



## International Journal of Research and Development in Pharmacy and Life Sciences



Rabbits were injected yeast at the dose of 0.5 ml/kg body weight, to induce pyrexia. Induction of fever was taken about one to two hours. (Grover, 1990)  
Negative control receiving 10ml distill water,

channel blocking agent verapamil (5mg/kg) and standard drug Aspirin(100 mg/kg) (Table 3). The extract reduced  $37.41 \pm 0.22$   $^{\circ}\text{C}$  of elevated rectal temperature compared to verapamil  $37.26 \pm 0.15$   $^{\circ}\text{C}$  and aspirin  $37.34 \pm 0.16$   $^{\circ}\text{C}$  after 3 hours in 2,4 dinitrophenol induced pyrexia rabbits while the mixture of verapamil and the extract (2mg/kg+50mg/kg) showed significant reduction in the temperature  $37.35 \pm 0.22$   $^{\circ}\text{C}$  as shown in Table 3.

Effect of Methanolic crude bark extract of plumeria rubra on E-coli induced pyrexia in rabbits.

The methanolic extract produced significant ( $P < 0.05$ ) antipyretic effect in E-coli induced pyretic rabbits. At a dose of 100 mg/kg body weight, plumeria rubra reduced  $(37.28 \pm 0.08$   $^{\circ}\text{C}$ ) of elevated rectal temperate compared to aspirin  $(37.34 \pm 0.12$   $^{\circ}\text{C}$ ) and ciprofloxacin  $(37.31 \pm 0.30$   $^{\circ}\text{C}$ ), while the combination of both extract and ciprofloxacin reduced the rectal temperature  $(37.22 \pm 0.16$   $^{\circ}\text{C}$

Table 2. Antipyretic effect of Pr.Cr on yeast induced pyrexia.

Dose

8. Surendra Kr. Sharma and Naresh Kumar (2012). Antimicrobial potential of Plumeria rubrabark Der PharmaChemica, 4(4):1591-1593
9. Al-Ghamdi MS (2001). The anti-inflammatory, analgesic and antipyretic activity of Nigella sativa. Journal of Ethnopharmacology. 76: 45 – 48.
- 10.

REFERENCES:

1. Wiart C (2002) Medicinal Plants of Southeast Asia, Kuala Lumpur; Pearson Malaysia Sdn. Bhd ;, pp 262.
2. Bobbarala V, Katikala PK, Naidu KC, Penumajji S (2009). Antifungal activity of selected plant extracts against phytopathogenic fungi Aspergillus niger F2723. Indian Journal of Science and Technology2(4): 87-90.
3. Williamson EM, Okpako DT, Evans FJ (1998). Selection, Preparation and Pharmacological Evaluation of Plant Material. John Wiley & Sons, Chichester, pp. 15–23.
4. Grover JK (1990). Experiments in Pharmacy and Pharmacology. 1st ed., Vol. 2, India, pp: 155.
5. Blackhouse N, Delporte C, Negrete R, Munoz O, Ruiz R. (1994). Anti inflammatory and antipyretic activities of Maytenusboaria. International Journal of Pharmacognosy 32: 239 -244.
6. Dardi MS, Sharma SK, Srivastava AK. (2005). Pharmacokinetics and dosage regimen of ceftriaxone in E. coli lipopolysaccharide induced fever in buffalo calves. J Vet Sci. 6(2):147-50.
7. Kirtikar KR, Basu BD (1935). Indian medicinal plants. 3rd edition, part II, International Book Distributors, Dehradun, pp. 1561-1564.