

# Molecular and Cellular Biomedical Sciences

Volume 6, Number 2, July 2022

## RESEARCH ARTICLES

**The Effects of *Moringa oleifera* Leaves on Complete Blood Count, Renal and Liver Functions as Potential Therapy for Malnutrition**

*Gamar Musa Kodi, Howeida Abdullah Mustafa, Alkhair Abd Almahmoud Idris; p.55-62*

**Association of CYP2A6 Genetic Polymorphism and Lung Cancer in Female Never Smokers**

*R.A Henny Anggriani, Noni Novisari Soeroso, Setia Putra Tarigan, Putri Chairani Eyanoer, Hidayat; p.63-9*

**Genetic Variant of Vascular Endothelial Growth Factor (VEGF)-A rs699947 is Associated with Preeclampsia**

*Anggelia Puspasari, Rina Nofri Enis, Herlambang; p.70-6*

**Development of Recombinant Immunoblot Assay Diagnostic Test Based on HIV-1 in Indonesia**

*Jeanne Elvia Christian, Silvia Tri Widyaningtyas, Budiman Bela; p.77-84*

**An Experimental Study on the Healing Effect of Water to Traditional Sudanese Liquor (Aragi)-induced Stomach Peptic Ulcers**

*Entisar Kuku Yousif, Howeida Abdullah Mustafa, Alkhair Abd Almahmoud Idris; p.85-8*

**Correlation between Genetic Polymorphism of CYP2A13 Genotype and Lung Cancer in Female Passive Smokers**

*Nurul Ramadhani, Noni Novisari Soeroso, Setia Putra Tarigan, Putri Chairani Eyanoer, Hidayat; p.89-95*

**Utilization of Expired Platelet Concentrate for Production of Human Platelet Lysate as a Medium for T47D Cell Propagation**

*Diani Mentari, Relita Pebrina, Diah Nurpratami; p.96-103*

Print ISSN: 2527-4384

Online ISSN: 2527-3442

<https://www.cellbiopharm.com/ojs/index.php/MCBS>

Cell and  
Biopharmaceutical  
Institute



# Molecular and Cellular Biomedical Sciences

## PRINCIPAL CONTACT

### MCBS OFFICE

Prodia Tower 8th Floor, Jl. Kramat Raya No.150, Jakarta Pusat 10430  
Email: mcbs\_office@cellbiopharm.com

## SUPPORT CONTACT

Nurrani Mustika Dewi  
Email: nurranimustika@gmail.com

## EDITOR IN CHIEF

Dr. Anna Meiliana  
Postgraduate Program in Clinical Pharmacy, Faculty of Pharmacy,  
Padjadjaran University, Indonesia

## EDITORIAL BOARD

Dr. Ahmad Faried  
Department of Neurosurgery, Faculty of Medicine  
Universitas Padjadjaran, Indonesia

Prof. Anak Iamaroon  
Department of Oral Biology and Diagnostic Sciences,  
Faculty of Dentistry, Chiang Mai University, Thailand

Dr. Bin Ren  
Division of Hematology and Oncology, Department of Medicine,  
Medical College of Wisconsin, United States of America

Dr. Ines Atmosukarto  
College of Medicine, Biology & Environment,  
Australian National University, Australia

Dr. Mutsumi Miyauchi  
Department of Oral and Maxillofacial Pathobiology, Basic Life Sciences,  
Institute of Biomedical and Health Sciences, Hiroshima University, Japan

Dr. Thai Yen Ling  
Department of Pharmacology,  
College of Medicine, National Taiwan University, Taiwan

Dr. Wahyu Widowati  
Department of Biology,  
Faculty of Medicine, Maranatha Christian University, Indonesia

## PEER-REVIEWERS

Prof. Akihiro Shimosaka  
Hematology Institute, Peking Union Medical College, China

Dr. Ditha Diana  
Anatomic Pathology Laboratory,  
Prodia Clinical Laboratory, Indonesia

Prof. Gerard Pals  
Department of Clinical Genetics  
Amsterdam University Medical Center, Netherlands

Prof. Hee Young Shin  
Department of Pediatrics, Cancer Research Institute,  
Seoul National University College of Medicine, South Korea

Prof. Hiroyuki Kumamoto  
Division of Oral Pathology, Department of Oral Medicine and Surgery,  
Graduate School of Dentistry, Tohoku University, Japan

Dr. Irawan Satriotomo  
Center for Translational Research in Neurodegenerative Disease (CTRND),  
University of Florida, United States of America

Dr. Laifa Annisa Hendarmin  
Section of Biology, Faculty of Medicine and Health Sciences,  
Syarif Hidayatullah State Islamic University, Indonesia

Prof. Yen Hua Huang  
Department of Biochemistry and Molecular Cell Biology,  
Graduate Institute of Medical Sciences College of Medicine,  
Taipei Medical University, Taiwan

Dr. Yudi Her Oktaviono  
Department of Cardiology and Vascular Medicine,  
Faculty of Medicine / Dr. Soetomo Hospital, Airlangga University, Indonesia

## FOCUS AND SCOPE

Molecular and Cellular Biomedical Sciences (MCBS) is an open access, peer-reviewed journal that supports all topics in Biology, Pathology, Pharmacology, Biochemistry, Histology and Biomedicine in the aspect of molecular and cellular.

MCBS is dedicated to publish review and research articles. The editors will carefully select manuscript to be delivered for peer-reviewing process. Therefore MCBS is committed to present only the valuable and recent scientific findings.

## SECTION POLICIES

### REVIEW ARTICLE

Review Article should consist of no more than 10,000 words, not including the words in abstract, references, table, figure, and figure legend. The manuscript should have no more than six figures and/or tables in total and no more than 200 references.

### RESEARCH ARTICLE

Research Article should consist of no more than 3,500 words, not including the words in abstract, references, table, figure, and figure legend. The manuscript should have no more than six figures and/or tables in total and no more than 40 references.

## PEER REVIEW PROCESS

All manuscripts submitted to Molecular and Cellular Biomedical Sciences will be selected and blind peer-reviewed by 2 or more reviewers when necessary, to present valuable and authentic findings. All details will also be reviewed, including appropriate title; content reflecting abstract; concise writing; clear purpose, study method and figures and/or tables; and summary supported by content. The reviewing process will take generally 2-3 months depends on sufficiency of information provided.

Peer-reviewers were selected based on their specialties that fit to the topic. Additional reviewer/s can also be pointed when necessary. Author can suggest reviewer/s that not having publication together within five years and should not be member/s of the same research institution.

## PUBLICATION FREQUENCY

Molecular and Cellular Biomedical Sciences is published triannually (in March, July, and November).

## OPEN ACCESS POLICY

This journal provides immediate open access to its content on the principle that making research freely available to the public supports a greater global exchange of knowledge.

# Molecular and Cellular Biomedical Sciences

## ARCHIVING

This journal utilizes the LOCKSS system to create a distributed archiving system among participating libraries and permits those libraries to create permanent archives of the journal for purposes of preservation and restoration.

## PLAGIARISM SCREENING POLICY

All manuscripts submitted to Molecular and Cellular Biomedical Sciences will be screened for plagiarism by using Grammarly.

## CONTENT LICENSING

All materials are free to be copied and redistributed in any medium or format. However, appropriate credit should be given. The material may not be used for commercial purposes. This content licensing is in accordance with a CC license: CC-BY-NC

## CONFLICT OF INTEREST POLICY

### AUTHOR'S CONFLICT OF INTEREST

At the point of submission, Molecular and Cellular Biomedical Sciences requires that each author reveal any personal and/or financial interests or connections, direct or indirect, or other situations that might raise the question of bias in the work reported or the conclusions, implications, or opinions stated. When considering whether you should declare a conflicting interest or connection, please consider the conflict of interest test: Is there any arrangement that would embarrass you or any of your co-authors if it was to emerge after publication and you had not declared it? Corresponding authors are responsible to confirm whether they or their co-authors have any conflicts of interest to declare, and to provide details of these. The statement includes any information regarding whether the manuscript is under consideration for other publication, or whether you have any patents that relevant to the manuscript. If the manuscript is published, any conflict of interest information will be written in the Conflict of Interest statement.

### AUTHOR'S ACKNOWLEDGEMENT

Authors whose manuscripts are submitted for publication must declare all relevant sources of funding in support of the preparation of a manuscript. Molecular and Cellular Biomedical Sciences requires full disclosure of financial support as to whether it is from government agencies, the pharmaceutical or any other industry, or any other source. Authors are required to specify sources of funding for the study and to indicate whether or not the manuscript was reviewed by the sponsor prior to submission. This information should be included in the Acknowledgements section of the manuscript. In addition to disclosure of direct financial support to the authors or their laboratories and prior sponsor-review of the paper, corresponding authors will be asked to disclose all relevant consultancies since the views expressed in the contribution could be influenced by the opinions they have expressed privately as consultants. This information should also be included in the Acknowledgments section of the manuscript.

### REVIEWER'S CONFLICT OF INTEREST

Reviewers must disclose to editors any conflicts of interest that could bias their opinions of the manuscript, and should recuse themselves from reviewing specific manuscripts if the potential for bias exists. As in the case of authors, silence on the part of reviewers concerning potential conflicts may mean either that such conflicts exist that they have failed to disclose, or that conflicts do not exist. Reviewers must not use information of the manuscript they are reviewing before it is being published, to further their own interests.

## PROTECTION OF HUMAN SUBJECT AND ANIMAL IN RESEARCH POLICY

When reporting experiments on human subjects, authors should indicate whether the procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the World Medical Association Declaration of Helsinki. If doubt exists whether the research was conducted in accordance with the said declaration, the authors must explain the rationale for their approach, and demonstrate that the institutional review body explicitly approved the doubtful aspects of the study.

When reporting experiments on animals, authors should be asked to indicate whether the institutional and national guide for the care and use of laboratory animals was followed. Further guidance on animal research ethics is available from the International Association of Veterinary Editors' Consensus Author Guidelines on Animal Ethics and Welfare.

## INFORMED CONSENT POLICY

Patients have a right to privacy that should not be violated without informed consent. Identifying information, including names, initials, or hospital numbers, should not be published in written descriptions, photographs, or pedigrees unless the information is essential for scientific purposes and the patient (or parent or guardian) gives written informed consent for publication. Authors should disclose to these patients whether any potential identifiable material might be available via internet as well as in print after publication. Nonessential identifying details should be omitted.

Molecular and Cellular Biomedical Sciences decides that patient confidentiality is better guarded by having the authors archive the consent, and instead providing us with a written statement in the manuscript attesting that they have received and archived written patient consent. When informed consent has been obtained, it should be indicated later in the published article.

## ROLE OF JOURNAL EDITOR

Editors of Molecular and Cellular Biomedical Sciences have responsibilities toward the authors who provide the content of the journals, the peer reviewers who comment on the suitability of manuscripts for publication, also toward the journal's readers and the scientific community. Editors are responsible for monitoring and ensuring the fairness, timeliness, thoroughness, and civility of the peer-review and other editorial processes.

Peer review by external reviewers with the proper expertise is the most common method to ensure manuscript quality. However, our editors may sometimes reject manuscripts without external peer review to make the best use of their resources. Reasons for this practice are usually that the manuscript is outside the scope of Molecular and Cellular Biomedical Sciences, does not meet our quality standards or lacks originality or novel information.

### Editor Responsibilities toward Authors

- Providing guidelines to authors for preparing and submitting manuscripts
- Providing a clear statement of the Journal's policies on authorship criteria
- Treating all authors with fairness, courtesy, objectivity, honesty, and transparency
- Establishing and defining policies on conflicts of interest for all involved in the publication process, including editors, staff, authors, and reviewers
- Protecting the confidentiality of every author's work
- Establishing a system for effective and rapid peer review
- Making editorial decisions with reasonable speed and communicating them in a clear and constructive manner
- Being vigilant in avoiding the possibility of editors and/or referees delaying a manuscript for suspect reasons
- Establishing a procedure for reconsidering editorial decisions
- Describing, implementing, and regularly reviewing policies for handling ethical issues and allegations or findings of misconduct by authors and anyone involved in the peer review process
- Informing authors of solicited manuscripts that the submission will be evaluated according to the journal's standard procedures or outlining the decision-making process if it differs from those procedures
- Clearly communicating all other editorial policies and standards

### Editor Responsibilities toward Reviewers

- Assigning papers for review appropriate to each reviewer's area of interest and expertise
- Establishing a process for reviewers to ensure that they treat the manuscript as a confidential document and complete the review promptly
- Informing reviewers that they are not allowed to make any use of the work described in the manuscript or to take advantage of the knowledge they gained by reviewing it before publication
- Providing reviewers with written, explicit instructions on the journal's expectations for the scope, content, quality, and timeliness of their reviews to promote thoughtful, fair, constructive, and informative critique of the submitted work

# Molecular and Cellular Biomedical Sciences

- Requesting that reviewers identify any potential conflicts of interest and asking that they recuse themselves if they cannot provide an unbiased review
- Allowing reviewers appropriate time to complete their reviews
- Requesting reviews at a reasonable frequency that does not overtask any reviewer
- Finding ways to recognize the contributions of reviewers, for example, by publicly thanking them in the journal; providing letters that might be used in applications for academic promotion; offering professional education credits; or inviting them to serve on the editorial board of the journal
- Making final decision regarding a submission status after receiving review result from reviewers

## Editor Responsibilities toward Readers and the Scientific Community

- Evaluating all manuscripts considered for publication to make certain that each provides the evidence readers need to evaluate the authors' conclusions and that authors' conclusions reflect the evidence provided in the manuscript
- Providing literature references and author contact information so interested readers may pursue further discourse
- Requiring the corresponding author to review and accept responsibility for the content of the final draft of each paper
- Maintaining the journal's internal integrity (e.g., correcting errors; clearly identifying and differentiating types of content, such as reports of original data, corrections/errata, retractions, supplemental data, and promotional material or advertising; and identifying published material with proper references)
- Ensuring that all involved in the publication process understand that it is inappropriate to manipulate citations by, for example, demanding that authors cite papers in the journal
- Disclosing all relevant potential conflicts of interest of those involved in considering a manuscript or affirming that none exist
- Working with the publisher to attract the best manuscripts and research that will be of interest to readers

## AUTHOR GUIDELINES

### 1. General Terms

Molecular and Cellular Biomedical Sciences welcomes articles covering all aspects of biomedical sciences. All submitted manuscripts must not be previously published and not under consideration for publication elsewhere. Papers may come from any country but must be written in English. The manuscript may be submitted as review articles, research articles, and short communications. There are no submission and processing charges for this journal.

All manuscripts are subjected to peer review. All submissions must be accompanied by abstracts of the authors' manuscripts on related subjects that are in press or under editorial review. Electronic reprints of related published papers by the author/s or manuscripts in the press also may be helpful to the reviewers.

All manuscripts must be accompanied by a covering letter signed by all author/s. Upon acceptance, author/s must transfer copyright to Cell and BioPharmaceutical Institute (CBPI). Accepted papers become the permanent property of CBPI and may be used according to copyright policy, or for particular purposes, please contact CBPI. It is the author/s' responsibility to obtain permission to reproduce illustrations, tables, etc. from other publication.

### 2. How to Submit

Authors are required to submit manuscripts electronically by using online journal system [cellbiopharm.com/ojs](http://cellbiopharm.com/ojs).

### 3. Requirements of Each Manuscript Type

**Review Article:** Review Article should consist of no more than 10,000 words, not including the words in abstract, references, table, figure, and figure legend. The manuscript should have no more than six figures and/or tables in total and no more than 200 references.

**Research Article:** Research Article should consist of no more than 3,500 words, not including the words in abstract, references, table, figure, and figure legend. The manuscript should have no more than six figures and/or tables in total and no more than 40 references.

### 4. Abstract

Provide an abstract of no more than 300 words (for Review Article) or 250 words (for Research Article). Structured-abstract should be followed in writing Research Article.

## 5. References

- References should be according to the Vancouver system.
- List all authors when there are six or fewer; when there are seven or more, list the first six, followed by "et al."
- A sequential number of references in the main text. Please follow in detail all examples below:

### Article:

Sandra F, Esposti MD, Ndebele K, Gona P, Knight D, Rosenquist M, et al. Tumor Necrosis Factor-Related Apoptosis-Inducing Ligand Alters Mitochondrial Membrane Lipids. *Cancer Res.* 2005; 65(18): 8286-97.

### Book

Murray PR, Rosenthal KS, Kobayashi GS, Pfaller MA. *Medical microbiology*. 4th ed. St. Louis: Mosby; 2002.

### Chapter in a book:

Rosenberg GA. Matrix metalloproteinase and proteolytic opening of the blood-brain-barrier in neuroinflammation. In: deVries E, Prat A, editors. *The Blood-brain Barrier and Its Microenvironment Basic Physiology To Neurological Disease*. New York: Taylor and Francis Group; 2005. p.335-58.

### Dissertation/Thesis/Essay:

Arlaukas SP. Near infrared fluorescent choline kinase alpha inhibitors for cancer imaging and therapy [Dissertation]. Philadelphia: University of Pennsylvania; 2015.

### Part of Website/Monograph:

Medline Plus [Internet]. Bethesda: US National Library of Medicine; ©2009. Diabetic Kidney Problems [update 2015 Nov 2; cited 2015 Nov 16]. Available from: <https://www.nlm.nih.gov/medlineplus/diabetickidneyproblems.html>.

### Conference Paper:

Fledelius HS. Myopia and significant visual impairment: global aspects. In: Lin LLK, Shin YF, Hung PT, editors. *Myopia Updates II: Proceedings of the 7th International Conference on Myopia 1998 Nov 17-20, Taipei*. Tokyo: Springer; 2000. p.3-17.

## 6. Unit of Measurement

- Authors can express all measurements in Conventional or International System (SI) units.
- Drug names must use generic names. When proprietary brands are used in research, include the brand name, the name and location (city & country) of the manufacturer in parentheses after the first mention of the generic name.

## SUBMISSION PREPARATION CHECKLIST

As part of the submission process, authors are required to check off their submission's compliance with all of the following items, and submissions may be returned to authors that do not adhere to these guidelines.

1. The submission has not been previously published, nor is it before another journal for consideration (or an explanation has been provided in Comments to the Editor).
2. The submission file is in OpenOffice, Microsoft Word, RTF, or WordPerfect document file format. Formatted as standard A4 page setup.
3. Where available, URLs for the references have been provided.
4. The text should be double-spaced with the 1-inch margin on the left and right sides. Use 12-point Times New Roman font.
5. The text adheres to the stylistic and bibliographic requirements outlined in the Author Guidelines, which is found in About the Journal.
6. Running title provided (not more than 8 words).
7. Proof of permission was obtained to reproduce illustrations, tables, etc. from other publication.
8. Complete information about author/s (first, middle, last name), author/s's affiliation, and email address of the corresponding author.
9. All pages are numbered at bottom right.

## COPYRIGHT NOTICE

For the submission of a manuscript to Molecular and Cellular Biomedical Sciences, I hereby certify that:

# Molecular and Cellular Biomedical Sciences

1. I have been granted authorization by my co-author/s to enter into these arrangements.
  2. I hereby declare, on behalf of myself and my co-author/s, that:
    - The manuscript submitted is an original work and has neither been published in any other peer-reviewed journal nor is under consideration for publication by any other journal. More so, the work has been carried out in the author/s' lab and the manuscript does not contravene any existing copyright or any other third party rights.
    - I am/we are the sole author/s of the manuscript and maintain the authority to enter into this agreement and the granting of rights to the publisher: The Cell and BioPharmaceutical Institute (CBPI), does not infringe any clause of this agreement.
    - The manuscript contains no such material that may be unlawful, defamatory, or which would, if published, in any way whatsoever, violate the terms and conditions as laid down in the agreement.
    - I/we have taken due care that the scientific knowledge and all other statements contained in the manuscript conform to true facts and authentic formulae and will not, if followed precisely, be detrimental to the user.
    - I/we permit the adaptation, preparation of derivative works, oral presentation or distribution, along with the commercial application of the work.
- No responsibility is assumed by Molecular and Cellular Biomedical Sciences (MCBS) and CBPI, its staff or members of the editorial boards for any injury and/or damage to persons or property as a matter of products liability, negligence or otherwise, or from any use or operation of any methods, products instruction, advertisements or ideas contained in a publication by MCBS.

**Copyright:**

Author/s who publish in any MCBS print & online journal will transfer copyright to their work to CBPI. Submission of a manuscript to the respective journals implies that all author/s have read and agreed to the content of the Covering Letter or the Terms and Conditions. It is a condition of publication that manuscripts submitted to this journal have not been published and will not be simultaneously submitted or published elsewhere. Plagiarism is strictly forbidden, and by submitting the manuscript for publication the author/s agree that the publishers have the legal right to take appropriate action against the author/s, if plagiarism or fabricated information is discovered. By submitting a manuscript, the author/s agree that the copyright of their manuscript is transferred to CBPI, if and when the manuscript is accepted for publication. Once submitted to the journal, the author/s will not withdraw their manuscript at any stage prior to publication. However, the copyright will be released to author/s when the manuscript is rejected.

# Molecular and Cellular Biomedical Sciences

## CONTENT

### RESEARCH ARTICLES

**The Effects of *Moringa oleifera* Leaves on Complete Blood Count, Renal and Liver Functions as Potential Therapy for Malnutrition**

*Gamar Musa Kodi, Howeida Abdullah Mustafa, Alkhair Abd Almahmoud Idris*  
p.55-62

**Association of CYP2A6 Genetic Polymorphism and Lung Cancer in Female Never Smokers**

*R.A Henny Anggriani, Noni Novisari Soeroso, Setia Putra Tarigan, Putri Chairani Eyanoer, Hidayat*  
p.63-9

**Genetic Variant of Vascular Endothelial Growth Factor (VEGF)-A rs699947 is Associated with Preeclampsia**

*Anggelia Puspasari, Rina Nofri Enis, Herlambang*  
p.70-6

**Development of Recombinant Immunoblot Assay Diagnostic Test Based on HIV-1 in Indonesia**

*Jeanne Elvia Christian, Silvia Tri Widyaningtyas, Budiman Bela*  
p.77-84

**An Experimental Study on the Healing Effect of Water to Traditional Sudanese Liquor (Aragi)-induced Stomach Peptic Ulcers**

*Entisar Kuku Yousif, Howeida Abdullah Mustafa, Alkhair Abd Almahmoud Idris*  
p.85-8

**Correlation between Genetic Polymorphism of CYP2A13 Genotype and Lung Cancer in Female Passive Smokers**

*Nurul Ramadhani, Noni Novisari Soeroso, Setia Putra Tarigan, Putri Chairani Eyanoer, Hidayat*  
p.89-95

**Utilization of Expired Platelet Concentrate for Production of Human Platelet Lysate as a Medium for T47D Cell Propagation**

*Diani Mentari, Relita Pebrina, Diah Nurpratami*  
p.96-103



# Molecular and Cellular Biomedical Sciences

## Abstract

DDC 615.321

Kodi GM, Mustafa HA, Idris AAA (School of Health Sciences, Ahfad University for Women, Omdurman, Sudan)

### **The Effects of *Moringa oleifera* Leaves on Complete Blood Count, Renal and Liver Functions as Potential Therapy for Malnutrition**

*Mol Cell Biomed Sci.* 2022; 6(2): 55-62

#### **Abstract (English)**

**Background:** *Moringa oleifera* which is available in many areas all over the world including Sudan is low-cost and traditionally used in the treatment of many disorders, including malnutrition. This study aimed to determine the effect of aqueous extract of *M. oleifera* leaves in renal, liver functions and complete blood count (CBC) parameters, and its potential as therapy for malnutrition.

**Materials and methods:** This was an experimental case control study using twenty-five Wistar albino rats. Rats were divided into three groups: normal protein diet group, low protein diet with or without *M. oleifera* extract groups. We determined rats' weight, CBC parameters, blood mineral concentrations, as well as liver and renal functions at day 0, 7, and 14.

**Results:** Our findings showed that rats' weight were significantly different between the three groups at day 0, 7, and 14. Rats' weight, blood sodium, potassium, calcium, and urea concentration, as well as Hb concentration, TWBCs count, total platelets count, and %lymphocyte showed significant differences between three groups at day 0, 7, and 14.

**Conclusion:** *M. oleifera* leaves can be used as potential therapy for malnutrition because they have some effects on weight, blood mineral concentrations, renal and liver function, as well as CBC parameters.

**Keywords:** ALP, AST, ALT, creatinine, *Moringa oleifera*

DDC 616.9940231

Anggriani RAH, Soeroso NN, Tarigan SP, Eyanoe PC, Hidayat (Department of Pulmonology and Respiratory Medicine, Faculty of Medicine, Universitas Sumatera Utara/Adam Malik General Hospital, Medan, Indonesia)

### **Association of CYP2A6 Genetic Polymorphism and Lung Cancer in Female Never Smokers**

*Mol Cell Biomed Sci.* 2022; 6(2): 63-9

#### **Abstract (English)**

**Background:** The major significant factor that affected lung cancer development among female passive smokers is environmental tobacco smoke. Nicotine can be found in a never smoker population, such as a child whose father is a smoker. Lung carcinogenesis in never smoker populations is affected by nicotine metabolism by CYP2A6 gene, which encodes the main nicotine metabolizing-enzyme. The aim of this study was to assess the genetic polymorphism of CYP2A6 and its association with secondhand smokers among females who have suffered from lung cancer in North Sumatra population.

**Materials and methods:** This study was a case-control study, composed of 53 case subjects and 46 control subjects that were involved through a purposive sampling technique from two hospitals in Medan. PCR-RFLP was used for the examination of CYP2A6 gene to determine the genotype. The data were analyzed with conditional logistic regression test using Epi Info 7.0 software.

**Results:** The most common genotype of CYP2A6 detected in this study was \*1B/\*1B (40.4%), while \*1B allele had the highest prevalence (55.5%). There was no significant association between CYP2A6 genotype ( $p$ -value=0.61) or alleles ( $p$ -value=0.25) and the incidence of lung cancer.

**Conclusion:** There was no association between CYP2A6 polymorphism and the incidence of lung cancer in secondhand smoker females.

**Keywords:** CYP2A6, PCR-RFLP, female secondhand smokers, lung cancer

DDC 599.935

Puspasari A, Enis RN, Herlambang (Department of Medical Biology and Biochemistry, Faculty of Medicine and Health Sciences, Universitas Jambi, Jambi, Indonesia)

### **Genetic Variant of Vascular Endothelial Growth Factor (VEGF)-A rs699947 is Associated with Preeclampsia**

*Mol Cell Biomed Sci.* 2022; 6(2): 70-6

#### **Abstract (English)**

**Background:** Preeclampsia remains as the leading cause of maternal-neonatal mortality and morbidity worldwide. Vascular endothelial growth factor A (VEGF-A) is a proangiogenic factor related to endothelial dysfunction and plays an important role in the preeclampsia pathophysiology. Genetic variants of VEGF-A are associated with VEGF-A expression and preeclampsia risk, however there are still

# Molecular and Cellular Biomedical Sciences

inconsistent results between different populations. The aim of this study was to determine the association of this genetic variant as preeclampsia risk factor.

**Materials and methods:** A cross-sectional study was performed with 76 pregnant women (29 preeclampsia and 47 normotensive) Jambi-Malay ethnic subjects. Sample DNA was extracted from subject's blood. To determine the genotype, one-step tetra amplification refractory mutation system (ARMS) polymerase chain reaction (PCR) method for VEGF-A rs699947 C/A was used.

**Results:** We found that pregnant woman with AC genotype ( $p$ -value=0.045; OR=2.76; 95% CI=1.01-7.58) and AA genotype ( $p$ -value=0.026; OR=12.44; 95% CI=1.23-126.18) had higher risk of preeclampsia than the CC genotype.

**Conclusion:** Genetic variant VEGF-A rs699947 C/A is associated with preeclampsia. The AC and AA genotype is the risk genotype for preeclampsia in Jambi-Malay ethnics.

**Keywords:** preeclampsia, VEGF-A, genetic variant, Jambi-Malay, Indonesia

DDC 616.075

Christian JE, Widyningtyas ST, Bela B (Master Program of Biomedical Science, Faculty of Medicine, Universitas Indonesia, Jakarta, Indonesia)

## Development of Recombinant Immunoblot Assay Diagnostic Test Based on HIV-1 in Indonesia

*Mol Cell Biomed Sci.* 2022; 6(2): 77-84

### Abstract (English)

**Background:** High mutation rates in HIV-1 could affect the accuracy of diagnostic tests. Therefore, recombinant antigen that has an immunodominant and conserved region from HIV-1 need to be developed to detect HIV-1 infection in Indonesia.

**Materials and methods:** The recombinant antigens comprise of Gag (p24), Pol and Env (gp41). Each antigens was expressed in the *Escherichia coli* expression system and purified using Ni-NTA chromatography. The reactivity of purified antigen against HIV antibodies was tested against a group of 50 HIV-positive plasma samples and 45 HIV-negative plasma samples in a recombinant immunoblot assay (RIBA) platform test. Moreover, 21 of 50 HIV-positive samples and 3 of 45 HIV-negative samples were also tested using HIV blot 2.2 to compare RIBA with a commercial western blot kit. Ten HBV-positive and 10 HCV-positive plasma samples were used to check cross-reactivity with HIV recombinant proteins in RIBA.

**Results:** All HIV-positive samples (100%) tested with RIBA were reactive towards Gag (p24), Pol, Env (gp41). Otherwise, 3 of 21 HIV-positive samples assayed with HIV blot 2.2 were not reactive to Pol protein. All HIV-negative samples tested with RIBA and 3 HIV-negative samples tested with HIV blot 2.2 did not produce any bands of HIV antigens. Few HBV and HCV samples showed reactivity towards HIV recombinant proteins.

**Conclusion:** Each recombinant protein, Gag (p24), Pol, Env (gp41), could be expressed and purified, as well as had reactivity to HIV-positive samples in RIBA test. Therefore, RIBA can be used as a diagnostic test to detect HIV-1 infection in Indonesia.

**Keywords:** diagnostic, HIV-1, immunodominant, recombinant immunoblot assay (RIBA)

DDC 616.343

Yousif EK, Mustafa HA, Idris AAA (School of Health Sciences, Ahfad University for Women, Omdurman, Sudan)

## An Experimental Study on the Healing Effect of Water to Traditional Sudanese Liquor (Aragi)-induced Stomach Peptic Ulcers

*Mol Cell Biomed Sci.* 2022; 6(2): 85-8

### Abstract (English)

**Background:** Peptic ulcer is one of the most common gastrointestinal tract diseases which affect the stomach. This study aimed to determine the effect of aragi on the adult rats' stomach and investigate the effect of water as a therapeutic agent on aragi-induced ulcerations.

**Materials and methods:** Thirty-five adult Wistar albino rats were used in this experimental study. Five rats were sacrificed on day 0, 5 rats were used as a control group, and 25 rats were treated with aragi. On day 15, all rats in the control group and five aragi-treated rats were sacrificed for histological examination of the stomachs. The remaining 20 rats were stopped from aragi intake and 10 of them were treated with water for 15 days. After 15 day, all rats were sacrificed for histopathological examination of their stomachs. Stomach tissues were stained using hematoxylin and eosin (H&E) and documented under a microscope.

**Results:** Our research showed that aragi-treated rats had different severity of peptic ulcers after 15 days of continuous aragi intake, while the control group showed normal stomach histology. Nine out of 10 rats treated by water after aragi treatment also showed normal stomach histology.

**Conclusion:** Aragi is a causative agent for peptic ulcer and water can be used as potential natural therapy for treating ulcerative stomach.

**Keywords:** aragi, water, stomach, peptic ulcer



# Molecular and Cellular Biomedical Sciences

DDC 616.9940231

Ramadhani N, Soeroso NN, Tarigan SP, Eyanoe PC, Hidayat (Department of Pulmonology and Respiratory Medicine, Faculty of Medicine, Universitas Sumatera Utara/Adam Malik General Hospital, Medan, Indonesia)

## **Correlation between Genetic Polymorphism of CYP2A13 Genotype and Lung Cancer in Female Passive Smokers**

*Mol Cell Biomed Sci.* 2022; 6(2): 89-95

### **Abstract (English)**

**Background:** Nicotine is metabolized to cotinine by cytochrome P450 enzyme, and this enzyme is involved in the activation of toxic and carcinogenic substances. The aim of this research was to assess the relationship between genetic polymorphism of CYP2A13 and lung cancer incidence in female passive smokers.

**Materials and methods:** This research was a case-control study that involved 104 research subjects. Subjects were recruited through purposive sampling technique from 2 hospitals in Medan, North Sumatra, Indonesia. The case population consisted of female passive smokers with lung cancer and the control population consisted of female passive smokers without lung cancer. All research subjects underwent blood sampling for genomics DNA extraction and CYP2A13 genotyping by polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP). Data was analyzed by conditional logistic regression by Epi Info 7.0 software.

**Results:** Among 104 subjects, 26 (25%) individuals were heterozygous, 76 (73%) individuals were wild type, and 2 (2%) were mutant for the 257Cys allele. There was a significant correlation between CYP2A13 genotype and lung cancer incidence ( $p$ -value<0.05). Female passive smokers with CT genotype had 2.7 greater risk of developing lung cancer than those with CC genotype (wild type). The C allele had more frequency and 1.6 times higher risk of lung cancer compared to T allele with a wide confidence range (0.73–3.52).

**Conclusion:** There was a significant correlation between CYP2A13 polymorphism and lung cancer incidence in female passive smokers.

**Keywords:** polymorphism, CYP2A13, PCR-RFLP, female passive smoker, lung cancer

DDC 616.0277

Mentari D, Pebrina R, Nurpratami D (Faculty of Biology, Universitas Jenderal Soedirman, Purwokerto, Indonesia)

## **Utilization of Expired Platelet Concentrate for Production of Human Platelet Lysate as a Medium for T47D Cell Propagation**

*Mol Cell Biomed Sci.* 2022; 6(2): 96-103

### **Abstract (English)**

**Background:** Platelet concentrate (PC) has a short shelf life (5 days). Expired PC cannot be used for clinical purposes. PC is used for human platelet lysate (HPL) production, which was found to be more effective than FBS at increasing T47D cell proliferation. HPL production using expired PC has not been reported. This study aimed to investigate whether the use of HPL produced from expired PC (storage duration >5 days) can increase the proliferation of T47D cells in vitro.

**Materials and methods:** Expired PC samples with a shelf life of 7 and 11 days were used to produce HPL via freeze/thaw method. pH, total protein content, glucose and albumin levels were measured. The 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay was used to measure proliferation rate and doubling time of HPL-treated T47D cells.

**Results:** After HPL production, the glucose level was influenced by the pH ( $p=0.003$ ), and albumin level was influenced by total protein content ( $p=0.030$ ). HPL stored for 7 and 11 days increased cell proliferation rate by 1.41 and 1.80 times higher than 10% FBS, respectively. HPL produced from expired PC did not cause morphological abnormality of the cells. In this study, the glucose levels affected cell proliferation ( $p=0.030$ ). High glucose levels inhibited T47D cell proliferation.

**Conclusion:** Expired PC can be used as a potential material for HPL production, since HPL produced from expired PC increases cell proliferation rate and shortens cell doubling time.

**Keywords:** cell proliferation, human platelet lysate, platelet concentrate, thrombocyte, T47D

# Molecular and Cellular Biomedical Sciences

## Thank to Reviewers

We thank the following reviewers for their contributions in this number:

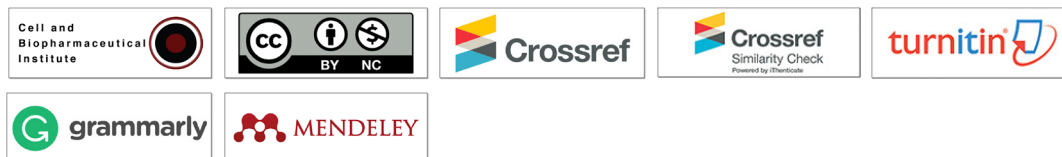
**Anna Suraya**  
**Bayu Winata Putera**  
**Dina Keumala Sari**  
**Dona Arlinda**  
**Enos Tangke Arung**  
**Erizal Sugiono**  
**Gianni Yosephine**  
**Haerani Rasyid**  
**Juminten Saimin**  
**Made Putra Semadhi**  
**Riesa Rohmat**  
**Sri Adi Sumiwi**  
**Tutik Harjianti**  
**Yusrawati**



# Molecular and Cellular Biomedical Sciences

Volume 6, Number 2, July 2022

Information of this journal can be accessed at: <https://CellBioPharm.com/ojs/index.php/MCBS>



Print ISSN: 2527-4384



9 772527 438000

Online ISSN: 2527-3442



9 772527 344004