**WHO Medical Devices Information system (MEDEVIS) including electronic Essential In Vitro Diagnostics list (eEDL)**

**Request for Proposals (RFP)**

Bid Reference

**MVP/EMP/eEDL2019\_1**

Unit Name

**EMP**

**Purpose of the RFP:**

Establishment of a web-based business solution for the WHO Medical devices lists (MEDEVIS) including the Essential In vitro diagnostics list (EDL)

**Closing Date:**

14 August 2019 12:00 hours (noon)   
Geneva, Switzerland

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# Introduction

* 1. Objective of the RFP

The purpose of this Request for Proposals (RFP) is to enter into a contractual agreement with a successful bidder and select a suitable contractor to design and establish a modern digital business solution for online management and web publishing of the WHO Priority Medical devices Lists including the WHO Model List of Essential in vitro diagnostics (EDL).

WHO is an Organization that is dependent on the budgetary and extra-budgetary contributions it receives for the implementation of its activities. Bidders are, therefore, requested to propose the best and most cost-effective solution to meet WHO requirements, while ensuring a high level of service.

* 1. About WHO

### WHO Mission Statement

The World Health Organization was established in 1948 as a specialized agency of the United Nations. The objective of WHO (www.who.int) is the attainment by all peoples of the highest possible level of health. “Health”, as defined in the WHO Constitution, is a state of complete physical, mental and social well being and not merely the absence of disease or infirmity. WHO's main function is to act as the directing and coordinating authority on international health work.

### Structure of WHO

The World Health Assembly (WHA) is the main governing body of WHO. It generally meets in Geneva in May of each year and is composed of delegations representing all 194 Member States. Its main function is to determine the policies of the Organization. In addition to its public health functions, the Health Assembly appoints the Director-General, supervises the financial policies of the Organization, and reviews and approves the proposed programme budget. It also considers reports of the WHO Executive Board, which it instructs with regard to matters upon which further action, study, investigation or report may be required.

The Executive Board is composed of 34 members elected for three-year terms. The main functions of the Board are to give effect to the decisions and policies of the WHA, to advise it and generally to facilitate its work. The Board normally meets twice a year; one meeting is usually in January, and the second is in May, following the World Health Assembly.

The WHO Secretariat consists of some 7,900 staff at the Organization's headquarters in Geneva, in the six regional offices and in countries. The Secretariat is headed by the Director-General, who is appointed by the WHA on the nomination of the Executive Board. The head of each regional office is a Regional Director. Regional directors are appointed by the Executive Board in agreement with the relevant regional committee.

### Description of Cluster/Service/Unit

The department of Essential Medicines and Health Products (EMP) works with Member States and partners to improve access to essential medicines and health products of assured quality for patients, and to promote their rational use. The department works within the Access to Medicines, Vaccines and Health Products division within WHO and contributes significantly towards Universal Health Coverage (UHC), especially in improving access to medicines and health products. Among many, the EMP department has two main areas of work: (1) Regulation of medicines and other health technologies (medicines, vaccines, other biologicals, blood products, medical devices, and diagnostics) and (2) Access, innovation and use of medicines and health products

* 1. Definitions, Acronyms and Abbreviations

**APL:** Priority Assistive Products List. Link: <https://www.who.int/phi/implementation/assistive_technology/EMP_PHI_2016.01>

**DCP:** Disease Commodity Packages: critical medical supplies and consumables and their related technical specifications to directly support emergency responders’ abilities to save lives during an outbreak. Link: <https://www.who.int/emergencies/what-we-do/prevention-readiness/disease-commodity-packages/en/>

**HFA:** Health family assessment tools. Link: <https://www.who.int/healthinfo/topics_standards_tools_data_collection/en/>

**SARA:** Service availability and readiness assessment. Link: <https://www.who.int/healthinfo/systems/sara_introduction/en/>

**ICD:** The International Classification of Diseases is the standard diagnostic tool for epidemiology, health management, and clinical purposes. This includes the analysis of the general health situation of population groups. It is used to monitor the incidence and prevalence of diseases and other health problems, providing a picture of the general health situation of countries and populations. Links: <https://www.who.int/classifications/en/> and <https://www.who.int/classifications/icd/en/>

**ICD-11:** International Classification of Diseases 11th Revision. Link: <https://icd.who.int/en/>

**ICMD:** International Classification of Medical Devices (under development , codes available by 4Q)

**ICHI:** International Classification of Health Interventions. Link: <https://www.who.int/classifications/ichi>

**Medical device (MD):** any article, apparatus, instrument, machine, appliance, implant, reagent for in vitro use, software, material or other similar related articles, intended to be used, alone or in combination, for human beings, for one or more of the specific medical purpose(s) of:

• diagnosis, prevention, monitoring, treatment or alleviation of disease,

• diagnosis, monitoring, treatment, alleviation of or compensation for an injury,

• investigation, replacement, modification, or support of the anatomy or of a physiological process, supporting

or sustaining life,

• control of conception,

• disinfection of medical devices, or

• providing information by means of in vitro examination of specimens derived from the human body;

and does not achieve its primary intended action by pharmacological, immunological or metabolic means, in

or on the human body, but which may be assisted in its intended function by such means”.

**MEDEVIS (****Medical Devices Information System)**: WHO Acronym for the digital business solution and electronic platform for managing all WHO medical devices .

**In vitro diagnostics (IVD):** a subset of medical devices, defined as devices which, whether used alone or in combination, are intended by the manufacturer for the in-vitro examination of specimens derived from the human body solely or principally to provide information for diagnosis or monitoring for compatibility purposes. They include reagents, calibrators, control material and test kits.

**EDL:** WHO Model list of **E**ssential in vitro **D**iagnostics

**eEDL:** **e**lectronic Model list of **E**ssential in vitro **D**iagnostics

**EDL Secretariat**: WHO technical team in charge of EDL and eEDL projects

**EML:** WHO Model Lists of Essential Medicines. Link: <https://www.who.int/medicines/publications/essentialmedicines/en/>

**eEML:** soon to be released electronic version of EML

**EMP:** WHO Essential Medicines and Health Products Department**RMNCH:** Interagency List of Medical Devices for Essential Interventions for Reproductive, Maternal, Newborn and Child Health <https://www.who.int/medical_devices/publications/interagency_med_dev_list/en/>

**LPMD:** WHO list of priority medical devices for cancer management

<https://www.who.int/medical_devices/publications/priority_med_dev_cancer_management/en/>

**MVP:** WHO division for Vaccines and Health products

**UHC:** Universal health coverage. General info: <https://www.who.int/news-room/fact-sheets/detail/universal-health-coverage-(uhc)>

**UHC menu:** WHO repository of guidelines and interventions – not published yet.

# BACKGROUND: DESCRIPTION OF PRESENT ACTIVITIES

Description of the existing activities currently undertaken by **EMP** and related to the objective of this Request for Proposals.

* 1. Overview and current approach

During the World Health Assembly (WHA60.29, 2007), Member States required the WHO Director Genera to: “..(6) establish and update regularly an evidence and web-based health technologies database to serve as a clearing house which will provide guidance on appropriate Medical devices according to levels of care, setting, environment, and intended health intervention, tailored to the specific needs of country or region”.

In 2018, the World Health Organization (WHO) has established a Strategic Advisory Group of Experts on In Vitro Diagnostics (SAGE IVD) to act as an advisory body to matters of global policies and strategies related to in vitro diagnostics (IVDs). The SAGE IVD meets annually to make recommendations on the content, format and implementation of the Model List of Essential In Vitro Diagnostics (EDL) and on any matters related to IVDs. The EDL is published in May every year with new tests and modification of current tests. The main list and the changes have to be reflected in the platform.

* 1. Objectives of the present activity

The Medical Devices lists including EDL are recommended by WHO for use in countries. The lists are not intended to be prescriptive; rather, countries should decide which medical devices and IVDs to select and where to use them, depending on their epidemiology, human resources and infrastructure. They are expected to provide guidance and serve as a reference for Member States (programme managers, laboratory managers, procurement officers and reimbursement officers) that are developing and/or updating national lists for interventions within universal health coverage and for selecting and using medical devices and IVDs. They are also informative for United Nations agencies and nongovernmental organizations that support selection, procurement, supply, donation or provision of medical devices and inform the private medical technology sector about the priorities to address global health issues.

This process would be highly facilitated by a free of access and easy to use electronic platform containing all IVDs in the EDL.

* 1. Activity coordination

Medical Devices team aims to ensure improved access, quality and use of safe and appropriate medical devices to advance universal health coverage, address health emergencies, and promote healthier populations. And advisory group of experts for medical devices, will support the development of the electronic platform and will be consulted for input as needed. The main responsible team in WHO is the medical devices team that includes the EDL secretariat.

# requirements

* 1. Introduction

WHO requires the successful bidder, the Contractor, to carry out the task of developing and successfully implementing the business solution for WHO Medical Devices lists including the Essential in vitro diagnostics list (EDL) as described and underlying including platform setup/configuration and data migration/pre-population with the relevant content

* 1. Characteristics of the provider

### Status

The provider shall be a for profit institution operating in the field of developing, implementing and maintaining secure online business and workflow solutions with proven expertise in health or non-health related areas.

### Accreditations

An accreditation (and certification related to advanced digital information management, online e-communication, business analysis/engineering and software development by Microsoft, IBM, SalesForce, SAP, Oracle; as well as international bodies such as ISO, PM Institute or equivalent etc.) or an on-going accreditation process by a certified accreditation body will be an asset.

### Previous experience

Proven experience in implementing web based software solutions which efficiently handle online workflow processes, online submissions and scientific reviews, as well as business intelligence/data queries and reporting for global audiences is required.

Previous work with WHO, other international organizations and/or major institutions in the field of online business solution design and information management for health and life sciences in an international context /for a muntinational organization is desirable.

### Staffing

The bidder should indicate how they intend to staff the project and provide the profiles of key resources proposed for this project. All resources that will have contact with WHO should be fully fluent in English (spoken and written). Continuity of resources across the duration of the project is expected.

***Please refer to Annex 4: Information about Bidder for structuring your response to each of the above aspects of 3.2 Characteristics of the contractor***

The successful bidder will need to demonstrate its personnel’s capability and the capacity to conduct this project. Details of staff (including CV’s of key personnel) which the bidder shall assign to the project, together with approximate proportions of the time allocated to the work, and the phases of the project in which they will be active, shall be provided as part of the RFP response. A contingency plan should be described in case of absence of key personnel..

* 1. Work to be performed

The work to be performed by the successful bidder is to establish a web-based business solution for the WHO Medical Devices Lists including the List of Essential in vitro diagnostics. It comprises the detailed design, development and full implementation of a web based (online) workflow and process management solution for the Medical Devices list presented in a phased approach.

WHO requires the successful bidder(s) to fully understand the current and forward-looking scope of our needs, identify the most suitable technical solution and to support WHO beyond the successful go-live of the business solution.

The scope of this project and activities is summarized as follows:

1. It shall allow for editing and updating the related information online through an admin and data entry/maintenance interface for WHO and external (non-WHO) users and greatly enhance efficiency and effectiveness of the review and publishing process, and establish highly automated processes.

2. Secondly, the new online platform shall allow a global audience of an estimated 100,000 daily

users to conduct simplified and at the same time efficient and browser independent information

retrieval (searching, filtering, viewing and downloading relevant information), on a variety of devices

including computers, smart phones, and tablets.

3. The proposed online platform shall comprise a wealth of additional data via aggregation of WHO managed databases information such as EML, WHO guidelines, UHC menu, ICHI, ICD-11 to provide additional reference information for the user. Linkages to other sources from UN organization, partners or approved agencies (UNICEF, UNFPA, UNOPS, IAEA, etc) shall be included also. Access to the information resources provided in and via the platform shall be granted free of charge to a public audience. The use of the platform and its information content shall be subject to the acceptance of the respective WHO corporate and Partners’ Copyright and licensing schemes.

The new online portal shall therefore become and serve as a WHO Public knowledge repository and clearing house for Medical Device information, offering seamless integration of data shared with partner organizations.

The planned project activities comprise the initial design workshops, compilation and approval of the design proposal, development, testing towards full implementation and go-live, as well and handover to and training of the business users. This RFP also expects to include an appropriate service delivery proposal for expanding the tool to creating new categories or fields, uploading information to expand to all medical devices, maintaining and supporting the business solution and ensuring its operational sustainability.

At a minimum, WHO envisions that this work will entail the following:

1. Implementation and integration of the proposed technology within the WHO hosting and network infrastructure
2. Development and delivery of the Medcial devices project including (non-exhaustive):
   * Design and specifications
   * Configuration and development
   * Data Migration
   * User Acceptance Testing (UAT) support
   * Documentation and training materials
   * Change management and user adoption
   * Launch, post launch hyper-care, and warranty service
3. Following go-live the service and support of the platform and finished project for a minimum of 2 years (with possibility of extension)
4. Readiness to undertake new projects using this platform.

### Key requirements

The key requirement is to establish a modern online business solution and workflow application for managing all aspects of WHO Medical Devices workflow processes. This project and its activities propose to transform the current lists (LPMD, RMNCH and EDL) and the process of their creation/maintenance from manually editing Excel,Word, PDF documents into a digital workflow and publishing solution.

Key work processes includes online data submission, online commenting and reviewing as well publishing of approved data; moreover, the sought business solution is expected to provide advanced and user friendly tools for online data mining, digital reporting and features for programmatic data exchange with external partners (via web api etc.).

The product delivered by the contractor shall offer all qualities and all required features as outlined in the attached “Detailed Requirements specifications” document (Appendix 1).

**A major organisational requirement of the project is developing and establishing the business solution while prioritizing the needs for managing the electronic Essential In vitro Diagnostics List (eEDL) first.**

The project therefore comprises two major components which the bidder can deliver in two phases and which allow grouping activities according to the following stages:

1. **Deliverable 1: WHO** **Medical Devices information system (MEDEVIS) Online Platform Infrastructure implemented for eEDL**
   1. “MEDEVIS” Online Platform build and administrative components implemented
   2. Portal for Online submissions designed and released for testing
   3. eEDL workflow process designed and released for testing
   4. eEDL List update and publishing module released for testing
   5. data connectivity/Integration with IVD types (currently 250) achieved and released for testing
   6. Data queries and reports designed and released for testing
   7. UAT carried out and passed
   8. Data migration or entry carried out as applicable
   9. Go-live of the eEDL business solution – eEDL operational
   10. Support contract issued for post-go-live service delivery
2. **Deliverable 2: WHO Medical Devices (MEDEVIS) Online Platform Infrastructure extended/enhanced for additional medical devices lists and categories** 
   1. MEDEVIS portal extended for medical devices lists with multiple level and flexible categories released (currently 1500, available on the Interagency List of Medical Devices for Essential Interventions for Reproductive, Maternal, Newborn and Child Health or the WHO list of priority medical devices for cancer management)
   2. workflow process design updated/enhance and released
   3. MEDEVIS List update and publishing module released
   4. data connectivity/Integration achieved and integrated between the MEDEVIS Online portal and external partners
   5. UAT carried out and passed
   6. Data migration or entry carried out as applicable
   7. Go-live of the MEDEVIS business solution and operational
   8. Support contract amended and issued for post-go-live Service delivery

*Please refer to the attached “Detailed Requirements Specifications” document which describes in more detai. the overall objectives, concept and detailed business requirements.*

**Project management and timelines:**

As outlined above, the project is set to comprised of 2 major project phases with major outputs. The major organisational requirement for project design and management is the ability to develop and establish the overarching MEDEVIS business solution’s platform while

* establishing the electronic Essential **In vitro Diagnostics** List (eEDL) first **and within 5 months after contract issuance**
* thereafter expanding MEDEVIS to comprise all remaining medical devices lists and their management

The project plan has tight timelines and requires close collaboration between the contractor and WHO.

* The project kick-off is expected to occur 1-2 days after signature of the contract, and
* the contractor is expected to commence work 2 weeks after issuance of the contractual agreement and
* deliver the fully functional business solution 5 months after project kick-off.

WHO suggests as Project management methodology of Waterfall and Agile project management principles. (see also below under reporting requirements).

**Functional and non-functional requirements:**

Please refer to Appendix 1 “Detailed Requirements Specifications” document which describes in full the detailed functional and non-functional requirements.

The following is to highlight selected aspects of the functional and non-functional requirements:

*Access and user authentication:*

The platform shall offer seamless Internet based access to the different components of the business solution via a unified interface for internal and external users. It shall provide:

- secure and restricted access to restricted areas via user authentication; providing access for authenticated and WHO authorized users to restricted areas (for example access to the portal administration, advanced management features including on-line submissions of EDL applications, etc. and other advanced data access).

- public access to public areas for anonymous/public users - i.e. for the retrieval of public data only.

User authentication, user management and access authorizations are to be handled by the portal and shall NOT be dependent on WHO Domain (WIMS) authentication.

*Hosting:*

The business solution and its underying software platform are to be hosted on the WHO network (on-premise Drupal or IIS) or preferably WHO Cloud (MS O365, MS Azure).

*Business solution, releases and IT platforms:*

With respect to the proposed approach and functional requirements, the software solution and underlying platform must:

* conform to WHO guidelines for developing, deploying and hosting IT applications in particular (best project management and software development practices, internet security,etc.).
* follow a standard release process for versions: The process of designing and releasing version shall bebased upon
  + The bidder maintaining its Development platform
  + The bidder configuring the WHO User acceptance test platform (UAT) and ensuring proper functioning of the replication process from the development to the UAT platform
* successfully pass the WHO Internet vulnerability test.

Once the UAT version of the software solution has been accepted, the UAT platform is deployed by the WHO IMT department to the WHO production environment.

The proposed business solution

-shall NOT require installation of any software component on client computers or client devices for use of its complete feature set; furthermore, it

- Shall operate as expected on most commonly web browsers (IE, Firefox, Google Chrome) and NOT require a specific web browser for use of its complete feature set;

- Shall be conforming to industry standard best practices in

* web application development, data management
* Ux interface design/usability
* seamless integration with external applications for data exchanges etc. (for example XML data exchange via Web API (Restfull API etc.)

- Shall allow for regular support updates as and when announced by the manufacturer/license holder of required components such as the operating system, database software, application development software etc.) and

* be error free and self-reliant and easily maintainable;
* be adaptable/extensible for future enhancements such as addition of changes to product categorIes/types, the the integration of new WHO reference data/libraries or changes in the workflow process, etc.

The contractor is furthermore expected to offer a guarantee period and propose a comprehensive service delivery model which includes business solution enhancements, platform upgrades and maintenance/user support to WHO for a minimum of two years after successful implementation of the project

### Place of performance

The execution of the work of this bid is not limited to a specific geographic location and much of the work can and will be conducted by the selected contractor remotely. However:

* The selected contractor(s) is expected to have regular contact with the key WHO resources in Geneva, Switzerland conforming to our office hours and holiday schedule.
* On-site interaction and meeting participation is expected for key vendor personnel where face to face engagement is required or advantageous. Such engagement could relate to project management, requirements gathering, acceptance test coordination, etc.
* The selected contractor(s) is expected to make all arrangements for their resources (visas, travel, meals and lodging, etc.).

### Timelines

The bidder is expected to put forth a schedule of work that addresses the timeline expectations indicated in the business requirement annexes. Lacking such information, the bidder should propose a timeline that is optimal for executing the project in an effective manner while allowing for any indicated risks or constraints

### Project supervision & Reporting requirements

Detailed project reporting requirements will be negotiated in the contract made between WHO and the successful bidder(s). As a baseline, WHO expects the selected bidder(s) to issue a bi-weekly progress report, subject to WHO designated project manager approval, covering the achievement of the past week in term of project execution as well as the forecast tasks to be performed in the following weeks along with risks, issues, delays, etc.

WHO will require regular reporting according to set milestones, expected delivery of outputs, and at all relevant project stages. As a general rule, reporting shall be aligned with the chosen Project management methodology (e.g. Prince2, etc.). As appropriate, a mix of Waterfall and Agile project management techniques and principles shall be applied and managed using collaboration tools (MS Teams, Atlassian/Jira etc.).

In addition to the communication and meetings recommended by the adopted Project management methodology, the WHO project lead will require a weekly teleconference and a monthly written report on progress, highlighting any issues and including, but not restricted to, emerging and existing risks, timelines, staffing and budget implications.

A closing report following completion of each phase of the project is also required highlighting barriers encountered, best practice identified and lessons learnt.

Either party can request additional reporting, teleconferences or face to face meetings as required during the course of the project:

**Table 3.3.4: Minimum reporting requirements and timelines for MEDEVIS activities under component (phase) 1**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Activity items (Phase 1)** | **MEDEVIS platform and eEDL**  **Activity** | **What to report & How to report** | **Timeline/ frequency** | **Type of report** |
| 0 | Project kick-off | Upon signature of contract | Max. 2 days following signature | Note for the record |
| 1 | Review detailed Solution approach and design document | 1. Report from the contractor on their current status, documenting the detailed design requirements, deliverables and planned delivery relating to the agreed workplan  2. Review and Approval from IAU | 14 days after kick-off  2 days after receipt of contactor’s report | Single report with proposed detailed design  Single approval |
| 2 | Business solution **design and development** (MEDEVIS platform setup and eEDL) | Summary report outlining   * he current status and level of achieving the deliverable/quality goal of the current deliverable/project step * Timelines and activities pending * Exceptions, Constraints, Support/Changes requested | bi-weekly after completion of Item1 **and until go-live is achieved** | Routine Progress reports |
| 3 | Release of MEDEVIS platform and eEDL prototype and **testing** using data for/from one IVD fully functioning components for at least the data entry/management component | Summary Milestone report outlining   * The current status and level of achieving the deliverable/quality goal of the current deliverable/project step * Timelines and activities pending * Exceptions, Constraints, Support/Changes requested | 2 months after completion of Phase 1 | Milestone report |
| 4 | **UAT Testing** of full eEDL solution which needs to have fully functioning components for its three components 1) data entry/management and 2) web based search  3) integration/federation with external sources such ICD. eEML)  Data migration/entry | **1 Contractor:**  Summary Milestone report for **release to User Acceptance Testing (UAT) including test of data migration and each following iteration** *(If any)*outlining   * The current status and level of achieving the deliverable/quality goal of the current deliverable/project step * Timelines and activities pending * Exceptions, Constraints, Support/Changes requested   **2 WHO**  Review report outlining the results of this phase, i.e. based on WHO’s review of the UAT or following iteration   * Acceptance, partial/conditional acceptance or rejection with or without requests to be fulfilled by contractor | 3 months after completion after Phase 1 | Milestone report  WHO Review report |
| 6 | Go-live and post-go service delivery | 1 Certificate of QA testing issued at Go-live (full system release and handover to WHO) assuring the application operates and can be handled/handle and safeguard information as intended  2 Technical Solution Handover Guide for System administrators and IT support contractors  3 End-user manual  4 Agreement on service delivery model and Signature of support contract | 5 months after kick-off | Reports |

### Performance monitoring

Performance will be monitored based on the milestones and targets committed for delivery of specified pieces of work such as those proposed in Section 4. These may be negotiated further while creating a contract between WHO and the successful bidder(s)

### Finance and accounting requirements

For fixed price contracts (e.g. project execution), payments will be issued against acceptance of agreed deliverables and subsequent receipt of invoice.

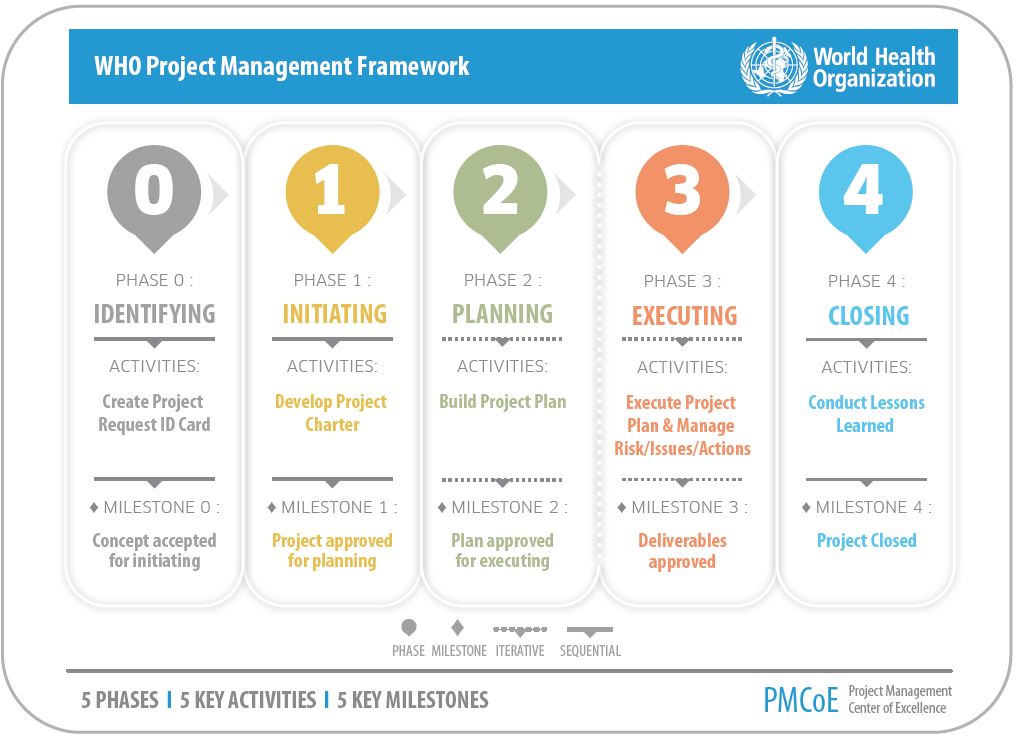
### Further capacities

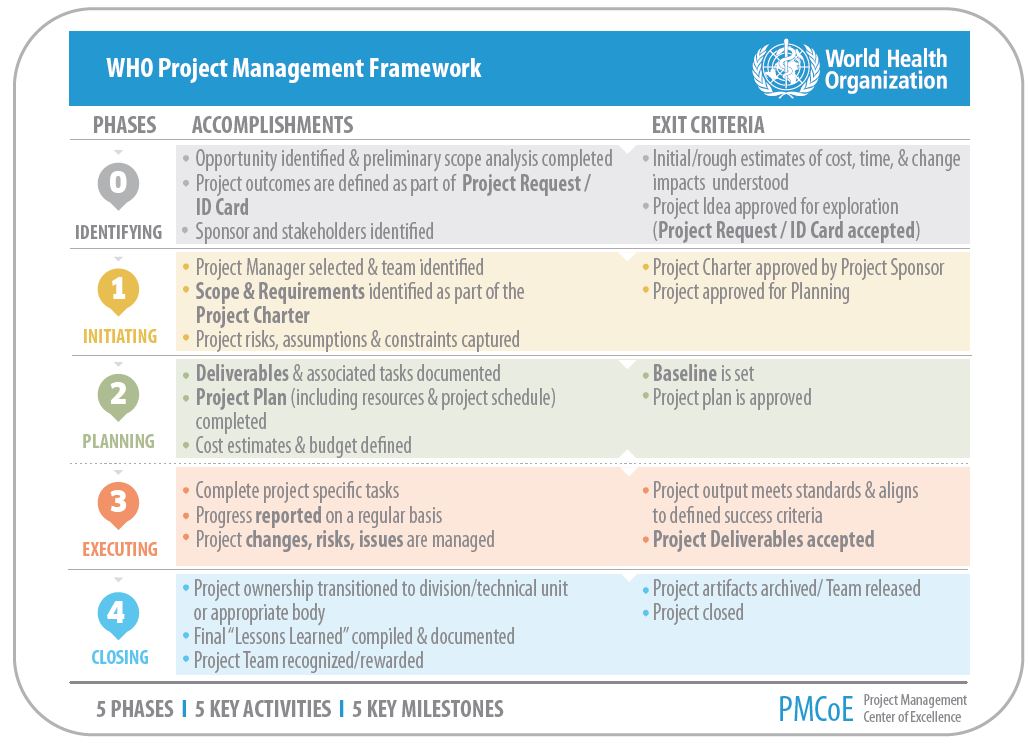
**Project Management**

Bidders are requested to detail their project management methodology which must remain compliant with the WHO Project Management framework. The bidder may propose additional steps if needed, and may propose a Waterfall, Agile, or hybrid approach.

Compliance with WHO Project Management framework is mandatory.

The following graphics outline the WHO Project Management framework used by the WHO IMT Department

*:* 



# Instructions To Bidders

Bidders should follow the instructions set forth below in the submission of their proposal to WHO:

**WHO will not be responsible for any proposal which does not follow the instructions in this RFP, including this Section 4, and may, at its discretion, reject any such non-complaint proposal.**

* 1. Language of the Proposal and other Documents

The proposal prepared by the bidder, and all correspondence and documents relating to the proposal exchanged by the bidder and WHO shall be written in the English language.

* 1. Intention to Bid

**No later than 26 July 2019** the bidder shall complete and return by email to WHO to the following address: [edlsecretariat@who.int](mailto:edlsecretariat@who.int) Subject: Bid Ref. MVP/EMP/eEDL2019\_1 ).

1. The RFP MVP/EMP/eEDL2019\_1 Acknowledgement form, attached hereto as Annex 1, signed as confirmation of the bidder's intention to submit a bona fide proposal and designate its representative to whom communications may be directed, including any addenda; and
2. The RFP MVP/EMP/eEDL2019\_1 Confidentiality Undertaking form, attached hereto as Annex 2, signed;
3. The Self-Declaration form, attached hereto as Annex 6, signed.

These forms are confirming the bidder’s intention to submit a bona fide proposal and designating a representative to whom communications may be directed, including any addenda.

WHO reserves the right to reject proposals from bidders who have not submitted the above-listed forms in accordance with this section.

* 1. Cost of Proposal

The bidder shall bear all costs associated with the preparation and submission of the proposal, including but not limited to the possible cost of discussing the proposal with WHO, making a presentation, negotiating a contract and any related travel.

WHO will in no case be responsible or liable for those costs, regardless of the conduct or outcome of the selection process.

* 1. Contents of the Proposal

Proposals must cover the total requirement. Proposals offering only part of the requirement may be rejected.

The bidder is expected to follow the proposal structure described in paragraph “Proposal Structure” below and otherwise comply with all instructions, terms and specifications contained in, and submit all forms required pursuant to, this RFP. Failure to follow the aforesaid proposal structure, to comply with the aforesaid instructions, terms and specifications, and/or to submit the aforesaid forms will be at the bidder’s risk and may affect the evaluation of the proposal.

* 1. Joint Proposal

Two or more entities may form a consortium and submit a joint proposal offering to jointly undertake the work. Such a proposal must be submitted in the name of one member of the consortium - hereinafter the “lead organization". The lead organization will be responsible for undertaking all negotiations and discussions with, and be the main point of contact for, WHO. The lead organization and each member of the consortium will be jointly and severally responsible for the proper performance of the contract.

* 1. Communications during the RFP Period

A prospective bidder requiring any clarification on technical, contractual or commercial matters may notify WHO via email at the following address no later than **seven (7)** working days prior to the closing date for the submission of offers:

**Email for submissions of all queries:** [edlsecretariat@who.int](mailto:edlsecretariat@who.int)

*(use subject: Bid Ref. MVP/EMP/eEDL2019\_1 )*

The EMP Team at WHO will respond in writing (via email only) to any request for clarification of the RFP that it receives by the deadline indicated above. A consolidated document of WHO's responses to all questions (including an explanation of the query but without identifying the source of enquiry) will be sent to all prospective bidders who have received the RFP.Questions are to be submitted following the format of the form "Questions from Bidders", attached hereto as Annex 4.

There shall be no individual presentation by or meeting with bidders until after the closing date for submission of proposals. From the date of issue of this RFP to the final selection, contact with WHO officials concerning the RFP process shall not be permitted, other than through the submission of queries and/or through a possible presentation or meeting called for by WHO, in accordance with the terms of this RFP.

* 1. Submission of Proposals

The bidder shall submit the complete proposal to WHO **no later than** 14/08/2019 at **12:00 (noon)** hours , Geneva, Switzerland time (“the Closing Date for Submission of Proposals”), by E-mail at the following address: [edlsecretariat@who.int](mailto:edlsecretariat@who.int) *(use subject: Bid Ref. MVP/EMP/eEDL2019\_1 )*

Each proposal should be prepared in two distinct parts: the technical proposal and the financial offer.

Each proposal must include the signed Proposal Completeness Form (attached hereto as Annex 3) and supporting documents, as well as the signed Acceptance Form (attached hereto as Annex 5).

Each proposal shall be marked Bid Ref: MVP/EMP/eEDL2019\_1 and be signed by a person or persons duly authorized to represent the bidder, submit a proposal and bind the bidder to the terms of the RFP.

A proposal shall contain no interlineations, erasures, or overwriting except, as necessary to correct errors made by the bidder, in which case such corrections shall be initialled by the person or persons signing the proposal.

It shall be the Bidder’s responsibility to obtain a confirmation of receipt by WHO of the signed Acknowledgement form (see section “Intention to Bid” 4.24.2 above) and the proposal, marking in particular the Bid Reference number and the date and time of receipt by WHO.

WHO may, at its own discretion, extend the closing date for the submission of proposals by notifying all bidders thereof in writing.

Any proposal received by WHO after the closing date for submission of proposals will be rejected.

**WHO may, at its discretion, reject late bids. Bidders are therefore advised to ensure that they have taken all steps to submit their proposals in advance of the above closing date and time.**

* 1. Period of Validity of Proposals

The offer outlined in the proposal must be valid for a minimum period of **180** calendar days after the closing date for submission of proposals. A proposal valid for a shorter period may be rejected by WHO. In exceptional circumstances, WHO may solicit the bidder’s consent to an extension of the period of validity. The request and the responses thereto shall be made in writing. Any bidder granting such an extension will not, however, be permitted to otherwise modify its proposal.

* 1. Modification and Withdrawal of Proposals

The bidder may withdraw its proposal any time after the proposal’s submission and before the closing date for submission of proposals, provided that written notice of the withdrawal is received by WHO via email or mail as provided in section 4.7 above, prior to the Closing Date for Submission of Proposals.

No proposal may be modified after the closing date for submission of proposals, unless WHO has issued an amendment to the RFP allowing such modifications (see section 4.11 “Amendment of the RFP”).

No proposal may be withdrawn in the interval between the closing date and the expiration of the period of proposal validity specified by the bidder in the proposal in accordance with section 4.8 “Period of Validity of Proposals”.

* 1. Receipt of Proposals from Non-invitees

WHO may, at its own discretion, if it considers this necessary and in the interest of the Organization, extend the RFP to bidders that were not included in the original invitation list.

* 1. Amendment of the RFP

WHO may, at any time before the closing date, for any reason, whether on its own initiative or in response to a clarification requested by a (prospective) bidder, modify the RFP by written amendment. Amendments could, inter alia, include modification of the project scope or requirements, the project timeline expectations and/or extension of the closing date for submission of proposals.

All prospective bidders that have received the RFP will be notified in writing of all amendments to the RFP and will, where applicable, be invited to amend their proposal accordingly.

* 1. Proposal Structure

The contents of the bidder's proposal should be concisely presented and structured in the following order to include, but not necessarily be limited to, the information listed in sections 4.12.1to 4.12.7.

Any information which the bidder considers confidential, should be clearly marked confidential.

### Acceptance Form

The bidder's proposal must be accompanied by the Acceptance Form (see Annex 5, attached) signed by a duly authorized representative of the bidder and stating:

* That the bidder undertakes on its own behalf and on behalf of its possible partners and contractors to perform the work in accordance with the terms of the RFP;
* The total cost of the proposal, indicating the United Nations convertible currency used[[1]](#footnote-2) (preferably US Dollars);
* The number of days the proposal is valid (from the date of the form) in accordance with section 4.8 “Period of Validity of Proposals”.

### Executive Summary

The bidder's proposal must be accompanied by an Executive Summary introducing the proposed solution and approach / methodology.

### Information about Bidders

Bidders should include the following information in their bids.

|  |  |
| --- | --- |
| **RFP Ref.** If applicable | **Information required** |
|  | **1. Company Information** |
|  | **1.1 Corporate information** |
| 3.2.1 | 1.1.1 Company mission statement *(including profit or not for profit status)* |
|  | 1.1.2 Service commitment to customers and measurements used |
| 3.2.2 | 1.1.3 Accreditations |
|  | 1.1.4 Organization structure |
|  | 1.1.5 Geographical presence |
|  | 1.1.6 Declared financial statements for the past (3) three years[[2]](#footnote-3) |
|  | **1.2 Legal Information** |
|  | 1.2.1 History of Bankruptcy |
|  | 1.2.2 Pending major lawsuits and litigations in excess of USD 100,000 at risk |
|  | 1.2.3 Pending Criminal/Civil lawsuits |
| 3.2.3 | **2. Experience and Reference Contact Information** |
|  | **2.1 Relevant Contractual relationships** |
|  | 2.1.1 Relevant Contractual projects (with other UN agencies or Contractors) |
|  | **2.2 Relevant Project Names** *(list and provide detailed examples of relevant experience gained within the past five years of the issuance of this RFP that demonstrate the Contractor’s ability to satisfactorily perform the work in accordance with the requirements of this RFP).* |
|  | 2.2.1 Project Description |
|  | 2.2.2 *Status (under development / implemented)* |
|  | 2.2.3 Reason for relevance *(provide reason why this project can be seen as relevant to this project)* |
|  | 2.2.4 Roles and responsibilities *(list and clearly identify the roles and responsibilities for each participating organization)* |
|  | 2.2.4.1 Client’s Role and Responsibility: Inputs from beneficiary |
|  | 2.2.4.2 Contractor’s Role and Responsibility: role in project |
|  | 2.2.4.3 Third party Contractors’ Role and Responsibility: previously specified 3rd party role in project |
|  | 2.2.5 Team Members *(indicate relevant members of the team that will also be used for this project)* |
| 3.2.4 | **3. Staffing information** |
|  | **3.1 Number and Geographical distribution of staff** |
|  | 3.1.1 Staff turnover rate for the past three years |
|  | **3.2 Staff dedicated to the Project** |
|  | 3.2.1 Name and CV of each team member |
|  | 3.2.2 Structure of the team, and role of each member in the project |
|  | 3.2.3 Time dedicated to the project |
|  | 3.2.3 Contingency plans in the event of a vacancy |
| 4.5 | **4. Proposed sub-contractor arrangements including sub-contractor information** *(as above for each sub-contractor)* |

### Proposed Solution

Please provide details of the proposed solution(s) **together with any assumptions** **you (the Bidder) have made**. This information must accord with the requirements confirmation (e.g. the Appendixes submitted as part of your proposal).

The topics of set-up, implementation work, warranty, post go-live support, and future project services should be individually addressed in this section of your proposal.

Please note that the contractor will be responsible for quantifying and describing requirements for any integration with other systems and any technology and platform requirements within WHO to implement the solution successfully (infrastructure requirements, user interface devices supported etc.)

Bidders are also requested to structure their proposals for the business solution in alignment with the suggested two-phases approach to this RFP.

***The proposal must include the confirmation of achieving the Business Processes, Functional Requirements and Non-Functional Requirements as set forth in the Appendices.***

Please provide the full list of 3rd party solution needed for the proposed solution and why each one been proposed over competing products.

### Approach/Methodology

Please provide details of the proposed Approach to implementation / configuration.

This should include:

* Your proposed Approach for organizing and performing the work, including all assumptions made and requirements for interaction with WHO staff (IT department/Information Management staff and business/subject matter experts) or provision of WHO facilities.
* Project organizational structure indicating key personnel by name and title/role, and their location e.g. offsite or onsite at WHO premise; along with the CVs for key staff.
* A list of work products/deliverables/documentation.
* Key milestone dates.

With respect to the proposal for the business solution to be delivered,

• All deliverables, cost and timelines should be understandable, transparent and stated clearly;

• The contractor will handover design documents, source code, technical and user documentation of the system to WHO. Rights for the source code and documentation of the system and its distribution will remain with WHO;

• The contractor can however, with the permission of WHO, use in full/ part the source code of developed application for further enhancing or modifying the application for other national governments and national and subnational health programs and use the developed application and pictorial, electronic or documented depiction of the developed application in full or part(s) for demonstration of its work or as an example to other organizations/ companies.

• Must contain the recommendations for requirements and cost estimates for cloud hosting; excellent platform and business solution performance is vital (experienced at end-user level)

### Proposed Timeline

Please provide a more detailed schedule/timeline for performing the work which elaborates on the information provided in your Approach. The proposed Timeline project plan should be presented either in MS Project MPP, XLS, or PDF format

### Financial Proposal

The Financial proposal is intended to provide WHO with multiple perspectives on the costs associated with different phases of the project. As a minimum, the bidder must provide the overall cost for the project, as well as a detailed cost breakdown based staff, feature components and system functionality for each phase and major sub-components of the project.

The Financial Proposal has two elements:

1) The cost for your proposal should be documented using the template available in Annex 5-Acceptance Form. The work should be broken into its key components, based on the objectives and outputs described in Section 3.3 (Work to be performed), and indicate the effort assumed for each to provide transparency. Such transparency should include the costs and assumptions made in connection with design, development, and user acceptance test to facilitate our decision making.

2) The bidder also is required to provide a staff rate table to be used for change control purposes and as the basis for future work. The table should be broken down by roles, experience levels, and location. The rates offered must remain valid for 2 years. The rate table can be **appended to the Acceptance Form or submitted separately** **as a Financial document** (do not submit with the Technical Proposal).

In addition to Annex 5, bidders are encouraged to submit additional financial information to facilitate the evaluation process.

* 1. Conduct and Exclusion of Bidders

All bidders must adhere to the UN Supplier Code of Conduct, which is available on the WHO procurement website at the following link: <http://www.who.int/about/finances-accountability/procurement/en/>

In addition, bidders must submit a signed Self Declaration form, attached hereto as Annex 6.

Bidders will be excluded if:

- they are bankrupt or being wound up, are having their affairs administered by the courts, have entered into an arrangement with creditors, have suspended business activities, are the subject of proceedings concerning those matters, or are in any analogous situation arising from a similar procedure provided for in national legislation or regulations;

- they or persons having powers of representation, decision making or control over them have been the subject of a final judgment or of a final administrative decision for fraud, corruption, involvement in a criminal organization, money laundering, terrorist-related offences, child labour or trafficking in human beings;

- they or persons having powers of representation, decision making or control over them have been the subject of a final judgment or of a final administrative decision for financial irregularity(ies);

- it becomes apparent to WHO that they are guilty of misrepresentation in supplying, or if they fail to supply, the information required under this RFP and/or as part of the bid evaluation process;

- they have a conflict of interest, as determined by WHO in its sole discretion; or

- they are, or have found to be, in violation of any standard of conduct as described in the WHO Policies, referred to in section 7.33 of this RFP.

WHO may decide to exclude bidders for other reasons.

# Evaluation Of Proposals

After the closing date for submission of proposals, WHO will open the proposals received in a timely manner.

There will be no public bid opening.

* 1. Preliminary Examination of Proposals

WHO will examine the proposals to determine whether they are complete, whether any computational errors have been made, whether the documents have been properly signed, and whether the proposals are generally in order. Proposals which are not in order as aforesaid may be rejected.

**Please note that WHO is not bound to select any bidder and may reject all proposals.** Furthermore, since a contract would be awarded in respect of the proposal which is considered most responsive to the needs of the project concerned, due consideration being given to WHO’s general principles, including economy and efficiency, WHO does not bind itself in any way to select the bidder offering the lowest price.

* 1. Clarification of Proposals

WHO may, at its discretion, ask any bidder for clarification of any part of its proposal. The request for clarification and the response shall be in writing. No change in price or substance of the proposal shall be sought, offered or permitted during this exchange.

* 1. Evaluation of Proposals

The following procedure will be utilized in evaluating the proposals, with technical evaluation of the proposal being completed prior to any focus on or comparison of price.

The evaluation panel will evaluate the technical merits of all the proposals which have passed the Preliminary Examination of proposals based on the following weighting:

|  |  |
| --- | --- |
| Technical Weighting: | 60 % of total evaluation |
| Financial Weighting: | 40 % of total evaluation |

The technical evaluation of the proposals will include:

A minimum of 60 points is required to pass the technical evaluation.

The bids will be scored according to the table below:



Adherence to Requirements will be the heaviest weighted criteria. The Proposed Approach and Execution criteria will be heavily weighted as well.

During the financial evaluation, the price proposal of all bidders who have passed the technical evaluation will be compared.

* 1. Bidders' Presentations

WHO may, during the evaluation period, at its discretion, invite selected bidders to supply additional information on the contents of their proposal (at such bidders' own cost). Such bidders will be asked to give a presentation of their proposal (possibly with an emphasis on a topic of WHO's choice) followed by a question and answer session. If required, the presentation will be held at WHO or by videoconference.

NOTE: Other presentations and any other individual contact between WHO and bidders is expressly prohibited both before and after the closing date for submission of proposals.

# Award Of Contract

* 1. Award Criteria, Award of Contract

WHO reserves the right to

1. Award the contract to a bidder of its choice, even if its bid is not the lowest;
2. Award separate contracts for parts of the work, components or items, to one or more bidders of its choice, even if their bids are not the lowest;
3. Accept or reject any proposal, and to annul the solicitation process and reject all proposals at any time prior to award of contract, without thereby incurring any liability to the affected bidder or bidders and without any obligation to inform the affected bidder or bidders of the grounds for WHO's action;
4. Award the contract on the basis of the Organization’s particular objectives to a bidder whose proposal is considered to be the most responsive to the needs of the Organization and the activity concerned;
5. Not award any contract at all.

WHO has the right to eliminate bids for technical or other reasons throughout the evaluation/selection process. WHO shall not in any way be obliged to reveal, or discuss with any bidder, how a proposal was assessed, or to provide any other information relating to the evaluation/selection process or to state the reasons for elimination to any bidder.

NOTE: WHO is **acting in good faith** by issuing this RFP. However, **this** **document does not oblige WHO to contract for the performance of any work, nor for the supply of any products or services.**

* 1. WHO's Right to modify Scope or Requirements during the Evaluation/Selection Process

At any time during the evaluation/selection process, WHO reserves the right to modify the scope of the work, services and/or goods called for under this RFP. WHO shall notify the change to only those bidders who have not been officially eliminated due to technical reasons at that point in time.

* 1. WHO's Right to Extend/Revise Scope or Requirements at Time of Award

WHO reserves the right at the time of award of contract to extend, reduce or otherwise revise the scope of the work, services and/or goods called for under this RFP without any change in the base price or other terms and conditions offered by the selected bidder.

* 1. WHO's Right to enter into Negotiations

WHO also reserves the right to enter into negotiations with one or more bidders of its choice, including but not limited to negotiation of the terms of the proposal(s), the price quoted in such proposal(s) and/or the deletion of certain parts of the work, components or items called for under this RFP.

* 1. Signing of the Contract

Within 30 days of receipt of the contract, the successful bidder shall sign and date the contract and return it to WHO according to the instructions provided at that time. If the bidder does not accept the contract terms without changes, then WHO has the right not to proceed with the selected bidder and instead contract with another bidder of its choice.

* 1. Publication of Contract

WHO reserves the right, subject to considerations of confidentiality to acknowledge the existence of the Contract to the public and publish and/or otherwise publicly disclose the Contractor’s name and country of incorporation, general information with respect to the work described herein and the Contract value. Such disclosure will be made in accordance with WHO’s Information Disclosure Policy and shall be consistent with the terms of the Contract.

# General And Contractual Conditions

The contract between WHO and the selected bidder ("the Contract") will, unless otherwise explicitly agreed in writing, include the provisions as set forth in this section, and will otherwise inter alia address the following issues:

* responsibilities of the selected bidder(s) ("the Contractor(s)") and WHO;
* clear deliverables, timelines and acceptance procedures;
* payment terms tied to the satisfactory performance and completion of the work;
* notices.

The prices payable by WHO for the work to be performed under the Contract shall be fixed for the duration of the Contract and shall be in a UN convertible currency (preferably US Dollars), based on the UN exchange rate of the date of invoice. The total amount payable by WHO under the Contract may be either a lump sum or a maximum amount. If the option for payment of a lump sum applies, that lump sum is payable in the manner provided, subject to satisfactory performance of the work. If the option for payment of a maximum amount applies:

1. the Contract shall include a detailed budget;
2. the Contractor shall be held to submit a financial statement together with each invoice;
3. any advance payments by WHO shall be used by the Contractor exclusively for the work in accordance with the budget and any unspent balance shall be refunded to WHO;
4. payment by WHO shall be subject to satisfactory performance and the acceptance of the Contractor's financial statements;
5. to the extent the Contractor is required to purchase any goods and/or services in connection with its performance of the Contract, the Contractor shall ensure that such goods and/or services shall be procured in accordance with the principle of best value for money. "Best value for money" means the responsive offer that is the best combination of technical specifications, quality and price; and
6. consistent with section 7.3,(Audit and Access), all financial reports shall be subject to audit by or on behalf of WHO, including examination of supporting documentation and relevant accounting entries in the Contractor's books. In order to facilitate financial reporting and audit, the Contractor shall keep systematic and accurate accounts and records in respect of the work.

Unless otherwise specified in the Contract, WHO shall have no obligation to purchase any minimum quantities of goods or services from the Contractor, and WHO shall have no limitation on its right to obtain goods or services of the same kind, quality and quantity as described in the Contract, from any other sources at any time.

Unless otherwise specified in the Contract, in the event that the Contract is a Long-Term Agreement (“LTA”), the Contractor shall offer the same prices and terms as those agreed with WHO under the Contract to other interested United Nations system agencies and to organizations eligible to purchase through WHO, it being understood that each such agency and organization will be responsible for independently entering into and administering its own contract with the Contractor. The Contractor shall take into account the additional quantities of services purchased by all United Nations system agencies and other organizations as aforesaid to further reduce the prices for WHO and such other agencies and organizations.

* 1. Conditions of Contract

Any and all of the Contractor's (general and/or special) conditions of contract are hereby explicitly excluded from the Contract, i.e., regardless of whether such conditions are included in the Contractor's offer, or printed or referred to on the Contractor's letterhead, invoices and/or other material, documentation or communications.

* 1. Responsibility

The Contractor will be responsible to ensure that the work performed under the Contract meets the agreed specifications and is completed within the time prescribed.

* 1. Audit and Access

WHO may request a financial and operational review or audit of the work performed under the Contract, to be conducted by WHO and/or parties authorized by WHO, and the Contractor undertakes to facilitate such review or audit. This review or audit may be carried out at any time during the implementation of the work performed under the Contract, or within five years of completion of the work. In order to facilitate such financial and operational review or audit, the Contractor shall keep accurate and systematic accounts and records in respect of the work performed under the Contract.

The Contractor shall make available, without restriction, to WHO and/or parties authorized by WHO:

1. the Contractor’s books, records and systems (including all relevant financial and operational information) relating to the Contract; and
2. reasonable access to the Contractor’s premises and personnel.

The Contractor shall provide satisfactory explanations to all queries arising in connection with the aforementioned audit and access rights.

WHO may request the Contractor to provide complementary information about the work performed under the Contract that is reasonably available, including the findings and results of an audit (internal or external) conducted by the Contractor and related to the work performed under the Contract.

* 1. Source of Instructions

The Contractor shall neither seek nor accept instructions from any authority external to WHO in connection with the performance of the work under the Contract. The Contractor shall refrain from any action which may adversely affect WHO and shall fulfil its commitments with the fullest regard to the interests of WHO.

* 1. Warranties

The Contractor warrants and represents to WHO as follows:

1. The deliverables shall meet the specifications called for in the Contract and shall be fully adequate to meet their intended purpose. The Contractor furthermore warrants that the deliverables shall be error-free. The Contractor shall correct any errors in the deliverables, free of charge, within fifteen days after their notification to the Contractor, during a period of at least one year after completion of the work. It is agreed, however, that errors and other defects which have been caused by modifications to the deliverables made by WHO without agreement of the Contractor are not covered by this paragraph.
2. The deliverables shall, to the extent they are not original, only be derived from, or incorporate, material over which the Contractor has the full legal right and authority to use it for the proper implementation of the Contract. The Contractor shall obtain all the necessary licenses for all non-original material incorporated in the deliverables (including, but not limited to, licenses for WHO to use any underlying software, application, and operating deliverables included in the deliverables or on which it is based so as to permit WHO to fully exercise its rights in the deliverables without any obligation on WHO’s part to make any additional payments whatsoever to any party.
3. The deliverables shall not violate any copyright, patent right, or other proprietary right of any third party and shall be delivered to WHO free and clear of any and all liens, claims, charges, security interests and any other encumbrances of any nature whatsoever.
4. The Contractor, its employees and any other persons and entities used by the Contractor shall not violate any intellectual property rights, confidentiality, right of privacy or other right of any person or entity whomsoever.
5. Except as otherwise explicitly provided in the Contract, the Contractor shall at all times provide all the necessary on-site and off-site resources to meet its obligations hereunder. The Contractor shall only use highly qualified staff, acceptable to WHO, to perform its obligations hereunder.
6. The Contractor shall take full and sole responsibility for the payment of all wages, benefits and monies due to all persons and entities used by it in connection with the implementation and execution of the Contract, including, but not limited to, the Contractor’s employees, permitted subcontractors and suppliers.

Contractor furthermore warrants and represent that the information provided by it to WHO in response to the RFP and during the bid evaluation process is accurate and complete. Contractor understands that in the event Contractor has failed to disclose any relevant information which may have impacted WHO's decision to award the Contract to Contractor, or has provided false information, WHO will be entitled to rescind the contract with immediate effect, in addition to any other remedies which WHO may have by contract or by law.

* 1. Legal Status

The Contractor shall be considered as having the legal status of an independent contractor vis-à-vis WHO, and nothing contained in or relating to the Contract shall be construed as establishing or creating an employer/employee relationship between WHO, on the one hand, and the Contractor or any person used by the Contractor in the performance of the work, on the other hand.

Thus the Contractor shall be solely responsible for the manner in which the work is carried out. WHO shall not be responsible for any loss, accident, damage or injury suffered by the Contractor or persons or entities claiming under the Contractor, arising during or as a result of the implementation or execution of the Contract, including travel, whether sustained on WHO premises or not.

The Contractor shall obtain adequate insurance to cover such loss, accident, injury and damage, before commencing work on the Contract. The Contractor shall be solely responsible in this regard and shall handle any claims for such loss, accident, damage or injury.

* 1. Relation Between the Parties

Nothing in the Contract shall be deemed to constitute a partnership between the Parties or to constitute either Party as the agent of the other.

* 1. No Waiver

The waiver by either Party of any provision or breach of the Contract shall not prevent subsequent enforcement of such provision or excuse further breaches.

* 1. Liability

The Contractor hereby indemnifies and holds WHO harmless from and against the full amount of any and all claims and liabilities, including legal fees and costs, which are or may be made, filed or assessed against WHO at any time and based on, or arising out of, breach by the Contractor of any of its representations or warranties under the Contract, regardless of whether such representations and warranties are explicitly incorporated here in or are referred to in any attached Appendices.

* 1. Assignment

The Contractor shall not assign, transfer, pledge or make any other disposition of the Contract or any part thereof, or any of the Contractor's rights, claims or obligations under the Contract except with the prior written consent of WHO.

* 1. Indemnification

The Contractor shall indemnify and hold WHO harmless, from and against the full amount of any and all claims and liabilities, including legal fees and costs, which are or may be made, filed or assessed against WHO at any time and based on, or arising out of, the acts or omissions of the Contractor, or the Contractor's employees, officers, agents, partners or sub-contractors, in the performance of the Contract. This provision shall extend, inter alia, to claims and liabilities in the nature of workmen's compensation, product liability and liability arising out of the use of patented inventions or devices, copyrighted material or other intellectual property by the Contractor, its employees, officers, agents, servants, partners or sub-contractors.

* 1. Contractor's Responsibility for Employees

The Contractor shall be responsible for the professional and technical competence of its employees and will select, for work under the Contract, reliable individuals who will perform effectively in the implementation of the Contract, respect the local laws and customs, and conform to a high standard of moral and ethical conduct.

* 1. Subcontracting

Any intention to subcontract aspects of the Contract must be specified in detail in the proposal submitted. Information concerning the subcontractor, including the qualifications of the staff proposed for use must be covered with same degree of thoroughness as for the prime contractor. No subcontracting will be permitted under the Contract unless it is proposed in the initial submission or formally agreed to by WHO at a later time. In any event, the total responsibility for the Contract remains with the Contractor.

The Contractor shall be responsible for ensuring that any and all subcontracts shall be fully consistent with the Contract, and shall not in any way prejudice the implementation of any of its provisions.

* 1. Place of Performance

The place of performance of the work under the Contract shall be as mentioned in section 3.3.2 above.

* 1. Language

All communications relating to the Contract and/or the performance of the work thereunder shall be in English.

* 1. Confidentiality

1. Except as explicitly provided in the Contract, the Contractor shall keep confidential all information which comes to its knowledge during, or as a result of, the implementation and execution of the Contract. Accordingly, the Contractor shall not use or disclose such information for any purpose other than the performance of its obligations under the Contract. The Contractor shall ensure that each of its employees and/or other persons and entities having access to such information shall be made aware of, and be bound by, the obligations of the Contractor under this paragraph. However, there shall be no obligation of confidentiality or restriction on use, where: (i) the information is publicly available, or becomes publicly available, otherwise than by any action or omission of the Contractor, or (ii) the information was already known to the Contractor (as evidenced by its written records) prior to becoming known to the Contractor in the implementation and execution of the Contract; or (iii) the information was received by the Contractor from a third party not in breach of an obligation of confidentiality.
2. The Contractor, its employees and any other persons and entities used by the Contractor shall furthermore not copy and/or otherwise infringe on copyright of any document (whether machine-readable or not) to which the Contractor, its employees and any other persons and entities used by the Contractor have access in the performance of the Contract.
3. The Contractor may not communicate at any time to any other person, Government or authority external to WHO, any information known to it by reason of its association with WHO which has not been made public except with the authorization of WHO; nor shall the Contractor at any time use such information to private advantage.
   1. Title Rights
4. All rights pertaining to any and all deliverables under the Contract and the original work product leading thereto, as well as the rights in any non-original material incorporated therein as referred to in section 7.5 2) above, shall be exclusively vested in WHO.
5. WHO reserves the right to revise the work, to use the work in a different way from that originally envisaged or to not use the work at all.
6. At WHO's request, the Contractor shall take all necessary steps, execute all necessary documents and generally assist WHO in securing such rights in compliance with the requirements of applicable law.
   1. Termination and Cancellation

WHO shall have the right to cancel the Contract (in addition to other rights, such as the right to claim damages):

1. In the event the Contractor fails to begin work on the date agreed, or to implement the work in accordance with the terms of the Contract; or
2. In the event the progress of work is such that it becomes obvious that the obligations undertaken by the Contractor and, in particular, the time for fulfilment of such obligations, will not be respected.

In addition, WHO shall be entitled to terminate the Contract (or part thereof), in writing:

* At will with the provision of thirty (30) days prior notice in writing; and
* With immediate effect (in addition to other rights, such as the right to claim damages), if, other than as provided above, the Contractor is:
  1. In breach of any of its material obligations under the Contract and fails to correct such breach within a period of thirty (30) days after having received a written notification to that effect from WHO; or
  2. Adjudicated bankrupt or formally seeks relief of its financial obligations.
  3. Force Majeure

No party to the Contract shall be responsible for a delay caused by force majeure, that is, a delay caused by reasons outside such party's reasonable control it being agreed, however, that WHO shall be entitled to terminate the Contract (or any part of the Contract) forthwith if the implementation of the work is delayed or prevented by any such reason for an aggregate of thirty (30) days. Such termination shall be subject to payment of an equitable part of the Contract sum and/or other reasonable charges. In the event of such termination, the Contractor shall, in accordance with the ownership rights referred to in section 7.17 (Title Rights), deliver to WHO all work products and other materials so far produced.

In the event of and as soon as possible after the occurrence of any cause constituting force majeure, the Contractor shall give notice and full particulars in writing to WHO, of such occurrence or change if the Contractor is thereby rendered unable, wholly or in part, to perform its obligations and meet its responsibilities under the Contract. The Contractor shall also notify WHO of any other changes in conditions or the occurrence of any event which interferes or threatens to interfere with its performance of the Contract. The notice shall include steps proposed by the Contractor to be taken including any reasonable alternative means for performance that is not prevented by force majeure. On receipt of the notice required under this section, WHO shall take such action as it, in its sole discretion, considers to be appropriate or necessary in the circumstances, including the granting to the Contractor of a reasonable extension of time in which to perform its obligations under the Contract.

* 1. Surviving Provisions

Those rights and obligations of the Parties as set forth in sections 7 and 8 that are intended by their nature to survive the expiration or earlier termination of the Contract shall survive indefinitely. This includes, **but is expressly not limited to**, any provisions relating to WHO's right to financial and operational audit, conditions of contract, warranties, legal status and relationship between the parties, breach, liability, indemnification, subcontracting, confidentiality, title rights, use of the WHO name and emblem, successors and assignees, insurance and liabilities to third parties, settlement of disputes, observance of laws, privileges and immunities, no terrorism or corruption, foreign nationals and compliance with WHO policies.

* 1. Use of WHO name and emblem

Without WHO’s prior written approval, the Contractor shall not, in any statement or material of an advertising or promotional nature, refer to the Contract or the Contractor’s relationship with WHO, or otherwise use the name (or any abbreviation thereof) and/or emblem of the World Health Organization.

* 1. Publication of Contract

Subject to considerations of confidentiality, WHO may acknowledge the existence of the Contract to the public and publish and/or otherwise publicly disclose the Contractor’s name and country of incorporation, general information with respect to the work described herein and the Contract value. Such disclosure will be made in accordance with WHO’s Information Disclosure Policy and shall be consistent with the terms of the Contract.

* 1. Successors and Assignees

The Contract shall be binding upon the successors and assignees of the Contractor and the Contract shall be deemed to include the Contractor’s successors and assignees, provided, however, that nothing in the Contract shall permit any assignment without the prior written approval of WHO.

* 1. Payment

Payment will be made against presentation of an invoice in a UN convertible currency (preferably US Dollars) in accordance with the payment schedule contained in the Contract, subject to satisfactory performance of the work. The price shall reflect any tax exemption to which WHO may be entitled by reason of the immunity it enjoys. WHO is, as a general rule, exempt from all direct taxes, custom duties and the like, and the Contractor will consult with WHO so as to avoid the imposition of such charges with respect to this contract and the goods supplied and/or services rendered hereunder. As regards excise duties and other taxes imposed on the sale of goods or services (e.g. VAT), the Contractor agrees to verify in consultation with WHO whether in the country where the VAT would be payable, WHO is exempt from such VAT at the source, or entitled to claim reimbursement thereof. If WHO is exempt from VAT, this shall be indicated on the invoice, whereas if WHO can claim reimbursement thereof, the Contractor agrees to list such charges on its invoices as a separate item and, to the extent required, cooperate with WHO to enable reimbursement thereof.

* 1. Title to Equipment

Title to any equipment and supplies that may be furnished by WHO shall remain with WHO and any such equipment shall be returned to WHO at the conclusion of the Contract or when no longer needed by the Contractor. Such equipment, when returned to WHO, shall be in the same condition as when delivered to the Contractor, subject to normal wear and tear. The Contractor shall be liable to compensate WHO for equipment determined to be damaged or degraded beyond normal wear and tear.

* 1. Insurance and Liabilities to Third Parties

The Contractor shall provide and thereafter maintain:

(i) insurance against all risks in respect of its property and any equipment used for the execution of the Contract;

(ii) all appropriate workmen's compensation insurance, or its equivalent, with respect to its employees to cover claims for personal injury or death in connection with the Contract; and

(iii) liability insurance in an adequate amount to cover third party claims for death or bodily injury, or loss of or damage to property, arising from or in connection with the performance of the work under the Contract or the operation of any vehicles, boats, airplanes or other equipment owned or leased by the Contractor or its agents, servants, employees, partners or sub-contractors performing work in connection with the Contract.

Except for the workmen's compensation insurance, the insurance policies under this section shall:

1. Name WHO as additional insured;
2. Include a waiver of subrogation to the insurance carrier of the Contractor's rights against WHO;
3. Provide that WHO shall receive written notice from the Contractor's insurance carrier not less than thirty (30) days prior to any cancellation or material change of coverage.

The Contractor shall, upon request, provide WHO with satisfactory evidence of the insurance required under this section.

* 1. Settlement of Disputes

Any matter relating to the interpretation of the Contract which is not covered by its terms shall be resolved by reference to Swiss law. Any dispute relating to the interpretation or application of the Contract shall, unless amicably settled, be subject to conciliation. In the event of failure of the latter, the dispute shall be settled by arbitration. The arbitration shall be conducted in accordance with the modalities to be agreed upon by the parties or, in the absence of agreement, with the rules of arbitration of the International Chamber of Commerce. The parties shall accept the arbitral award as final.

* 1. Authority to Modify

No modification or change of the Contract, no waiver of any of its provisions or any additional contractual relationship of any kind shall be valid and enforceable unless signed by a duly authorized representative of both parties.

* 1. Privileges and Immunities

Nothing in or relating to the Contract shall be construed as a waiver of any of the privileges and immunities enjoyed by WHO under national or international law, and/or as submitting WHO to any national court jurisdiction.

* 1. Anti-Terrorism and UN Sanctions; Fraud and Corruption

The Contractor warrants for the entire duration of the Contract that:

(i) it is not and will not be involved in, or associated with, any person or entity associated with terrorism, as designated by any UN Security Council sanctions regime, that it will not make any payment or provide any other support to any such person or entity and that it will not enter into any employment or subcontracting relationship with any such person or entity;

(ii) it shall not engage in any illegal, corrupt, fraudulent, collusive or coercive practices (including bribery, theft and other misuse of funds) in connection with the execution of the Contract; and

(iii) the Contractor shall take all necessary precautions to prevent the financing of terrorism and/or any illegal corrupt, fraudulent, collusive or coercive practices (including bribery, theft and other misuse of funds) in connection with the execution of the Contract.

Any payments used by the Contractor for the promotion of any terrorist activity or any illegal, corrupt, fraudulent, collusive or coercive practice shall be repaid to WHO without delay.

* 1. Ethical Behaviour

WHO, the Contractor and each of the Contractor’s partners, subcontractors and their employees and agents shall adhere to the highest ethical standards in the performance of the Contract. .In this regard, the Contractor shall also ensure that neither the Contractor nor its partners, subcontractors, agents or employees will engage in activities involving child labour, trafficking in arms, promotion of tobacco or other unhealthy behaviour, or sexual exploitation and abuse.

* 1. Officials not to Benefit

The Contractor warrants that no official of WHO has received or will be offered by the Contractor any direct or indirect benefit arising from the Contract or the award thereof.

* 1. Compliance with WHO Codes and Policies

By entering into the Contract, the Contractor acknowledges that it has read, and hereby accepts and agrees to comply with, the WHO Policies (as defined below).

In connection with the foregoing, the Contractor shall take appropriate measures to prevent and respond to any violations of the standards of conduct, as described in the WHO Policies, by its employees and any other persons engaged by the Contractor to perform any services under the Contract.

Without limiting the foregoing, the Contractor shall promptly report to WHO, in accordance with the terms of the applicable WHO Policies, any actual or suspected violations of any WHO Policies of which the Contractor becomes aware.

For purposes of the Contract, the term “WHO Policies” means collectively: (i) the WHO Code of Ethics and Professional Conduct; (ii) the WHO Policy on Sexual Exploitation and Abuse Prevention and Response; (iii) the WHO Code of Conduct for responsible Research; (iv) the WHO Policy on Whistleblowing and Protection Against Retaliation; and (v) the UN Supplier Code of Conduct, in each case, as amended from time to time and which are publicly available on the WHO website at the following links: http://www.who.int/about/finances-accountability/procurement/en/ for the UN Supplier Code of Conduct and at http://www.who.int/about/ethics/en/ for the other WHO Policies.

* 1. Zero tolerance for sexual exploitation and abuse

WHO has zero tolerance towards sexual exploitation and abuse. In this regard, and without limiting any other provisions contained herein:

1. each legal entity Contractor warrants that it will: (i) take all reasonable and appropriate measures to prevent sexual exploitation or abuse as described in the WHO Policy on Sexual Exploitation and Abuse Prevention and Response by any of its employees and any other persons engaged by it to perform any services under the Contract; and (ii) promptly report to WHO and respond to, in accordance with the terms of the Policy, any actual or suspected violations of the Policy of which the contractor becomes aware; and
2. each individual Contractor warrants that he/she will (i) not engage in any conduct that would constitute sexual exploitation or abuse as described in the WHO Policy on Sexual Exploitation and Abuse Prevention and Response; and (ii) promptly report to WHO, in accordance with the terms of the Policy, any actual or suspected violations of the Policy of which the Contractor becomes aware.
   1. Tobacco/Arms Related Disclosure Statement

The Contractor may be required to disclose relationships it may have with the tobacco and/or arms industry through completion of the WHO Tobacco/Arms Disclosure Statement.  In the event WHO requires completion of this Statement, the Contractor undertakes not to permit work on the Contract to commence, until WHO has assessed the disclosed information and confirmed to the Contractor in writing that the work can commence.

* 1. Compliance with applicable laws, etc.

The Contractor shall comply with all laws, ordinances, rules, and regulations bearing upon the performance of its obligations under the terms of the Contract. Without limiting the foregoing or any other provision of these General and Contractual Conditions, the Contractor shall at all times comply with and ensure that each of its partners, subcontractors and their employees and agents comply with, any applicable laws and regulations, and with all WHO policies and reasonable written directions and procedures from WHO relating to: (i) occupational health and safety, (ii) security and administrative requirements, including, but not limited to computer network security procedures, (iii) sexual harassment, (iv) privacy, (v) general business conduct and disclosure, (vi) conflicts of interest and (vii) business working hours and official holidays.

In the event that the Contractor becomes aware of any violation or potential violation by the Contractor, its partners, subcontractors or any of their employees or agents, of any laws, regulations, WHO policies or other reasonable written directions and procedures, the Contractor shall immediately notify WHO of such violation or potential violation. WHO, in its sole discretion, shall determine the course of action to remedy such violation or prevent such potential violation, in addition to any other remedy available to WHO under the Contract or otherwise.

* 1. Breach of Essential Terms

The Contractor acknowledges and agrees that each of the provisions of section 7.30 (Anti-Terrorism and UN Sanctions; Fraud and Corruption), section 7.31 (Ethical Behaviour), section 7.32 (Officials not to Benefit), section 7.33 (Compliance with WHO Codes and Policies), and section 7.36 (Zero tolerance for sexual exploitation and abuse), section 7.35 (Tobacco/Arms Related Disclosure Statement) and section 7.36(Compliance with applicable laws, etc.) hereof constitutes an essential term of the Contract, and that in case of breach of any of these provisions, WHO may, in its sole discretion, decide to:

(i) terminate the Contract, and/or any other contract concluded by WHO with the Contractor, immediately upon written notice to the Contractor, without any liability for termination charges or any other liability of any kind; and/or

(ii) exclude the Contractor from participating in any ongoing or future tenders and/or entering into any future contractual or collaborative relationships with WHO.

WHO shall be entitled to report any violation of such provisions to WHO’s governing bodies, other UN agencies, and/or donors.

# Personnel

* 1. Approval of Contractor Personnel

WHO reserves the right to approve any employee, subcontractor or agent furnished by the Contractor and Contractor's consortium partners for the performance of the work under the Contract (hereinafter jointly referred to as "Contractor Personnel"). All Contractor Personnel must have appropriate qualifications, skills, and levels of experience and otherwise be adequately trained to perform the work. WHO reserves the right to undertake an interview process as part of the approval of Contractor Personnel.

The Contractor acknowledges that the qualifications, skills and experience of the Contractor Personnel proposed to be assigned to the project are material elements in WHO’s engaging the Contractor for the project. Therefore, in order to ensure timely and cohesive completion of the project, both parties intend that Personnel initially assigned to the project continue through to project completion. Once an individual has been approved and assigned to the project, such individual will not, in principle, thereafter be taken off the project by the Contractor, or reassigned by the Contractor to other duties. Circumstances may arise, however, which necessitate that Personnel be substituted in the course of the work, e.g. in the event of promotions, termination of employment, sickness, vacation or other similar circumstances, at which time a replacement with comparable qualifications, skills and experience may be assigned to the project, subject to approval of WHO.

WHO may refuse access to or require replacement of any Contractor Personnel if such individual renders, in the sole judgment of WHO, inadequate or unacceptable performance, or if for any other reason WHO finds that such individual does not meet his/her security or responsibility requirements. The Contractor shall replace such an individual within fifteen (15) business days of receipt of written notice from WHO. The replacement will have the required qualifications, skills and experience and will be billed at a rate that is equal to or less than the rate of the individual being replaced.

* 1. Project Managers

Each party shall appoint a qualified project manager (“Project Manager”) who shall serve as such party’s primary liaison throughout the course of the project. The Project Manager shall be authorized by the respective party to answer all questions posed by the other party and convey all decisions made by such party during the course of the project and the other party shall be entitled to rely on such information as conveyed by the Project Manager.

The Project Managers shall meet on a monthly basis in order to review the status of the project and provide WHO with reports. Such reports shall include detailed time distribution information in the form requested by WHO and shall cover problems, meetings, progress and status against the implementation timetable.

* 1. Foreign Nationals

The Contractor shall verify that all Contractor Personnel is legally entitled to work in the country or countries where the work is to be carried out. WHO reserves the right to request the Contractor to provide WHO with adequate documentary evidence attesting this for each Contractor Personnel.

Each party hereby represents that it does not discriminate against individuals on the basis of race, gender, creed, national origin, citizenship.

* 1. Engagement of Third Parties and use of In-house Resources

The Contractor acknowledges that WHO may elect to engage third parties to participate in or oversee certain aspects of the project and that WHO may elect to use its in-house resources for the performance of certain aspects of the project. The Contractor shall at all times cooperate with and ensure that the Contractor and each of its partners, subcontractors and their employees and agents cooperate, in good faith, with such third parties and with any WHO in-house resources.

# List Of Annexes & APPENDICES

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| **Appendix 1** | Detailed Requirements Specifications |
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**Request for Proposals:** MVP/EMP/eEDL2019\_1

**Annex 1: Acknowledgement Form** (Ref. Paragraph 4.2)

|  |
| --- |
| **Please check the appropriate box (see below) and email this acknowledgement form immediately upon receipt to** [edlsecretariat@who.int](mailto:edlsecretariat@who.int)  Please include bid reference in your subject line: MVP/EMP/eEDL2019\_1 |
| **Intention To Submit A Proposal**  We hereby acknowledge receipt of the RFP. We have perused the document and advise that we intend to submit a proposal **on or before** 14/08/2019 **at 12:00 (noon) hours** , Geneva, Switzerland  **time**. |
| **Non-Intention To Submit A Proposal**  We hereby acknowledge receipt of the RFP. We have perused the document and advise that we do not intend to submit a proposal for the following reasons:  Insert reason here: |
| **Bidder's Contact Information is as follows**: |

|  |  |
| --- | --- |
| **Entity Name:** | ………………………………………………………………………………………………… |
| **Mailing Address:** | …………………………………………………………………………………………………  …………………………………………………………………………………………………  ………………………………………………………………………………………………… |
| **Name and Title of duly authorized representative:** | ………………………………………………………………………………………………… |
| **Signature:** |  |
| **Date:** | ………………………………………………………………………………………………… |

**Request for Proposals:** MVP/EMP/eEDL2019\_1

**Annex 2: Confidentiality Undertaking** (Ref. Paragraph 4.6)

1. The World Health Organization (WHO), acting through its Department of EMP, has access to certain information relating to Medical Devices List and Essential Diagnostics List which it considers to be proprietary to itself or to entities collaborating with it (hereinafter referred to as “the Information”).
2. WHO is willing to provide the Information to the Undersigned for the purpose of allowing the Undersigned to prepare a response to the Request for Proposals (RFP) for the Establishment of web-based business solution for the WHO Medical Devices List and List of Essential in vitro diagnostics Project ("the Purpose"), provided that the Undersigned undertakes to treat the Information as confidential and proprietary, to use the Information only for the aforesaid Purpose and to disclose it only to persons who have a need to know for the Purpose and are bound by like obligations of confidentiality and non-use as are contained in this Undertaking.
3. The Undersigned undertakes to regard the Information as confidential and proprietary to WHO or parties collaborating with WHO, and agrees to take all reasonable measures to ensure that the Information is not used, disclosed or copied, in whole or in part, other than as provided in paragraph 2 above, except that the Undersigned shall not be bound by any such obligations if the Undersigned is clearly able to demonstrate that the Information:
   1. was known to the Undersigned prior to any disclosure by WHO to the Undersigned; or
   2. was in the public domain at the time of disclosure by WHO; or
   3. becomes part of the public domain through no fault of the Undersigned; or
   4. becomes available to the Undersigned from a third party not in breach of any legal obligations of confidentiality to WHO.
4. At WHO's request, the Undersigned shall promptly return any and all copies of the Information to WHO.
5. The obligations of the Undersigned shall be of indefinite duration and shall not cease on termination of the above mentioned RFP process.
6. Any dispute relating to the interpretation or application of this Undertaking shall, unless amicably settled,   
   be subject to conciliation. In the event of failure of the latter, the dispute shall be settled by arbitration.   
   The arbitration shall be conducted in accordance with the modalities to be agreed upon by the parties or, in the absence of agreement, with the rules of arbitration of the International Chamber of Commerce. The parties shall accept the arbitral award as final.

|  |  |
| --- | --- |
| **Entity Name:** | ………………………………………………………………………………………………… |
| **Mailing Address:** | …………………………………………………………………………………………………  …………………………………………………………………………………………………  ………………………………………………………………………………………………… |
| **Name and Title of duly authorized representative:** | ………………………………………………………………………………………………… |
| **Signature:** |  |
| **Date:** | ………………………………………………………………………………………………… |

**Request for Proposals:** MVP/EMP/eEDL2019\_1

**Annex 3: Proposal Completeness Form** (Ref. Paragraphs 4.4 & 4.6)

|  |  |  |
| --- | --- | --- |
| **Section** | **Requirement** | **Completed in full (Yes/No)** |
| Annex 2 | Confidentiality undertaking form | Yes  No |
| Annex 3 | Proposal completeness form | Yes  No |
| Annex 5 | Acceptance form | Yes  No |
| Annex 6 | Self-Declaration Form | Yes  No |
| 4.12.2 to 0 | **Technical Proposal,** *including*  a)Executive Summary, proposed solution, approach/methodology, project plan and timelines;  b)Post-go-live service delivery model and support/maintenance proposal | Yes  No |
| 4.12.7 | **Financial Proposal** (for Techncal proposal including post go-live support/maintenance proposal) | Yes  No |

**The enclosed Proposal is valid for \_\_\_\_\_\_\_\_\_\_\_\_\_ days from the date of this form** (Ref. Paragraph 4.8)**.**

Agreed and accepted on **\_\_\_\_\_\_\_\_\_\_\_\_\_**

|  |  |
| --- | --- |
| **Entity Name:** | ………………………………………………………………………………………………… |
| **Mailing Address:** | …………………………………………………………………………………………………  …………………………………………………………………………………………………  ………………………………………………………………………………………………… |
| **Name and Title of duly authorized representative:** | ………………………………………………………………………………………………… |
| **Signature:** |  |
| **Date:** | ………………………………………………………………………………………………… |

**Request for Proposals:** MVP/EMP/eEDL2019\_1

**Annex 4: Questions from Bidders** (Ref. Paragraph 4.6)

|  |  |  |
| --- | --- | --- |
| **No.** | **RFP Section reference** | **Question** |
| 1 | Enter Text | Enter Text |
| 2 | Enter Text | Enter Text |
| 3 | Enter Text | Enter Text |
| 4 | Enter Text | Enter Text |
| 5 | Enter Text | Enter Text |
| 6 | Enter Text | Enter Text |
| 7 | Enter Text | Enter Text |
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| 18 | Enter Text | Enter Text |
| 19 | Enter Text | Enter Text |
| 20 | Enter Text | Enter Text |

**Request for Proposals:** MVP/EMP/eEDL2019\_1

**Annex 5: Acceptance Form** (Ref. Paragraph 4.6)

**The Undersigned, ……………………….., confirms to have read, understood and accepted the terms of the Request for Proposals (RFP) No**. MVP/EMP/eEDL2019\_1**, and its accompanying documents. If selected by WHO for the work, the Undersigned undertakes, on its own behalf and on behalf of its possible partners and contractors, to perform** MVP/EMP/eEDL2019\_1 **in accordance with the terms of this RFP** **and any corresponding contract between WHO and the Undersigned, for the following sums *(an Excel detailed/itemized proposal following the same structure can be attached to the below) - For more information please refer to Appendix 1***

|  |  |
| --- | --- |
| **Item** | **Cost**  *(indicate* ***CUR****rency)* |
| **One time costs** |  |
| **Deliverable 1: MEDEVIS for EDL** | |
| **Output 1.1: Convert EDL into web based eEDL** | |
| **Project Manager** costs | 0.00 |
| **Team members** costs *(please itemize by function)* | 0.00 |
| Other **technical costs** *(please itemize and specify whether there are one-time or recurring costs)*:  Operating System, database, application, license, etc. | 0.00 |
| **Other** Costs *(please itemize and specify whether there are one-time or recurring costs)* | 0.00 |
| **Proposed Output 1.1 Costs** | **0.00** |
| **Output 1.2: Develop structure for submission, review and publishing of EDL** | |
| **Project Manager** costs | 0.00 |
| **Team members** costs *(please itemize by function)* | 0.00 |
| Other **technical costs** *(please itemize and specify whether there are one-time or recurring costs)*:  Operating System, database, application, license, etc. | 0.00 |
| **Other** Costs *(please itemize and specify whether there are one-time or recurring costs)* | 0.00 |
| **Proposed Output 1.2 Costs** | **0.00** |
|  |  |
| **Deliverable 2: MEDEVIS** | |
| **Output 2.1: MEDEVIS- remaining lists implemented as web based eLists** | |
| **Project Manager** costs | 0.00 |
| **Team members** costs *(please itemize by function)* | 0.00 |
| Other **technical costs** *(please itemize and specify whether there are one-time or recurring costs)*:  Operating System, database, application, license, etc. | 0.00 |
| **Other** Costs *(please itemize and specify whether there are one-time or recurring costs)* | 0.00 |
| **Proposed Output 2.1 Costs** | **0.00** |
| **Output 2.2: MEDEVIS extending digital platform** |
| **Project Manager** costs | 0.00 |
| **Team members** costs *(please itemize by function)* | 0.00 |
| Other **technical costs** *(please itemize and specify whether there are one-time or recurring costs)*:  Operating System, database, application, license, etc. | 0.00 |
| **Other** Costs *(please itemize and specify whether there are one-time or recurring costs)* | 0.00 |
| **Proposed Output 2.1 Costs** | **0.00** |
| **Project Management** | |
| **Project Manager** costs for the management of the global project *(if any)* | 0.00 |
| Other **technical** costs *please itemize and specify whether there are one-time or recurring costs)*:  Operating System, database, application, license, etc. | 0.00 |
| Proposed **Travel** Costs | 0.00 |
| **Other** Costs *(please itemize and specify whether there are one-time or recurring costs)* | 0.00 |
| **Project management Costs** | **0.00** |
| **Other one time general costs (platform configuration, etc)** | **0.00** |
| **User and Admin Training** | **0.00** |
| **Recurring Costs** |  |
| **Support and maintenance** | **0.00** |
| **License fees** | **0.00** |
| **Other recurring costs (please detail)** | **0.00** |
| **TOTAL PROJECT COSTS** | **0.00** |

**The enclosed Proposal is valid for \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ days from the date of this form** (Ref. Paragraph 4.8)**.**

Agreed and accepted on **\_\_\_\_\_\_\_Date\_\_\_\_\_\_\_\_**

|  |  |
| --- | --- |
| **Entity Name:** | ………………………………………………………………………………………………… |
| **Mailing Address:** | …………………………………………………………………………………………………  …………………………………………………………………………………………………  ………………………………………………………………………………………………… |
| **Name and Title of duly authorized representative:** | ………………………………………………………………………………………………… |
| **Signature:** |  |

**Annex 6: Self Declaration Form**

**Applicable to private and public companies**

<**COMPANY**> (the “Company”) hereby declares to the World Health Organization (WHO) that:

1. it is not bankrupt or being wound up, having its affairs administered by the courts, has not entered into an arrangement with creditors, has not suspended business activities, is not the subject of proceedings concerning the foregoing matters, and is not in any analogous situation arising from a similar procedure provided for in national legislation or regulations;
2. it is solvent and in a position to continue doing business for the period stipulated in the contract after contract signature, if awarded a contract by WHO;
3. it or persons having powers of representation, decision making or control over the Company have not been convicted of an offence concerning their professional conduct by a final judgment;
4. it or persons having powers of representation, decision making or control over the Company have not been the subject of a final judgment or of a final administrative decision for fraud, corruption, involvement in a criminal organization, money laundering, terrorist-related offences, child labour, human trafficking or any other illegal activity;
5. it is in compliance with all its obligations relating to the payment of social security contributions and the payment of taxes in accordance with the national legislation or regulations of the country in which the Company is established;
6. it is not subject to an administrative penalty for misrepresenting any information required as a condition of participation in a procurement procedure or failing to supply such information;
7. it has declared to WHO any circumstances that could give rise to a conflict of interest or potential conflict of interest in relation to the current procurement action;
8. it has not granted and will not grant, has not sought and will not seek, has not attempted and will not attempt to obtain, and has not accepted and will not accept any direct or indirect benefit (finanical or otherwise) arising from a procurement contract or the award thereof;
9. it adheres to the UN Supplier Code of Conduct;
10. it has zero tolerance for sexual exploitation and abuse and has appropriate procedures in place to prevent and respond to sexual exploitation and abuse.

The Company understands that a false statement or failure to disclose any relevant information which may impact upon WHO's decision to award a contract may result in the disqualification of the Company from the bidding exercise and/or the withdrawal of any proposal of a contract with WHO. Furthermore, in case a contract has already been awarded, WHO shall be entitled to rescind the contract with immediate effect, in addition to any other remedies which WHO may have by contract or by law.

|  |  |
| --- | --- |
| **Entity Name:** | ………………………………………………………………………………………………… |
| **Mailing Address:** | …………………………………………………………………………………………………  …………………………………………………………………………………………………  ………………………………………………………………………………………………… |
| **Name and Title of duly authorized representative:** | ………………………………………………………………………………………………… |
| **Signature:** |  |
| **Date:** | ………………………………………………………………………………………………… |

1. https://treasury.un.org/operationalrates/default.php [↑](#footnote-ref-2)
2. For companies in existence less than two years, please provide the available audited financial statements. [↑](#footnote-ref-3)