# Phytochemical investigation and pharmacological screening of leaves of *Achyranthus aspera* Linn. as analgesic and antipyretic

#### N.G. SUTAR\*, U.N. SUTAR, Y.P. SHARMA., I.K. SHAIKH and S.S. KSHIRSAGAR

S.N.D. College of Pharmacy, Babhulaon, Tal-Yeola, District Nashik (India).

(Received: July 20, 2008; Accepted: October 21, 2008)

### ABSTRACT

Achyranthus aspera Linn. leaves have antimicrobial, anti-inflammatory, venereal affection, analgesic, antipyretic activity, in treatment of eyes, arthritis as antiirlityetc. Methanolic extract of leaves of *Achyranthus aspera* Linn. reported as analgesic activity but leaves till not reported, the effect of alcoholic extract and its various fraction as Petroleum ether, Ethyl acetate, Diethyl ether, n-Butanol were tested for phytochemically which contain glycosides, flavonoids, tannins, amino acids. As analgesic activity by using hot plate method and antipyretic by using brewers yeast induced method and compare with Aspirin as a standard in a dose of [25mg/kg] for analgesic and [125mg/kg] for antipyretic effect.

Key words: Leavess of *Achyranthus aspera* Linn., Analgesic activity by hot plate, and Antipyretic activity by Brewers yeast suspension, Aspirin.

#### INTRODUCTION

Achyranthus aspera Linn. [Amaranthaceae] is grown in India, in sub tropical parts of India. Western Himalayas. It is known as Prickly chaff flower in English, Chirchira in Hindi, Aghada in Marathi<sup>1</sup>. Dried leaves used in venereal affection, anti-inflammatory, purgative<sup>2</sup>, as tonic, The methanolic extract of leaves of Achyranthus aspera Linn.. reported to have analgesic activity but leaves till not reported the effect of alcoholic extract and its various fraction as Petroleum ether, Ethyl acetate, Diethyl ether, n-Butanol The aim of study was to screen effect of leaves of *Achyranthus aspera* Linn. as Analgesic and Antipyretic<sup>4</sup>.

#### MATERIAL AND METHODS

The fresh leaves of *Achyranthus aspera* Linn. were collected. They were shade dried and ground to obtained coarse particles size. The powdered material was extracted with (95%) alcohol in a continuous hot extractor at 40°-50°c temps. Some part of the extract was kept aside and remaining was fractionated with Pet. ether, Ethyl acetate, Diethyl ether, and n-Butanol what ever the fraction collected was wash with water air dry and kept separately with Na<sub>2</sub>So<sub>3</sub> ad dehydrating agent.<sup>5</sup> Qualitative test were performed for the alcohol extract and its fractions alcohol extract showed the presence of glycosides, amino acids, and sterols. Pet. Ether extract showed presence of fats and oils <sup>6,7</sup>

# Evaluation of analgesic and antipyretic activity Hot plate method

In this method Wister male albino rats (180-200gm) were used for the study. The animals were segregated into seven groups of six animals each.

Group 1 - Normal saline solution,

Group 2 - Aspirin as standard (25mg/kg),

|                                  |   | Table<br>fracti                           | <ul> <li>1: Showing resources</li> <li>ons of leaves of</li> </ul> | ult of analgesic<br>Achyranthus a       | Table 1: Showing result of analgesic effect of Alcohol extract and its fractions of leaves of <i>Achyranthus aspera</i> Linn. on Hot plate method | ol extract and it<br>Hot plate metho | qv                            |                            |
|----------------------------------|---|---|--|---|---|--------------------------------------|-------------------------------|----------------------------|
| S.<br>No.                        | Time Co<br>in min (ve   | Control As<br>(vehicle)                   | Aspirin Alc<br>ext   | Alcohol Pe<br>extract ext               | Pet.ether So<br>extract fra   | Solvent ether<br>fraction            | Ethy lacetate<br>fraction     | n-Buthanol<br>fraction     |
| -                                | 0 2.0   | 2.00±0.15 2.80                            | 15   | 2.6±0.18 2.4                            | 2.4±0.21 2.5  | 2.3±0.17<br>/> >>>/                  | 2.4±0.21                      | 2.4±0.13                   |
| 0                                | 20 2.2  | 2.25±0.11 7.2                             | m  | 35                                      | 00  | (2.22 %)<br>4.30±0.60                | 5.00±1.23                     | 6.20±0.70                  |
| ო                                | 60 1.5  | (28.<br>1.50±0.07 8.19                    | (28.38%) (28.38%)<br>8.19±0.74 7.9                                 | (25.63%) (25<br>7.99±0.22 6.4           | (23.84%) (1 <sup>-</sup><br>6.49±0.60 6.1   | (11.54%)<br>6.14±0.55                | (15.49%)<br>5.30±1.25         | (22.25%)<br>7.49±0.71      |
| 4                                | 90 1.5  | (36.<br>1.50±0.22 10.9                    | (36.21%) (35<br>10.9±1.16 8.9                                      | (35.13%) (27<br>8.93±0.63 7.5           | (27.02%) (29<br>7.58±0.61 7.1   | (25.13%)<br>7.17±0.70                | (20.54%)<br>5.40±1.07         | (32.43%)<br>7.46±1.14      |
|                                  |   |   | (50.8%) (40  | (40.17%) (34                            | (34.32%) (30  | (30.65%)                             | (21.08%)                      | (34.25%)                   |
| Group                            |   | Rect                                      | Rectal Temp °C   |   | Time after  | Time after medication in min         | min                           |                            |
|                                  |   | Initial                                   | 18 hr after<br>Yeast injection                                     | 30 min                                  | 60min   | 90min                                | 120min                        | 180min                     |
| Control<br>Asnirin 1             | Control<br>Asnirin 150 ma/ka  | 38.30±0.031<br>38.30+0.073                | 39.35±0.025<br>39.35±0.027   | 39.30± 0.015<br>38.37+ 0.025            | 39.23± 0.025<br>38.09+ 0.015  | 5 39.20± 003<br>5 37.67+0.053        | 39.15±0.038<br>3 37.58+ 0.058 | 39.00±0.02<br>8 37.40+0.47 |
| Alcohol<br>Pet ethe              | Alcohol 30mg/kg<br>Pet ether 100ma/ka                                 | 38.28±0.022<br>38.25+0.058                | 39.37±0.038  | 38.68+ 0.051                            | 37.87± 0.065<br>38.00+ 0.065  |                                      |                               |                            |
| Solvent<br>Ethyl eta<br>n-Butane | Solvent ether300mg/kg<br>Ethyl etate. 300 mg/kg<br>n-Butanol 300mg/kg | 38.27±0.065<br>38.24±0.080<br>38.26±0.065 | 39.40±0.012<br>39.38±0.10<br>39.40±0.051                           | 38.80±0.17<br>38.67±0.13<br>38.75± 0.11 | 38.60± 0.11<br>38.50± 0.069<br>38.61± 0.060   |                                      |                               |                            |
|                                  |   |   |  |   |   |                                      |                               |                            |

Sutar et al., Biosci., Biotech. Res. Asia, Vol. 5(2), 841-844 (2008)

842

- Group 3 Alcohol extract (30mg/kg)
- Group 4 Pet ether fraction (100mg/kg),
- Group 5 Ethyl acetate fraction (300mg/kg),
- Group 6 Diethyl ether fraction (300mg/kg),
- Group 7 -n-Butanol fraction (300mg/kg).

The dried extract and its fraction were formulated as a suspension in distilled water. Alcoholic extract and its various fraction were administered orally using intragastic tube. The pain threshold (No. of liking of paw/jumping) were measured at 20, 60, 90 min after administration of standard and test solution <sup>6,710.</sup>

#### Antipyretic activity

In this method Wister male albino rats (180-200gm) were used for the study. The animals were segregated into seven groups of six animals each. The standard and test group were fevered by brewers yeast suspension in propylene glycol [15%] at a dose of 10 ml/kg.

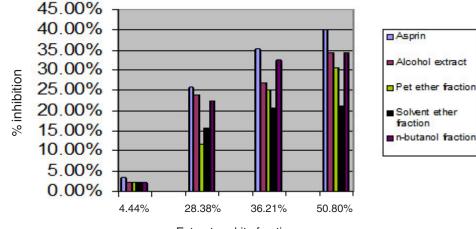
- Group 1 Normal saline solution, Group 2 - Aspirin as standard (25mg/kg), Group 3 - Alcohol extract (30mg/kg) Group 4 - Pet ether fraction (100mg/kg), Group 5 - Ethyl acetate fraction (300mg/kg),
- Group 6 Diethyl ether fraction (300mg/kg),
- Group 7 -n-Butanol fraction (300mg/kg).

The dried alcoholic extract and its various fractions were formulated as a suspension in distilled water. Alcoholic extract and its various fractions were administered orally using intragastic tube. The rectal temperatures were measured at 30, 60, 90, 120 and 180 min. after administration of standard and test solution<sup>11</sup>.

#### RESULTS

## Analgesic activity and Antipyretic activity

Amongst alcoholic extract and its various fractions of leaves of *Achyranthus aspera* Linn.



Extract and its fraction

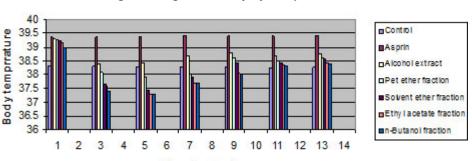


Fig. 1: Analgesic activity by Hot plate method

Fig. 2: Antipyretic activity by Brewers yeast induced supension method

alcoholic extract showed potent analgesic and antipyretic activity which is comparable to that of aspirin at the dose of 25mg/kg. For analgesic and 125mg/kg. for antipyretic activity the peak effect in alcohol extract and fraction of leaves of *Achyranthus aspera* Linn... were seen after 1h.in case of analgesic and 0.5h.in antipyretic treatment This could be due to more availability of saponin as active principle's in leaves.

#### DISCUSSION

From this study, it can be concluded that the leaves of *Achyranthus aspera* Linn.. possesses marked analgesic and antipyretic activity and is equipotent to standard drugs. The present study establishes effectiveness and pharmacological screening, rational for use of leaves of *Achyranthus aspera* Linn. in folklore medicine as analgesic drug. Thus the plant can be further explored for its phytochemical profile to identify the active constituent saponin is responsible for the above mention activities.

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844