

## ABSTRACT

**ANALYSIS OF DIFFERENCES OF SERUM THROMBOXANE B<sub>2</sub> LEVEL AFTER TAKING ASETOSAL IN ACUTE THROMBOTIC STROKE WITH DIABETES MELLITUS AND NON DIABETES MELLITUS**

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**Background:** The underlying pathophysiology of thrombotic stroke is a formation of thrombus in cerebrovascular, causing decrease in blood flow to certain cerebral region and thus manifested as neurological deficit. Acetosal as a secondary prevention therapy for thrombotic stroke, exert its pharmacological effect by inhibiting COX-1 enzyme so it decrease Thromboxane A<sub>2</sub> (TxA<sub>2</sub>) synthesis, a potent agonist for platelet activation. Diabetes Mellitus known as one of the major risk factor for thrombotic stroke, shows a platelet hyperactivity. It is thought that the higher rate of recurrence in thrombotic stroke is due to platelet hyperactivity. To analyze the pharmacological effect of ASA in thrombotic stroke, especially in diabetic patients, we measure the level of thromboxane A<sub>2</sub> (TxA<sub>2</sub>) through the measurement of its stabile Metabolite, TxB<sub>2</sub>.

**Objectives:** To analyze the difference of serum TxB<sub>2</sub> levels in acute thrombotic stroke patients with diabetes mellitus (DM) and non DM.

**Method:** Data were collected from August – October 2016. Venous blood sample were obtained to extract serum TxB<sub>2</sub> level before and after 5-7 days taking asetosal 100 mg. Protocol of this study was approved by Ethical Committee of Dr. Soetomo Teaching Hospital Surabaya. The informed consents must be signed as a proof of participation. The measured data of serum TxB<sub>2</sub> were analyzed using paired t-test to compare parametric data within group and independent t-test to compare parametric data between DM and non DM group.

**Result:** Total 27 inpatients were participated in this study divided into 15 and 12 patients in DM and non DM group. Sixty percent (60%) and 67% subject in diabetic and non diabetic group were men. Ages and sex between two groups were not significantly different ( $p > 0.05$ ). Serum TxB<sub>2</sub> before taking ASA (pre ASA) in diabetic and non diabetic group was 0.56 – 55.77 ng/mL and 4.24 – 60.31 ng/mL with mean value was  $16.43 \pm 16.08$  ng/mL and  $27.36 \pm 21.04$  ng/mL respectively. After taking ASA 5-7 days (post ASA), we measure serum TxB<sub>2</sub> level in diabetic and non diabetic group was 0.90 – 6.58 ng/mL and 0.95 – 16.23 ng/mL with mean value was  $2.93 \pm 1.83$  ng/mL and  $5.36 \pm 4.06$  ng/mL respectively. Serum TxB<sub>2</sub> level after taking ASA 100 mg decrease significantly in both group ( $p = 0.006$  in diabetic group and  $p = 0.005$  in non diabetic group). Mean reduction of serum TxB<sub>2</sub> level in diabetic ( $13.49 \pm 15.98$  ng/mL) is less than non diabetic group ( $22.00 \pm 21.65$  ng/mL). Statistical analysis shows that the mean reduction of serum TxB<sub>2</sub> level between two groups were not significantly different ( $p > 0.05$ ).

**Conclusion:** ASA 100 mg could decrease serum TxB<sub>2</sub> level significantly in both Diabetic and non Diabetic group. Mean reduction of serum TxB<sub>2</sub> level in non diabetic group was greater than in diabetic group although it is not significantly different.

**Keyword:** Thrombotic Stroke, Acetosal, platelet activation, Diabetes Mellitus, Thromboxane B<sub>2</sub>, TxB<sub>2</sub>, Thromboxane A<sub>2</sub>, TxA<sub>2</sub>.