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

闽医大附一伦理医技审[2015]084-1 号 MICA,ECFAH 01 FWO [2015]064-1				
项目名称 Protocol Title	住院病人医疗信息和生物标本的二次利用 Reutilization of Medical Information and Biological Specimens of the Inpatients			
项目来源 Protocol Source	福建医科大学附属第一医院 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			
审查类别 Type of Review	□初次审查 Initial Review ☑ 修正案审查 Review on Amendments □跟踪审查 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 □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		
送审材料 Documents for Reviewing	福建医科大学附属第一医院患者住院须知(V4.0 2019.4.3) Instructions for hospitalization in the First Affiliated Hospital of Fujian Medical University(V4.0 2019.4.3)	审查委员 Review Person	见附件 2 See Attachment 2	
出席人员 Attendees	实到 Attendance: 9人, 回避 Avoidance: 0人			
投票情况 Voting Results	1.同意 Apptoval (9) 人 2.作必要修正后同意 Approval after necessary amendment (0) 人 3.作必要修正后再审 Check Review after necessary amendmen (0) 人 4.不同意 Disapproval (0) 人 5.终止或暂停已批准的技术 Terminating / Suspending the Approved Technology (0) 人			
审查意见 Comments	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》、《赫尔辛基宜言》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 GC 成和开展工作,其审查和工作过程不受伦理委员	P、中国相关法: 会以外的任何组	规及指南的要求组 l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			
	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.			
备注	Requirements on researches given by the IRB to the project are as			
Remarks	follows: 1. Submitting a concluding report to the IRB when completing			
	researches on the project; 2. Submitting at least one "annual progress			
	report" to the IRB each year; 3. If there i	s any modificat	tion 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.			
		主任委员签字:	953	
	Signature of the chairman of committee: 日期,2019年4月3月 Date: 初初的)			
		Date: William		
批件有效期			ž.	
Validity Period of	2019年4月3日——2022年4月2日			
Ethical Review Approval	3rd,April,20192nd,April,2022			
Document		and the second second		
伦理委员会地址	福建省福州市茶中路 20 号	联系电话	0001 07001000	
Ethics Committee	No.20, Chazhong Road, Fuzhou, Fujian Province	Contact Number	0591-87981028	
Address	,	Number		