Phytochemical Investigation and In-Vitro Anti-Deppressant Activity of Albizia Lebbeck

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Abstract: The plant Albizia lebbeck is a species of the family Fabaceae. Native to Indian subcontinent and Myanmar. It is widely cultivated and naturalized in other tropical and subtropical regions, including Australia. Albizia lebbeck plant was selected because of its therapeutic uses, the part used was leaves. The leaves of Albizia lebbeck were extracted using maceration and soxhlet extraction process and yield obtained was 2mg and 86.23mg respectively, so soxhlet extraction gives better yield when compared with maceration. By invitro assay method the methanolic extract of Albizia lebbeck leaves produced better Anti depressant activity against PC12 and C6 cell lines, which is supported by increased cell viability from inducing leaves extract of Albizia lebbeck on PC12and C6 cell lines.

Keywords: Albizia lebbeck, Fabaceae, Invitro, viability, cell lines

1. Introduction

Depression is an affective mental disorder characterized by extreme exaggeration and mood disturbance. Decreased brain cells of monoamines like noradrenaline, dopamine, and serotonin leads to depression.1 Drugs that increase the level of these neurotransmitters in the CNS show antidepressant activity. Many of currently available antidepressant drugs have proven to be effective but they are burdened with some disadvantages such as various adverse effects, problematic interactions and relatively low response. On the other hand, drugs obtained from natural sources have good efficacy, least risk and low side effects profile. Therefore, herbal therapies should be considered as alternative medicines. In recent years, there has been growing interest in developing new antidepressant drugs from natural sources, one such source can be Albizia lebbeck, ²The plant Albizia lebbeck is a species of the family Fabaceae. Native to Indian subcontinent and Myanmar.3 It is widely cultivated and naturalized in other tropical and subtropica regions, including Australia. Common name sinenglish include Siris, Indian siris, East Indian walnut, Broomerain tree, Lebbeck tree, Frywood, Koko and woman's tongue tree. The leaves are bipinnate7.5 - 15 cm long with one to four pairs of pinnae. Each pinna with 6 - 8 le aflets. General chemical constituents in Albezialebbeck include alkaloids, anthraquinones, essential oils, flavanoids, glycosides, saponins, steroids and triterpenoids.4 The plant possesses many therapeutic activities such as treatment of leprosy, ulcers, ophthalmic and skin eruptions, skin diseases. It is a astringent also used by some cultures to treat boils, cough, eye flu, gingivitis, lung problems, pectoral problems. It is used as a tonic and is used to treat abdominal tumors, the bark is used medicinally treat inflammation and also effective in migraine

2. Materials and Methods

Albizia lebbeck plant was selected because of its therapeutic uses, the part used was leaves.

The leaves of Albizia lebbeck was collected Locally from Tumkur and It was identified and authenticated. The leaves of Albizia lebbeck was cleaned and shade dried at 25 degree Celsius. The dried leaves were coarsely powdered by a Mixer grinder and the powder was stored in

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airtight container, and this was used throughout the investigation. In this process, 10g of coarsely powdered leaves of albizialebbeck drug was dissolved in 150ml of methanol and was allowed to stand at room temperature for 3 days with frequent agitation. Then the mixture was strained and the marc was pressed, the combined liquids were clarified by filtration. After boiling, the filtrate was evaporated and thick mass was collected.5

1) LCMS analytical technique

Sample Preparation: 10mg of the sample extract is dissolved in 2mL of Methanol. Filtered and Injected. Instrumentation: The Acquity H - class UPLC (Waters Corporation, Milford, MA, USA) was employed, which had an integrated vacuum degasser, automatic sample manager (Serial C10UPA554M, Waters Corporation, Singapore), ultra performance binary solvent manager (Serial C10UPB081A, Waters Corporation, Singapore), and injection volume range of up to 100 µLwith an optional extension loop. A C18 stationaryphase (BEH C18, 50 x 1.0mm, 1.7µ) was used for chromatographic separation. A photodiode array detector (DAD) was employed in conjunction with a Xevo G2 - XS QToF (Serial # YFA1548, Waters Corporation, Wilmslow, UK) for mass spectrometric (MS) detection

2) In - vitro study of Albizia lebbeck leaves by MTTAssay Method

MTT assay is a colorimetric assay used for the determination of cell proliferation and cytotoxicity, basedonreductionofthe

yellowcolouredwatersolubletetrazoliumdyeMTTto formazan crystals. Mitochondrial lactate dehydrogenase produced by live cells reduces MTT to insoluble formazan crystals, which upon dissolution into an appropriate solvent exhibits purple colour, the intensity of which is proportional to the number of viable cells and can be measured spectrophotometrically at 570nm.

3. Result and Discussion

PC12 and C6 cell lines are used to determine the antidepressant activity, by MTT assay we have evaluated the

extract antidepressant activity of methanolic of albizialebbeck leaves depending on the percentage cell viability. The result have been summarised in Table No 1 and 2. It is seen that the methanolic extractof albizialebbeck leaves at 10ug/ml concentration showed antidepressant activity near to that of a Standard drug Fluoxetine at 10ug/ml concentration. By inducing cytotoxic drug Doxorubicin at 0.3ug/ml concentration preliminarily to PC12 and C6 cell lines the cell viability is decreased then the Standard drug Fluoxetine and Methanolic extract of albizialebbeck leaves is added to estimate the percentage increase in cell viability.

From this study we have determined that increase in the percentage of cell viability will increase the number of live and healthy cells which directly promotes the normal functioning of cells. On phytochemical screening of methanolic extract of albizialebbeck leaves by general test and by LCMSMS test we determined the presence of Alkoloids, flavonoids, saponins, phenolic acids, carbohydrates and proteins. Therefore, from this study we can presume that out of many phytoconstituents determined by general test and also by LCMSMS test, one or few constituents present in methanolic extract of albizia lebbeck leaves are responsible for the exhibited Antidepressant activity.

S. No	Concentration (ug/ml)	% cell viability
1)	Untreated	100
2)	Doxorubicin induced 0.3ug/ml	40.67
3)	Standard drug - 10ug/ml	91.21
4)	Testsample - 10ug/ml	84.28

1) % cell viability of Methanol extract against C6 cells

2) %Cell viability of methanol extract against PC12 cell

S. No	Concentration (ug/ml)	% cell viability
1)	Untreated	100
2)	Doxorubicin induced0.3ug/ml	41.63
3)	Standard drug - 10ug/ml	94.12
4)	Test sample - 10ug/ml	91.32

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4. Conclusions

The leaves of Albizia lebbeck were extracted using maceration and soxhlet extraction process and yield obtained was 2mg and 86.23mg respectively, so soxhlet extraction gives better yield whencompared with maceration. General chemical tests indicated presence of carbohydrates, alkaloids, proteins, saponins, and flavonoids. Whereas on further analysis by LCMS it showed presence of flavonoid glycosides, polyphenols, terpinoids and steroidal saponins, among this some of the constituents may be effective in showing Antideppressant activity. ByInvitro assay method the methanolic extract of Albizia lebbeck leaves produced better Antideppresant activity against PC12 and C6 cell lines, which is supported by increased cell viability from inducing leaves extract of Albizia lebbeck on PC12and C6 cell lines. From this invitro assay performed and observations noted, it can be conclude that Albizia lebbeck, Benth (Fabaceae) possess significant Antideppressant activity when compared with that of the standard marketed drug,

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