

Analgesic Activity of Ethanol Extract of Temu Giring Rhizome (*Curcuma heyneana* Val & Zijp) in Mice

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Abstract. Temu giring (*Curcuma heyneana* Val & Zijp) is a traditional medicinal plant that is believed in community as an analgesic. The objective of this research was to determine the analgesic activity of the *C. heyneana* rhizome by using infra red (IR) thermal induction method in mice. Mice were divided into 7 groups. Group 1 served as negative control, group 2,3,4,5 served as treatment groups which were given ethanol extract of *C. heyneana* rhizome at dose of 5, 25, 125, and 625 mg/kg respectively, group 6 and 7 served as comparable groups, given antalgin 65 mg/kg and morphine sulphate 1.3 mg/kg, respectively. The observation has been done, pain stimulus of mice by exposing to infra red (IR) every 10 minutes for 80 minutes. The data were analyzed by ANOVA at the significance level of 95%. Ethanol extract of *C. heyneana* at the doses of 25, 125, and 625 mg/kg had significant effect to reduce the pain compared to the negative control ($p < 0.05$). Ethanol extract of *C. heyneana* rhizome at dose of 125 mg/kg, had the same effect to antalgin 65 mg/kg ($p \geq 0.05$), while the ethanol extract of *C. heyneana* at the dose of 625 mg/kg had the same effect as morphine sulfate 1.3 mg/kg ($p \geq 0.05$). It can be concluded that ethanol extract of *C. heyneana* rhizome has analgesic activity.

Keywords: temu giring, analgesic, *Curcuma heyneana*, rhizome

Abstrak. Temu giring (*Curcuma heyneana* Val & Zijp) adalah tanaman obat tradisional yang diyakini berkhasiat sebagai penghilang rasa nyeri. Tujuan dari penelitian ini adalah untuk membuktikan aktivitas analgesik dari rimpang *C. heyneana* dengan menggunakan metode induksi termal infra merah (IR). Mencit dibagi menjadi 7 kelompok. Kelompok 1 adalah kontrol negatif, kelompok 2, 3, 4 adalah kelompok perlakuan yang diberikan ekstrak etanol rimpang *C. heyneana* dengan dosis 5, 25, 125, dan 625 mg/kg, kelompok 5 dan 6 adalah kelompok pembanding yang diberikan antalgin 65 mg/kg serta morfin sulfat 1,3 mg/kg. Pengamatan meliputi daya tahan mencit untuk menahan nyeri yang ditimbulkan oleh paparan infra merah (IR) pada telapak kaki yang diinduksi setiap 10 menit dan berlangsung selama 80 menit. Data dianalisis dengan menggunakan ANOVA pada tingkat signifikansi 95%. Ekstrak etanol *C. heyneana* pada dosis 25, 125, dan 625 mg/kg memiliki efek yang signifikan untuk mengurangi nyeri dibandingkan dengan kontrol negatif ($p < 0,05$). Pada dosis 125 mg/kg, ekstrak etanol rimpang *C. heyneana* memiliki efek yang sama seperti antalgin 65 mg/kg ($p \geq 0,05$). Sedangkan ekstrak etanol *C. heyneana* pada dosis 625 mg/kg memiliki efek yang sama seperti morfin sulfat 1,3 mg/kg ($p \geq 0,05$). Dapat disimpulkan bahwa ekstrak etanol rimpang *C. heyneana* dosis 25, 125 dan 625 mg/kg memiliki aktivitas analgesik.

Kata kunci: temu giring, analgesik, *Curcuma heyneana*, rimpang

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1. Introduction

NSAIDs or nonsteroidal anti-inflammatory drugs are the most common medicine as pain relief in the community. It is used to relieve symptoms of arthritis, sprains, headaches and other daily discomforts [1]. Long periode of NSAIDs utilization could increase its side effect such as gastrointestinal irritation, bleeding, blurred vision, and decreased kidney function [2]. Utilization of paracetamol in the long period could induce hypersensitivity reactions, blood disorders, and liver damage [3]. Opioid analgesics also have the potential effect in tolerance, dependence and addiction [4].

Herbal medicine treatment is currently popular as an alternative treatment to heal various disease including as pain reliefer. One of the herbal medicine that widely used by Indonesian people is temu giring (*Curcuma heyneana* Val & Zijp). Empirically, local people use temu giring rhizome to treat various disease such as rheumatism, irregular menstruation, stomach aches, fever and as anthelmintic. Temu giring rhizome has various phytochemical properties of essential oils, starch, resin, fat, tannins, saponins, and flavonoids [5]. It is necessary to conduct research in determining analgesic activity of ethanol extract of temu giring (EETG). The purpose of this study was to determine the analgesic activity of the ethanol extract of temu giring on male mice using a infra red plantar test.

2. Material and Method

2.1. Materials

Temu giring rhizome was obtained from PT. Sumatera Busan, Deli Serdang, North Sumatra, Indonesia. Sodium Carboxymethyl Cellulose, distilled water were obtained from PT. Bratachem, Indonesia . Morphine sulfate and antalgin were obtained from PT. Kimia Farma, Indonesia.

2.3. Animals

Male mice of 3 months old were obtained from the Animal House of Faculty of Pharmacy Universitas Sumatera Utara. Mice were acclimatized for 2 weeks under standard laboratory conditions at room temperature condition 25-30°C. The mice were fed with standard laboratory diet and allowed to drink water. The studies were carried out in accordance with the institutional ethical guidelines.

2.4. Research Methods

The research had obtained approval ethical clearance with No. 785 / KEPH-FMIPA / 2016. The extract was obtained by maceration method using ethanol 96%. The analgesic test used a plantar test with infra red (IR). Mice were devided into 7 groups. Group 1 served as negative control,

group 2, 3, 4, 5 served as treatment groups which were given ethanol extract of temu giring rhizome at dose of 5, 25, 125, and 625 mg/kg respectively, group 6 and 7 served as comparable groups, given antalgin 65 mg/kg and morphine sulphate 1.3 mg/kg, respectively. The observation had been done, included the pain stimulus of mice which exposed by infra red (IR) every 10 minutes for 80 minutes.

2.5. Statistical Analysis

Analysis of all results were performed using ANOVA. *P* values for significance were set at 0.05.

3. Result and Discussion

The results of analgesic activity of ethanol extract of temu giring rhizome by the infra red (IR) plantar test method can be seen in Figure 1.

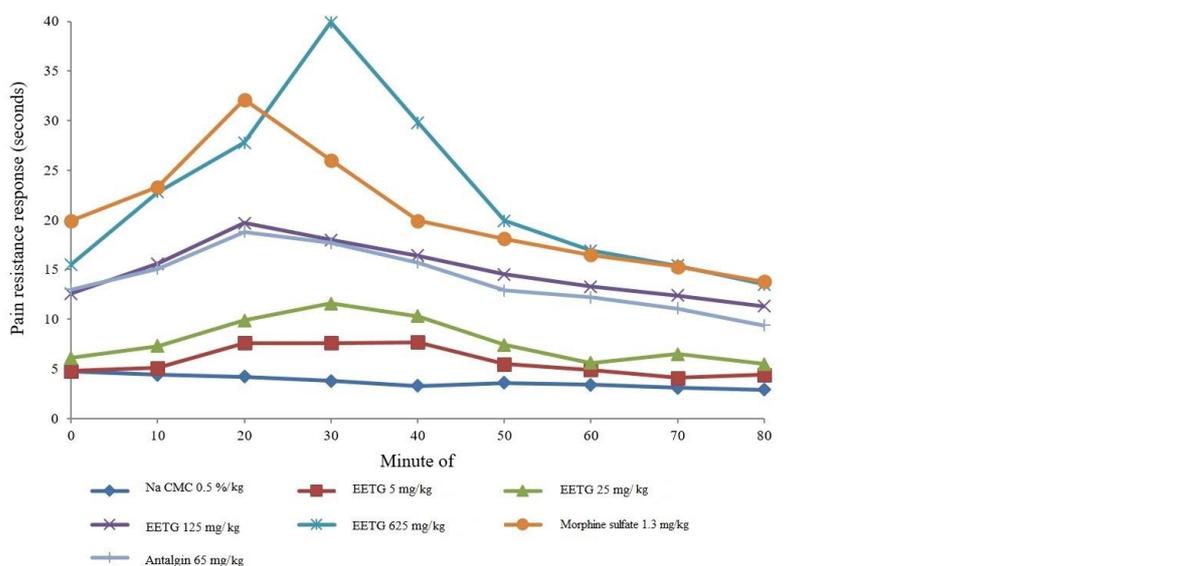


Figure 1. Analgesic effect ethanol extract of temu giring rhizome on mice induced by infra red (IR).

Figure 1 shows that negative control which only given a vesicle (sodium carboxymethyl cellulose), there was no analgesic effect when it exposed by infrared for 5 seconds, the mice appeared raise their legs. Ethanol extract at dose of 5 mg/kg showed no analgesic effect. Based on observations the mice were not able to resist pain induction, the condition of mice were same as to the negative controls. Dose of 25 mg/kg, showed ability as pain reliefer to infra red compare to negative control group ($p < 0.05$). However, the pain relief effect was not as strong as the antalgin and morphine groups ($p < 0.05$).

Dose of 125 mg/kg were able as pain reliefer due to infra red induction, it had longer activity than the effect caused by dose of 25 mg/kg and showed similar effects with antalgin ($p > 0.05$) but not to morphine ($p < 0.05$). Mice given a dose of 625 mg/kg showed the strongest analgesic effect and comparable with the analgesic effects of morphine ($p \geq 0.05$). The plantar test method

(Hargreaves method) was an appropriate model to determine peripheral response of thermal stimuli in animals.

The results showed that animals that had been given ethanol extract of temu giring rhizome were able to resist pain induction longer than animals that were not given an extract. Temu giring rhizome has secondary metabolite content of flavonoids. Previous research stated that flavonoids content in plants had an analgesic effect in mice [6,7]. Other studies stated that flavonoids were known to inhibit the formation of cyclooxygenase (COX) or lipoxygenase [8]. Temu giring rhizome also had curcuminoid content which were composed of curcumin, demethoxycurcumin and bismethoxycurcumin. Curcumin also had an effect in inhibiting the metabolism of arachidonic acid, cyclooxygenase, lipoxygenase and cytokines [9]. It was stated that curcumin significantly inhibited mRNA expression and COX-2, but not in COX-1 [9]. Cyclooxygenase (COX) is known as substances of pain mediators, phytochemical properties of temu giring could inhibit cyclooxygenase (COX) and it will reduce pain. In addition, curcumin also has a high selectivity to COX-2 which has lower side effect on gastrointestinal than non selective analgesics drug.

4. Conclusion

It can be concluded that the ethanol extract of temu giring rhizome at dose of 25, 125 and 625 mg/kg have analgesic effect.

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