

Association between serum PGE2 levels and degree of acid- fast bacilli positivity in sputum of pulmonary tuberculosis patients

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Cross-sectional Study

Association between serum PGE₂ levels and degree of acid-fast bacilli positivity in sputum of pulmonary tuberculosis patients



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ABSTRACT

Keywords:

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Background: *Mycobacterium tuberculosis* that infected apoptotic macrophages is triggered by PGE₂. Apoptosis suppresses the growth of *Mycobacterium tuberculosis* bacteria, which is shown in the results of acid-fast bacilli (AFB) in the sputum that becomes a marker of the number of bacteria.

Objective: Analyzing the association between serum PGE₂ levels and the positivity of AFB in the sputum of tuberculosis patients.

Methods: A cross-sectional study was carried out from August 2019–July 2020. Serum PGE₂ levels and AFB levels in sputum were collected from participants. Data analysis used the Chi-square test and Spearman's correlation with $p < 0.05$.

Results: The average participants' serum PGE₂ levels were 446.37 ± 510.27 pg/ml, with a median value of 216.95 pg/ml. Most participants had normal serum PGE₂ levels (62.9%). Most participants had a high positivity of AFB in sputum (58.1%). Analysis of the association between serum PGE₂ levels and the degree of AFB positivity in sputum obtained $r = -0.036$ and p -value = 0.780.

Conclusion: There is a weak negative association between serum PGE₂ levels and the degree of AFB positivity in sputum but not statistically significant.

1. Introduction

Tuberculosis (TB) is still a global health problem [1]. The increase in TB cases is accompanied by an increase in drug-resistant TB (DR TB) cases. In the Global Tuberculosis Report, WHO reported that 10 million people were suffering from TB, both new and relapsed cases, with 558,000 of whom had DR TB [2]. Indonesia ranks third in the country with the highest TB incidence globally, both new and relapse cases. The number of new and relapsed TB cases in Indonesia in 2017 was 442,172, and 54% of them were confirmed bacteriologically by either acid-fast bacilli (AFB) sputum staining or sputum culture [3].

The pathogenesis of TB is an interaction between *Mycobacterium tuberculosis* and the host [4]. The process begins with alveolar macrophages and dendritic cells as the first cells facing *Mycobacterium tuberculosis* bacteria. Macrophages' response as the mainline in dealing with *Mycobacterium tuberculosis* infection is influenced by various inflammatory mediators [5]. The failure of macrophages to control the number of *Mycobacterium tuberculosis* will result in the significant growth of

bacteria [6,7].

This condition emphasizes the important role of the host immune system in determining the susceptibility of TB to relapse. Several studies pointed out that Prostaglandin E₂ (PGE₂) affects macrophages as the main cells in the innate immune system. PGE₂ induces apoptosis and inhibits necrosis of macrophages infected with *Mycobacterium tuberculosis* [5,8,9]. Macrophage apoptosis is reported to reduce the growth rate of *Mycobacterium tuberculosis*, which is very important in the elimination mechanism of bacteria that infects the lungs, whereas necrosis plays the opposite role [5,8,10]. When the growth of *Mycobacterium tuberculosis* cannot be inhibited, the number of bacteria will increase. The high number of bacteria is reflected in the degree of phlegm AFB positivity. The higher the value of positivity for AFB in sputum, the greater the number of *Mycobacterium tuberculosis* bacteria contained in each ml of sputum [11]. The higher the number of bacteria, the easier it is to transmit, broader lung damage, and an increased risk of resistance [12,13].

Based on the facts above, this study further revealed the association

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between PGE₂, which represents the innate immune system, and the degree of phlegm AFB positivity, which represents the number of bacteria. This research is important because no similar study was conducted in humans, so it is hoped that this research could provide further research.

2. Methods

2.1. Participants

Participants in this study were both new and relapsed patients with pulmonary tuberculosis. The inclusion criteria were patients diagnosed with pulmonary tuberculosis [3,14], positive sputum examination results for AFB, aged 21–65 years, who cooperated during the research procedure. Meanwhile, the exclusion criteria included patients with risk factors for immunocompromised (AIDS, malignancy, and systemic lupus erythematosus), patients having received anti-tuberculosis drug therapy for their current illness, patients taking non-steroidal anti-inflammatory drugs and/or corticosteroids in the past week.

2.2. Ethical clearance

Participants and their families filled out the consent form before the study. Participants filled out the consent form consciously and without coercion. This study received ethical approval based on the Declaration of Helsinki and obtained the registry of research at the Health Research Ethics Committee in the Hospital.

2.3. Study design

A cross-sectional study was carried out from August 2019–July 2020. The number of participants in this study was 62 patients that were obtained using Ronald Fisher’s classic z transformation formula. The sample collection used a consecutive sampling technique (Fig. 1). Serum PGE₂ levels and levels of AFB sputum were taken from the participants. This study report is by the Strengthening the Reporting of Cohort Studies in Surgery (STROCSS) 2019 guideline [15].

2.4. Measurement of serum PGE₂ level

Serum PGE₂ level is the total concentration of PGE₂ in the blood of pulmonary tuberculosis patients. This examination was carried out by taking 3–5 ml of the patient’s venous blood and analyzed using the Elisa

Kit PGE₂ (pg/ml). Serum PGE₂ level is categorized into high if the value is more than 400 pg/ml, normal if the value is 200–400 pg/ml, and low if the value is less than 200 pg/ml [16].

2.5. Acid-fast bacilli test

Sputum examination was conducted to determine the degree of the participant’s AFB positivity. Sputum collection for participants is carried out by the patient independently in the morning [17] which the participant gets an explanation from a pulmonary specialist regarding effective deep breathing and coughing techniques [18]. The sputum is put into a tube that has been prepared previously and then taken to the laboratory for analysis. The examination of AFB in the participant’s sputum used the acid-fast staining method (Ziehl Nielsen) or the rapid molecular test of sputum with the GeneXpert machine [19]. The degree of phlegm AFB positivity was assessed based on the International Union Against Tuberculosis Lung Disease (IUATLD) standards which were categorized into 2: low (1+ and scanty) and high (2+ and 3+) [19,20].

2.6. Statistical analysis

The analysis in this study used descriptive analysis and bivariate analysis. Descriptive analysis included the presentation of the results descriptively using the distribution table, mean, median, standard deviation, maximum value, and minimum value. Meanwhile, bivariate analysis was used to assess the association between two variables. The association between variables was analyzed using the Chi-Square test and assessed the association strength using the Spearman correlation test. The analysis was declared significant if $p < 0.05$. The analysis was assisted by IBM SPSS Statistics software version 21.0 (IBM Corp., Armonk, NY, USA).

3. Results

3.1. Characteristic of participant

Most participants were male who was 43.37 ± 12.58 years old. Meanwhile, the median of participants’ age was 44.5 years, with the lowest age being 21 years and the highest being 64 years. Some patients had a smoking habit (56.5%) and comorbidity of diabetes mellitus (32.3%). A total of 37 participants were new tuberculosis patients and the rest were relapsed, tuberculosis patients. Most participants had a body mass index (BMI) in the skinny category as much as 53.2%

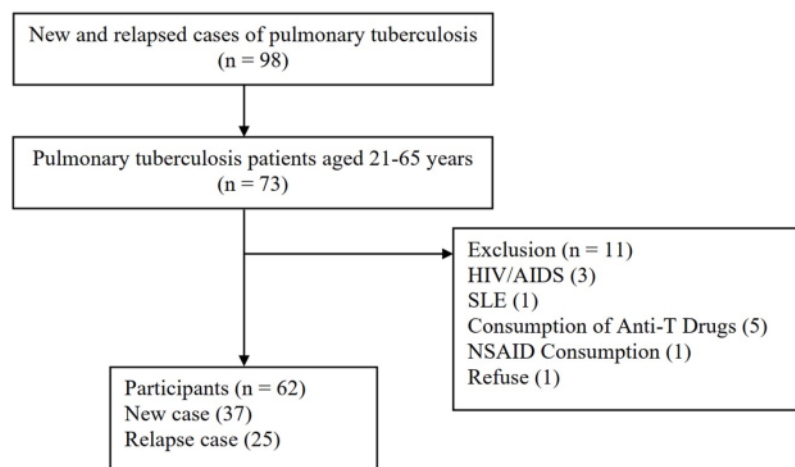


Fig. 1. Participant requirement process.

Table 1
Characteristic of participant.

Variable	n (%)
Sex	
Male	36 (58.1)
Female	26 (41.9)
Education	
Elementary School	8 (12.9)
Junior High School	12 (19.4)
Senior High School	34 (54.8)
College	7 (11.3)
Not attending school	1 (1.6)
History of Diabetes Mellitus	
Yes	20 (32.3)
No	42 (67.7)
History of Tuberculosis Treatment	
New case	37 (59.7)
Relapse	25 (40.3)
Smoking Habit	
Smoking	35 (56.5)
No smoking	27 (43.5)
Degree of Acid-Fast Bacilli Positivity	
Low	26 (41.9)
High	36 (58.1)
Serum PGE ₂ Level	
Low	9 (14.5)
Normal	39 (62.9)
High	14 (22.6)
Body Mass Index	
Skinny (<18.5 kg/m ²)	33 (53.2)
Normal (18.5–25.0 kg/m ²)	24 (38.7)
Fat (>25.0 kg/m ²)	5 (8.1)

(Table 1). The average BMI value was 19.46 ± 4.05 kg/m², with a value range of 14.20–38.28 kg/m².

3.2. Distribution of serum PGE₂ levels in tuberculosis patients

Most participants had normal serum PGE₂ levels (62.9%; Table 1). The average participants had serum PGE₂ levels of 446.37 ± 510.27 pg/ml, with a median value of 216.95 pg/ml. The lowest and highest value of the participants' serum PGE₂ levels were 191.00 pg/ml and 2374.00 pg/ml, respectively. The serum PGE₂ levels of smoking and non-smoking participants was 228.80 (191.0–2,3374.0) pg/ml and 214.40 (198.3–1724.0) pg/ml, respectively. Most serum PGE₂ levels of smoking participants were normal (50%), while the serum PGE₂ levels of non-smoking participants were mostly normal (78%; $p = 0.053$). The median value of serum PGE₂ levels for participants with and without diabetes mellitus was 217.30 (191.0–1986.0) pg/ml and 216.80 (193.0–2374.0) pg/ml, respectively. The value of serum PGE₂ levels of participants with and without diabetes mellitus were 45% and 71%, respectively, indicating that most participants had normal values ($p =$

Table 2
Distribution of serum PGE₂ levels in tuberculosis patients.

Variable	Serum PGE ₂ Levels			p
	Low	Normal	High	
Pulmonary Tuberculosis				
New case	6 (16)	23 (62)	8 (22)	0.292
Relapse case	3 (12)	16 (64)	6 (24)	
Diabetes mellitus				
Yes	4 (20)	9 (45)	7 (35)	0.118
No	5 (12)	30 (71)	7 (17)	
BMI				
Skinny	1 (3)	24 (73)	8 (24)	0.058
Normal	7 (29)	12 (50)	5 (21)	
Fat	1 (20)	3 (60)	1 (20)	
Smoking				
Yes	6 (18)	17 (50)	11 (32)	0.053
No	3 (11)	22 (78)	3 (11)	

Abbreviation: BMI = body mass index.

0.118; Table 2).

Most of the participants' serum PGE₂ levels were normal in both groups of participants with a new diagnosis of pulmonary tuberculosis (62%) and relapsed (64%; $p = 0.292$). The median value of serum PGE₂ levels for participants diagnosed with new pulmonary tuberculosis was 215.70 (191.0–1724.0) pg/ml and participants diagnosed with relapsed pulmonary tuberculosis was 224.40 (193.2–2374.0) pg/ml. Participants' serum PGE₂ levels that were categorized by BMI were mostly normal, with 73% of skinny participants, 50% of normal participants, and 60% of fat participants ($p = 0.058$; Table 3). The median value of serum PGE₂ levels of participants with BMI in the skinny category was 222.60 (194.3–1986.0) pg/ml, normal was 210.30 (191.0–2374.0) pg/ml, and fat was 216.40 (199.0–1497.0) pg/ml.

3.3. Distribution of positivity of acid-fast bacilli in sputum of tuberculosis patients

Most participants had a high degree of AFB positivity in sputum as much as 58.1% (Table 1). Most participants who were diagnosed with new cases of pulmonary tuberculosis had a high degree of AFB positivity (68%). Meanwhile, most participants diagnosed with relapsed pulmonary tuberculosis had a low positivity degree (56%; $p = 0.065$). Some participants had a high degree of AFB positivity in participants with and without a history of diabetes mellitus of 65% and 55%, respectively ($p = 0.455$). Participants' BMI was categorized into 3, namely skinny, normal, and fat, in which some participants had a high degree of AFB positivity ($p = 0.561$). Most smoking (56%) and non-smoking (61%) participants had high positivity of AFB ($p = 0.798$; Table 3).

4. Association between serum PGE₂ levels and positivity of acid-fast bacilli in sputum of tuberculosis patients

The results showed that most participants with low (89%) and high (71%) serum PGE₂ levels had a high positivity of AFB in sputum as much as 89%. Meanwhile, participants with normal serum PGE₂ levels had a low positivity degree of AFB in sputum as much as 54% ($p = 0.036$). The strength of the association between serum PGE₂ levels and the degree of AFB positivity in sputum obtained $r = -0.036$ and p -value = 0.780 (Table 4).

5. Discussion

PGE₂ is a derivative of arachidonic acid produced by various inflammatory cells, especially macrophages. PGE₂, as an inflammatory mediator, plays a role in regulating various cell functions, namely macrophages, T cells, etc. In addition, PGE₂ plays a role in various body functions such as blood pressure regulation, temperature regulation, gastric protection, and childbirth [21]. Under various conditions such as

Table 3
Distribution of positivity of acid-fast bacilli in sputum of tuberculosis patients.

Variable	Degree of Acid-Fast Bacilli Positivity		p
	Low (%)	High	
Pulmonary Tuberculosis			
New case	12 (32)	25 (68)	0.065
Relapse case	14 (56)	11 (44)	
Diabetes Mellitus			
Yes	7 (35)	13 (65)	0.455
No	19 (45)	23 (55)	
BMI			
Skinny	15 (45)	18 (55)	0.561
Normal	10 (42)	14 (58)	
Fat	1 (20)	4 (80)	
Smoking			
Yes	15 (45)	19 (56)	0.798
No	11 (39)	17 (61)	

Abbreviation: BMI = body mass index.

Table 4
Association between PGE₂ levels and positivity of acid-fast bacilli in the sputum of tuberculosis patients.

Variable	Tuberculosis Positivity		p ^a	r	p ^b
	Low	High			
PGE ₂ Levels					
Low	1 (11)	8 (89)	0.036	-0.036	0.780
Normal	21 (54)	18 (46)			
High	4 (29)	10 (71)			

Note: p^a = Chi-square test; p^b = Spearman's correlation test.

changes in environmental temperature, hunger conditions, stress, PGE₂ will be produced so that levels in the body will rise and fall in various ways [22].

Schoenberger et al. reported an increase in serum PGE₂ levels in patients with diabetic retinopathy [23]. A study conducted by Lo et al. showed that the increase in serum PGE₂ levels was due to the upregulation of the cyclooxygenase-2 (COX₂) enzyme in patients with diabetes mellitus [24]. Kumar et al. reported differences in plasma PGE₂ levels in TB patients compared to TB-DM [16]. These results are inconsistent with various studies that reported increased levels of PGE₂ in smokers. Amadio et al. reported an increase in PGE₂ production in smokers due to the modulation of expression of tissue factors exposed to cigarette smoke [25]. Chen et al. in their study also reported the role of cigarette smoke in increasing PGE₂ production [26].

The condition obtained in this study seemed to occur because of the patient's experience factor. In patients with relapse cases, the experience of suffering from TB in the past will make the patient who has a cough immediately come to the health facility. Meanwhile, new case patients ignore the cough complaint that leads to accompanying complaints such as weight loss, hemoptysis, or fever. When these accompanying complaints occur, the course of TB disease would be long enough to increase the number of bacteria [1].

The profile of serum PGE₂ levels showed that the average participants had 446.23 pg/ml, with a standard deviation of 510.27 pg/ml. According to some literature, normal serum PGE₂ levels range from 200 to 400 pg/ml [16]. PGE₂ is a derivative of arachidonic acid produced mainly by inflammatory cells to face invading pathogens from outside. The effect of PGE₂ will trigger apoptosis of macrophages infected with *Mycobacterium tuberculosis* [4]. Macrophage apoptosis will have an elimination effect because *Mycobacterium tuberculosis* bacteria can be destroyed. PGE₂ also suppresses macrophage necrosis which can lead to bacterial dissemination. Increased levels of PGE₂ are associated with a decrease in the number of bacteria in the lung [7].

The negative association between serum PGE₂ levels and the degree of phlegm AFB positivity is by a study conducted by Dietzold and Amaral. Dietzold et al. reported that high levels of PGE₂ and low levels of LXA₄ suppress the growth of *Mycobacterium tuberculosis* [7]. Amaral et al. also reported that PGE₂ is associated with macrophage apoptosis in vitro. Apoptotic macrophages infected with *Mycobacterium tuberculosis* will increase the elimination of these bacteria [4]. The two studies above reported a significant association between PGE₂ and the growth of *Mycobacterium tuberculosis*. The statistical analysis results of this study showed that the association between serum PGE₂ levels and the degree of AFB positivity was not statistically significant. The main difference between this study and the two studies above is that both were carried out on mice and in vitro, whereas this study was conducted on pulmonary TB patients with various complications and uncontrollable comorbidities.

The results of this study can be used as consideration for conducting further research on the predictor factors for the positivity of AFB in pulmonary TB patients. The use of PGE₂ together with LXA₄ is expected to be able to assist clinicians in predicting the level of AFB positivity in pulmonary TB patients with specific chest X-ray images but difficulty in expectorating phlegm. In addition, in the future study it can be

considered to analyze the comparison of PGE₂ in TB patients, smokers patients, smokers with tuberculosis, etc.

Nevertheless, this study has several limitations. First, extreme serum PGE₂ levels were found in some research subjects. This can be caused by various factors that can increase PGE₂ levels that cannot be controlled. Second, this study only examined PGE₂ levels in TB patients without comparing them with PGE₂ levels in healthy persons, so it cannot be used as a predictor factor for the degree of positivity of AFB with sputum.

6. Conclusion

The average age of new and relapsed pulmonary TB patients is 43.37 years, mostly male, have a high school education, have a smoking habit, have a low BMI, and have no history of DM. The median serum PGE₂ level of new and relapsed pulmonary TB patients was 216.95 pg/ml. The majority of new pulmonary TB patients have a high degree of positivity for AFB in sputum, but relapsed pulmonary TB patients have a low degree of positivity for AFB. This study finds a weak negative association between serum PGE₂ levels and the degree of phlegm AFB positivity but not statistically significant.

4

Please state any sources of funding for your research

None.

Ethical approval

We have conducted an ethical approval base on Declaration of Helsinki at Ethical Committee in Dr. Soetomo General Academic Hospital, Surabaya, Indonesia.

Consent

Written informed consent was obtained from the patient.

5

Author contribution

All authors contributed toward data analysis, drafting and revising the paper, gave final approval of the version to be published and agree to be accountable for all aspects of the work.

3

Registration of research studies

1. Name of the registry: Health Research Ethics Coommittee in the Dr. Soetomo General Academic Hospital, Surabaya, Indonesia.
2. Unique Identifying number or registration ID: 1355/KEKP/VII/2019.
3. [Perlink to your specific registration \(must be publicly accessible and will be checked\):-.](#)

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1

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Declaration of competing interest

The authors declare that they have no conflict of interest.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.amsu.2021.103008>.

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