# **Registry Profile for [**Acupuncture-Moxibustion Clinical Trial Registry**]**

Official Name of Registry (list abbreviation, if relevant)	Acupuncture-Moxibustion Clinical Trial Registry (AMCTR)
Address of Registry	No. 16, Nan Xiao Jie Street, Dong Zhi Men Nei , Beijing, China No. 16, Dongzhimendistrict, Beijing, China
Registry URL (Register's web site address)	www.acmctr.org
Registration URL (Web site where users go to register a trial)	www.acmctr.org
Affiliated Primary Registry	Chinese Clinical Trial Registry

#### Content

1.	Does the Registry accept prospective registration of interventional clinical trials submitted by Responsible Registrants?	⊠ Yes □ No
2.	Is the Registry open to all prospective registrants ( <i>ICMJE</i> ), either internationally, or within one or more specific countries	⊠ Yes □ No
	<ol><li>2.1. From which countries will the Registry accept trials for registrati Worldwide, No limits.</li></ol>	ion?
	2.2. If the Registry is not open to ALL registrants (that is, accepts tri particular sponsor, health care condition (eg cancer), int specify which trials the Registry is open to: The Registry is open to those trials which focus on acupuncture.	ervention, etc) please
3.	Is the Registry able to collect and publicly display the WHO Trial Registration Data Set ( <i>ICMJE</i> )	⊠ Yes □ No
4.	Does the Registry endeavour to keep registered information up-to-date?	⊠ Yes □ No
	4.1. Does the Registry have a reminder system to facilitate the submission of updated information by the responsible registrant?	⊠ Yes □ No
	4.2. Does the Registry highlight or flag records that are not current (have not been updated in more than 12 months) so readers are aware that information contained in such records may not be current?	⊠ Yes □ No
5.	Does the Registry never remove a trial once it has been registered?	⊠ Yes □ No

## **Quality and Validity**

<ol><li>Does the Registry have a mechanism in place to ensure the validity of the registered data? (ICMJE)</li></ol>	⊠ Yes □ No	
C4. Describe assistant contest Description Description to the contest	2110	
6.1. Does the registry contact Responsible Registrants when one	× Yes	
or more items in the Trial Registration Data Set are		
incomplete or potentially inaccurate?	□ No	
6.2. Does the registry comply with the International Standards for		
Clinical Trial Registries (ISCTR)?	× Yes	
Olimbai Thai Registries (100 TR):	□No	
C 0 1 If the angular to C 0 is "mall them places describe which	-	
6.2.1. If the answer to 6.2 is "no", then please describe which standards are a problem and your plans for achieving these standards in future.		
6.3. Does the registry have documented Standard Operating		
Procedures (SOPs)?	× Yes	
1 100edules (501 3):	□ No	
6.3.1. If the answer to 6.1 is "no", then please describe your plan for completing your SOPs and the date by which you expect them to be complete		
6.3.2. If the answer to 6.1 is "yes" then please let us know the languages in which the SOPs are written:		
× English		
6.4. Are mechanisms in place to ensure compliance with these	T_	
	×  Yes	
SOPs (eg staff training)?	□ No	
6.5. Please briefly describe these compliance mechanisms The full-time and partly-time staff for Acupuncture-Moxibustion Clinical Trial Registry will be trained by Chinese Clinical Trial Registry; The web, Registry system and databases of Acupuncture-Moxibustion Clinical Trial Registry will be designed by the IT engineer of Chinese Clinical Trial Registry.		
7. Does the registry maintain a publicly accessible audit trail so	W Voc	
changes made to the WHO Trial Registration Data Set for an		
individual trial can be tracked.?	│ □ No	
ilidividual tilai cati be tracked.:		
7.1. If yes, then please describe how users can access the audit trail.		
The audit trial will be flagged as "Audit trial" and could be search by using a key word "audit".		
7.2. If no, then please describe your plan for creating an audit tra you expect it to be complete and publicly available	ail and the date by which	

#### Accessibility

•		
8. Is the WHO Trial Registration Data Set accessible to the public at no charge? (ICMJE)	⊠ Yes □ No	
9. Is the WHO Trial Registration Data Set electronically searchable? (ICMJE)	⊠ Yes □ No	
10. Is the WHO Trial Registration Data Set available in English?	⊠ Yes □ No	
11. Can registrants submit a trial for registration at any time of day on any day of the week (24 hours a day, seven days a week)?	⊠ Yes □ No	
12. Can the register be searched at any time of day on any day of the week (24 hours a day, seven days a week)?	⊠ Yes □ No	
13. It is desirable that the WHO Trial Registration Data Set be available countries served by the registry. Please indicate the languages in winformation is displayed Chinese and English		
13.1. If trials are registered in more than one language, does the registry check the quality of translation?	⊠ Yes □ No	
Note: If the trial is submitted in both languages by the Responsible Registrant then the registry should check the translation. If the trial is submitted in only one language and the registry translates into the second language, then other registry staff should check the translation.		
<ul> <li>14. What quality control measures does the registry have in place to make sure that all items in the WHO Trial Registration Data Set are as complete and accurate as possible?</li> <li>We have double checking system:</li> <li>1. AMCTR staff will review, check and edit the data in the registration form, make sure the language is correct and understandable, the study design described is acceptable, then submit the form to ChiCTR;</li> <li>2. ChiCTR will check again.</li> </ul>		

# Unambiguous Identification

15. Does the Registry have processes in place to prevent the registration of a single trial more than once on their register?	⊠ Yes □ No
15.1. Please briefly describe these processes: Our SOP include checking the trial for duplicate registration by searching WHO ICTR Search Portal.	
16. Does the Registry agree to facilitate the retrospective linking (or bridging) on the WHO Search Portal of a single trial registered with more than one registry by encouraging Responsible Registrants to enter secondary identifiers, including the UTN?	⊠ Yes □ No
17. Does the Registry agree, as part of the registration process, to search the WHO Search Portal to ascertain if the trial has already been registered on another WHO Primary Registry?	⊠ Yes □ No

## Technical Capacity

18. Does the Registry agree to submit the WHO Trial Registration Data Set for all records on their registry to the Primary Registry so it can be included in the ICTRP's Central Repository?		yes □ No
Note: In the case of dual lan language version	nguage registries the ICTRP requires the English	
	registry already successfully submitted data to ted Primary Registry?	○ X Yes No, not yet
19. Does the Registry have access to a database that is used to store and manage the submitted data?		Yes D No
	ease provide details of the database that will and where it is based	
20. Does the Registry ha technology support?	ve access to adequate information	⊠ Yes □ No
21. Does the Registry ha against data corruption	ve adequate security and other provisions on/loss?	⊠ Yes □ No

#### Administration and Governance

22. Does the Re	egistry have a national remit?	☐ Yes ☒ No
	From which countries does the Registry have the remit to	o act as their Primary
	Registry? gistry of ChiCTR, AMCTR will have a national remit beca	ause of ChiCTR have
the national rem		adde di Cime i i i i i i i i i i i i i i i i i i
	egistry have the support of a relevant agency such as	× Yes
	of Health or similar? supported by the <b>China Association of</b>	□ No
	re-Moxibustion and the World Federation of	
•	re-Moxibustion Societies.	
	Please specify the names of the government agencies the support.	nat have given their
As a Partne	r Registry of ChiCTR, AMCTR could be considered supp	oorted by MOH.
24. Does the Re	egistry publicly disclose ownership, governance	
	d not/for-profit status?	× Yes
24.1.	Please indicate the web address where this information	∐ No
www.acmc		is displayed
	try managed by a not-for-profit agency? (ICMJE)	₩ Vaa
		X Yes
26 Should the	register cease to function, does the register agree that	□ No
	WHO Trial Registration Data Set for all trial records will	× Yes
be transferre	ed to a WHO Primary Register?	□ No
27. Does the registry agree to inform it's affiliated Primary Registry of		
	s that may be relevant to its status as a WHO Partner	□ No
Registry?	le that Partner Registries have a strategy in place	_
	the medium to long term sustainability of the registry.	× Yes
	egistry have such a strategy in place?	□ No
	rnance information:	
	What is the name of the agency (or agencies) that fund the Chine Aggregation of Aggregative Mayibustian and	
	The China Association of Acupuncture-Moxibustion and exibustion Societies.	the world rederation of
	What is the name of the agency that manages the regist	rv?
	The China Association of Acupuncture-Moxibustion and	
	exibustion Societies.	
	Is the agency that manages the registry a for-profit	□Yes
	agency?	⊠ No
29.4.	If the agency is for-profit, is the data on the register	
	(database) also available on a Registry that is	□Yes
	managed by a not-for-profit agency?	□No
29.5.	If the answer to 27.4 is "yes", then please specify the na	× Not applicable
	Registry	ino oi trie additional
	· ,	

29.6. Please describe any other ownership or governance issues of relevance to the register. (eg Does the register have an Advisory Board. If so, what is its role?) The register will have an Advisory Board, which will solute some problem for Registrants.