

Text S₁: Quality assessment rules of included interventional studies.

Note: The minimum 20 items of WHO Trial Registration Data Set (TRDS, V.1.2.1) plus optional additional 3 items recommended by ICTRP, and some specific items for TCM information (including TCM intervention, diagnosis, outcome and rationale) were adopted to assess the reporting quality of registered interventional trials included in this study. Aim to better assessment, the total 23-item was divided into two parts: (1) simple items (i.e. item 1-12, 16-18, 22) and (2) complex items (i.e. item 13-15, 19-21, 23), which contained some subitems that needed to be evaluated individually.

Part 1. Simple items

	Item/Label	Assessment rules: “1” (reported) or “0” (not reported)
1	Primary Registry and Trial Identifying Number	It should contain a name of primary registry and the unique ID number assigned by the Primary Registry to this trial.
2	Date of Registration in Primary Registry	A clear date when trial was officially registered in the Primary Registry.
3	Secondary Identifying Numbers	<p>These include: 1) The Universal Trial Number (UTN); 2) Identifiers assigned by the Sponsor (record Sponsor name and Sponsor-issued trial number (e.g., protocol number)); 3) Other trial registration numbers issued by other Registries (both Primary and Partner Registries in the WHO Registry Network, and other Registries); 4) Identifiers issued by funding bodies, collaborative research groups, regulatory authorities, ethics committees / institutional review boards, etc.</p> <p>All secondary identifiers will have 2 elements: an identifier for the issuing authority (eg CTN, ISRCTN, ACTRN) plus a number. There is no limit to the number of secondary identifiers that can be provided. Also, some trials do not have the secondary ID number, which recorded as “0”.</p>
4	Source(s) of Monetary or Material Support	Major source(s) of monetary or material support for the trial (e.g., funding agency, foundation, company, institution).
5	Primary Sponsor	It should contain the individual, organization, group or other legal entity which takes responsibility for initiating, managing and/or financing a study.
6	Secondary Sponsor(s)	Additional individuals, organizations or other legal persons , if any, that have agreed with the Primary Sponsor to take on responsibilities of sponsorship. Also, some trials do not have this item, which recorded as “0”.
7	Contact for public queries	Contact details should include the name, email address, telephone number, affiliation and postal address of the contact person. If all details or main details (i.e. name plus one contact information) were provided, the record was denoted

		to be “1”. However, if the valid contact information was missing or not reported, the record was denoted to be “0”.
8	Contact for scientific queries	<p>Contact details of the Principal Investigator (PI) should include: i) Name and title, email address, telephone number, postal address and affiliation of the PI, and; ii) Email address, telephone number, postal address and affiliation of the contact for scientific queries about the trial (if applicable). The record was denoted to be “1” as some valid contact information were reported.</p> <p>If some registries had one field for two contact persons, the information of person(s) responsible for the overall scientific leadership (including the PI) was extracted from that field and classified as the scientific contact item, and the other one was recorded as public contact.</p>
9	Public title	Title intended for the lay public in easily understood language. An informative public title will describe the participants, the intervention, the comparator and the main outcome of the study. Only absence in this item will denoted to be “0”.
10	Scientific title	Scientific title of the study as it appears in the protocol submitted for funding and ethical review. Include trial acronym if available. It is desirable that the scientific title contain all elements of PICO (Participants, Intervention, Comparator and Outcomes). Only absence in this item will denoted to be “0”.
11	Countries of Recruitment	It should contain the name of countries from which participants will be, are being, or have been recruited at the time of registration.
12	Health condition(s) or problem(s) studied	<p>List the name of primary health condition(s) or problem(s) studied (e.g., depression, breast cancer, medication error).</p> <p>If the study is conducted in healthy human volunteers belonging to the target population of the intervention (e.g. preventive or screening interventions), enter the particular health condition(s) or problem(s) being prevented.</p>
16	Date of first enrolment	Anticipated or actual date of enrolment of the first participant . Some Registries may use the label " trial start date ". If so, it should be made clear to Registrants (via ‘Help’ text or other documents) that this field must contain the date the first participant was enrolled. Specifically, if the date only contains the year and month, it also scored as “1”.
17	Target sample size	Number of participants that this trial plans to enroll in total.
18	Recruitment status	Recruitment status include: Pending, Recruiting, Suspended, Complete, etc.
22	Approvals	This item includes ethics committees and regulatory authorities. For each approving entity the name of the entity, the date and status of the approval should be reported.

Part 2. Complex items

Note: The following 7 complex items are important information for a trial, containing some subitems in each item according to the explanatory text in TRDS. Each subitem with detailed criteria need to be evaluated separately and the overall quality of each item was assessed comprehensively. Compared with WHO standard framework, different registries presented some variation in registration formats for some items. Take ClinicalTrials.gov for example, the field of 'detailed description' could not be ignored because it provided lots of additional information contributed to the assessment of intervention or study design item, although 'detailed description' was not specified in a particular item according to TRDS framework. Therefore, it was essential to **scan overall record information** in order to evaluate complex items objectively. Focus on the contents only with same name according to TRDS items is not enough.

Besides, according with the characteristics of TCM trials, some of key information related with TCM are specified in the data set, thus to assess whether these items are fully reported, including: 1) Details description of TCM intervention as a subitem included in item 13, 2) TCM diagnosis and its criteria as two subitems included in item 14, 3) TCM outcome(s) as a subitem included in item 19/20, 4) TCM background and rationale as an subitem included in item 21.

	Item/Label	Assessment rules: "1" (reported) or "0" (not reported)
13	Interventions	<p>This item contained four subitems:</p> <p>13.1 Intervention Name: For drugs use generic name; for other types of interventions provide a brief descriptive name. For investigational new drugs that do not yet have a generic name, a chemical name, company code or serial number may be used on a temporary basis. As soon as the generic name has been established, update the associated registered records accordingly. For non-drug intervention types, provide an intervention name with sufficient detail so that it can be distinguished from other similar interventions.</p> <p>Name was assessed as reported ("1") if a specific TCM name was provided, such as Si-Ni-Tang/ Sini decoction (a Chinese Herbal Formula), Acupunctue of Point Hegu, etc. No name or too generalized name was denoted as "0", such as Chinese medicine, TCM treatment, Drug group, etc. These names of TCM interventions were too confused to figure out what specific technology adopted in a trial.</p> <p><u>13.2 TCM Intervention Description:</u> Must be sufficiently detailed for it to be possible to distinguish between the arms of a study (e.g., comparison of different dosages of drug) and/or among similar interventions (e.g., comparison of multiple implantable cardiac defibrillators). For example, interventions involving drugs may include dosage form, dosage, frequency and duration. If the intervention is one or</p>

	<p>more drugs, then use the International Non-Proprietary Name for each drug if possible (not brand/trade names). For an unregistered drug, the generic name, chemical name, or company serial number is acceptable.</p> <p>Given the considerable variability in the types of TCM interventions, the more detailed information should be provided. For example, the ingredients in compound formula, and the operating methods of acupuncture, cupping, or other non-pharmacy therapies should be reported.</p> <p>13.3 More Interventions: If the intervention consists of several separate treatments, list them all in one line separated by commas (e.g., "low-fat diet, exercise"). Or use "and", etc. This point was assessed according to the correct format noted in TRDS, and was collected to understand the proportion of integrated traditional Chinese and conventional medicine.</p> <p>13.4 Controlled Intervention(s): For controlled trials, the identity of the control arm should be clear. The control intervention(s) is/are the interventions against which the study intervention is evaluated (e.g., placebo, no treatment, active control). If an active control is used, be sure to enter in the name(s) of that intervention or enter "placebo" or "no treatment" as applicable. For each intervention, describe other intervention details as applicable (dose, duration, mode of administration, etc).</p>
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14	Key Inclusion and Exclusion Criteria	<p>This item contained six subitems:</p> <p>14.1 Age: Clearly age range. Absence in this point was recorded as “0”.</p> <p>14.2 Sex: Male, Female or Both. Absence in this point was recorded as “0”.</p> <p>14.3 Conventional medicine criteria relate to clinical diagnosis and co-morbid conditions; exclusion criteria are often used to ensure patient safety.</p> <p>14.4 Healthy volunteers: If the study is conducted in healthy human volunteers not belonging to the target population (e.g., a preliminary safety study), enter "healthy human volunteer". If the trial included healthy volunteers, the record was denoted to be “1”. If not, the record was denoted to be “0”.</p> <p>14.5 Adopted TCM diagnosis: include name of TCM disease(s) or “zheng”/ syndrome/pattern or symptom and sign, etc. The record was denoted to be “1” as the trial adopted TCM diagnosis and reported its name.</p> <p>14.6 TCM diagnosis with clearly criteria: A clearly criteria should contain the specific descriptions (i.e. specific symptoms belonging to the pattern mentioned) or guideline references. If the diagnosis details were not provided, this information was denoted as “0”, although it had a name of TCM diagnosis. For example, the TCM diagnosis criteria reported as “TCM syndrome differentiation of depression” or “qi and blood stagnation pattern or cold and dampness stagnation pattern” without any specific descriptions or guideline references.</p>
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15	Study type	<p>This item contained six subitems:</p> <p>15.1 Type of study: Interventional or observational, etc.</p> <p>15.2 Randomized/Non-randomized: For randomized trials: the randomize numbers/sequence generation will be documented (i.e. the software, operator, etc). The completely reported of randomized methods and correctly reported non-randomized methods were both considered as “1”. The missing records or incomplete reported of randomized were both considered as “0”.</p> <p>15.3 Allocation concealment mechanism: For randomized trials: the detailed information of allocation concealment should be clarified (i.e. central randomized/controlled, opaque envelope, etc).</p> <p>15.4 Masking/Open label: Masking (is masking used and, if so, who is masked). The completely reported of masking methods and correctly reported open label were both considered as “1”. The missing records or incomplete reported of masking methods were both considered as “0”. For example, if the trial reported masking was adopted but we did not find any information of who was blinded.</p> <p>15.5 Assignment: single arm, parallel, crossover or factorial, etc.</p> <p>15.6 Phase: if applicable. Presented as “1”; Absence as “0”.</p>
19	Primary Outcome(s)	<p>This item contained three subitems:</p> <p>19.1 Name: The name of the outcome (do not use abbreviations).</p> <p>19.2 Measurement: The metric or method of measurement used (be as specific as possible).</p> <p>19.3 Time point(s) of primary interest: The time point is not the same as study duration or period of follow-up.</p> <p>Most trials should have only one primary outcome.</p>
20	Key Secondary Outcome(s)	<p>This item contained three subitems:</p> <p>19.1 Name: The name of the outcome (do not use abbreviations).</p> <p>19.2 Measurement: The metric or method of measurement</p>

		<p>used (be as specific as possible).</p> <p>19.3 Time point(s) of interest</p> <p>A secondary outcome may involve the same event, variable, or experience as the primary outcome, but measured at timepoints other than those of primary interest. Some trials may not have secondary outcome.</p>
	19/20 Outcome(s)	<p>Two subitems were included in the part of outcome(s):</p> <p><u>19/20.1 TCM related outcome(s):</u> In primary or secondary outcome(s), if TCM related outcome(s), such as “zheng”/symptom scoring, etc., was adopted in the trial, it would be considered as “1”.</p> <p>19/20.2 Adverse events outcome(s): In primary or secondary outcome(s), if any adverse events outcome(s) was reported in the trial, it would be considered as “1”.</p>
21	Lay Summary / Synopsis	<p>This item contained three subitems:</p> <p>21.1 Objective: short description of the primary purpose of the study.</p> <p><u>21.2 Study hypothesis/background with TCM theory:</u> a brief statement of the study hypothesis or short description of the background of the study. This should be written in language intended to be read and understood by the lay public. For better understanding the rationale of a TCM clinical trial, the rational/background of TCM interventions should be clarified and provided as much as possible based on TCM theory. In this part, a description with TCM theories in background of disease or intervention studied, was denoted to be “1”. However, the assessment criteria in this subitem was according to whether it included TCM related rational/background contents, no matter more or less would be given “1”.</p> <p>21.3 Description of PIO: a description of the included participants, interventions to be tested and outcomes to be measured.</p> <p>Do not include the entire protocol; do not duplicate information recorded in other data elements.</p>
23	Results links	<p>Citations to any publications (including results) related to the trial, if available. Provide either the unique citation database identifier of an article (e.g., Pubmed or Embase), the Digital Object Identifier (DOI) or enter the full bibliographic citation.</p> <p>Links to (abstracts of) articles, conference proceedings, results</p>

		<p>databases, registry results sections, trial websites with results and other links to results can also be provided here.</p> <p>Only available publication website/link was presented as “1”. Following situations were denoted to be “0”: No field of results or absence in this item or results/publication unavailable/ cannot be downloaded, etc.</p>
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Text S₂: Main ID of all included registration records.

ChiCTR (1480)

ChiCTR-DCD-15007073	ChiCTR-EOC-15007674	ChiCTR-ICC-15006335	ChiCTR-ICR-15005795
ChiCTR-ICR-15005823	ChiCTR-ICR-15006004	ChiCTR-ICR-15006138	ChiCTR-ICR-15006315
ChiCTR-ICR-15006657	ChiCTR-ICR-15006787	ChiCTR-ICR-15006850	ChiCTR-IIR-15007567
ChiCTR-CCH-13003487	ChiCTR-CCH-13003756	ChiCTR-INR-15007417	ChiCTR-INR-15007502
ChiCTR-CNC-12002214	ChiCTR-INR-15007539	ChiCTR-IOR-14005387	ChiCTR-IOR-14005542
ChiCTR-ECS-13003786	ChiCTR-IOR-14005609	ChiCTR-IOR-14005645	ChiCTR-EOC-15007621
ChiCTR-IOR-14005744	ChiCTR-ICR-15006257	ChiCTR-ICR-15006359	ChiCTR-IOR-15006313
ChiCTR-ICR-15006373	ChiCTR-ICR-15006639	ChiCTR-IOR-15006876	ChiCTR-ICR-15006988
ChiCTR-IOR-15007061	ChiCTR-ICR-15007004	ChiCTR-IOR-15007206	ChiCTR-IDR-15007540
ChiCTR-IOR-15007240	ChiCTR-IOR-15007309	ChiCTR-IOR-15007471	ChiCTR-IOR-15007551
ChiCTR-IIR-15007492	ChiCTR-IIR-15007683	ChiCTR-IPR-14005340	ChiCTR-IPR-14005341
ChiCTR-IPR-14005381	ChiCTR-IOC-15006309	ChiCTR-IOR-14005408	ChiCTR-IOR-14005679
ChiCTR-IOR-15005902	ChiCTR-IOR-15006144	ChiCTR-IPR-14005429	ChiCTR-IOR-15006149
ChiCTR-IPR-14005441	ChiCTR-IPR-14005665	ChiCTR-IPR-15005899	ChiCTR-IPR-15006694
ChiCTR-IPR-15006758	ChiCTR-IPR-15006759	ChiCTR-IPR-15006945	ChiCTR-IPR-15007208
ChiCTR-IOR-15006402	ChiCTR-IPR-15007396	ChiCTR-IOR-15006418	ChiCTR-IPR-15007457
ChiCTR-IPR-15007643	ChiCTR-IOR-15006419	ChiCTR-IOR-15006626	ChiCTR-IOR-15006706
ChiCTR-OCC-12002329	ChiCTR-IOR-15006768	ChiCTR-OCH-11001546	ChiCTR-IOR-15006812
ChiCTR-OCH-12003138	ChiCTR-IOR-15006973	ChiCTR-OCH-13003043	ChiCTR-OCH-13004245
ChiCTR-OCH-14004683	ChiCTR-IOR-15007366	ChiCTR-OCH-14004724	ChiCTR-OCH-14005137
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ChiCTR-IOR-15007545	ChiCTR-ONC-13003230	ChiCTR-IOR-15007672	ChiCTR-ONC-13003247
ChiCTR-ONC-13003662	ChiCTR-IOR-16007719	ChiCTR-ONRC-08000260	ChiCTR-ONRC-09000511
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ChiCTR-OCB-15006981	ChiCTR-TRC-07000010	ChiCTR-OCC-12002013	ChiCTR-TRC-07000033
ChiCTR-OCH-10000780	ChiCTR-TRC-07000047	ChiCTR-TRC-08000048	ChiCTR-TRC-08000069
ChiCTR-OCH-14004799	ChiCTR-TRC-08000083	ChiCTR-OCS-14004354	ChiCTR-TRC-08000098
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ChiCTR-TRC-08000156	ChiCTR-ONC-11001624	ChiCTR-ONC-12002009	ChiCTR-TRC-08000169
ChiCTR-TRC-08000209	ChiCTR-ONC-13003714	ChiCTR-TRC-08000261	ChiCTR-ONC-13003787
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ChiCTR-TRC-13003716	ChiCTR-TRC-14004180	ChiCTR-TRC-14004181	ChiCTR-TRC-13003717
ChiCTR-TRC-14004182	ChiCTR-TRC-13003780	ChiCTR-TRC-14004235	ChiCTR-TRC-14004293
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ChiCTR-TRC-13003936	ChiCTR-TRC-13004010	ChiCTR-TRC-14004613	ChiCTR-TRC-13004038
ChiCTR-TRC-13004040	ChiCTR-TRC-14004739	ChiCTR-TRC-13004072	ChiCTR-TRC-14004899
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ChiCTR-TRC-14005182	ChiCTR-TRC-14004244	ChiCTR-TRC-14004435	ChiCTR-TRC-14005185
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JPRN-UMIN000013967	JPRN-UMIN000015346	JPRN-UMIN000016625	JPRN-UMIN000017147
JPRN-UMIN000017983	JPRN-UMIN000000909	JPRN-UMIN000000994	JPRN-UMIN000001441
JPRN-UMIN000002140	JPRN-UMIN000003115	JPRN-UMIN000004401	JPRN-UMIN000005645
JPRN-UMIN000005737	JPRN-UMIN000007773	JPRN-UMIN000011349	JPRN-UMIN000011704
JPRN-UMIN000011757	JPRN-UMIN000014899	JPRN-UMIN000015645	JPRN-UMIN000016171
JPRN-UMIN000017520	JPRN-UMIN000018925	JPRN-UMIN000001277	JPRN-UMIN000001659
JPRN-UMIN000002254	JPRN-UMIN000004228	JPRN-UMIN000005332	JPRN-UMIN000005644
JPRN-UMIN000008934	JPRN-UMIN000009676	JPRN-UMIN000010139	JPRN-UMIN000011692
JPRN-UMIN000012683	JPRN-UMIN000013695	JPRN-UMIN000015729	JPRN-UMIN000017936
JPRN-UMIN000018797	JPRN-UMIN000005863	JPRN-UMIN000005899	JPRN-UMIN000015574
JPRN-UMIN000015920	JPRN-UMIN000016632	JPRN-UMIN000017609	JPRN-UMIN000018815
JPRN-UMIN000018929	JPRN-UMIN000022259	JPRN-UMIN000021377	JPRN-UMIN000021065
JPRN-UMIN000022211	JPRN-UMIN000023937	JPRN-UMIN000023856	JPRN-UMIN000023672
JPRN-UMIN000025309	JPRN-UMIN000025301	JPRN-UMIN000025297	JPRN-UMIN000025226
JPRN-UMIN000024994	JPRN-UMIN000024092	JPRN-UMIN000024351	JPRN-UMIN000024008
JPRN-UMIN000030588	JPRN-UMIN000030578	JPRN-UMIN000030273	JPRN-UMIN000027525
JPRN-UMIN000028737	JPRN-UMIN000028604	JPRN-UMIN000028266	JPRN-UMIN000028218
JPRN-UMIN000028216	JPRN-UMIN000028222	JPRN-UMIN000028044	JPRN-UMIN000028017
JPRN-UMIN000027474	JPRN-UMIN000026995	JPRN-UMIN000026141	JPRN-UMIN000025602
JPRN-UMIN000022018	JPRN-UMIN000022009	JPRN-UMIN000025962	JPRN-UMIN000025691
JPRN-UMIN000025559			

DRKS (27)

DRKS00004302	DRKS00004448	DRKS00005474	DRKS00005977
DRKS00003767	DRKS00004857	DRKS00009803	DRKS00007491
DRKS00003116	DRKS00008562	DRKS00000164	DRKS00009249
DRKS00010191	DRKS00010122	DRKS00010625	DRKS00006876
DRKS00011281	DRKS00011079	DRKS00011038	DRKS00013160
DRKS00013158	DRKS00012915	DRKS00012578	DRKS00012623
DRKS00012664	DRKS00012464	DRKS00011525	

REBEC (33)

RBR-3xktv5	RBR-58yq52	RBR-6w25zv	RBR-8whr7r
RBR-9tcn2x	RBR-4qxt3z	RBR-5g7xqh	RBR-5x69x2
RBR-2t9t7q	RBR-42kz9z	RBR-59f4yr	RBR-3w2p32
RBR-42c8wp	RBR-7cfctd	RBR-4n73hw	RBR-5vrgch
RBR-4msydx	RBR-84jx98	RBR-3dmsbk	RBR-5ndq22
RBR-45zj7b	RBR-69fnqt	RBR-4ndjtp	RBR-5t27sb
RBR-9h77k7	RBR-8f5mny	RBR-53bmfx	RBR-43byv6
RBR-9tk8z	RBR-4y8vd2	RBR-77y2sp	RBR-9kwjgx
RBR-5q9svp			

CTRI (12)

CTRI/2017/12/010901	CTRI/2017/09/009806	CTRI/2017/09/009589	CTRI/2017/08/009541
CTRI/2017/08/009384	CTRI/2017/08/009346	CTRI/2017/08/009317	CTRI/2017/08/009238
CTRI/2017/07/009046	CTRI/2017/06/008939	CTRI/2017/06/008782	CTRI/2017/01/007716

TCTR (16)

TCTR20131003001	TCTR20140903004	TCTR20141111001	TCTR20150504002
TCTR20131105001	TCTR20150629002	TCTR20160609004	TCTR20160204002
TCTR20171214001	TCTR20170823002	TCTR20170822002	TCTR20170818002
TCTR20170928004	TCTR20170612002	TCTR20170508002	TCTR20170215003

EU-CTR (6)

EUCTR2006-000947-24-GB	EUCTR2006-000810-18-GB	EUCTR2008-002343-16-IT	EUCTR2015-004250-18-SE
EUCTR2013-004657-24-GB	EUCTR2017-000145-40-ES		

NTR (5)

NTR1987	NTR3132	NTR4059	NTR4325
NTR4787			

PACTR (5)

PACTR201501000986244	PACTR201504001033387	PACTR201711002771340	PACTR201705002261398
PACTR201701001957330			

SLCTR (2)

SLCTR/2017/039	SLCTR/2017/029		
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Text S₃: Descriptive information of recruitment countries/regions.

Countries/Regions	Number of records	Percentage of records (%)
China mainland	1918	57.44
United States	308	9.22
Korea	191	5.72
Iran	181	5.42
Japan	93	2.79
Taiwan	88	2.64
Australia	82	2.46
German	70	2.10
Brazil	67	2.01
United Kingdom	65	1.95
Hong Kong	39	1.17
Spain	29	0.87
Israel	23	0.69
Canada	19	0.57
Sweden	15	0.45
France	13	0.39
Thailand	13	0.39
India	13	0.39
Denmark	11	0.33
Italy	11	0.33
Austria	9	0.27
Norway	9	0.27
Saudi Arabia	8	0.24
New Zealand	8	0.24
Egypt	7	0.21
Singapore	6	0.18
The Netherlands	6	0.18
Switzerland	5	0.15
Portugal	5	0.15
Tunisia	4	0.12
Mexico	3	0.09
Ireland	3	0.09
Belgium	2	0.06
Turkey	2	0.06

Nigeria	2	0.06
Greece	1	0.03
Uganda	1	0.03
Malaysia	1	0.03
Czech Republic	1	0.03
Ecuador	1	0.03
Bangladesh	1	0.03
Croatia	1	0.03
Ukraine	1	0.03
Poland	1	0.03
Nepal	1	0.03
Sri Lanka	1	0.03