

Uji potensi hepatoprotektif madu ps pollen substitute terhadap gambaran histologis hati mencit mus musculus l jantan galur ddy = Hepatoprotective potency study of ps pollen substitute honey on liver histology of ddy strain male mice mus musculus l

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Abstrak

Penelitian telah dilakukan untuk mengetahui potensi hepatoprotektif madu PS (Pollen Substitute) terhadap gambaran histologis hati mencit (*Mus musculus*) jantan galur DDY. Dua puluh empat ekor mencit jantan dibagi ke dalam 4 kelompok, yaitu kelompok kontrol normal (KK1) yang diberikan akuades dan minyak kelapa; kelompok kontrol perlakuan (KK2) yang diberikan akuades dan CCl₄; serta 2 kelompok perlakuan (KP1 dan KP2) yang diberikan madu PS dosis 0,04 ml/20 g bb dan 0,08 ml/20 g bb selama 14 hari berturut-turut, kemudian diinjeksikan CCl₄ 2 jam setelah pemberian madu PS terakhir. Organ hati diisolasi 24 jam setelah injeksi CCl₄. Hasil uji statistik menunjukkan tidak ada pengaruh pemberian madu PS terhadap berat basah organ hati (KK1 [1,57±0,34] gram, KK2 [1,92±0,21] gram, KP1 [1,73±0,34] gram, KP2 [1,75±0,30] gram) dan diameter vena sentralis (KK1 [37,91±2,44] μm, KK2 [73,39±3,06] μm, KP1 [70,03±9,65] μm, KP2 [67,61±6,33] μm), serta terdapat pengaruh pemberian madu PS terhadap warna organ hati, persentase derajat kerusakan lobulus hati (derajat 0, 1, 2, dan 3 KK1 [55,70%; 40,00%; 4,33%; dan 0,00%], KK2 [4,56%; 29,00%; 18,67%; dan 47,78%], KP1 [14,22%; 43,11%; 15,22%; dan 27,44%], KP2 [6,34%; 41,33%; 24,67%; dan 27,67%]), dan gambaran histologis hati. Hasil pengamatan menunjukkan pemberian madu PS dosis 0,04 ml/20 g bb memberikan efek hepatoprotektif yang lebih baik dibandingkan madu PS dosis 0,08 ml/20 g bb terhadap gambaran histologis hati.

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The aim of this study was to know hepatoprotective potency of PS (Pollen Substitute) honey to liver histology of DDY strain male mice (*Mus musculus*). Twenty four male mice were divided into 4 groups, which were normal control group (KK1) which was given aquadest and palm oil, treatment control group (KK2) which was given aquadest and CCl₄, as well as 2 treatment groups (KP1 and KP2) which were given PS honey dose 0,04 ml/20 g bb and 0,08 ml/20 g bb in 14 days. Then KP1 and KP2 were induced by CCl₄ 2 hours after the last administration of honey PS. The liver then was isolated 24 hours after CCl₄ injection. The result of statistic test showed that there were no effects of PS honey administration to liver wet weight (KK1 [1,57±0,34] gram, KK2 [1,92±0,21] gram, KP1 [1,73±0,34] gram; KP2 [1,75±0,30] gram) and central vein diameter (KK1 [37,91±2,44] μm, KK2 [73,39±3,06] μm, KP1 [70,03±9,65] μm, KP2 [67,61±6,33] μm), and there were effects of PS honey administration to liver color, percentage of liver lobulus damage level (level 0, 1, 2, and 3 of KK1 [55,70%; 40,00%; 4,33%; and 0,00%], KK2 [4,56%; 29,00%; 18,67%; and 47,78%], KP1 [14,22%; 43,11%; 15,22%; and 27,44%], KP2 [6,34%; 41,33%; 24,67%; and 27,67%]), and liver histology. The result of the observation showed that the hepatoprotective effect of administration of PS honey dose 0,04 ml/20 g bw was better than dose 0,08 ml/20 g to liver histology.;