LYMPHOCYTE-T TYPE TH1 AND TH2 ACTIVITY DIFFERENCE OF LUNG TISSUE ON Heligmosomoides nematode and Mycobacterium tuberculosis SEQUENTIAL CO-INFECTION

Laksni Wulandari¹, Muhammad Amin¹, Soedarto², Gatot Soegiarto³
¹Pulmonology and Respiratory Medicine Department, ²Parasitology Department, Dr. Soetomo Hospital, ³Internal Medicine Department, Faculty of Medicine, Airlangga University, Surabaya, Indonesia

ABSTRACT
Tuberculosis (TB) is a chronic infectious disease caused by Mycobacterium tuberculosis that are often associated with uneffectiveness of the BCG vaccine and high worm infection. The objective of this study was to determine the differences in the activity of lymphocytes T type Th1 (IFN-γ) and Th2 (IL-4) in lung tissue on Heligmosomoides nematode and Mycobacterium tuberculosis sequential co-infection. This research using 49 mice were divide into 7 groups treated with infection by Mycobacterium tuberculosis inhaled and Heligmosomoides polygyrus orally within 8 and 16 weeks. The levels of IFN-γ in peripheral blood serum (89.929 ± 3.533 pg/mL) resemblesthe pattern of the percentage of lymphocytes T CD4+ Th1 in lung tissue (3.246 ± 0.519%) and peripheral blood (4.930 ± 0.237%), while the levels IL-4 in the peripheral blood serum (20.782 ± 4.043%) resembles the pattern of the percentage of lymphocytes T CD4+ Th2 in intestinal tissue (1.048 ± 0.359%) and peripheral blood (1.916 ± 0.537%). In conclusion, there is difference in the activity of lymphocytes T type Th1 and Th2 but it does not affect the immune response to Mycobacterium tuberculosis infection. (FMI 2017;53:124-130)

Keywords: Mycobacterium tuberculosis, Heligmosomoides polygyrus, IFN-γ, IL-4

INTRODUCTION
Tuberculosis (TB) is a chronic infection caused by Mycobacterium tuberculosis with a mortality rate of around 1.4 million per year, especially in developing and low-income countries (WHO 2012). The high incidence rate of TB in most areas of the country is closely related to the effectiveness of low BCG vaccination and the high prevalence of worm infections (Fine et al 1995, Lipner et al 2006, Elias et al 2007).

Mycobacterium tuberculosis, a facultative intracellular paracitic bacillus bacterium (Todar 2009) can be eliminated by cellular immunity played by macrophages; CD4+ T-lymphocytes that secrete IFN-γ; CD8+ T lymphocytes that kill macrophages infected with TB germs; as well as T lymphocytes γδ (Schluger et al 2005) in which the immune response requires a strong Th1 type of cytokine (van Crewel et al 2002). In contrast, worm infections stimulate the activation of eosinophil cells, mast cells, basophile cells, and IgE formation, which is a Th2-type immune response (Anthony et al 2007).

The dominant Th2-type immune response suppresses the Th1 immune response through suppression by IL-4 which proves that worm infections can suppress the immune response to the TB germs (Resende et al 2006, Potian et al 2007, Bhatt et al 2007). In contrast, Erb et al (2002) and Frantz et al (2007) reported that worm or
parasite infections did not alter immunity to mycobacteria in mice-treated animals, in which dendritic cells mutually stimulated using bacterial antigens and worm antigens were able to undergo maturation and induce both Th1-type immune responses and Th2-type immune responses. The existence of differences in the results of these studies to date have not been able to be drawn conclusions that are certain and satisfactory.

Research on the co-infection of Nematoda *Heligmosomoides polygyrus* worm in mice previously infected with *Mycobacterium tuberculosis* as a sequential standard model is very necessary to be done to determine the difference of T1 and Th2 type T lymphocyte cell activity so that TB can be effectively controlled effectively.

**MATERIALS AND METHODS**

The research was conducted for 6 (six) months with the location of the research is at Animal Cage Try the Parasitology Division of Clinic Faculty of Medicine Universitas Brawijaya and in Bacteriology Laboratory of Tuberculosis Infection Study Group of Tropical Diseases Institution Airlangga University Surabaya.

The sample size or replication is the number of treatments in an experiment (r). Replication affects the number of replicates (r) of a study. Using the large formula of Steel and Torrie samples (1980) and Higgins correction factor formulas to anticipate a drop out with an estimated 55%. The total sample size was 49 male (Mus musculus) mice of wild type type which was 8-12 mingggu with body weight 30-35 gram.

Mice were divided into 7 groups consisting of: *Mycobacterium tuberculosis*-infected group for 8 weeks (M.tb8), tuberculosis infected group (*Mycobacterium tuberculosis*) for 16 weeks (M.tb16), a group infected with a worm (*Heligmosomoides polygyrus*) for 8 weeks (H.pg8), a worm infected group (*Heligmosomoides polygyrus*) for 16 weeks (H.pg16), a group of mice treated with helminths (*Heligmosomoides polygyrus*) followed by TB infection (*Mycobacterium tuberculosis*) (H.pg + M.tb), a group of mice treated with TB coinfection (*Mycobacterium tuberculosis*) followed by a helminth infections (*Heligmosomoides polygyrus*) (M.tb + H.pg), as well as infectious groups as controls. At the end of the 16th week (ie 8 weeks after the last infection treatment sequence), the study subjects of all treatment groups will be evaluated by taking peripheral blood specimens for the measurement of some IL-4 and IFN-cytokines with ELISA method.

**Mycobacterium tuberculosis infection method**

Mice Balb/c presented 10mL PBS-Tween 80 solution containing 106 *Mycobacterium tuberculosis* inhalation using a modified nose only inhalation system or Middlebrook Inhalation Exposure System (GLAC-Col) for 30 minutes in inhalation chamber placed in a large box filtered HEPA.

**Method of Heligmosomoides polygyrus Infection**

Balb/c mice were inoculated with oral *Heligmosomoides polygyrus* larvae with a blunt-specially-sided blistering syringe containing 100 µL PBS solution containing 2000 L3/mL. Gavage needle is inserted into the esophagus and the contents are sprayed slowly to avoid regurgitation (Camberis et al., 2003).

**Processing and analysis of data**

The data of the research variables were analyzed for distribution by group with Shapiro-Wilk test. Data that is not normally distributed will be tested by Kruskal-Wallis, while the normal distributed data is tested by homogeneity of variance. The homogeneous variance will be analyzed by ANOVA test, while the non homogeneous variance will be analyzed by Brown-Forsythe test.

**RESULTS**

**Effect of treatment on Th-1 lymphocyte activity**

Th1 T lymphocyte activity is characterized by the production of Interferon-γ (IFN-γ) cytokines in peripheral blood measured by the ELISA method and the percentage of CD4+ T lymphocytes expressing the IFN-γ intracellular as measured by flowisotometry in lung tissue, intestinal tissue, and peripheral blood. In the coinfection group of *M. tuberculosis* and *H. Polygyrus* there is a tendency that high levels of IFN-γ determined by the last sequence of infection, in which the group treated with the last co-infection was *M. tuberculosis* infection IFN-γ level (89,929± 3,533 pg/mL) was significantly higher than the group treated with the last co-infection of *H. Polygyrus* infection (46,168,7,821 pg/mL) seen in Table 1 and Figure 1.
Table 1. IFN-γ level in peripheral blood serum

<table>
<thead>
<tr>
<th>Treatment groups</th>
<th>Peripheral blood serum ELISA (pg/mL)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean ± SD</td>
<td>Min</td>
</tr>
<tr>
<td>H.pg 8</td>
<td>8.564 ± 1.414 *</td>
<td>7.278</td>
</tr>
<tr>
<td>H.pg 16</td>
<td>29.463 ± 6.279 b</td>
<td>22.203</td>
</tr>
<tr>
<td>H.pg + M.tb</td>
<td>89.929 ± 3.533 c</td>
<td>85.127</td>
</tr>
<tr>
<td>M.tb + H.pg</td>
<td>46.168 ± 7.821 c</td>
<td>38.852</td>
</tr>
<tr>
<td>M.tb 16</td>
<td>62.975 ± 7.824 d</td>
<td>51.203</td>
</tr>
<tr>
<td>M.tb 8</td>
<td>106.481 ± 5.446 f</td>
<td>99.620</td>
</tr>
<tr>
<td>Control</td>
<td>7.207 ± 2.034 e</td>
<td>4.747</td>
</tr>
</tbody>
</table>

Notes: H.pg: H. Polygyrus infection; M.tb: M. tuberculosis infection; 8 and 16: infection for 8 and 16 weeks; The letters a, b, c, d, e, f: indicate that groups with the same letter marks have insignificant differences, whereas groups with different letter marks have significant differences.

Figure 2. Flowcytometry results of CD4+ T lymphocytes in lung tissue
Percentage of CD4+ T lymphocytes expressing the IFN-γ molecule intracellular in lung tissue, intestinal tissue, and peripheral blood were measured by flow cytometry technique using antibodies against CD4 and INF-γ simultaneously with the FACSCalibur tool and the CellQuest software obtained the number of cell percentages in question and the dot plot visualization in the upper right quadrant of each graph as shown in Fig. 2.

CD4+ T lymphocytes were identified with anti-CD4 antibodies that were conjugated with fluoro- chrome fluorescein isothiocyanate (FITC) (top right). Furthermore, permeabilization and identification of CD4+ T lymphocytes expressing IFN-γ intracellular with anti-IFN-γ antibodies conjugated with phycoerythrin (PE) (bottom left) or expressing IL-4 with anti-IL-4 antibodies conjugated with peridinin chlorophyll protein (perCP) (bottom right). In the table at the bottom of the dot plot can be read the percentage of events in the upper right quadrant (UR). This figure represents the measurement in the treatment group no. 1 member no. 4.

This technique is a technique of calculating intracellular cytokines directly without in vitro stimulation. CD4+ T lymphocytes expressing the IFN-γ molecule intracellular percentage is very low (less than 6%), but the percentage pattern in each type of treatment group can still be seen in Figures 3, 4, and 5.

The highest percentage of CD4+ T-1 lymphocytes in lung tissue (Fig. 5.3) was highest in *M. tuberculosis* infection for 8 weeks (4.508 µ.947) and then decreased in *M. tuberculosis* infection for 16 weeks (2.058 × 0.845). *H. Polygyrus* infection for 8 weeks had a low percentage of CD4+ T1 lymphocytes (equivalent to control) and then increased in infection for 16 weeks. The group treated with the last co-infection in the form of *M. tuberculosis* infection had a significantly higher percentage of CD4+ T1 T lymphocytes than the opposite co-infection.

The calculated percentage of CD4+ T1 lymphocyte T lymphocytes in intestinal tissue showed no significant intergroup differences (Brown-Forsythe test, p = 0.109; p> 0.05) as shown in Figure 4.

The calculated percentage of CD4+ Th1 lymphocyte T lymphocytes in peripheral blood showed that the group treated with the last co-infection of *M. tuberculosis* infection had a significantly higher percentage of CD4+ T1 T lymphocytes than the opposite co-infection, as shown in Figure 5.
Lymphocyte CD4+ T1 lymphocyte graph pattern shows a tendency that IFN-γ in peripheral blood serum has a picture that more closely resembles the CD4+ Th1 lymphocyte T lymphocyte pattern in lung tissue and peripheral blood, in contrast to the CD4+ T1 lymphocyte T lymphocyte pattern in the intestinal tract (Fig. 1 to Fig. 5).

The Effect of treatment on T-type Th2 lymphocyte activity

Th2-type T lymphocyte activity is characterized by the production of the interleukin-4 (IL-4) cytokine in peripheral blood measured by the ELISA method and the percentage of CD4+ T lymphocytes expressing intracellular IL-4 molecules as measured by flow-citometry in lung tissue, intestinal tissue and peripheral blood.

In the co-infection group M. tuberculosis and H. Polygyrus it appears that significantly higher levels of IL-4 in peripheral blood (p = 0.009; p <0.05) were higher when the latter co-infection was H. Polygyrus infection (66,625 ± 13,937 pg/mL) than when the last co-infection treatment of M. tuberculosis infection (20.782 ± 4,043 pg/mL) can be seen in Table 2 and Figure 6.

Table 2. Peripheral blood ELISA in treatment groups

<table>
<thead>
<tr>
<th>Treatment groups</th>
<th>Peripheral blood ELISA (pg/mL)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean ± SD</td>
<td>Min</td>
</tr>
<tr>
<td>H.pg 8</td>
<td>93.887 ± 7.279 *</td>
<td>83.647</td>
</tr>
<tr>
<td>H.pg 16</td>
<td>78.964 ± 12.376 a</td>
<td>66.912</td>
</tr>
<tr>
<td>H.pg + M.tb</td>
<td>20.782 ± 4.043 a</td>
<td>17.070</td>
</tr>
<tr>
<td>M.tb + H.pg</td>
<td>66.625 ± 13.937 a</td>
<td>49.982</td>
</tr>
<tr>
<td>M.tb 16</td>
<td>16.961 ± 5.209 a</td>
<td>11.035</td>
</tr>
<tr>
<td>M.tb 8</td>
<td>14.007 ± 4.418 a</td>
<td>9.632</td>
</tr>
<tr>
<td>Control</td>
<td>5.014 ± 1.357 b,c</td>
<td>3.140</td>
</tr>
</tbody>
</table>

Notes: H.pg: H. Polygyrus infection; M.tb: M. tuberculosis infection; 8 and 16: infection for 8 and 16 weeks; The letters a, b, c,: indicate that groups with the same letter marks have insignificant differences, whereas groups with different letter marks have significant differences.

The results of the CD4+ Th2 lymphocyte T lymphocyte count in lung tissue showed no significant differences between the treatment groups of infections, but all the treatment group data were significantly different than the control group (Games-Howell Test, p <0.05) as shown in Fig. 7.

Figure 7. CD4+ T-lymphocytes expressing IL-4 in lung tissue

The calculated percentage of CD4+ Th2 lymphocyte T lymphocytes in the intestinal tract showed M. tuberculosis infection for 8 weeks and for 16 weeks had low CD4+ T2 lymphocyte percentage (equivalent to control group). The group that received the last co-infection treatment in the form of H. Polygyrus infection had a significantly higher percentage of CD4+ Th2 lymphocyte T than those treated with the last co-infection with M. tuberculosis infection (2,712,0,502 vs. 1,048 ± 0.359; p = 0.005, p <0.05) as shown in Figure 8.

Figure 8. CD4+ T-lymphocytes expressing IL-4 in intestinal tissue

The CD4+ Th2 lymphocyte T lymphocyte count in peripheral blood showed M. tuberculosis infection for 8 weeks had a low percentage of lymphocytes of T-CD4+ Th2 (0.844 ± 0.178), equivalent to 16 weeks of infection of M. tuberculosis (0.964 ± 0.273) and control group (0.334 ± 0.155). The group that received the last co-
infection treatment in the form of \textit{M. tuberculosis} infection had a significantly lower percentage of CD4+ Th2 lymphocytes than the group treated with the last coinfection of \textit{H. Polygyrus} infection as shown in Figure 9.

**DISCUSSION**

Lymphocyte CD4+ Th2 lymphocyte pattern pattern shows a tendency that the pattern of IL4 levels in peripheral blood serum has a picture that more closely resembles the CD4+ Th2 lymphocyte T lymphocyte pattern in intestinal and peripheral blood tissue, in contrast to the CD4+ Th2 lymphocyte T lymphocyte pattern in lung tissue (Figure 9).

The role of T lymphocytes in the immune response to \textit{M. tuberculosis} infection is undoubtedly, where a specific new immune response forms 2-3 weeks after the onset of infection. Tissue lymphocytes accumulate at the site of infection, proliferate and secrete cytokines, especially IFN-\(\gamma\). Protective immune responses to \textit{M. tuberculosis} are more necessary for the role of Th1 type cytokines (van Crevel et al., 2002). Th1-type cytokines, among others, IFN-\(\gamma\), are needed not only to activate macrophages (Shi et al., 2003) but also to assist the activity of CD8+ T lymphocytes (Salgame 2005; Vesosky et al., 2006).

In this study, there was an increase in the percentage of Th1 lymphocytes in lung tissue and in peripheral blood that correlated strongly with IFN-\(\gamma\) cytokine levels in peripheral blood serum. The increase occurred in the infection of \textit{M. tuberculosis} for 8 weeks which subsequently ‘subside’ at the time of infection lasting up to 16 weeks, which is consistent with the results of several other researchers who found that elevated levels of IFN-\(\gamma\) primarily in the early stages of infection, especially after the second week after infection (Shi et al., 2003; Vesosky et al., 2006). In contrast the role of Th2 lymphocytes is very limited, as evidenced by the low percentage of Th2 lymphocytes in lung tissue and peripheral blood and levels of IL-4 cytokine in peripheral blood serum in \textit{M. tuberculosis} infection for 8 weeks. Levels of IL-4 in new blood increased in the treatment of \textit{H. Polygyrus} infection. Increased levels of IL-4 in peripheral blood serum correlated with the percentage of Th2 lymphocytes in the intestinal and peripheral blood tissue but did not affect the percentage of Th2 lymphocytes in lung tissue.

In the co-infected group, the levels of IL-4 in peripheral blood serum only increased when the last treatment of infection was \textit{H. Polygyrus} infection. Nematode worm infection in the intestine does have a systemic effect on peripheral blood circulation but does not affect the percentage of Th1 or Th2 lymphocytes in lung tissue, so it can be concluded that chronic worm infection has no effect on \textit{M. tuberculosis} infection.

**CONCLUSION**

Sequential co-infection of \textit{Heligmosomoides polygyrus} and \textit{Mycobacterium tuberculosis} leads to differences in T1 and Th2 type T lymphocyte activity in intracellular expressing IFN-\(\gamma\), IFN-\(\gamma\) level in peripheral blood serum, intracellular IL-4 molecule and IL-4 levels in peripheral blood serum, but the changes do not affect immune response to \textit{Mycobacterium tuberculosis} infection.

**REFERENCES**


not interfere with efficient elimination of *Mycobacterium bovis* BCG from the lungs of mice. Clin Diagn Lab Immunol 9, 727-730


