ICTRP Registry Profile

When complete, this form should be returned to the ICTRP Secretariat by emailing to ictrpinfo@who.int.

Instructions

This form is to be completed by all registries applying for the status of Primary Registry in the WHO Registry Network. Please refer to the ICTRP web site for detailed guidance on how to submit an application (http://www.who.int/ictrp/en/).

Before completing this form it is strongly recommended that the Registry Administrator obtain a copy of the International Standards for Clinical Trial Registries. These Standards can be obtained by asking the ICTRP Secretariat for access to the Registry Network Sharepoint. The Sharepoint contains a number of useful documents including the Standards, which provide detailed information on the expectations of Primary Registries. The numbering of each item in the Standards is directly compatible with the numbering of each item on this profile form.

ALL questions on this form must be answered before the form is submitted. Incomplete applications will result in a delay in the consideration of your application.

The answer to all questions on this form (except those marked with an asterisk *) must be *"yes"* if a registry is to be considered potentially eligible for Primary Registry status.

Questions marked with an asterisk (*) provide valuable background information and must be completed.

If a registry achieves Primary Registry status then the profile will be published on the ICTRP web site.

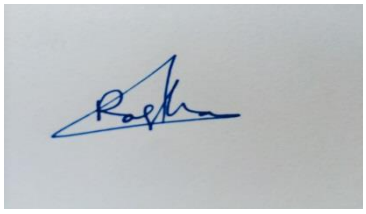
Updating

All Primary Registries are required to keep their profile up-to-date. Once each year, each Primary Registry will be required to resubmit an updated profile. This annual profile will be used to monitor registries and ensure continued compliance with ICTRP requirements. All profiles will be published on the ICTRP web site.

Approval Process for new applications

The completed Registry Profile should be submitted to the ICTRP Secretariat, along with the letter/s of support from the relevant ministry and/or other relevant national or regional agency/ies demonstrating that the registry has at least a national remit. The application will be reviewed by the ICTRP Primary Registry Panel and the registry will receive formal notification of the result of their application within 12 weeks of the application deadline. Submission deadlines and more details on the approval process are available on the ICTRP web site: http://www.who.int/ictrp.

Basic Information

I.	Name of person completing this form	**Rasha Hamra**
II.	Role of the person completing this form (that is, the role of this person in the registry making this application)	**Main Administrator** **Focal Point for Regulating Clinical Trials in Lebanon**
III.	Date this form was completed	____17__ / ___09___ / ___2020_____ *dd* *mm* *yyyy*
IV.	Signature of person completing this form	
V.	Official Name of Registry *(list abbreviation used, if relevant)* Eg Australian and New Zealand Clinical Trials Registry (ANZCTR)	**Lebanese Clinical Trial Registry** **LBCTR**
VI.	Registry postal address	**Ministry of Public Health** **Beir Hassan Highway, next to Ogero** **Beirut, Lebanon**
VII.	Registry street address (if different to postal address)	**Same as above**
VIII.	Registry URL *(Registry's web site address)*	**https://lbctr.moph.gov.lb**
IX.	Registration URL *(Web site where users go to register a trial)*√	**https://lbctr.moph.gov.lb**
X.	Application type	Application for Primary Registry status √ Application for Partner Registry status ☐
XI.	What is the name of the agency (or agencies) that funds the registry?	**Lebanese Ministry of Public Health**
XII.	What is the name of the agency that manages the registry?	**Lebanese Ministry of Public Health**

XIII.	Does the registry have an Advisory Board? *	Yes ☐ No √
XIV.	If the registry has an Advisory Board, please describe its terms of reference, which organizations are represented on the Board, and how often it meets. If necessary, please attach this information to the application as a separate document.	NA
XV.	Is registration of a clinical trial a legal requirement in the country (or countries) covered by the registry? * **If yes**, please provide the title of the relevant law and information on how a copy of the law can be obtained (including the relevant web address)	Yes ☐ No √
XVI.	Is registration of a clinical trial a requirement to obtain ethics approval in the country (or countries) covered by the registry? * **If yes**, please provide the title of the relevant document describing this requirement and information on how this document can be obtained (including the relevant web address)	Yes ☐ No √
XVII.	Is the registry currently accepting clinical trials for registration?	Yes √ No ☐
XVIII.	How many trials are on your database? On __17___/__09___/___2020_____ there were _74_ trials on the registry database. <u>Note</u>: The ICTRP cannot consider a registry for Primary Registry status until it contains at least 10 trials submitted directly by Responsible Registrants. It is not acceptable for a registry to only include trial information that has been downloaded and imported from another registry.	
XIX.	Does the registry agree in principle to comply with the International Standards for Clinical Trial Registries?	Yes √ No ☐

1. Content

1.1. The Registry will accept prospective registration of interventional clinical trials submitted by Responsible Registrants

1.1.1. Does the Registry register trials before the first participant has been recruited?	Yes √ No ☐
a) Does the Registry query submissions where registration is being sought after the first participant has been recruited?	Yes √ No ☐
b) Does the Registry make it clear to Responsible Registrants that prospective registration means that a trial must complete the registration process, **and** have a trial registration number issued, **before** the recruitment of the first participant.	Yes √ No ☐
1.1.2. Does the Registry register trials that have already recruited the first participant? (Also referred to as retrospective registration)	Yes √ No ☐
1.1.3. Does the Registry register other types of studies, including observational studies	Yes √ No ☐
1.1.4. Does the Registry register all trials submitted by Responsible Registrants	Yes √ No ☐
a) Does the Registry accept trials submitted by Responsible Registrants?	Yes √ No ☐
b) Does the Registry ask the person submitting a trial for registration to verify that they meet the terms and conditions for being a Responsible Registrant before being able to proceed to trial registration.	Yes √ No ☐
c) Does the Registry use the contact details provided by the Responsible Registrant to verify that those details are correct?	Yes √ No ☐
d) As a minimum, Registries must send an email to the address given and receive a reply from that same address. When possible, the telephone number and/or surface mail address will also be verified in a similar fashion.	Yes √ No ☐
e) Does the Registry ensure that all Responsible Registrants are associated with an institution or organization?	Yes √ No ☐
f) Does the Registry obtain institutional contact details for the Responsible Registrant, including name and telephone number of the institution?	Yes √ No ☐
1.1.5. Does the Registry accept studies for registration when the data is submitted as an electronic data file (eg as an xml file) Note: This is not a requirement and is being asked for information only.	Yes ☐ No √

1.2. The registry will be open to all prospective registrants (either internationally or within one or more specific countries) *(ICMJE requirement)*

a) Which types of study does the Registry accept for registration? (*tick all that apply*) √ Interventional studies √ Observational studies
b) If registration is restricted in some way (eg only accepts trials from a particular sponsor, or in a particular health care condition (eg cancer) or intervention) please specify how it is restricted: **Not restricted**
c) From which countries does the Registry accept trials for registration: **Only from Lebanon for the time being**

1.3. The Registry will be able to collect and publicly display the WHO Trial Registration Data Set (TRDS) *(ICMJE requirement)*.

a) Does the Registry collect and display, on a publicly accessible web site, all of the items in the WHO Trial Registration Data Set	Yes √ No ☐
b) Does the Registry have quality control procedures in place to ensure all items in the TRDS contain meaningful data?	Yes √ No ☐
c) Does the Registry collect the optional TRDS data items? If yes, please specify: **Lay summary** **Ethical Approvals**	Yes √ No ☐

1.4. The Registry will make an effort to keep registered information up-to-date.

a) Does the Registry permit Responsible Registrants to update information about their trial?	Yes √ No ☐
b) Does the Registry have a reminder system to facilitate the submission of updated information by the Responsible Registrant. If yes, please state how often the Registry reminds Responsible Registrants to update their data (eg once ever 6 months; once every year): **Once per year** Note: This is not a requirement and is being asked for information only.	Yes √ No ☐
c) Does the Registry display the date the trial record was last updated? Note: This is not a requirement and is being asked for information only.	Yes √ No ☐
d) Does the Registry request updates from Responsible Registrants at least annually until the Registrant has recorded meaningful information about the publication of the trial results (e.g. has listed a citation in a "Publications" field) Note: This is not a requirement and is being asked for information only.	Yes √ No ☐

1.5. The Registry will never remove a trial once it has been registered.

a) Does the Registry ever delete a trial record from their database, or remove it from public view, once a registration number has been issued? If yes, please explain the circumstances under which a record would be deleted:	Yes ☐ No √
b) Does the Registry inform Responsible Registrants at the time of registration that a trial cannot be deleted once it has been registered?	Yes √ No ☐
c) Does the Registry have clear and transparent processes for dealing with requests to remove a trial record from public view, and have these processes been documented in the SOPs?	Yes √ No ☐

2. Quality and Validity

2.1. The Registry will have processes in place to make sure that registered data is complete and accurate.

a) Do Registry staff routinely check all data submitted about a trial for completeness and meaningfulness to ensure that all TRDS fields are populated and comply with the minimum standards?	Yes √ No ☐
b) If one or more items in the TRDS submitted for registration are incomplete or not meaningful, do Registry staff contact the Responsible Registrant and attempt to obtain complete and meaningful data?	Yes √ No ☐
c) Does the Registry's database system apply automated checking procedures (e.g. range checks, logic rules) to data items to facilitate validity checking?	Yes √ No ☐
d) Does the Registry have processes in place for deciding whether to register trials where the Responsible Registrant remains non-compliant with requests to provide complete and meaningful data?	Yes √ No ☐
e) Does the Registry undertake regular internal quality control audits to assess the level of completeness and accuracy of the data collected?	Yes √ No ☐

2.2. The Registry will have documented Standard Operating Procedures (SOPs). These SOPs will be aligned with the International Standards for Clinical Trial Registries.

a) Does the Registry have written standards for all procedures and processes employed by the registry? ○ *These written standards are known as Standard Operating Procedures (SOPs)*	Yes √ No ☐
b) Does the Registry use these SOPs to train all staff processing trial registrations to ensure that common standards for ensuring data completeness and meaningfulness are adhered to?	Yes √ No ☐
c) Are the internal registry-specific SOPs aligned with the WHO International Clinical Trial Registry Platform International Standards for Clinical Trial Registration?	Yes √ No ☐

2.3. The Registry will have processes in place to make sure that people and trials exist

2.3.1. Does the Registry make sure that the person registering the trial exists and that they are the appropriate Responsible Registrant?	Yes √ No ☐
2.3.2. Does the Registry make sure that the trial exists? If yes, please briefly describe what the registry does to make sure that the trial exists. **We ask for ethical approvals and for annual updates, SUSARs**	Yes √ No ☐
a) Does the Registry obtain written third-party confirmation that a trial exists? If yes, please specify the method of confirmation: **Ethical approvals from authorized ethical committees in the country** **Registration of CT in other countries**	Yes √ No ☐
b) Does the Registry display in the trial record: a. if the registry has obtained written third party confirmation of the trial's existence, and b. the name of the third party from whom confirmation was received (eg the name of the ethics committee)	Yes √ No ☐

2.4. The Registry will have a publicly accessible audit trail so that changes made to the WHO TRDS for an individual trial can be tracked.

a) Does the Registry allow Responsible Registrants to update their registered trial records?	Yes √ No ☐
b) Does the Registry make available a publicly accessible audit trail of any changes to any TRDS items?	Yes √ No ☐
c) Does the have quality control procedures in place to ensure any updated information continues to fulfil the standards for each TRDS item.	Yes √ No ☐
d) Does the Registry use the most up-to-date information as the default display?	Yes √ No ☐
e) Can the TRDS, as originally registered, be accessed at all times?	Yes √ No ☐

2.5. The Registry agrees to comply with the International Standards for Clinical Trial Registries (ISCTR).

a) Does the Registry Administrator have a thorough working knowledge of the operational aspects of their registry?	Yes √ No ☐
b) Is the Registry Administrator committed to ensuring that all Registry staff are familiar with the standards described in the ISCTR?	Yes √ No ☐
c) Are all Registry staff familiar with the contents of the ISCTR?	Yes √ No ☐

3. Accessibility

3.1. The Registry will make the WHO TRDS for all registered trials accessible to the public at no charge *(ICMJE requirement)*.

a)	Does the Registry make the WHO TRDS items for all studies in their register (ie the registry database) accessible online at no charge to the end user?	Yes √ No ☐

3.2. The Registry will make it possible for the WHO TRDS for all registered trials to be searched electronically *(ICMJE requirement)*.

a)	Is it possible to search the Registry online using electronic searches of text words and phrases via a simple, single search box?	Yes √ No ☐
b)	Does the online search allow users to search in at least the condition field and the intervention field?	Yes √ No ☐
c)	When the results of a trial identified by a search are displayed, are all items in the WHO TRDS visible?	Yes √ No ☐
d)	Does the online search have an advanced search option? Note: This is not a requirement and is being asked for information only.	Yes ☐ No √

3.3. The Registry will allow Responsible Registrants to submit a trial for registration at any time of day on any day of the week (24 hours a day, seven days a week).

a)	Is it possible to submit a trial to the registry 24 hours a day, seven (7) days a week?	Yes √ No ☐
b)	If the Registry is planning downtime, does it publish advance notice of downtime at least one (1) week beforehand?	Yes √ No ☐

3.4. The Registry will allow their register database to be searched at any time of day on any day of the week (24 hours a day, seven days a week).

a)	Is it possible to search the register online 24 hours a day, 7 days a week?	Yes √ No ☐

3.5. It is desirable that Registries in the WHO Registry Network also make the WHO TRDS available in the language(s) of the country or countries served by the registry.

a) Does the Registry accept and/or display trial information in languages others than English? If yes, please specify the languages used:	Yes ☐ No √
Only answer the remaining questions in this section if the answer to the above question is *yes*. ALL the below is Not Applicable	
b) Does the Registry have quality control procedures in place to ensure that all translations are accurate?	Yes ☐ No ☐
c) Are all TRDS items for all records also available in English?	Yes ☐ No ☐
d) Are all trial records translated into English by the Responsible Registrant checked by registry staff against the non-English submission before being accepted for registration?	Yes ☐ No ☐
e) Are all trial records translated by Registry staff checked by at least one other staff member?	Yes ☐ No ☐
f) If there is a discrepancy in a translation, is the translation checked by a third person?	Yes ☐ No ☐
g) Does the Registry make users of the Registry aware of who performed the translation (the Responsible Registrant or Registry staff) of a registered record?	Yes ☐ No ☐
h) If a trial is registered in more than one language then will the Registry submit the "Scientific Title", and a language identifier, to the ICTRP Search Portal for each language used?	Yes ☐ No ☐

4. Unambiguous Identification

4.1. The Registry will have in place processes to prevent the registration of a single trial more than once on their database.

a)	Does the Registry make sure that a trial that has been submitted for registration is not already been included in their register by first searching and checking their own database?	Yes √ No ☐
b)	Does the Registry have policies and procedures in place to deal with inadvertent duplicate registration of the same trial within their own register?	Yes ☐ No √

4.2 The Registry will facilitate the retrospective linking (or bridging) on the WHO Search Portal of a single trial registered with more than one registry by entering secondary identifiers. This includes the UTN, and the unique identifiers allocated by other registries in the WHO Registry Network.

a)	Does the Registry require responsible Registrants to make an entry in the *Secondary Identifiers* field?	Yes √ No ☐
b)	If there are no known secondary identifiers, does the Registry require Responsible Registrants to enter 'Nil known' in the *Secondary Identifiers* field?	Yes √ No ☐
c)	Does the Registry require Responsible Registrants to enter a UTN? a. *The UTN may be entered into either the Secondary Identifiers field or a field designated specifically for collection of the UTN* Note: This is not a requirement and is being asked for information only.	Yes √ No ☐

4.3. It is desirable that Primary Registries will search the ICTRP Search Portal and attempt to determine if the trial has already been registered by another Primary Registry in the WHO Registry Network or an ICMJE approved registry.

a)	Does the Registry attempt to determine whether a submitted trial has been registered in another Primary Registry or an ICMJE approved registry before registration? Note: This is not a requirement and is being asked for information only.	Yes ☐ No √

5. Technical Capacity

5.1. The Registry will submit the WHO TRDS for all records on their register, in English, to the WHO ICTRP Central Repository.

a) Will (or does) the Registry submit the WHO TRDS items for **all records** on their register, in English, to the WHO ICTRP Central Repository? ○ **Note**: If a registry accepts study types other than interventional trials (i.e. observational studies) these must be provided as well.	Yes √ No ☐
b) Will (or does) the Registry submit records in the format requested by the WHO ICTRP (e.g. xml file) at least once per month?	Yes √ No ☐
c) Will (or does) the Registry, after the initial data transfer of all records, only submit new or updated records each time (rather than the entire data set every time)?	Yes ☐ No √

5.2. The Registry will have access to a database that is used to store and manage the submitted data.

a) Does the Registry have access to a database that is used to store and manage the submitted data?	Yes √ No ☐
b) Does the Registry use database software and hardware that guarantees reliable access to registered data and data safety at all times?	Yes √ No ☐

5.3. The Registry will have access to adequate information technology support.

a) Does the Registry have access to reliable information technology support?	Yes √ No ☐
b) Does the Registry have access to all of the following? a. reliable application, database, backup and mail servers b. good internet connectivity speed c. sound operating systems d. appropriate software for servers, desktops and laptops e. database and web development and maintenance personnel f. other skilled information technology personnel to support these systems, as required	Yes √ No ☐
c) Please briefly describe the Registry's information technology infrastructure. (A separate document may be submitted separately if necessary) **Attached separately**	

5.4. The Registry will have adequate security and other provisions against data corruption and loss.

a) Does the Registry have documented procedures for ensuring adequate data security and other provisions to prevent data corruption and loss?	Yes √ No ☐
b) Does the Registry issue alerts in advance of website downtime? If yes, please briefly describe how these alerts are circulated and who is responsible for circulating them.	Yes ☐ No √

6. Administration and Governance

6.1. The Registry will have at least a national remit, and the support of government within the country (or region) to act as the Primary Registry for that country or region (defined as a group of countries and not a group of states within a country).

a) Does the Registry have at least a national remit?	Yes √ No ☐
b) Does the Registry have a letter of support, or other appropriate documentation, from the Ministry of Health or other relevant national or regional agencies? Note: The letters of support must be submitted to the ICTRP Secretariat as part of the application	Yes √ No ☐
c) From which country (or countries) does the Registry have the remit to act as the national (or regional) clinical trial registry? **Lebanon**	
d) Please specify the name of the national agencies that have given their support to the Registry: **Lebanese Ministry of Public Health**	

6.2. The Registry will publicly disclose ownership, governance structure and not-for-profit status.

a) Is the Registry managed by a not-for-profit agency?	Yes √ No ☐
b) Does the Registry publicly disclose its ownership, governance structures and not-for-profit status in a prominent place on the registry's website?	Yes √ No ☐
c) What is the web address for the page where the ownership, governance and not-for-profit information is displayed? **www.moph.gov.lb**	
d) Will the Registry inform the ICTRP immediately if their ownership, governance structures or not-for-profit status change in any way?	Yes √ No ☐

6.3. The Registry agrees that, should it cease to function, at least the WHO TRDS (original and updated) for all trial records will be transferred to a Primary Registry in the WHO Registry Network.

a) Will the Registry transfer at least the WHO TRDS (original and updated) for all trial records to another Primary Registry in the WHO Registry Network if it ceases to function?	Yes √ No ☐

6.4. The Registry will have a strategy in place ensure the medium to long term sustainability of the registry

a) Does the Registry have a documented business plan?	Yes ☐ No ☐ NOT Relevant

b) Does the Registry's business plan include strategies to ensure its medium to long term sustainability?	Yes ☐ No ☐ Not Relevant

7. The Trial Registration Data Set (TRDS)

7.1. The Registry will collect and publicly display all items in the WHO Trial Registration Data Set (TRDS)

	Item/Label	Does the registry collect this data item?	Does the registry publicly display this data item?
1	Primary Registry and Trial Identifying Number	Yes √ No ☐	Yes √ No ☐
2	Date of Registration in Primary Registry	Yes √ No ☐	Yes √ No ☐
3	Secondary Identifying Numbers	Yes √ No ☐	Yes √ No ☐
4	Source(s) of Monetary or Material Support	Yes √ No ☐	Yes √ No ☐
5	Primary Sponsor	Yes √ No ☐	Yes √ No ☐
6	Secondary Sponsor(s)	Yes √ No ☐	Yes √ No ☐
7	Contact for public queries	Yes √ No ☐	Yes √ No ☐
8	Contact for scientific queries	Yes √ No ☐	Yes √ No ☐
9	Public title	Yes √ No ☐	Yes √ No ☐
10	Scientific title	Yes √ No ☐	Yes √ No ☐
11	Countries of Recruitment	Yes √ No ☐	Yes √ No ☐
12	Health condition(s) or problem(s) studied	Yes √ No ☐	Yes √ No ☐
13	Interventions	Yes √ No ☐	Yes √ No ☐
14	Key Inclusion and Exclusion Criteria	Yes √ No ☐	Yes √ No ☐
15	Study type	Yes √ No ☐	Yes √ No ☐
16	Date of first enrolment	Yes √ No ☐	Yes √ No ☐
17	Target sample size	Yes √ No ☐	Yes √ No ☐
18	Recruitment status	Yes √ No ☐	Yes √ No ☐

	Item/Label	Does the registry collect this data item?	Does the registry publicly display this data item?
19	Primary Outcome(s)	Yes √ No ☐	Yes √ No ☐
20	Key Secondary Outcome(s)	Yes √ No ☐	Yes √ No ☐

7.2. The Registry may choose to collect and publicly display other data items. It is recommended that registries consider the following optional, additional data items:

	Item/Label	Does the registry collect this data item?	Does the registry publicly display this data item?
21	Lay Summary / Synopsis	Yes √ No ☐	Yes √ No ☐
22	Approvals	Yes √ No ☐	Yes √ No ☐
23	Results links	Yes √ No ☐	Yes √ No ☐

8. Partner Registries

8.1. Primary Registries in the WHO Registry Network will have the capacity to partner with other Registries.

a) Is the Registry willing and able to form partnerships with other Registries that do not themselves fulfil the criteria for a Primary Registry in the WHO Registry Network?	Yes √ No ☐
b) Does the Registry currently have any Partner Registries? If yes, please provide the name(s) of these partners:	Yes ☐ No √
c) If the registry has partners, are they listed on the proposed Primary Registry's web site? If yes, please provide the address of this web page: Note: A completed Registry Profile form is required for all Partner Registries. This profile will be published on the ICTRP's web site.	Yes ☐ No ☐ **Not Applicable**

8.2. Primary Registries in the WHO Registry Network will ensure that potential Partner Registries meet WHO minimum standards requirements.

a) Does the proposed Primary Registry agree that, before agreeing to accept a Partner Registry and their trial registration records, they will make sure that the Partner Registry meets all the WHO minimum standards listed in the International Standards for Clinical Trial Registries?	Yes √ No ☐
b) Has a Registry Profile form been completed and submitted for all of the Registry's Partner Registries?	Yes √ No ☐

8.3. Primary Registries will have procedures in place to enable exchange of data with Partner Registries.

ALL Not Applicable for Now

a) Is the Registry able to accept data (that is, as electronic data files) from Partner Registries or other appropriate data providers?	Yes ☐ No ☐
b) Does the Registry agree to establish a Memorandum Of Understanding (MOU) or other such agreement with each Partner Registry or other data providers, as per the requirement described in the International Standards for Clinical Trial Registries?	Yes √ No ☐
c) Does the Primary Registry agree the area of coverage/responsibility of their Partner Registries or other data providers (such as geographical location, health condition, intervention type, etc) and incorporate this into their SOPs and instructions to Registrants to avoid any confusion or unintentional duplicate registration?	Yes √ No ☐
d) Does the Primary Registry record the identification number and date of registration in the Partner Registry within the trial record on the Primary Registry?	Yes √ No ☐
e) Does the Primary Registry identify records that have been sourced from Partner Registries or other data providers so users are aware of the data source?	Yes ☐ No ☐
f) Before announcing Partner Registries, Primary Registries must have successfully imported data into the Primary Registry?	Yes ☐ No ☐

Contact Information

Administrator
The Administrator is the person employed to manage the Registry and will be the primary point of contact between the Registry and the ICTRP Secretariat.

Title *(Dr/Prof/Mr/Mrs/Ms/Miss)*	Dr.
Given Name	Rasha
Family Name	Hamra
Telephone number	00961-1-830300. ext 295
Fax	00961-1-843762
Email	rashahamra@yahoo.com

Secondary Contact
The Secondary Contact must not be the same as the Administrator. Secondary contact details are requested for circumstances when the Administrator is unavailable.

Title *(Dr/Prof/Mr/Mrs/Ms/Miss)*	Mrs.
Given Name	Marie
Family Name	Bou Saadah
Telephone number	00961-1-830300, ext 212
Fax	00961-1-843762
Email	tonielie@hotmail.com

Information Technology Officer
The Information Technology Officer should be the person who will be responsible for submitting the data for inclusion on the ICTRP Central Repository. They should have a good understanding of all of the IT issues relevant to the registry.

Title *(Dr/Prof/Mr/Mrs/Ms/Miss)*	Mrs.
Given Name	Lina
Family Name	Abou Mrad
Telephone number	00961-1-830300
Fax	
Email	laboumrad@moph.gov.lb