الجامعة الإسلامية العالمية ماليزيا
INTERNATIONAL ISLAMIC UNIVERSITY MALAYSIA
يونيورسيتي إسلام انتارابغسا ملسيا
(Company No 101067-P)

RESEARCH MANAGEMENT CENTRE

Our Ref. : IIUM/504/14/11/2/ IREC 2017-009
Date : 3 July 2017

Asst. Prof. Dr. Norafiza Zainuddin (Principal Investigator)
Department of Biomedical Sciences
Kulliyyah of Allied Health Sciences
International Islamic University Malaysia
Indera Mahkota Campus
25200 Kuantan, Pahang

Dear Asst. Prof. Dr.,

The IIUM Research Ethics Committee (IREC) has reviewed your study protocol as mentioned below:-

ID NO.	:	**IREC 2017-009**
TITLE	:	**Knowledge and Perception of IIUM Kuantan Students on Lung Cancer and Its Screening**
REGISTRATION DATE	:	**31 May 2017**
INVESTIGATOR	:	**Nil**
STUDENT	:	**Humairah Thabit (Undergraduate Student)**
NAME OF SITE	:	**IIUM Kuantan Campus, Kuantan, Pahang**
DURATION	:	**9 Feb 2017 - 31 Jul 2017**

The IIUM Research Ethics Committee (IREC) operates in according to the Declaration of Helsinki, International Conference of Harmonization Good Clinical Practice Guidelines (ICH-GCP), Malaysia Good Clinical Practice Guidelines and Council for International Organizations of Medical Sciences (CIOMS) International Ethical Guidelines.

The following documents have been received and reviewed to the above study:-

1. Study Proposal/Protocol: Version 1, revision 00, dated 6 Jun 2017
2. Informed Consent Form (ICF):-
   i. Information Sheet – English: Version 1, revision 00, dated 6 Jun 2017
   ii. Consent Form – English: Version 1, revision 00, dated 6 Jun 2017
3. Questionnaire: Version 1, revision 00, dated 6 Jun 2017
4. Approval Letter from Kulliyyah
5. Principal Investigator's CV

Decision by IIUM Research Ethics Committee (IREC):

(√) Approved
( ) Disapproved

Date of Approval: **21 June 2017**

The investigator(s) are required to:

a) register the study with National Medical Research Register (NMRR) for any study done in Ministry of Health (MOH) facilities.
b) notify IREC of any change in protocol and obtaining further ethical approval as appropriate.
c) report any adverse incident during the course of a study to IREC even if the incident is not directly related to the study.
d) report serious adverse event (SAE) within 24 hours.
e) report minor adverse event within 2 weeks.
f) provides information of minor protocol deviation in Progress Report or End Report whichever necessary.
g) report any major protocol deviation occurs within 5 working days.
h) submit Progress report form before the end of six (6) month given by IREC.
i) complete and submit the End of Project Report Form to the IREC Secretariat's Office.

Thank you.

Yours sincerely,

PROF. DR. NASSER MUHAMMAD AMJAD
Chairman,
IIUM Research Ethics Committee (IREC)

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