ABSTRACT
The present investigation showed that the extracts of E. hirta Linn. displayed antibacterial activities. The powdered plant material of Euphorbia hirta was extracted in three solvents ethanol, benzene and water using the Soxhlet extraction apparatus 241 (PSAWINDIA). The percentage yields of the extracts were 58 (water), 32 (ethanol), and 15 (benzene). Phytochemical screening of the crude extracts revealed the presence of saponins, flavonoids, cardiac glycosides, cyanogenic glycosides, anthraquinones, and alkaloids. The presence of these constituents was linked to the antibacterial activity of the plant, using the agar well-diffusion method. The results suggested that the use of the extracts of E. hirta Linn. as an oral medication, both on short-term and long-term was not safe, due to the presence of some toxic constituents.

Keywords: Euphorbia hirta Linn., Ethanol extract, Benzene extract, phytochemical, water extract.

1. INTRODUCTION
Euphorbia hirta belongs to the family Euphorbiaceae. It is a small annual herb common to tropical countries [1]. The plant has been used for female disorders but is now more important in treating respiratory ailments, especially cough, coryza, bronchitis and asthma. In India it is used to treat worm infestations in children and for dysentery, gonorrhea, jaundice, pimples, digestive problems and tumors [2]. The plant is also widely used in Angola against diarrhea and dysentery, especially amoebic dysentery. In Nigeria extracts or exudates of the plant are used as ear drops and in the treatment of boils, sore and promoting wound healing [3, 4]. The alcoholic extract of Euphorbia hirta leaves is highly effective against gram positive bacteria and moderately effective against gram negative bacteria. This work was carried out to determine the in-vitro activities of Euphorbia hirta extracts on some clinical isolates.

2. MATERIALS AND METHODS
Collection of Plant Material
The fresh plant of E. hirta Linn. was collected from Diara land of Hajipur, Vaishali district in Bihar, India.

Preparation of Plant Material
The fresh plant was rinsed with tap water and air-dried under shade for a week, and then reduced to coarse powder using a pestle and mortar. Subsequently, it was ground into fine powder using the Kenwood electric blender (Kenwood Limited, Havant, United Kingdom). The powder was then stored in an airtight bottle for further use.

Preparation of the Extracts

Two hundred grams of the powdered sample (whole plant) was soaked in 100 ml of solvent in a Soxhlet extractor 241(PSAWINDIA) and left to stand for 24 hours, to extract. The filtrate was then evaporated to dryness using a rotary evaporator attached to a vacuum pump (Model type 349/2, Corning Limited). The percentage yield of the crude extract was determined for each solvent; for water it was 59%, for methanol 33%, and for benzene 16%. The percentage extract yield was estimated as dry weight/dry material weight × 100 [5]. For the preparation of the dilutions of crude extracts for antibacterial assay, the extracts was reconstituted by dissolving in the respective extracting solvents and further diluted with distilled water to obtain 400, 200, 100, 50, 25, 12.5, 6.25, 3.085, and 1.03 mg/ml. The reconstituted extracts were maintained at a temperature between 2 and 8°C.

Phytochemical Screening of the Plant Material

Phytochemical screening was carried out on the powdered plant material for the presence of bioactive components, such as, tannins, phenols, alkaloids, cardiac glycosides, anthraquinones, cyanogenic glycosides, saponins, and flavonoids [6].

3. RESULTS AND DISCUSSIONS

Percentage yield of the powdered plant *Euphorbia hirta* Linn. crude extracts, obtained using various solvents are shown Table 1. Out of the 200 g of powdered plant material, the percentage yield obtained for water was 58%, for ethanol 32%, and for benzene 15%. Phytochemical screening of the crude extracts of *E. hirta* Linn. revealed the presence of some bioactive components as shown in Table 2. It contained tannins, terpenes, cardiac glycosides, anthraquinones, saponins, cyanogenic glycosides, flavonoids, and alkaloids.

The percentage yield obtained for water was 58%, for ethanol 32%, and for benzene 15% (Table 1). This yield was not far lower than that obtained by Doughari et al. [7] for *Sennaangustifolia* as they reported a yield of 52% for water extract, 50% for benzene extract, and 28% for dichloromethane, and also that of Owolabi et al. [8] who reported a yield of 10.74% for water extract and 3.78% for their ethanol extracts. Ogbolie et al. [9] also reported a yield of 9.1% for water extracts of *E. hirta* Linn.. Factors like the age of the plant and the polarity of the solvent used affected the yield. Thus, in this study, water seemed to be the best solvent for this plant material, thus supporting the use of water as a solvent of choice in traditional practice.

The results obtained in this study support the methods used by the traditional healers. It is evident from the results that water extract has some significantly low antibacterial activity, suggesting that the active principles are more soluble in water, and that water is the appropriate solvent for the extraction of the bioactive constituents present in *E. hirta* Linn.[10, 11].
4. CONCLUSIONS

The present investigation showed that the extracts of *E. hirta Linn.* displayed antibacterial activities. However, long term use of the extract was slightly toxic in experimental animals, even though it was potent on the bacteria, which points to its toxicity in the vital organs such as liver, lungs, kidneys, spleen, and heart. It is therefore suggested that detoxification of the constituents responsible for its toxicity be researched upon for effective ethno-medicinal prescriptions of the plant.

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Table 1: Percentage yields of extracts of *E. hirta* Linn.

| Extraction Solvent | Yields (%) |
|--------------------|-----------|
| Water              | 58.0      |
| Ethanol            | 32.0      |
| Benzene            | 15.0      |

Table 2: Phytochemical screening and chromatographic resolution of crude extracts

| Class of compounds | RF values of spots |
|--------------------|--------------------|
|                    | Aqueous | Ethanol | Benzene |
| Alkaloids          | 0.51    | 0.79    | -       |
| Flavonoids.        | 0.72    | 0.82    | 0.81    |
| Saponins           | 0.79    | 0.63    | 0.76    |
| Tannins            | 0.83    | 0.88    | 0.84    |
| Anthraquinones     | 0.89    | 0.61    | -       |
| Cyanogenicgly      | 0.70    | 0.65    | 0.63    |
| Terpenes           | 0.88    | 0.62    | 0.66    |
| Cardiac gly        | 0.80    | -       | -       |