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 - *Aloe vera* extract reduces 8-oxo-2'-deoxyguanosine levels and improves total antioxidants in streptozotocin-induced diabetic rats
- Blood cadmium levels increase prostate specific antigen and insulin-like growth factor-1 among cadmium exposed workers
- Increase in neutrophil count after repeated exposure of *Plasmodium berghei*-infected mice to artemisinin
 - Contact with poultry and animals increases risk of *Campylobacter* infections in adults of Ardabil province, Iran
- Combination of five clinical data as prognostic factors of mortality after ischemic stroke

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Abstract / Indexing



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Campbell NRC, Gilbert RE, Leiter LA, et al. Hypertension in people with type 2 diabetes: update on pharmacologic management. *Can Fam Physician* 2011;57:997-1002.

2. Corporate author

Diabetes Prevention Program Research Group. Ten year follow-up of diabetes incidence and weight loss in the Diabetes Prevention Program Outcomes Study. *Lancet* 2009;374:1677-86. doi:10.1016/S0140-6736(09)61457-4.

3. Volume with supplement

Maeshiro R, Koo D, Keck CW. Patients and populations: public health in medical education. *Am J Prev Med* 2011; 41(Suppl 3):S145-S318. DOI: <http://dx.doi.org/10.1016/j.amepre.2011.07.010>.

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Santos CAST, Fiaccone RL, Oliveira NF, et al. Estimating adjusted prevalence ratio in clustered cross-sectional epidemiological data. *BMC Med Res Method* 2008;8:80. doi:10.1186/1471-2288-8-80.

Books and Other Monographs

1. Editor(s), compiler(s) as author

Gilstrap LC, Cunningham FG, VanDorsten JP, editors. Operative obstetrics. 4th ed. New York: McGraw-Hill;2010.

2. Chapter in a book

Meltzer PS, Kallioniemi A, Trent JM. Chromosome alterations in human solid tumors. In: Vogelstein B, Kinzler KW, editors. The genetic basis of human cancer. New York: McGraw-Hill;2010.p.93-113.

3. Conference paper

Christensen S, Oppacher F. An analysis of Koza's computational effort statistic for genetic programming. In: Foster JA, Lutton E, Miller J, et al. editors. Genetic programming. EuroGP 2002: Proceedings of the 5th European Conference on Genetic Programming; 2002 Apr 3-5; Kinsdale, Ireland. Berlin: Springer;2002.p.182-91.

4. Dissertation

Hos J. Mechanochemically synthesized nanomaterials for intermediate temperature solid oxide fuel cell membranes [dissertation]. Crawley, Western Australia: University of Western Australia;2005.

Electronic Material

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Murray G. A duty of care to children and young people in Western Australia: Report on the quality assurance and review of unsubstantiated allegations of abuse in care;2005. Available at: <http://www.community.wa.gov.au/NR/rdonlyres/851183A4-A822-4592-AB66-C1etsC410E453AEEC/0/DCDRPTGwennMurrayreportwithcover2006.pdf> Accessed April 12, 2008.

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Sillick T J, Schutte NS. Emotional intelligence and self-esteem mediate between perceived early parental love and adult happiness. *Applied Psychol* 2006;2:38-48. Available at: <http://ojs.lib.swin.edu.au/index.php/ejap/article/view/71/100>. Accessed June 10, 2010.

3. Monograph on the internet

Foley KM, Gelband H, editors. Improving palliative care for cancer [monograph on the Internet]. Washington: National Academy Press; 2008. Available at: <http://www.nap.edu/books/0309074029/html/>. Accessed July 9, 2010.

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ORIGINAL ARTICLE

Increase in neutrophil count after repeated exposure of *Plasmodium berghei*-infected mice to artemisinin

Lilik Maslachah* and Rahmi Sugihartuti*

ABSTRACT

BACKGROUND

Leukocytes play an important role in the elimination of malaria infection. The leukocyte profile upon elimination of the malaria parasites that have been exposed to antimalarials and are subsequently capable of faster growth has not been researched. The aim of this research was to evaluate the role of mouse leukocytes in the elimination of parasites as shown by the leukocyte profile.

METHODS

An experimental research with post test only control group design was conducted involving 24 male mice of the Swiss Albino strain weighing 20 g -30 g, and 2.5 months old. They were randomized into four groups: two control groups (K1, KP) and two treatment groups (P1, P4). Artemisinin at a dose of 0.04 mg/g body weight was given to the mice for 3 days, starting 2 days after infection. The leukocyte profile was observed on the 2nd, 5th, 8th, and 10th day after infection. The results were analyzed by two-way Anova.

RESULTS

As shown in treatment control group KP and treatment group P4, *P. berghei* that had been passaged in the mice and were still viable after repeated exposure to artemisinin, may cause changes in leukocyte profile. On the 10th day of infection, the neutrophil percentage in group P1 showed a significantly different decrease when compared with the other groups (K1, KP and P4) ($p < 0.05$).

CONCLUSION

Repeated exposure to artemisinin of mice infected with *P. berghei* can cause changes in neutrophil profile in mice .

Keywords: Artemisinin, leukocytes, repeated passage, *P. berghei*

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INTRODUCTION

Malaria is still a public health problem in more than 95 countries around the world including Indonesia. In 2015 there were approximately 214 million cases of malaria with 438.000 deaths, mostly occurring in African children under 5 years of age.⁽¹⁾

The increased incidence of morbidity and mortality of malaria is rapid and widespread due to increased parasite resistance to antimalarial medications. The new drugs for malaria treatment which are used to date are artemisinin and its derivatives. The effect of these drugs is faster than that of other anti-malarial drugs because they have more complex mechanisms of action. However, there have been indications that the Plasmodium parasite has become resistant to these drugs.⁽²⁾ This was shown clinically in two patients in Cambodia who were infected with *P. falciparum*, in whom the parasites had become resistant to artesunate.⁽³⁾ The research results showed that there has been decreasing efficacy of the combination of artesunate-mefloquine in falciparum malaria in Cambodia.⁽⁴⁾ Currently, the resistance of *P. falciparum* to artemisinin has been detected to have occurred in five countries including Cambodia, Lao People's Democratic Republic, Myanmar, Thailand and Vietnam.⁽¹⁾

The results of an in vitro study of *P. falciparum* that were repeatedly exposed to artemisinin showed increased inhibitory concentration 50 % (IC_{50}), dormant phenotypic changes, and faster growth of viable Plasmodium evolving from dormant forms. Moreover, artemisinin exposure also can cause mutations in the *pfatpase6* gene.⁽⁵⁾ It is believed that the artemisinin resistance occurs because of mutations in *pfert*, *pfmdr1*, *pfatpase6* and *pfk13*.^(6,7)

The presence of parasite pressure on the use of medicines at subcurative doses will lead to the development of new parasite strains that can survive the medicine. The results of this research are to be viewed as indicating an emergency, because there will be a risk of the

development of in vivo resistance in humans, which will be one of the health problems in the world. Moreover, there is no substitute for artemisinin. Malaria treatment failure with artemisinin and its derivatives will open an era of untreatable malaria.

Experimental in vivo studies using rodent malaria are used to support the translation of laboratory studies into clinical studies. Since the pathologic mechanism for the spectrum of malaria in humans is not known clearly, these experimental studies can be used to explain the mechanism of artemisinin resistance in vivo by using mice that are infected with *P. berghei*. Because of the resilience and virulence of the malaria parasites in antimalarial resistance, there is a need to do research to develop effective strategies for malaria control. However, it is really difficult to do research in endemic areas because there are many confounding factors, such as the cloning of multiple infections from infective mosquito bites, which is impossible to do in humans because of ethical factors. So that this research is using rodent malaria as an in vivo drug resistance model of human malaria by exposing *P. berghei* to artemisinin. The effective dose of artemisinin that is used is 99% (ED_{99} : 200 mg/kg body weight of the mice), and is given repeatedly to the mice upon parasite passage.

P. berghei is one of the species of *Plasmodium* that infect rodents. Many studies of malaria use *P. berghei* with mice as hosts, because *P. berghei* has the same life cycle and morphology as the *Plasmodium* parasites that infect humans. Molecular analysis show that there are similarities between rodent malaria and *P. falciparum* malaria. In order to support in vivo studies on *P. falciparum* drug resistance, *P. berghei* can be used as a model.⁽⁸⁾

Results of research by Darlina et al.^(9,10) showed that leukocytes play a role in the protective mechanism during malaria infection, as is indicated by the increasing numbers of leukocytes, lymphocytes, and monocytes and decreasing rates of parasitemia in mice that have been infected with irradiated *P. berghei*.

One study showed that artemether, artesunate and artemisinin markedly decreased the leukocyte count after 24 h of treatment. Artemether and artesunate showed the highest efficacy in rescuing mice with late-stage cerebral malaria and rapidly decreased leukocyte accumulation in the brain.⁽¹¹⁾ The results of the present study can give an idea of the leukocyte profile in hosts infected with plasmodium that have been repeatedly exposed to artemisinin. This is not possible in humans because of ethical considerations, therefore the results of the animal studies can be used for improvements in the treatment of malaria infection.

Leukocytes are part of the blood that are vital to the body defenses against infection. The objective of the present study was to evaluate the effect of artemisinin on leukocyte profile in mice infected with *P. berghei*, so that this research was conducted to determine the changes in leukocyte profile during the development of artemisinin resistance in mice infected with *P. berghei* upon passage of *P. berghei* that had been exposed repeatedly to antimalarial drugs,

METHODS

Research design

This research was an experimental research with post test only control group design. The parasite that was used to infect the mice was *P. berghei* ANKA strain. The experimental animals were 2.5 months old male Swiss albino mice weighing 20g -30 g. The artemisinin was of analytical grade from Sigma Chemical Co. Leukocyte examinations were conducted at the Faculty of Veterinary Medicine, Airlangga University, from June to October 2015.

Research sample

This research used twenty four male mice of the Swiss Albino strain, weighing 20 g -30 g, and 2.5 months old. Each group consisted of 6 treated mice, with one mouse as the donor. The sample size was determined by Federer's formula. The animals were obtained from the

animal facility at the Veterinary and Pharmaceutical Center (Pusat Veteriner dan Farmasi, Pusvetma), Surabaya.

Research procedure

Mice were intraperitoneally infected with mouse red blood cells (RBC) containing 1×10^5 *P. berghei* per 0.2 ml. The infection was detected by daily microscopic examination of the erythrocytes in thin blood smears taken from the tail vein of the mice and stained with 20% Giemsa. The mice were randomized into four groups, consisting of two control groups (K1, KP) and two treatment groups (P1, P4).

In the treatment group P1, the mice were infected with *P. berghei* and 48 hours later treated with a single administration of artemisinin. Subsequently the parasites in the blood of P1 animals were passaged into another group of animals (P2), who were then given 2 artemisinin treatments. The procedure was repeated for groups P3 and P4, which were infected with blood from P2 and P3 donors, respectively, and received 3 and 4 artemisinin treatments, respectively. The control group K1 consisted of mice who were infected with *P. berghei* and were untreated (did not receive artemisinin), while treatment control group KP consisted of untreated mice who were infected with *P. berghei* that had previously been treated four times with artemisinin.

Details of the treatment are as follows: Treatment group P1 comprised six mice that had been inoculated with RBC containing 1×10^5 *P. berghei* in 0.2 ml (D_0) and were then treated with the artemisinin "4-day test" (4-DT) at the ED_{99} dose of 200mg/kgBW for 3 days, starting 48 hours after infection (D_2). After that, the parasitemia rate was monitored and calculated at 120 hours after infection. After the parasitemia rate was higher than 2 % of RBC, one mouse was used as donor of parasites to be passaged to 5 new mice. Then, 48 hours post infection, the 5 new mice were treated artemisinin at the same ED_{99} dose for 3 days and so on up to 4 passages (P2, P3, P4). The control group K1 contained

five mice inoculated with 0.2 ml RBC containing 1×10^5 *P. berghei* that had never been treated with artemisinin (D_0) and were left untreated. Treatment control group (KP) contained six mice inoculated with 0.2 ml RBC containing 1×10^5 *P. berghei* that had previously been treated 4 times with artemisinin (D_0) and were left untreated. Parasite development was followed in all treatment groups until 10 days after infection.^(12,13)

Leukocyte differential count

The leukocyte differential count was done to represent one of the immune system parameters that plays an important role in malaria infection. The leukocyte differential count was done at 48 hours, and on the 5th, 8th and 10th day post-infection, by making thin blood smears taken from the mouse tail vein, fixed with methanol for 5 minutes, put in a Giemsa staining solution for 20 minutes, washed with running water with the slide in a tilted position, and dried in the air. The leukocyte differential count was determined under the light microscope at 1000x magnification. Every 100 leukocytes that were found were counted and grouped into neutrophils, eosinophils, basophils, lymphocytes, and monocytes. The leukocytes were counted in a number of visual fields throughout the smear, the examination starting from the center of the smear, proceeding parallel to the edge of the smear, then moving back and forth until a total leukocyte count of 100 was reached. The proportions of the types of leukocytes was expressed in percentages.⁽¹⁰⁾

Data analysis

The data on the leukocyte profile was processed with ANOVA. If there were any significant differences, the multiple comparisons test was done using SPSS 17.0, with the level of significance set at 5% to determine differences in treatment.

Ethical clearance

This research has been approved by the ethics committee of the Faculty of Veterinary

Medicine, Airlangga University, on June 3rd 2015 with certificate number No. 464 KE.

RESULTS

Table 1 and Fig. 1A show the resulting neutrophil percentage in P4, which was highest among the other treatment groups, starting from the 2nd day until the 10th day post infection. The graph also shows that the infection on the 2nd day in the treatment control group KP (which was infected with *P.berghei* that had been treated 4 times with artemisinin and were then left untreated) and the treatment group P4 (that was infected with *P.berghei* and had been treated four times with artemisinin) increased at significantly different rates when compared with the control group K1 (mice that had been infected with *P.berghei* that had never been treated with artemisinin) and group P1 (infected with *P.berghei* and treated once with artemisinin). On the 5th day of infection, group KP showed significantly different increases in neutrophils when compared with the other three groups, namely groups K1, P1, and P4. On the 8th day of infection, the neutrophil percentage did not show any significant differences between all the groups (K1,KP,P1,P4). On the 10th day of infection, the neutrophil percentage in group P1 showed significantly different decreases when compared with the other groups (K1, KP and P4).

Table 1 and Figure 1B show that the highest monocyte percentage was in group P4 when compared with the other treatment groups. The monocytes started to increase on the 2nd - 8th days post infection and then decreased on the 10th day post infection, but the percentage was still higher than in the other treatment groups. These results show that on the 2nd day of infection the KP control group showed significantly different increases when compared with groups K1, P1, and P4. On the 5th and 8th days of infection the monocytes percentage did not show any significant differences between all groups (K,KP,P1,P4). On the 10th day of infection, group

Table 1. Distribution of mean leukocyte count by control and treatment groups

Leukocytes (%)	Treatment group				P
	K1 (n=5)	KP (n=5)	P1 (n=5)	P4 (n=5)	
Day 2					
Neutrophils	0.00 ± 0.00 ^a	20.80 ± 3.42 ^b	6.60 ± 8.53 ^a	27.40 ± 8.76 ^b	< 0.05
Monocytes	0.40 ± 0.89 ^a	2.00 ± 1.00 ^b	0.60 ± 1.34 ^b	4.00 ± 3.16 ^c	< 0.05
Lymphocytes	94.00 ± 6.04 ^a	75.00 ± 6.20 ^b	90.20 ± 11.69 ^a	64.60 ± 4.87 ^b	< 0.05
Basophils	0.00 ± 0.00 ^a	1.00 ± 1.00 ^b	0.00 ± 0.00 ^a	0.40 ± 0.84 ^{a,b}	< 0.05
Eosinophils	5.60 ± 6.34 ^b	1.20 ± 1.78 ^a	2.60 ± 3.71 ^b	3.60 ± 2.19 ^b	< 0.05
Day 5					
Neutrophils	12.80 ± 13.06 ^a	25.40 ± 3.71 ^o	13.20 ± 10.82 ^a	23.40 ± 4.87 ^a	< 0.05
Monocytes	3.40 ± 1.94 ^a	1.80 ± 1.30 ^a	2.80 ± 2.77 ^a	6.20 ± 5.49 ^a	> 0.05
Lymphocytes	78.00 ± 7.68 ^a	69.40 ± 6.65 ^a	82.00 ± 12.18 ^a	67.60 ± 7.40 ^a	> 0.05
Basophils	0.00 ± 0.00 ^a	0.20 ± 0.44 ^a	0.00 ± 0.00 ^a	2.00 ± 1.41 ^o	< 0.05
Eosinophils	5.60 ± 3.91 ^a	3.20 ± 4.54 ^{ab}	2.00 ± 3.93 ^{ab}	0.60 ± 1.34 ^b	< 0.05
Day 8					
Neutrophils	20.60 ± 9.71 ^a	26.40 ± 3.78 ^a	25.00 ± 14.69 ^a	25.00 ± 8.39 ^a	> 0.05
Monocytes	1.20 ± 1.0 ^a	3.60 ± 2.70 ^a	9.20 ± 4.20 ^a	4.20 ± 1.48 ^a	> 0.05
Lymphocytes	78.20 ± 9.73 ^a	69.60 ± 4.92 ^a	65.40 ± 15.12 ^a	69.60 ± 8.96 ^a	> 0.05
Basophils	0.00 ± 0.00 ^a	0.20 ± 0.44 ^b	0.00 ± 0.00 ^a	0.60 ± 0.54 ^b	< 0.05
Eosinophils	0.00 ± 0.00 ^a	0.20 ± 0.44 ^a	0.40 ± 0.89 ^a	0.80 ± 0.83 ^a	> 0.05
Day 10					
Neutrophils	35.00 ± 8.68 ^a	28.80 ± 5.89 ^a	15.20 ± 8.75 ^o	30.60 ± 9.83 ^a	< 0.05
Monocytes	3.00 ± 2.54 ^a	1.80 ± 0.83 ^{ab}	7.40 ± 7.60 ^c	2.80 ± 1.78 ^{ab}	< 0.05
Lymphocytes	60.80 ± 11.16 ^a	68.60 ± 5.54 ^a	77.40 ± 5.81 ^a	66.60 ± 9.53 ^a	> 0.05
Basophils	0.00 ± 0.00 ^a	0.40 ± 0.54 ^a	0.00 ± 0.00 ^a	0.40 ± 0.89 ^a	> 0.05
Eosinophils	1.20 ± 2.16 ^a	0.40 ± 0.54 ^a	0.00 ± 0.00 ^a	0.00 ± 0.00 ^a	> 0.05

Values are means ± SEM. Values for each day along the same rows with different superscript letters are significantly different ($p < 0.05$); Note: K1 = control group infected with *P.berghei* and untreated with artemisinin; KP = treatment controls (infected with *P.berghei*, treated 4 times with artemisinin and then untreated; P1 = treatment group infected with *Plasmodium berghei* and treated with a single dose of artemisinin; P4 = treatment group infected with *Plasmodium berghei* and treated 4 times with artemisinin

P1 showed significant differences when compared with the other three groups (K1, KP, and P4).

Table 1 and Figure 1C show that the highest lymphocyte percentage was in group P4 when compared with the other groups, starting on the 2nd day until the 10th day post infection. The results show that on the 2nd day of infection groups KP and P4 showed significant differences when compared with groups K1 and P1. On the 5th, 8th, and 10th days of infection, the lymphocyte percentage did not show any significant differences between all groups (K1, KP, P1, P4). Table 1 and Figure 1D show that the highest basophil percentage was in group P4 when compared with the other groups, starting on the 2nd day until the 5th day post infection then on the 8th day post infection, the basophil percentage decreased. These results show that on the 2nd day

of infection the KP control group showed significantly different increases when compared with groups K1 and P1. On the 5th day of infection, group P4 showed significantly different increases in percentages of basophils when compared with the other three groups (K1, KP, P1). On the 8th day of infection, group P4 showed significantly different increases in percentages of basophils when compared with groups K1 and P1, but did not show any significant differences with group KP. On the 10th day of infection, the basophil percentage did not show any significant differences between all control and treatment groups (K, KP, P1, P4).

Table 1 and Figure 1E show that the eosinophil percentage in group P4 was higher the 2nd day post infection. After that, the eosinophil percentage decreased. The image shows that on the 2nd day of infection the KP

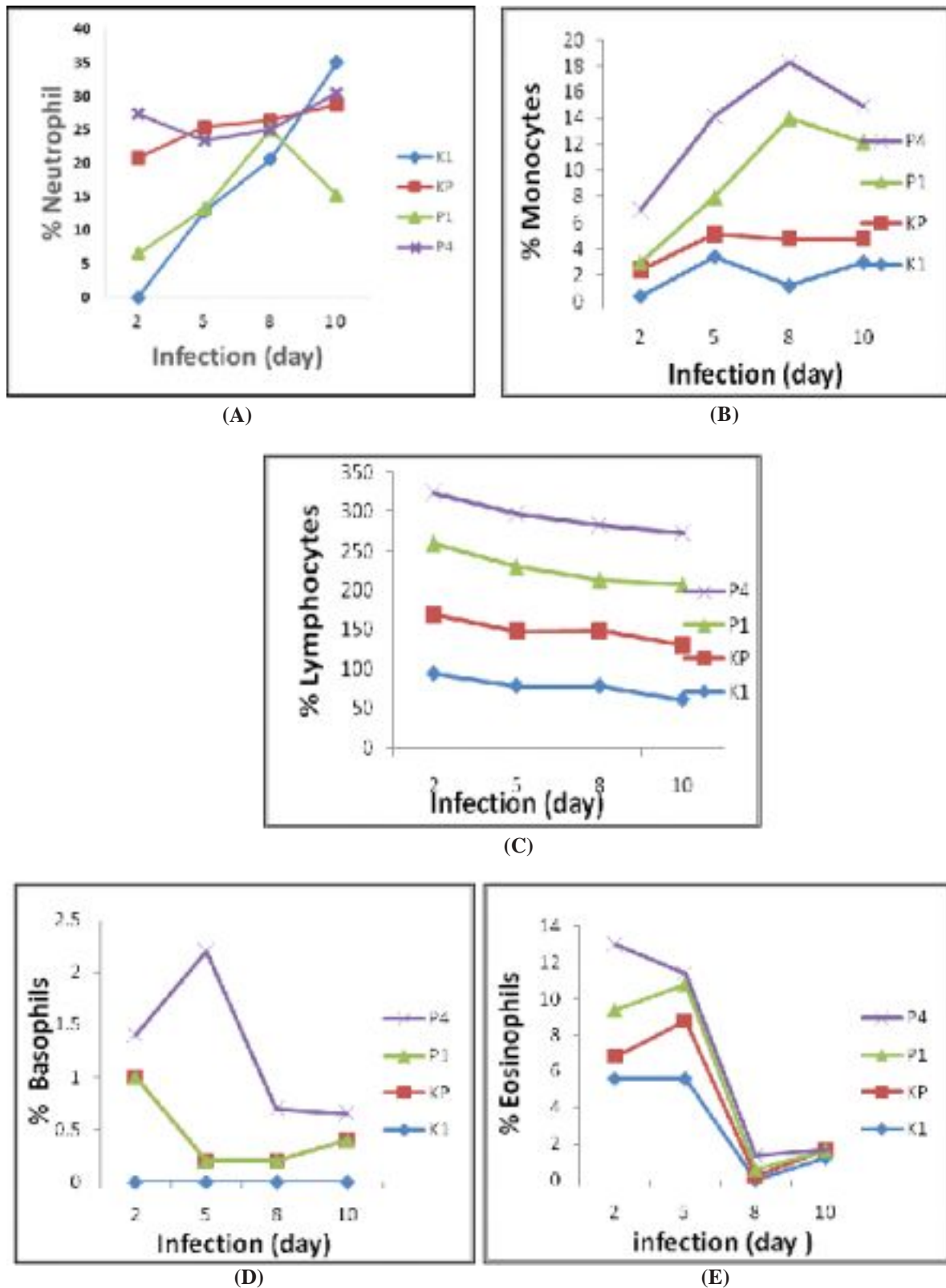


Figure 1. Leukocyte type percentage in control and treatment groups repeatedly exposed to artemisinin in mice infected with *P.berghei*. Note: K1 = control group infected with *P.berghei* and untreated with artemisinin; KP = treatment controls (infected with *P.berghei*, treated 4 times with artemisinin and left untreated); P1 = treatment group infected with *Plasmodium berghei* and treated with a single dose of artemisinin; P4 = treatment group infected with *Plasmodium berghei* and treated 4 times with artemisinin

control group showed significantly different increases when compared with groups P1 and P4. On the 5th and 8th days of infection, group P4 showed significantly different increases in eosinophil percentages when compared with the K1 control group, but did not show any significant differences with groups KP and P1. On the 10th day of infection, the eosinophil percentage did not show any significant differences between all control and treatment groups (K,KP,P1,P4).

DISCUSSION

In the present study we showed that neutrophil percentage increased from the 2nd day until the 5th day post infection with *P. berghei*. Repeat treatments with artemisinin showed that neutrophils are cells that function as the first line of defense of the body against infection or pathogens entering the body (to eliminate *P.berghei*). Mice infected with *P. berghei* and treated with a one-time dose of an antimalarial drug yielded the same results as mice treated with a one-time dose of artemisinin (P1), as is shown by the increasing number of neutrophils, monocytes and lymphocytes after more than 5 days of infection.⁽¹⁴⁾ Research with repeated exposure of artemisinin in mice as a model of in vivo in infections in humans has not been done before.

Cytokines and interleukins that are released after lysis of neutrophils will stimulate the bone marrow to release neutrophils so that the production of neutrophils is increased. Malaria infections can induce both humoral and cellular immune responses. In the humoral immune response, antibodies provide protection against malaria infection through a variety of mechanisms. Antibodies can block the invasion of merozoites into erythrocytes and inhibit the growth of parasites in the erythrocytes, while cellular immunity against Plasmodium plays a role in the erythrocytic stage. Cellular immunity is activated by T helper (Th) lymphocytes, which are a subset of T cells needed in the induction of

immune responses against an invasion of foreign antigens. According to the type of cytokine produced, Th lymphocytes are divided into Th1 cells, which will activate cellular immunity and Th2 cells, which will activate humoral immunity.⁽¹⁵⁾

The increase in neutrophil percentage on the 8th day post infection was not significantly different between groups. On the 10th day post infection, the group of mice infected with *P.berghei* and treated with a single dose of artemisinin (P1) showed a decrease in neutrophils when compared with groups K1, KP and P4. This shows that neutrophils function as the first line of defense to eliminate *P.berghei* by phagocytizing parasites and neutrophils on the first days of infection or when the *P.berghei* attacks. This suggests that infections of longer duration will increase the percentage of parasitemia in the mice so that the high numbers of parasites in their bodies will suppress the immune system (immunosuppression), causing the immune system to become inadequate and unable to respond properly. High parasitemia rates will improve the mechanism of parasite invasion and suppress the host immune system.⁽¹⁶⁾

Neutrophils play an important role in the immune response by modulating both cellular and humoral immunity via the synthesis and release of immunoregulatory cytokines. All neutrophil profiles in the groups showed that *P.berghei* which had been treated with repeated artemisinin showed differences when compared with *P.berghei* that had never been treated with artemisinin. This shows the infection had become more powerful because of the parasites' ability to develop faster. This is consistent with in vitro studies that were performed by Maslachah⁽¹⁷⁾ who found that Plasmodium which remained viable after repeated artemisinin exposure will increase the parasitemia rate and take a shorter time to become viable again when compared with the parasites in the treatment group which only received a one-time artemisinin exposure.

Monocytes are phagocytic cells that can destroy antigens by ingestion. Moreover,

monocytes may contribute to delivery of antigens to the lymphocytes (antigen presenting cells, APC). The relatively high increases in the number of monocytes in mice that had been infected with *P. berghei* and exposed repeatedly to artemisinin, resulted in increased phagocytosis of the parasites, this being the body's defense mechanism against *P. berghei*. Repeat artemisinin therapy in the treatment group causes faster growth of *P. berghei* that remained viable from medicine exposure, so increasing the parasitemia rate and also the number of cells contributing to the elimination of the parasites, which will affect the monocyte profile.

The treatment control group of mice that were infected *P. berghei* and had been treated four times with artemisinin and left untreated (KP) showed an increasing average percentage of monocytes that is higher when compared with the control mice that were infected with *P. berghei* and were not exposed to the medicine (K0). This significant difference is visible on the 2nd day post infection. This can be explained from the research results of Maslachah,⁽¹⁷⁾ who found that Plasmodium that have been exposed to artemisinin and are still viable and grow normally, have a faster rate of intra-erythrocytic development, so that the increase in the average percentage of monocytes is a mechanism for defense against *P. berghei*.

The monocyte profile from the 2nd to the 8th day post infection showed an increasing trend in the treated group of mice exposed to artemisinin, but after the 10th day of infection, there was a decrease, whereas in the group of control mice, the trend was decreasing or stable. This shows that after chronic infection, monocyte numbers will be decreased so that the monocytes function maximally in the first week of infection.

The average percentage of lymphocytes in the treated groups of mice showed that had been exposed to artemisinin, seemed to decrease when compared with the control groups. This is because artemisinin treatment for 3 days with a therapeutic dose in mice infected with *P. berghei* caused suppression of Plasmodium growth so

that the growth of malaria parasites in all treatment groups was inhibited, as compared with the untreated control group. Research conducted by Teuscher et al.⁽¹⁸⁾ in *P. falciparum* strains W2 and W2AL80 also showed a decrease in parasitemia rate to 1% up to 8 days after medicine exposure, when it began to show increases in parasitemia rate, reaching more than 10% after 12 days.

With regard to the percentage of lymphocytes on the 2nd day of infection, both the treatment control group KP and the treatment group P4 infected with *P. berghei* that had been repeatedly exposed to artemisinin, showed significant differences when compared with the control group K1 and the treatment group P1 infected with *P. berghei* and received no treatment and a one-time exposure to artemisinin, respectively. This can be explained by the fact that lymphocytes play a role in the body defense mechanism against *P. berghei*. At the beginning of infection, *P. berghei* will stimulate the formation of antigen presenting cells (APCs). APCs will stimulate the mice to produce T lymphocytes that play a role in the immune response to destroy the Plasmodium. However, after the 5th, 8th and 10th day of infection, the lymphocyte percentage showed no significant differences between all groups of mice. This shows that antigen presenting cells (APC) can affect the response of T lymphocytes during acute infection in the erythrocytic stage.

In the present study, basophils and eosinophils did not play a major role in infection by the parasite *P. berghei*. Basophils have an important role in the allergic process, while eosinophils play a major role in the process of infection by parasitic worms.⁽¹⁹⁾ The average value of basophils from this study ranged from 0 to 0.4%. These results do not show significantly different increases from the physiological value of basophils of 0 to 0.3%. The percentage of eosinophils was 0 to 5.6%, which did not show a significant increase from the physiological value of eosinophils of 0-4%. On the 2nd day post infection, the basophil and

eosinophil profiles in the treatment control group KP showed that the groups of mice infected with *P. berghei* and exposed 4 times to artemisinin had been able to stimulate the production of cells that play a role the immune system (basophils and eosinophils). This suggests that infection with *P.berghei* that have been exposed repeatedly can stimulate the immune system to work optimally by expressing all types of leukocytes in order to neutralize the infection. Derivatives of artemisinin (artemether and artesunate) are effective in rescuing mice with late state cerebral malaria and in decreasing leukocyte accumulation in the brain.⁽¹¹⁾ Many research studies of medicinal plants have shown them to have antimalarial activity by inducing the synthesis of red blood cells, white blood cells and platelets.^(20,21)

A limitation of this study is that the leukocyte profile data were obtained by using blood smear preparations only, whereas specialized tools are required such as heme analyzers to obtain results with better validity. The clinical implication of this study is that this exposure model of artemisinin with in vivo repeated passages in mice infected with *P.berghei* can be used as a basis to describe the functioning of the immune system against infections with *P. berghei* parasites which have been exposed to repeated artemisinin. The results of this study can provide a baseline leukocyte profile as one of the immune system's lines of defense against malaria infections caused by *P. berghei* that have repeatedly been exposed to antimalarial drugs, as an in vivo model of malaria in humans.

Further research is needed on the immunologic profile of cytokine-mediated tissue or organ damage. The leukocyte response occurs under normal physiological and pathological conditions. The manifestations of the leukocyte response to malaria infection are in the form of decreases or increases in one or several types of leukocyte, which can provide information and clinical clues that may help in the diagnosis of infections caused by Plasmodium.

CONCLUSION

Repeated exposure to artemisinin in mice infected with *Plasmodium berghei* will cause changes in leukocyte profile, comprising neutrophils, lymphocytes, monocytes, basophils and eosinophils.

CONFLICT OF INTEREST

The authors state that there is no conflict of interest with the other authors, publisher or study.

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CONTRIBUTION

LM contributed to drafting the manuscript and the design of the study. LM and RS contributes to data collection, analysis and interpretation. LM contributed revising manuscript for critically important content. All authors read and approved the final manuscript.



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