同意書編號 NUM	AS-IRB02-103087	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： 0226’2013
日期 Date	0522’2014		

申請案編號 Application No： AS-IRB01-11031（M3R4）

計畫名稱 Project title： 情緒穩定劑的藥效與藥物不良反應之分子遺傳研究
Molecular genetic study of drug response and adverse drug reactions of mood stabilizers

申請人 Project Investigator： 鄭泰安 Andrew Cheng

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IRB on Biomedical Science Research /IRB-BM
Academia Sinica

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執行期間 Project Duration： From 2011 / 05 / 01 / to 2017 / 04 / 30 / (yyyy / mm/ dd) (yyyy / mm/ dd)

核准日期 Approval Date： 2014 / 05 / 22 / (yyyy / mm/ dd)

有效期限 Due Date： From 2014 / 05 / 22 / to 2017 / 04 / 30 / (yyyy / mm/ dd) (yyyy / mm/ dd)

進度或成果報告 Progress Report：

1. 請於 2015 / 04 / 30 /(yyyy / mm/ dd)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 2015 / 04 / 30 / (yyyy/ mm/ dd) 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. 如研究對象有不良反應，應即時向主管機關報告並副知本委員會。
   All the adverse events are required to be reported to the official institute and IRB of Academia Sinica promptly.

附註意見 Additional Opinion：

若需與研究對象溝通（如與研究對象通信、致贈禮物等），與研究對象溝通之文件、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(AS-IRB-BM-11031 v.5)，包括計畫書、研究對象說明同意書、問卷，已經本委員會之審核，並同意此計畫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(AS-IRB-BM-11031 v.5) has been reviewed, along with the informed consent form, questionnaire, and found to conform with the guidelines set forth by IRB on Biomedical Science Research / IRB-BM Academia Sinica.

主任委員： 日期：
Chairman Date

中央研究院醫學研究倫理委員會
IRB on Biomedical Science Research /IRB-BM
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