation of the anti-inflammatory activity of *Aegle marmelos* Corr Root. *Indian Journal of Pharmacology*. 43(4):393-397.
22. Bhardwaj, R.L. 2012. Fruit juice: a novel functional food and its categories. *Agrobios News Letter*. 11(8):87-88.
23. Bhardwaj, R.L. 2014. Role of *bael*fruit juice in nutritional security of Sirohi tribals. *Benchmark Survey Report of Sirohi Tribals*, Krishi Vigyan Kendra, AU, Jodhpur. 45(8):11-37.
24. Bhardwaj, R.L. and Pandey, S. 2011. Juice blends-a way of utilization of underutilized fruits, vegetables and spices: a review. *Critical Review in Food Science and Nutrition*. 51(6):563-570.
25. Bhardwaj, R.L., Nandal, U., Pal, A. and Jain, S. 2014. Bioactive compounds and medicinal properties of fruit juice. *Fruits*. 69(5):391-421.
26. Bhaskara Rao, K.V., Sekar, D.K., Kumar, G. and Karthik, L. 2011. A review on pharmacological and phytochemical properties of *Aegle marmelos*L. Corr (Rutaceae). *Asian Journal of Plant Sciences and Research*. 1(2):8-17.
27. Biswas, K., Bandyopadhyay, U., Chattopadhyay, I., Varadaraj, A., Ali, E. and Banerjee, R.K. 2003. A novel antioxidant and antiapoptotic role of omeprazole to block gastric ulcer through scavenging of hydroxyl radical. *The Journal of Biology Chemistry*. 278(13):109-119.

28. Bramhachari, P.V. and Reddy, Y.K. 2010. Phytochemical examination: antioxidant and radical scavenging activity of *Aegle marmelos* L. Corr extracts. *Journal of Pharmacy Research*. 3(12):3023-3025.
29. Brijesh, S., Daswani, P., Tetali, P., Antia, N. and Birdi, T. 2009. Studies on the antidiarrhoeal activity of *Aegle marmelos* Corr unripe fruit: validating its traditional usage. *BMCComplementary and Alternative Medicine*. 9(1):47-52.
30. Capasso, R., Pinto, L., Vuotto, M.L. and Di Carlo, G. 2000. Preventive effect of eugenol on PAF and ethanol-induced gastric mucosal damage. *Fitoterapia*. 71(1):131-135.
31. Chaouki, W., Leger, D.Y., Liagre, B., Beneytout, J.L. and Hmamouchi, M. 2009. Citral inhibits cell proliferation and induces apoptosis and cell cycle arrest in MCF-7 cells. *Fundamental and Clinical Pharmacology*. 23(5):549-556.
32. Charoensiddhi, S. and Anprung, P. 2008. Bioactive compounds and volatile compound of *Thai baelfruit* (*Aegle marmelos* L. Corr) as a valuable source for functional food ingredients. *International Journal of Food Research*. 15(3):287-295.
33. Citarasu, T., Rajajeyasekar, R., Venkatmalingam, K., Dhandapani, P.S. and Peter Marian, M. 2003. Effect of wood apple *Aegle marmelos* Corr extract as an antibacterial agent on pathogens infecting prawn (*Penaeus indicus*) larvi-culture. *Indian Journal of Marine Science*. 32(2):156-161.
34. Costa-Lotufo, L.V., Khan, M.T.H., Ather, A., Wilke, D.V., Jimenez, P.C., Pessoa, C., Moaraes, M.E.A. and Moreas, M.O.D. 2005. Studies of anticancer potential of plants used in Bangladeshi folk medicine. *Journal of Ethnopharmacology*. 99(1):21-30.
35. Dhankhar, S. 2010. *Aegle marmelos* L. Corr: a source of phytomedicine. *Journal of Medicinal Plants Research*. 5(9):1497-1507.
36. Dhankhar, S., Ruhil, S., Balhara, M., Dhankhar, S. and Chhillar, A.K. 2011. *Aegle marmelos* L. Corr: a potential source of phytomedicine. *Journal of Medicinal Plants Research*. 5(9):1497-1507.
37. Dhar, M.L., Dhar, M.M., Dhawan, B.N., Mehrotra, B.N. and Ray, C. 1968. Screening of Indian plant for biological activity. *Indian Journal of Experimental Biology*. 7(4):232-239.
38. Dhiman, A.K. 2003. *Discussion of Plant, Sacred Plants and their Medicinal Uses*. Daya Publication House, New Delhi. 4(1):18-19.

39. Dhuley, J.N. 2003. Investigation on the gastroprotective and antidiarrhoeal properties of *Aegle marmelos* unripe fruit extracts. *Hindustan Antibiotics Bulletin*. 45(46):41-46.
40. Dhuley, J.N. 2007. Investigation on the gastroprotective and antidiarrhoeal properties of *Aegle marmelos* unripe fruit extracts. *Hindustan Antibiotics Bulletin*. 41(1):45-46.
41. Evans, W.C. and Saponin 2002. Cardioactive drugs and other stories. *Trease and Evans Pharmacognosy*, 15th ed., British Library Cataloguing in Publication Data. 23(8):294-298.
42. Fenner, F.J., Gibs, E.P.J., Murphy, F.A., Rott, R. and Studdart, M.J. 1993. *White D. Veterinary Virology*, 2nd ed., Academic Press, London. 45(8):301-309.
43. Geetha, T. and Varalakshmi, P. 2001. Anti-inflammatory activity of lupeol and lupeol linoleate in rats. *Journal of Ethnopharmacology*. 76(1):77-80.
44. Gheisari, H.R., Amiri, F. and Zolghadri, Y. 2011. A review on *Aegle marmelos*: a potential medicinal tree. *International Journal of Current Pharmaceutical Research*. 3:85-88.
45. Ghosh, S. and Playford, R.J. 2003. Bioactive natural compounds for the treatment of gastrointestinal disorders. *Clinical Science (London)*. 104(6):547-556.
46. Goel, R.K., Maiti, R.N., Manickan, M. and Ray, A.B. 2000. Antiulcer activity of naturally occurring pyrano-coumarin and isocoumarin and their effect on prostanoid synthesis using human colonic mucosa. *Indian Journal of Experimental Biology*. 35(10):1080-1083.
47. Gopalan, C., Ramashastri, B.V., Balasubramanian, S.C., Rao, N.B.S., Deosthale, Y.G. and Pant, K.C. 2010. *Nutritive Value of Indian Foods*, National Institute of Nutrition, ICMR, Hyderabad. 1-135.
48. Gupta, N., Agrawal, R., Shrivastava, V., Roy, A. and Prasad, P. 2012. Chemopreventive potential of *Aegle marmelos* fruit extract against 7, 12-Dimethylbenz (a) anthracene-induced skin papillomagenesis in mice. *Research Journal of Pharmacology Pharmacodynamics*. 4(2):87-90.
49. Haider, R., Khan, A.K., Aziz, K.M., Chowdhury, A. and Kabir, I. 1991. Evaluation of indigenous plants in treatment of acute shizellosis. *Tropical and Geographical Medicine*. 43(3):266-270.

50. Jafri, M.A., Farah, J.K. and Singh, S. 2001. Evaluation of the gastric antiulcerogenic effect of large cardamom. *Journal of Ethnopharmacology*, 23(75):89-94.
51. Jagetia, G.C., Venkatesh, P. and Baliga, M.S. 2003. Evaluation of the radioprotective effect of *Aegle marmelos* L. Corr in cultured human peripheral blood lymphocytes exposed to different doses of radiation: a micronucleus study, *Mutagenesis*. 4(18):387-393.
52. Jagetia, G.C., Venkatesh, P. and Baliga, M.S. 2004. Fruit extract of *Aegle marmelos* Corr protects mice against radiation-induced lethality. *Integrative Cancer Therapies*. 3(4):323-325.
53. Jagetia, G.C., Venkatesh, P. and Baliga, M.S. 2005. *Aegle marmelos* L. Corr inhibits the proliferation of transplanted Ehrlich ascites carcinoma in mice. *Biology and Pharmaceutical Bulletin*. 28(1):58-64.
54. Johnson, M. 2010. Biochemical variation studies in *Aegle marmelos* L. Corr: a medicinally important plant. *Journal of Chemical and Pharmaceutical Research*. 2(6):454-462.
55. Jyothi, K.S. and Rao, B.S. 2010. Antibacterial activity of extracts from *Aegle marmelos* Corr: against standard pathogenic bacterial strains. *International Journal of Pharm Tech Research*. 2(3):1824-1826.
56. Kala, C.P., Farooquee, N.A. and Majila, B.S. 2005. Indigenous knowledge and medicinal plants used by Vaidyas in Uttarakhand. *Indian Natural Product Radiance*. 4(3):195-204.
57. Kamalakkannan, N. and Prince, P.S.M. 2003. Hypoglycaemic effect of water extract of *Aegle marmelos* Corr: fruit in streptozotocin diabetic rats. *Journal of Ethnopharmacology*. 87(23):207-210.
58. Kamalakkannan, N. and Prince, P.S.M. 2005. Antihyperlipidemic effect of *Aegle marmelos* Corr fruit extract in streptozotocin induced diabetes in rats. *Journal of the Science of Food and Agriculture*. 85:569-573.
59. Kar, A., Choudhry, B.K. and Bandhopadhyay, N.G. 2002. Comparative evaluation of hypoglycemic activity of some Indian medicinal plants in alloxan diabetic rats. *Journal of Ethnopharmacology*. 84(1):105-108.
60. Karakaya, S. 2004. Bioavailability of phenolic compounds. *Critical Review in Food Science Nutrition*. 44:453-464.

61. Kaur, S., Kaur, P., Walia, A. and Kumar, S. 2009. Antigenotoxic activity of polyphenolic rich extracts from *Aegle marmelos* L. Corr in human blood lymphocytes and *E. coli* *Records of Natural Products*. 3(1):68-75.
62. Kaushik, R.A., Yamdagni, R. and Dhawan, S.S. 2000. Physico-chemical characteristics of baelfruit at green and ripe stage of maturity. *Haryana Journal Horticultural Sciences*. 29(12):44-45.
63. Khan, H.T. and Sultana, S. 2011. Effect of *Aegle marmelos* Corr on DEN initiated and 2-AAF promoted hepatocarcinogenesis: a chemo preventive study. *Toxicology Mechanisms and Methods*. 21(6):453-462.
64. Kokate, C.K., Purohit, A.P. and Gokhale, S.B. 2002. Drugs containing glycosides *Pharmacognosy*. 21st ed. Nirali Prakashan, Pune. 12(7):158-239.
65. Kour, K. and Singh, N.D. 2012. Fruits: a smart choice of antioxidants. *Agrobios News Letter*. 11(8):76-77.
66. Kruawan, K. and Kangsadalampai, K. 2006. Antioxidant activity, phenolic compound contents and antimutagenic activity of some water extract of herbs. *Thai Journal of Pharmaceutical Sciences*. 30:28-35.
67. Kumar, V., Ahmed, D., Verma, A., Anwar, F., Ali, M. and Mujeeb, M. 2013. Umbelliferone D-galactopyranoside from *Aegle marmelos* L. Corr an ethno medicinal plant with antidiabetic, antihyperlipidemic and antioxidative activity. *BMC Complementary and Alternative Medicine*. 13:273-275.
68. Kyle, J.A., Sharp, L., Little, J., Duthie, G.G. and McNeill, G. 2009. Dietary flavonoid intake and colorectal cancer: a case control study. *British Journal of Nutrition*. 7:1-8.
69. Lambole, V.B., Krishana, M., Upendra, K., Bhatt, S.K.P. and Vipul, G. 2010. Phytopharmacological properties of *Aegle marmelos* L. Corr as a potential medicinal tree: an overview. *International Journal of Pharmaceutical Sciences Review and Research*. 5(2):67-72.
70. Laphookhieo, S. 2011. Chemical constituent from *Aegle marmelos* L. Corr. *Journal of Brazilian Chemistry Society*. 22:176-178.
71. Leticia, V. and Costa, L. 2005. Evaluation of anticancer potential used in Bangladeshi folk medicine. *Journal of Ethnopharmacology*. 99(1):21-38.

72. Liu, E.S. and Cho, C.H. 2000. Relationship between ethanol-induced gastritis and gastric ulcer formation in rats. *Digestion*. 62(4):232-239.
73. Ma, Y. 2005. Association between dietary carbohydrates and body weight. *American Journal of Epidemiology*. 161(4):359-367.
74. Madhu, C., Hindu, K., Sudeepthi, C., Maneela, P., Reddy, K.V. and Sree, B.B. 2012. Anti-ulcer activity of aqueous extract of *Aegle marmelos* Corr leaves on rats. *Asian Journal of Pharmaceutical Research*. 2(4):132-135.
75. Maheshwari, V.L., Joshi, P.V. and Patil, R.H. 2009. In vitro anti diarrheal activity and toxicity profile of *Aegle marmelos* L. Corr dried fruit pulp. *Natural Product Radiance*. 8(5):498-502.
76. Maity, P., Biswas, K., Roy, S., Banerjee, R.K. and Bandyopadhyay, U. 2003. Smoking and the pathogenesis of gastroduodenal ulcer-recent mechanistic update. *Molecular and Cellular Biochemistry*. 253(12):329-338.
77. Maity, P., Hansda, D., Bandyopadhyay, U. and Mishra, D.K. 2009. Biological activities of crude extracts and chemical constituents of *Aegle marmelos* L. Corr. *Indian Journal of Experimental Biology*. 47(11):849-861.
78. Malviya, R., Kumar, A., Singh, A. and Kulkarni, G.T. 2012. Pharmacological screening, ayurvedic values and commercial utility of *Aegle marmelos* Corr. *International Journal of Drug Development and Research*. 4(1):28-37.
79. Marwat, S.K. and Khan, M.A. 2009. Fruit plant species mentioned in the holy Qura'n and ahadith and their ethnomedicinal importance. *American-Eurasian Journal of Agricultural and Environmental Sciences*. 5(2):284-295.
80. Mukharjee, B. and Ahmad, K. 1957. Riboflavin. *Pakistan Journal of Biological and Agricultural Sciences*. 4(5):47-51.
81. Nandal, U. 2013. Nutritive value and therapeutic uses of beverages. *A Handbook of Foods and Nutritional Biochemistry*, Agrobios, Jodhpur. 8(4):251-270.
82. Nandal, U. and Bhardwaj, R.L. 2012. Role of underutilized fruits in nutritional and economic security of tribals: a review. *Critical Review in Food Science*. 54(7):880-890.
83. Nandal, U. and Bhardwaj, R.L. 2013. Role of fruit juices in nutritional and health security. *Indian Food Pack*. 67(4):112-122.

84. Nandal, U. and Meena, R.P. 2012. Role of fruit juices in nutritional security: a concept. *Agrobios News Letter*. 11(2):89-90.
85. Nirupama, G.S., Padmasri, G., Ramesh, R.V. and Vasanthi, M. 2012. Comparative analysis of phytochemical constituents presents in various parts of *Aegle marmelos*L. *Corr. AsianPacific Journal of Tropical Disease*. 45(8):774-777.
86. Parekh, J. and Chanda, S.V. 2007. *In vitro* antimicrobial activity and phytochemical analysis of some Indian medicinal plants. *Turkey Journal of Biology*. 31(1):53-58.
87. Paricha, S. 2004. *Aegle Marmelos*Corr: nature's most natural medicinal fruit. *Orissa Review*. 46(2):16-17.
88. Parihar, N. and Kumar, S. 2013. Study of antifungal potential of *Aegle marmelos*L. *Corr: a medicinal plant*. *International Journal of Plant, Animal and Environmental Sciences*. 3(1):126-129.
89. Patel, P. and Mohammad Syed, A.B. 2010. Immunomodulatory activity of methanolic fruit extract of *Aegle marmelos*Corr in experimental animals. *Saudi Pharmaceutical Journal*. 18(3):161-165.
90. Patel, P.K., Jyoti, S., Lokesh, S., Narendra, K.P. and Dubey, B.K. 2012. *Aegle marmelos*L. *Corr: a review on its medicinal properties*. *International Journal of Pharmaceutical andPhytopharmacological Research*. 1(5):332-341.
91. Pawar, R.S. and Bhutani, K.K. 2004. Protobasic acid glycosides from *Madhuca indica* with inhibitory activity on free radical release from phagocytes. *Journal of Natural Product*. 67(5):668-671.
92. Prakash, D., Upadhyay, G., Pushpangadan, P. and Gupta, C. 2011. Antioxidant and free radical scavenging activities of some fruits. *Journal of Complementary and Integrative Medicine*. 8(2):1513-1515.
93. Prasad, S., Nigam, N., Kalra, N. and Shukla, Y. 2008. Regulation of signaling pathways involved in lupeol induced inhibition of proliferation and induction of apoptosis in human prostate cancer cells. *Molecular Carcinogenesis*. 47(12):916-924.
94. Purohit, S.S. and Vyas, S.P. 2004. *Aegle Marmelos Corr, Medicinal Plant Cultivation: A Scientific Approach*, Agrobios, Jodhpur. 25(1):280-285.
95. Rajadurai, M. and Prince, P.S. 2005. Comparative effects of *Aegle marmelos*Corr extract and alpha-tocopherol on serum lipids, lipid peroxides and cardiac enzyme levels in rats

- with isoproterenol-induced myocardial infarction. *Singapore Medical Journal*. 46(5):78-80.
96. Rajan, S., Gokila, M., Jency, P., Brindha, P. and Sujatha, R.K. 2011. Antioxidant and phytochemical properties of *Aegle marmelos* L. Corr fruit pulp. *International Journal of Current Pharmaceutical Research*. 3(2):65-70.
 97. Rangari, V.D. 2004. *Traditional Drugs of India, Pharmacognosy and Phytochemistry Part-II*, 1st ed., Carrier Publication, Nasik. 25(1):182-184.
 98. Rastogi, R.P. and Mehrotra, B.N. 1995. *In Compendium of India Medicinal Plants*, Lucknow Publication and Information Directorate, New Delhi. 15(1):15-18.
 99. Rathore, M. 2009. Nutrient content of important fruit trees from arid zone of Rajasthan. *Journal of Horticulture and Forestry*. 1(7):103-108.
 100. **Robbers, J.E.** and Tyler, V.E. 2002. Herbs of choice: the therapeutic use of phytochemicals. *International Journal of Pharmacy Science*. 3(2):199-203.
 101. Roy, S.K. and Singh, R.N. (1980), "Studies on changes during development and ripening of baelfruit", *Punjab Horticulture Journal*, Vol. 20 Nos 3/4, pp. 190-197.
 102. Ruxton, C.H.S., Gardner, E.J. and Walker, D. (2006), "Can pure fruit and vegetable juice protect against cancer and cardiovascular disease too: a review of the evidence", *International Journal of Food Science Nutrition*, Vol. 3, pp. 1-24.
 103. Sabu, M.C. and Kuttan, R. (2004), "Antidiabetic activity of *Aegle marmelos* L. Corr and its relationship with its antioxidant properties", *Indian Journal of Physiology and Pharmacology*, Vol. 48 No. 1, pp. 81-88.
 104. Sachdewa, A., Raina, D., Srivastava, A.K. and Khemani, L.D. (2001), "Effect of *Aegle marmelos* Corr and *Hibiscus rosa sinensis* leaf extract on glucose tolerance in glucose induced hyperglycemic rats", *Journal of Environmental Biology*, Vol. 22 No. 1, pp. 53-56.
 105. Sampathkumar, K.P., Madevi, M.U., Bhowmik, D., Singh, D.M. and Dutta, A.S. (2012), "Recent trends in medicinal uses and health benefits of Indian traditional herbs *Aegle marmelos* Corr", *The Pharma Innovation*, Vol. 1 No. 4, pp. 70-77.
 106. Sanchez-Moreno, C., Jimenez-Escrig, A. and Martin, A. (2009), "Stroke: roles of B vitamins, homocysteine and antioxidants", *Nutrition Research and Review*, Vol. 22 No. 1, pp. 49-67.

107. Sharma, G.N. and Dubey, S.K. (2011), "Ulcer healing potential of *Aegle marmelos* fruit seed", *Asian Journal of Pharmacy and Life Science*, Vol. 1 No. 2, pp. 172-178.
108. Sharma, G.N., Dubey, S.K., Sharma, P. and Sati, N. (2011), "Medicinal values of *bael* (*Aegle marmelos* L. Corr): a review", *International Journal of Current Pharmaceutical Review and Research*, Vol. 2 No. 1, pp. 12-22.
109. Sharma, P.C., Bhatia, V., Bansal, N. and Sharma, A. (2007), "A review on *bael*", *Natural Product Radiance*, Vol. 6 No. 2, pp. 171-178.
110. Shoba, F.G. and Thomos, M. (2001), "Study of antidiarrheal activity of four medicinal plants in castor-oil induced diarrhea", *Journal of Ethnopharmacology*, Vol. 76 No. 1, pp. 73-76.
111. Simmons, R.A. (2006), "Developmental origins of diabetes: the role of oxidative stress", *Free Radical Biology and Medicine*, Vol. 40 No. 6, pp. 917-922.
112. Singh, A., Sharma, H.K., Kaushal, P. and Upadhyay, A. (2014a), "*Bael* (*Aegle marmelos* L. Corr) products processing: A review", *African Journal of Food Science*, Vol. 8 No. 5, pp. 204-215.
113. Singh, A.K., Chakraborty, I. and Chaurasiya, A.K. (2014b), "*Bael* preserves syrup as booster of human health as a health drink", *The BioScan*, Vol. 9 No. 2, pp. 565-569.
114. Singh, D., Chaudhary, M., Chauhan, P.S., Prahalad, V.C. and Kavita, A. (2009), "Value addition to forest produces for nutrition and livelihood", *Indian Forester*, Vol. 135 No. 9, pp. 1271-1284.
115. Sivraj, R. and Balakrishana, A. (2011), "Preliminary phytochemical analysis of *Aegle marmelos* (L.) Corr", *International Journal of Pharmaceutical Science and Research*, Vol. 2 No. 1, pp. 146-150.
116. Slamenova, D., Horvathova, E., Wsolova, L., Sramkova, M. and Navarova, J. (2009), "Investigation of anti-oxidative, cytotoxic, DNA-damaging and DNA-protective effects of plant volatiles eugenol and borneol in human-derived HepG2, Caco-2 and VH10 cell lines", *Mutation Research*, Vol. 677 No. 1, pp. 46-52.
117. Srivastava, G.K., Prabhuji, S.K., Sinha, D., Srivastava, R. and Rao, V. (2012), "*Aegle marmelos* Corr: a potent medicinal tree of the tropics", *Medicinal Plants*, Vol. 4 No. 2, pp. 111-114.

118. Subramaniam, D., Giridharan, P., Murmu, N., Shankaranarayanan, N.P., May, R. and Houchen, C.W. (2008), "Activation of apoptosis by 1-hydroxy-5, 7- dimethoxy-2-naphthalene-carboxaldehyde, a novel compound from *Aegle marmelos* Corr", *Cancer Research*, Vol. 68 No. 20, pp. 8573-8581.
119. Suvimol, C. and Pranee, A. (2008), "Bioactive and volatile compounds of *Thaibaelfruit*", *International Food Research Journal*, Vol. 15 No. 3, pp. 1-9.
120. Tiwari, N.N. and Joshi, M.P. (1990), "Medicinal plants of Nepal II", *Journal of Nepal Medical Association*, Vol. 28, pp. 221-232.
121. Upadhyaya, S., Shanbhag, K.K., Suneetha, G., Naidu, B.M. and Upadhyaya, S. (2004), "A study of hypoglycemic and antioxidant activity of *Aegle marmelos* in alloxan induced diabetic rats", *Indian Journal Physiology and Pharmacology*, Vol. 48 No. 4, pp. 476-480.
122. Veerappan, A., Miyazaki, S., Kadarkaraisamy, M. and Ranganathan, D. (2007), "Acute and subacute toxicity studies of *Aegle marmelos* L. Corr, an Indian medicinal plant", *Phytomedicine*, Vol. 14 Nos 2/3, pp. 209-215.
123. Venkatesan, D., Karunakaran, M., Kumar, S.S., Palaniswamy, P. and Ramesh, G. (2009), "Antimicrobial activity of *Aegle marmelos* Corr against pathogenic organism compared with control drug", *Ethnobotanical Leaflets*, Vol. 13, pp. 968-974.
124. Vimal, V. and Devaki, T. (2004), "Linear furanocoumarin protects rat myocardium against lipidperoxidation and membrane damage during experimental myocardial injury", *Biomedical and Pharmacotherapy*, Vol. 58 Nos 6/7, pp. 393-400.
125. Yadav, N., Tyagi, G., Jangir, D.K. and Mehrotra, R. (2011), "Rapid determination of using reverse phase high performance liquid chromatography", *Journal of Pharmacy Research*, Vol. 4 No. 3, pp. 717-719.
126. Yadav, N.P. and Chanotia, C.S. (2009), "Phytochemical and pharmacological profile of leaves of *Aegle marmelos* L. Corr", *The Pharma Review*, pp. 144-149.