福建医科大学附属第一医院医学伦理委员会医学研究与临床技术应用分会审查批件

Ethics Review Form for Branch for Medical Research and Clinical Technology Application, Ethics Committee of First Affiliated Hospital of Fujian Medical University

闽医大附一伦理医技审[2015]084-1 号 MTCA,ECFAH of FMU [2015]084 -1

		H of FMU [2015]084	• -1
项目名称	住院病人医疗信息和生物标本的二次利用		
Protocol Title	Reutilization of Medical Information and Biological Specimens of the Inpatients		
项目来源	福建医科大学附属第一医院		
Protocol Source	The First Affiliated Hospital of Fujian Medical University		
伦理审查方式			
Mode of Review	☑ 会议审查 Meeting Review □ 快速审查 Quick Review		
	□初次审查 Initial Review ☑ 修正案审查 Review on Amendments		
	审查类别 pe of Review □ 国际审查 Tracing Review □ 违背方案审查 Review on Contrary to Protocol □ 严重不良事件审查 Review on Serious Adverse Events □ 暂停/终止研究审查 Suspending/Terminating Review on Research		
Type of Review			
	□结题审查 Review on End of Research □复审 Review		
会议地点	门诊十二楼大会议室	日 期	2019-4-3
Meeting Location	Conference room at the 12th floor	Date	2019-4-3
	福建医科大学附属第一医院患者住院须知		
送审材料	(V4.0 2019.4.3)	审查委员	见附件 2
Documents for	Instructions for hospitalization in the First	Review Person	See Attachment
Reviewing	Affiliated Hospital of Fujian Medical		2
山市人品	University(V4.0 2019.4.3)		
出席人员 Attendees	实到 Attendance: 9人, 回避 Avoidance: 0人		
Attenuces	1.同意 Apptoval(9)人		
In The late was	2 作必更修正后同音 Approval after pecessary amendment (0) 人		
投票情况	3.作必要修正后再审 Check Review after necessary amendmen (0) 人		
Voting Results	4.不同意 Disapproval(0)人		
5.终止或暂停已批准的技术 Terminating / Suspending the Approved Technology			nology(0)人
	2019年4月3日,伦理委员会根据国家《药物临床试验质量管理规范》、《赫		
	尔辛基宣言》及《涉及人的生物医学研究伦理审查办法》等伦理原则,对本方案进		
审查意见 Comments			
	行伦理跟踪审查,同意在我院继续实施。		
	On Apr. 3rd, 2019, According to the national ethical principles of Good		
	Clinical Practice, Declaration of Helsinki and Measures for the Ethical		
	Review of Biomedical Research Involving Humans, a meeting ethical review		
	of the project was conducted, and the project is approved to be carried out		
	in our hospital.		

本审查委员会声明: 遵从 ICH-GCP、中国 GCP、中国相关法规及指南的要求组 成和开展工作,其审查和工作过程不受伦理委员会以外的任何组织及个人的影响。 本伦理委员会对本项目研究要求: 1. 完成项目研究时, 向伦理委员会提交结题 报告; 2. 每年度至少向伦理委员会提交1次"年度进展报告"; 3. 如研究方案、知 情同意书等文件的任何修改, 应及时通知伦理委员会, 必要时重新审查, 获批准后 执行。4. 在批件有效期届满1个月前向伦理委员会提出申请。 The Institutional Review Board (IRB) declares that: In accordance with the requirements of ICH-GCP, China GCP and relevant laws and regulations or guidelines in China, the review and other working processes are free from any influence of any organization or individuals other than this Ethics Committee. Requirements on researches given by the IRB to the project are as follows: 1. Submitting a concluding report to the IRB when completing Remarks researches on the project; 2. Submitting at least one "annual progress report" to the IRB each year; 3. If there is any modification on documents such as research plan or informed consent, notify to the IRB in time and do Check Review if necessary for further execution after approval; 4. Please submit an application within 30 days before the expiry date. Signature of the chairman of committee: Date: 批件有效期 Validity Period of 2019年4月3日——2022年4月2日 **Ethical Review** 3rd, April, 2019----2nd, April, 2022 Approval Document 联系电话 伦理委员会地址 福建省福州市茶中路 20 号 Contact 0591-87981028 **Ethics Committee** No.20, Chazhong Road, Fuzhou, Fujian Province Number Address