INTRODUCTION
Depression is a psychiatric disorder, which affects 21% of the world population and is considered as disorders of mood rather than disturbances of thought or cognition. It is a chronic illness associated with low mood and a range of associated emotional, cognitive, physical, and behavioral symptoms [1]. There are two types of depression: the first one is unipolar depression, in which the mood swings are always in the same direction and caused due to stressful events of life. The second one is bipolar depression, also termed as endogenous depression, in which the mood swings are bidirectional and unrelated to external stress [2]. Although there are numerous synthetic drugs available for the treatment of depression, they are associated with various adverse effects and drug–drug interactions that can compromise the therapeutic treatment. In the traditional systems of medicine, many plants and formulations have been used to treat depression for thousands of years. The results of the study indicate that CC can be used as an antidepressant herb.

MATERIALS AND METHODS
Collection of plant material and extraction
The aerial parts of CC were collected at NH-5 near Guntur. The plant was authenticated by the Department of Botany, Acharya Nagarjuna University, Guntur, and voucher specimen was preserved. The leaves and stems were separated, dried, powdered, and then extracted with alcohol as solvent using soxheletation for 4 cycles. Then, the extracted drug was further evaporated using simple distillation apparatus to obtain the concentrate. To this extract, aliquots of water were added and then fractionated successively using petroleum ether and chloroform.
by mother liquor method [13]. To this extract, 250 ml of water was added and shaken thoroughly, and then to this, 100 ml of petroleum ether was added to separate the non-polar constituents. This procedure was repeated until the appearance of colorless petroleum ether layer. All the fractions of petroleum ether layer were collected and evaporated to a concentrated residue. After separation of petroleum ether fraction, 100 ml of chloroform was added to the hydroalcoholic extract and this procedure is repeated until the chloroform layer becomes colorless. All the fractions of chloroform layer were collected and evaporated to a concentrated residue. The leftover portion is considered as hydroalcoholic fraction.

**Preliminary phytochemical screening**

Various qualitative tests were performed for the detection of phytochemical constituents present in all three fractions, for the presence of carbohydrates, tannins, flavonoids, steroids, glycosides, alkaloids, and saponins [14].

**Animals**

Swiss albino mice of either sex 3-4 months old and weighing around 20-30 g were selected. The animals had free access to food and water and were housed in an animal room with alternating light-dark cycle of 12 hrs each. The animals were acclimatized for at least 5 days to the laboratory conditions before the commencement of behavioral experiments. Experiments were carried out between 9:00 am and 11:00 am. The Institutional Animal Ethics Committee (IAEC) approved the experimental protocol, and the care of laboratory animals was taken as per the guidelines of CPCSEA, Ministry of Forests and Environment, Government of India (1529/P0/A/11/CPCSEA/IAEC/ PRO-03/2015-2016).

**Preparation of standard**

Imipramine was procured from Baroda Pharmaceuticals as a gift sample. Stock solution was prepared by dissolving 10 mg in 10 ml of distilled water, and then diluted to required dilutions.

**Measurement of antidepressant activity in albino mice**

In the present investigation, tail suspension test (TST) and forced swim test (FST) were selected as animal models for the evaluation of antidepressant activity in albino mice.

**Experimental protocol**

Animals were divided into 16 groups, and each group consists of 6 mice.

**TST**

The total duration of immobility induced by tail suspension was measured according to the method described by Steru [15] and Thierry [16]. Mice were suspended on the edge of a table 50 cm above the floor by the adhesive tape placed approximately 1 cm from the tip of the tail. Immobility time was recorded during a 6-minute period. Animal was considered to be immobile when it did not show any movement of the body and hanged passively. Each animal was used only once. The experimental protocol and treatment schedule were given in Table 1.

**FST**

FST was proposed as a model for antidepressant activity by Porsolt et al. [17,18]. Mice were forced to swim individually in a glass jar containing fresh water of 15 cm height and maintained at 25°C (±3°C). Mice were considered to be immobile when it remained floating in the water without struggling, making no or minimum movements of its limbs necessary to keep its head above water. The total duration of immobility was recorded during a 6-minute test. The changes in immobility duration were studied in all groups of animals. The experimental protocol and treatment schedule were similar to TST. On the 14th day, 90 minutes after administration of extracts, immobility period was recorded in all groups.

Statistical analysis

All the results were expressed as mean±standard error. Data were analyzed using one-way ANOVA followed by Dunnett’s test. In all the tests, the criterion for statistical significance was p<0.05.

**RESULTS AND DISCUSSION**

Depression is a psychiatric disorder that affects individuals’ quality of life, emotional and social relations directly. Most of the therapies require several weeks of treatment; there are various side effects caused by conventional antidepressant drugs before improvement of signs and symptoms is observed. To overcome these limitations of available antidepressant drugs, attempts are underway to explore medicinal plants with antidepressant activity. Many single and compound drug formulations of plant origin are being used in the treatment of psychiatric disorders and proved to have a better acceptance due to lower incidence of side effects [19,20]. Nowadays, plants with chemical constituents such as flavonoids, phenols, alkaloids, and triterpenoid saponins were reported to have antidepressant activity. Preliminary phytochemical screening was done for petroleum ether, chloroform, and alcoholic fractions of CC using various qualitative tests. The results of phytochemical screening are presented in Tables 2 and 3.

**Table 1: Experimental protocol for TST and FST**

| Group | Treatment schedule for TST |
|-------|---------------------------|
| Control | Only distilled water was administered orally for 14 days |
| Standard | Imipramine (10 mg/kg) was administered orally for 14 days |
| CCA1 | CCA (100 mg/kg) was administered orally for 14 days |
| CCA2 | CCA (200 mg/kg) was administered orally for 14 days |
| CCC1 | CCC (100 mg/kg) was administered orally for 14 days |
| CCC2 | CCC (200 mg/kg) was administered orally for 14 days |
| CCP1 | PCC (100 mg/kg) was administered orally for 14 days |
| CCP2 | PCC (200 mg/kg) was administered orally for 14 days |

**Groups 9-16 were similar to 1-8, except, that the immobility time was recorded using FST. FST: Forced swim test, TST: Tail suspension test, CC: Callistemon citrinus, CCA: Callistemon citrinus with petroleum, CCC: Callistemon citrinus with chloroform, CCP: Callistemon citrinus with petroleum ether**

**Table 2: Percentage yield of different fractions of CC extract**

| Fraction | % Yield |
|----------|---------|
| Petroleum ether (CCP) | 0.9 |
| Chloroform (CCC) | 6.5 |
| Alcohol (CCA) | 7.0 |

**Table 3: Preliminary phytochemical analysis of fractions of CC extract**

| Phytochemical | CCP | CCC | CCA |
|---------------|-----|-----|-----|
| Alkaloids      | -ve | +ve | -ve |
| Glicosides     | -ve | +ve | +ve |
| Flavonoids     | -ve | +ve | +ve |
| Carbohydrates  | -ve | -ve | +ve |
| Tannins        | -ve | +ve | +ve |
| Steroids       | -ve | +ve | -ve |
| Fats and oils  | +ve | -ve | -ve |

**CC: Callistemon citrinus, CCA: Callistemon citrinus with alcohol, CCC: Callistemon citrinus with chloroform, CCP: Callistemon citrinus with petroleum ether**

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Table 4: Effect of CC on immobility time in FST

| Groups          | Mean±SEM of immobility time (seconds) | % Decrease in immobility time |
|-----------------|-------------------------------------|------------------------------|
| Control         | 97.6±5.5                            |                              |
| Standard        | 22.5±2.17                           | 77                           |
| CCA 100 mg/kg   | 38.5±6.02                           | 60.5                         |
| CCA 200 mg/kg   | 26.5±3.14                           | 72.9                         |
| CCC 100 mg/kg   | 31.6±2.30                           | 67.7                         |
| CCC 200 mg/kg   | 23.2±2.51                           | 76.3                         |
| CCP 100 mg/kg   | 78.5±5.02                           | 20                           |
| CCP 200 mg/kg   | 40.6±4.24                           | 49.5                         |

Values are expressed as mean±SEM, *p<0.05, **p<0.01 versus control group. CCA: Callistemon citrinus; CCA: Callistemon citrinus with alcoholic; CCC: Callistemon citrinus with chloroform; CCP: Callistemon citrinus with petroleum ether; SEM: Standard error of mean, FST: Forced swim test

Table 5: Effect of CC on immobility time in TST

| Groups          | Mean±SEM of immobility time (seconds) | % Decrease in immobility time |
|-----------------|-------------------------------------|------------------------------|
| Control         | 184.3±80.01                         |                              |
| Standard        | 69.2±5.71                           | 73.4                         |
| CCA 100 mg/kg   | 114.4±7.0                           | 30                           |
| CCA 200 mg/kg   | 85.3±5.12                           | 54                           |
| CCC 100 mg/kg   | 111.6±5.77                          | 40                           |
| CCC 200 mg/kg   | 78.5±4.18                           | 57.5                         |
| CCP 100 mg/kg   | 165.26.21                           | 11                           |
| CCP 200 mg/kg   | 121.25.21                           | 34                           |

Values are expressed as mean±standard error of mean (SEM), *p<0.05, **p<0.01 versus control group. CCA: Callistemon citrinus; CCC: Callistemon citrinus with alcoholic; CCC: Callistemon citrinus with chloroform; CCP: Callistemon citrinus with petroleum ether. TST: Tail suspension test

In this study, we used the animal models TST and FST, which are widely accepted behavioral models for assessing antidepressant activity. Reduction in immobility time is the characteristic behavior scored in these tests, reflecting behavioral despair as seen in human depression. In addition, it is well known that conventional antidepressant drugs are able to reduce the immobility time in rodents [21]. This decrease in duration of immobility is considered to have a good predictive value in the evaluation of potential antidepressant agents. The results of different extracts of CC on the immobility duration in TST are presented in Table 4. Administration of CC alcoholic (CCA), CC chloroform (CCC), and CC petroleum ether (CCP) fractions of CC 200 mg/kg for 14 successive days decreased the immobility time in TST by 54%, 75.7%, and 34%, respectively, as compared to control group. CCA, CCC, and CCP 200 mg/kg administration for 14 days decreased immobility time in FST by 72.9%, 76.3%, and 49.5%, respectively as compared to control group. Standard drug imipramine 10 mg/kg on 14 days treatment decreased the immobility time by 73.4% and 77%, respectively, in TST and FST as compared to control group (Tables 4 and 5). A dose of 200 mg/kg p.o. CCC extract showed a potent antidepressant-like effect in both TST and FST as indicated by highest decrease in immobility period. The effect was comparable to a standard drug in FST.

CONCLUSION

In the present study, the preliminary phytochemical analysis revealed that the presence of tannins and flavonoids in hydroalcoholic fraction. Chloroform fraction showed positive results toward flavonoids, alkaloids, and steroids. Petroleum ether fraction showed positive results toward sterols and lipids; CCA and CCC showed a significant decrease in immobility time in both FST and TST indicating their antidepressant activity.

Chloroform fraction of CC showed antidepressant activity, which was comparable to a standard drug, i.e., imipramine (10 mg/kg). As reported earlier, CC contains many bioactive compounds and majority of these compounds are terpenoids and flavonoids that are responsible for the health benefits. Therefore, this study explores the use of CC in the treatment of depression. Further study can be explored to isolate the active constituents and evaluate the mechanism of antidepressant activity of CC.

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