Ethics Committee of Shenzhen Centre for Disease Control and Prevention

Ethical Review Approval

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Approval		
The sample size of this study was 2.		
All samples in this study were isolates from hospitals after		
clinical diagnostic tests were done, and no original information		
was involved, while the information data used were anonymised,		
with no subsequent retrospective analyses, and did not involve the		
patients themselves.		
The study is a public interest exploration of scientific		
knowledge and does not involve treatment or intervention, nor		
does it involve genetic modification or hereditary issues, and the		
chance of additional harm to the subjects is low.		
Exemption from signing informed consent.		
Consent to conduct the study.		
1. If serious adverse events occur, they should be reported to		
the Ethics Committee in a timely manner.		
2. If any modification of human biomedical research trial		
protocol, informed consent, or change of principal investigator		
should be promptly notified to the Ethics Committee, re-examined,		
and implemented after approval.		
3. This approval is valid for one year. Please apply for follow-		
up review one month before the expiration of the approval.		
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 was involved, while the information data used were and with no subsequent retrospective analyses, and did not in patients themselves. The study is a public interest exploration of scientific knowledge and does not involve treatment or intervention does it involve genetic modification or hereditary issues chance of additional harm to the subjects is low. Exemption from signing informed consent. Consent to conduct the study. 1. If serious adverse events occur, they should be retted the Ethics Committee in a timely manner. 2. If any modification of human biomedical researce protocol, informed consent, or change of principal invest should be promptly notified to the Ethics Committee, retaind implemented after approval. 3. This approval is valid for one year. Please apply 		

Approval No.: Shen Zilun Zi [2023] No. 029A

Remarks: The conclusions of the review include four scenarios: approval, re-examination after revision and disapproval. Principle of numbering the approval number: the first four digits represent the year; in the middle is the acceptance sequence number; the letter A at the end represents the initial ethical review of the project, B represents the follow-up review, and C represents the revised review of the research protocol.

深圳市疾病预防控制中心伦理委员会 伦理审查批件

批件号: 深疾伦字 [2023] 第 029A 号

项目名称	中国深圳1例创伤弧菌死亡病例的病原学检测及溯源分析			
项目负责人	石晓路	0	1	
原审查批件号				
申请受理号	QS2023040068			
审查结论	批准			
审查意见	始信息,同时所使用的 析,不涉及患者本身。	医院作完临床诊断相信息数据均经匿名4 科学知识探索,不被 对受试者产生额外的	金测后的分离物,不涉及房 化处理,后续不需要溯源分 步及治疗和干预,也不涉及 的伤害的机会很低。	
注意事项	 1、如发生严重不良事件,应及时报告伦理委员会; 2、如人体生物医学研究试验方案、知情同意书的任何修改、主要研究者变更应及时通知伦理委员会,重新审查,获得批准后执行。 3、本批件有效期一年。请在批件到期前一个月提出跟踪审查申请。 			
跟踪审查委员	赵锦 刘建平	联系电话	13823311336 13823132646	
主任委员签字	352	SULUEIN CENTER FOR DIS SULUEIN CENTER FOR F	A A A A A A A A A A A A A A A A A A A	

备注:审查结论包括批准、修改后批准、修改后重审和不批准等四种情形。批件号编号原则:前四位代表 年份;中间是受理序列号;后面字母A代表项目初次伦理审查,B代表跟踪审查,C代表研究方案修订审 查。