

RESEARCH ARTICLE

**Folic Acid Usual Doses Decrease
the Buccal Micronucleus Frequency on Smokers**

Yuktiana Kharisma,¹ Meta Maulida Damayanti,² Fajar Awaliya Yulianto,³
Santun Bhekti Rahimah,¹ Winni Maharani,⁴ Meike Rachmawati,²
Herri S. Sastramihardja,¹ Muhammad Alief Abdul 'Aziiz,⁵ Muhammad Ilham Halim⁵

¹Department of Pharmacology, ²Department of Phatology, ³Department of Public Health,
⁴Department of Microbiology, ⁵Medical Undergraduate Study Program,
Faculty of Medicine, Universitas Islam Bandung, Bandung, Indonesia

Abstract

Cigarette contains toxic chemical compounds that trigger DNA instability. Initial genotoxic oral cavity characterized by the appearance of micronucleus (MN) in the buccal mucosa. Folate is needed in maintaining DNA stability. This study aimed to compare the effects of folic acid usual doses (400 mcg and 1.000 mcg) on the MN frequency of buccal mucosa in active smokers. It is a clinical trial conducted in November 2018 in the Laboratory of the Faculty of Medicine, Universitas Islam Bandung of 53 active smokers who divided into two treatment groups. Group A was administered by 400 mcg and group B 1,000 mcg folic acid supplementation within three weeks. The buccal mucosa smear stained with hematoxylin-eosin (HE) and observed through a light microscope with 100× and 400× magnification. Data were analyzed by the Wilcoxon test statistically. The results showed that there was a significant decrease ($p=0.00$) in MN frequency in folic acid supplementation for three weeks, namely group A=6.39±3.92 and group B=6.93±5.82 in pre-supplementation, and group A=3.80±2.66 and group B=3.31±2.71 post-supplementation of folic acid. Giving a dose of 400 mcg and 1,000 mcg for three weeks did not provide significant results ($p=0.94$) with Kruskal-Wallis test. In conclusion, administration of folic acid at usual dose give results to a decrease in the buccal mucosa MN frequency in active smokers.

Key words: Active smokers, folic acid, micronucleus

**Asam Folat Dosis Lazim Menurunkan
Frekuensi Mikronukleus Mukosa Bukal pada Perokok****Abstrak**

Asap rokok mengandung senyawa kimia toksik yang memicu ketidakstabilan DNA. Deteksi genotoksik awal rongga mulut ditandai dengan kemunculan mikronukleus (MN) pada mukosa bukal. Folat diperlukan dalam menjaga kestabilan DNA. Penelitian ini bertujuan mengetahui efek asam folat dosis lazim (400 mcg dan 1.000 mcg) terhadap frekuensi MN mukosa bukal pada perokok aktif. Penelitian ini merupakan uji klinis yang dilakukan pada bulan November 2018 di Laboratorium Fakultas Kedokteran, Universitas Islam Bandung terhadap 53 perokok aktif yang dibagi ke dalam dua kelompok perlakuan. Kelompok A mendapatkan suplementasi asam folat 400 mcg dan kelompok B mendapatkan suplementasi asam folat 1.000 mcg selama tiga pekan. Apus mukosa bukal diwarnai dengan *hematoxylin-eosin* (HE) dan diamati melalui mikroskop cahaya dengan pembesaran 100× dan 400×. Data dianalisis dengan uji Wilcoxon secara statistik. Hasil penelitian menunjukkan bahwa terdapat penurunan frekuensi MN yang signifikan ($p=0.00$) terhadap suplementasi asam folat selama tiga minggu, yaitu kelompok A=6,39±3,92 dan kelompok B=6,93±5,82 pada pre-suplementasi, serta kelompok A=3,80±2,66 dan kelompok B=3,31±2,71 pada pascasuplementasi asam folat. Pemberian dosis 400 mcg dan 1.000 mcg selama tiga minggu tidak memberikan hasil yang bermakna ($p=0,94$) berdasar atas Uji Kruskal-Wallis. Simpulan, pemberian asam folat dosis lazim memberikan hasil baik terhadap penurunan frekuensi MN mukosa bukal pada perokok aktif.

Kata kunci: Asam folat, mikronukleus, perokok aktif

Received: 1 February 2019; Revised: 10 August 2019; Accepted: 17 August 2019; Published: 31 August 2019

Correspondence: Yuktiana Kharisma. Department of Pharmacology, Faculty of Medicine, Universitas Islam Bandung. Jln. Tamansari No. 22, Bandung 40116, West Java, Indonesia. Phone: (+6222) 4203368. Facsimile: +(6222) 4231213. Mobile: +6287821360031. E-mail: yuktiana@gmail.com

Introduction

Smoking is one of the unethical lifestyles associated with genetic damage. Tobacco contains toxic chemical compounds and free radicals which can trigger cell damage and induces mutations in the deoxyribonucleic acid (DNA) which will increase the risk of malignancy in the oral cavity, lungs, and other non-communicable diseases.¹⁻³

Indonesia had the first largest number of smokers in the world in 2015; there were 76.2% of smokers who were aged 15 years and older. West Java contributes to the 4-6 times increase in the proportion of active smokers at 1% and passive smokers at 62.8%.³ A total of 10.9 million residents of the Bandung city became smokers by spending 12-29 cigarettes per day.⁴

The buccal mucosa (BM) is a covered flat epithelial layer that functions as a protection for the oral cavity against potential carcinogenic substances that can produce potential reactive products in the oral cavity. The cells in the BM layer used to detect the occurrence of initial genotoxins caused by carcinogenic inhalant substances such as cigarette smoke. Exfoliated buccal cells used as non-invasive indicators of genotoxic effects characterized by the appearance of MN in cells. The frequency of the micronucleus is proportional to the degree of exposure and the potential of carcinogenic substances.⁵ An examination of buccal MN cells can be done using HE staining through observation of a light microscope or specific DNA stainings such as acridine orange and Feulgen. This method can be used regularly as an initial screening in groups of individuals who are susceptible to exposure to carcinogenic substances. The presence of MN is a reliable indicator of chromosome damage due to the loss of the whole and partial chromosomes.⁶

Free radicals in cigarettes cause oxidative stress and can lead to atherosclerosis, diabetes, cancer, neurodegenerative disorders, cardiovascular disorders, and other chronic conditions.^{7,8} High free radicals will cause endogenous antioxidants in the body to decline and are not effective against the harmful effects of free radicals. Exogenous antioxidants or other micronutrient supplements are needed to prevent the harmful effects of free radicals for becoming a cofactor for antioxidants to maintain optimal gene health.^{9,10} One of the essential micronutrients is folic acid.⁸

Folate need in DNA stability by synthesizing with deoxythymidine monophosphate (dTMP) derived from deoxyuridine monophosphate

(dUMP). It plays a role in donating methyl in folate-methionine methylation. Vitamin B12 deficiency also produces high uracil, which causes restrictions on folate synthesis through the inhibition of the synthetic of 5,10 methylenetetrahydrofolate. The result is in DNA mutations, single-stranded DNA formation, DNA fragmentation, chromosomal damage, and MN formation. Folate and vitamin B12 need for the synthesis of methionine through remethylating of homocysteine (HCy) and the S-adenosylmethionine (SAM) synthetic; a typical methyl donor needed for the maintenance of DNA methylation patterns.^{6,7} Based on the description above, the researcher was interested to see the effects of folic acid at usual dosage (400 mcg and 1,000 mcg) to the frequency of buccal mucosa MN in active smokers.

Methods

The 60 participants selected regarding the duration of smoking habits; minimum for one year. The participants should have no alcoholic consumption habits, systemic diseases, allergy to folic acid, undergoing radiotherapy, performing oral x-ray examination procedures in the last month, amalgam dental fillings, and chronic infection, and inflammation of the oral cavity. Fifty-three participants follow the entire three weeks of the research period.

The samples swabbed from the buccal mucosa using a buccal brush at pre-supplementation and post-supplementation group of 400 mcg and 1,000 mcg folic acid. The buccal smeared on the glass object and stained by HE. The results were observed on 100 cells and evaluated with 100× and 400× magnification using a light microscope

The data analyzed using the Statistical Package for Social Sciences software (SPSS Inc., Chicago, IL, United States). Data presented as a mean and standard deviation. The differences between the variables were analyzed using the Wilcoxon and Kruskal-Wallis test. A p value of <0.05 was considered significant. This research had approved by the Health Research Ethics Committee of Faculty of Medicine Universitas Islam Bandung with the ethical clearance number: 379/Komite Etik.FK/X/2018.

Results

The MN in the buccal mucosa found in varying amounts between participants. Micronucleus

observations are carried out through a light microscope with 100× and 400× magnification with calculations performed on 100 cells in each buccal mucosa smear. The size of MN is 1/3–1/6 times than the central nucleus, but they have similar round shape and density. The micronucleus images found in this study presented in Figure 1 and 2 below.

The comparison of MN frequency between the participant who were provided 400 mcg and 1,000 mcg folic acid as pre-supplementation and post-supplementation are shown on Table.

In the group of participants who consumed 400 mcg (group A) and 1,000 mcg (group B) the frequency of the post-supplementation micronucleus was lower (group A=3.80±2.66; group B=3.31±2.71) than the pre-supplementation MN (group A=6.39±3.92; group B=6.93±5.82) respectively. There was a significant difference between the frequency of the MN pre-supplementation and post-supplementation of folic acid, both in the group provided with folic acid at 400 mcg and 1,000 mcg. Therefore the implementation of 400 mcg of folic acid was able to significantly reduce the buccal mucosa MN frequency of active smokers, while the dosage

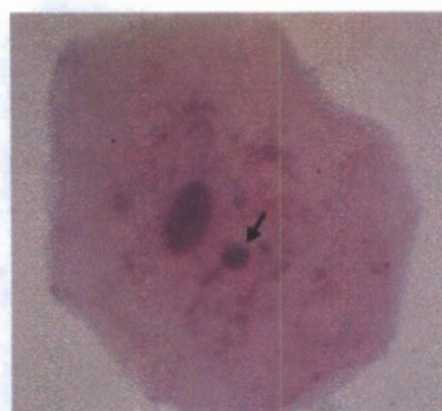


Figure 2 Micronucleus in Buccal Mucosa Cell (arrow head)

Observed through 400× magnification with HE stained

of the 400 mcg and 1,000 mcg did not give any significant effect.

Discussion

Tobacco use habits in various forms are associated with an increased risk of oral cancer.¹¹ Cigarettes

Table Comparison of MN Frequency Between Group

Supplementation	MN Frequency Group A	MN Frequency Group B	p Value
Pre-supplementation	6.39±3.92	6.93±5.82	0.00*
Post-supplementation	3.80±2.66	3.31±2.17	0.00*
Δ	2.59±1.26	3.32±3.65	0.94**

Group A: participants who were provided with 400 mcg folic acids; Group B: participants who were provided with 1000 mcg folic acids; *Wilcoxon test, significance if p<0.05; **Kruskal-Wallis test, significance if p<0.05

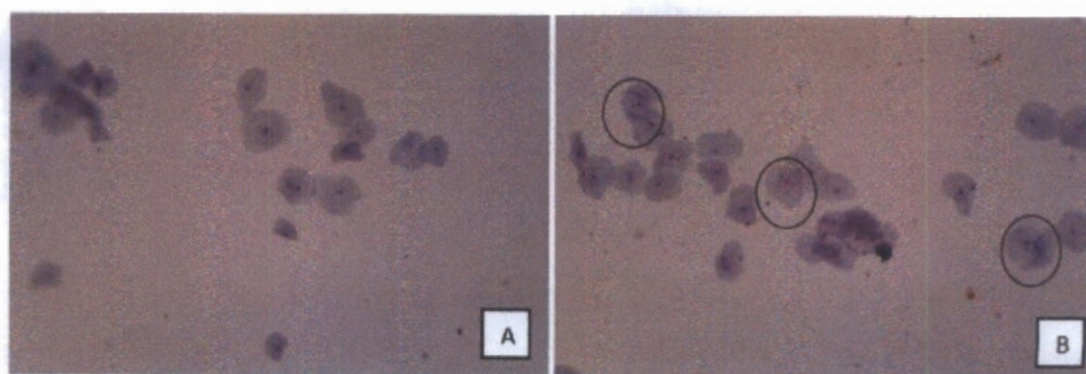


Figure 1 (A) Normal Cell, (B) Micronucleus in Buccal Mucosa

Observed through 100× magnification with HE stained

are complex mixtures of various substances that are genotoxic and carcinogenic to oral epithelial cells.¹² Micronucleus is the result of chromosome changes originating from fragments or whole chromosomes that lag in the anaphase during the break up of chromosomes induced by many genotoxic agents.

Several studies have shown different results regarding the association between cigarette consumption and increasing MN frequencies in buccal mucosa cells.^{13,14} In this study, the results of the increase of MN frequency was varied between individuals, with an average value of 6.39 ± 3.92 in group A and 6.93 ± 5.82 in group B. The study by Nersesyan showed that smoking induced MN and other chromosomal anomalies in buccal cells. The cytogenetic effects of cigarettes on exfoliated buccal cells are inconsistent, and these findings suggest that this mismatch may partly be due to variations in tar and nicotine levels of cigarettes smoked by participants.

Folic acid is a part of vitamin B which act as an antioxidant. Several studies have shown that folic acid supplementation can reduce the MN frequency in lymphocytes, buccal mucous epithelial cells significantly in high-risk groups (smokers, alcoholics, patients with chronic diseases). Folic acid provided in high dosage in order to protect cells from the occurrence of DNA damage due to the exposure to high oxidants. Supplementation of $2 \times 1,000$ mcg and 3×500 mcg of folic acid was provided for 30 days and was shown to decrease the buccal mucosa MN frequency in previous studies.¹⁶ Our study used folic acid supplementation once daily of 400 mcg, and 1,000 mcg dose indicates the improvement of MN frequency by 50% between the pre-supplementation and post-supplementation of folic acid. Nevertheless, the MN reduction not synchronized with the average MN frequency of healthy people about 0.5–2.5 MN per 1,000 cells according to the research implemented by Jois et al.¹⁷

Several factors that influence the form of anomalies include smoking habits, local and systemic infections, lifestyle, and the other chronic diseases underlying it. In this study, participants had the same risk factors for the emergence of MN based on data obtained through filling out questionnaires in participants at the beginning of the study. Some participants had an adequate to bad oral health index in the two treatment groups. It could be one of the supporting factors for the

high frequency of MN in participants. There is a strong relationship between the plaque index and the calculus with an increase in the frequency of micronucleus in the cell oral epithelium. Dental and calculus plaques are places for oral bacteria that produce chronic bacterial infections.¹² Chronic infections lead to chronic inflammatory processes that are carcinogenic and are associated with the formation of clastogenic genetic damage in oral epithelial cells.¹⁸

Conclusion

This study concluded that administration of folic acid at a dose of 400 mcg and 1,000 mcg give similar results to a decrease in the buccal mucosa MN frequency in active smokers.

Conflict of Interest

The authors declare no conflict of interests.

Acknowledgments

The author's thanks to all the Biomedical Laboratory staff of the Faculty of Medicine of Universitas Islam Bandung for support of this study.

References

1. World Health Organization (WHO). Prevalance of tobacco smoking [Internet]. 2015 [cited 2017 February 9]. Available from: http://gamapserver.who.int/gho/interactive_charts/tobacco/use/atlas.html.
2. Valavanidis A, Vlachogianni F, Fiotakis K. Tobacco smoke: involvement of reactive oxygen species and stable free radicals in mechanisms of oxidative damage, carcinogenesis and synergistic effects with other respirable particles. *Int J Environ Res Public Health*. 2009;6(2):445–62.
3. Badan Penelitian dan Pengembangan Kesehatan, Kementerian Kesehatan Republik Indonesia. Riset kesehatan dasar (Riskesdas) 2013. Jakarta: Badan Litbang Kesehatan, Kemenkes RI; 2013.
4. Dio. 30 persen warga Bandung perokok [Internet]. Tobacco Control Support Centre Indonesia. 2011 [cited 2017 February 9]. Available from: <http://www.tcsc-indonesia.org/30-persen-warga-bandung-perokok/>.

5. Devi P, Thimmarasa VB, Melhotra V, Arora P. Micronucleus assay for evaluation of genotoxicity in potentially malignant and malignant disorders. *J Indian Acad Oral Med Radiol.* 2011;23(2):97-100.
6. Fareed M, Afzal M, Siddique YH. Micronucleus investigation in buccal mucosal cells among pan masala/gutkha chewers and its relevance for oral cancer. *Biol Med.* 2011;3(2):8-15.
7. Thomas P, Wu J, Dhillon V, Fenech M. Effect of dietary intervention on human micronucleus frequency in lymphocytes and buccal cells. *Mutagenesis.* 2011;26(1):69-76.
8. Noori S. An overview of oxidative stress and antioxidant defensive system [Internet]. 3 November 2012 [cited 2017 February 9]. Available from: <http://dx.doi.org/10.4172/scientificreports.413>.
9. Yuan G, Sun B, Yuan J, Wang Q. Effect of 1-methylcyclopropene on shelf life, visual quality, antioxidant enzymes and health-promoting compounds in broccoli florets. *Food Chem.* 2010;118(3):774-81.
10. Fenech M. Genome health nutrigenomics and nutrigenetics – diagnosis and nutritional treatment of genome damage on an individual basis. *Food Chem Toxicol.* 2008;46(4):1365-70.
11. Giustarini D, Dalle-Donne I, Tsikas D, Rossi R. Oxidative stress and human diseases: origin, link, measurement, mechanisms, and biomarkers. *Crit Rev Clin Lab Sci.* 2009;46(5-6):241-81.
12. Saeed SH, Younis WH. A cytopathological study of the effect of smoking on the oral epithelial cells in relation to oral health status by the micronucleus assay. *J Bagh College Dentistry.* 2012;24(3):67-70.
13. Pradeep MR, Guruprasad Y, Jose M, Saxena K, Deepa K, Prabhu V. Comparative study of genotoxicity in different tobacco related habits using micronucleus assay in exfoliated buccal epithelial cells. *J Clin Diagn Res.* 2014;8(5):ZC21-4.
14. Lazalde-Ramos BP, Zamora-Perez A, Sosa-Macías M, Guerrero-Velázquez C, Zúñiga-González GM. DNA and oxidative damages decrease after ingestion of folic acid in patients with type 2 diabetes. *Arch Med Res.* 2012;43(6):476-81.
15. Nersesyan A, Muradyan R, Kundi M, Knasmueller S. Impact of smoking on the frequencies of micronuclei and other nuclear abnormalities in exfoliated oral cells: a comparative study with different cigarette types. *Mutagenesis.* 2011;26(2):295-301.
16. Fenech M, Bonassi S. The effect of age, gender, diet and lifestyle on DNA damage measured using micronucleus frequency in human peripheral blood lymphocytes. *Mutagenesis.* 2011;26(1):43-9.
17. Jois HS, Kale AD, Mohan Kumar KP. Micronucleus as a potential biomarker of oral carcinogenesis. *IJDA.* 2010;2(2):197-202.
18. Bloching M, Reich W, Schubert J, Grummt T, Sandner A. Micronucleus rate of buccal mucosal epithelial cells in relation to oral hygiene and dental factors. *Oral Oncol.* 2008;44(3):220-6.