同意書 編號 AS-IRB02-99080 NUM 日期

Date

0715'2010

中央研究院醫學研究倫理委員會 研究計畫審查通過證明 Certificate of Approval

IRB02 修訂日期 Revised Date: 0730'2009

申請案編號 Application No: AS-IRB01-10054 (08007)

免疫基因多型性與人類乳突病毒持續感染及子宮頸癌之長期追

human papillomavirus, and cervical cancers: a long-term follow-up

study

申請人 Project Investigator: 陳建仁 Chien-Jen Chen

合作機構 Collaborating Institute(s): 無

From 2008 / 08 / 01 / to 2011 / 07 / 31 /

執行期間 Implement Duration:

(yyyy / mm/ dd) (yyyy / mm/ dd)

 核准日期 Approval Date:
 2010 / 07 / 15 / (yyyy / mm/ dd)

 有效期限 Due Date:
 2011 / 07 / 31 / (yyyy / mm/ dd)

進度或成果報告 Progress Report:

1. 請於 <u>2010 / 07 / 14 /(yyyy / mm/ dd)</u>前繳交進度或成果報告至本委員會追蹤審查。
The progress report should be submitted to the IRB at Academia Sinica before <u>2010 / 07 / 14 / (yyyy/ mm/ dd)</u> for continuing review.

- 2. 如向衛生署定期提出進度或成果報告,需將完整報告內容副知本委員會。
 If Progress reports are submitted to the Department of Health, a copy should be submitted to the IRB at Academia Sinica.
- 3. 如受試者有不良反應,應即時向主管機關報告並副知本委員會。 All the adverse events are required to report to the official institute and IRB of Academia Sinica promptly.

附註意見 Additional Opinion:

若需與受試者溝通(如與受試者通信、致贈禮物等),與受試者溝通之文件、信件或禮物,須經本委員會審核通過後方可進行。

Any materials (correspondence, brochure, compensation, etc.) provided to subjects, must be approved by the IRB of Academia Sinica.

茲 證明上項醫學研究計畫,包括其受試者之說明同意書,已經本委員會之審核,並同意此計畫書之內容符合本委員會所訂定之醫學研究倫理標準。

The above named study proposal has been reviewed, along with the informed consent, by the committee and found to conform with the guidelines set forth by this committee.

主任委員: 日期:

Chairman Date

中央研究院醫學研究倫理委員會

Human Subject Research Ethics Committee / IRB Academia Sinica