

**RFP-NIAID-DMID-NIHAI2008026
Amendment #1**

**“Tuberculosis Clinical Diagnostics Research
Consortium (CDRC)”**

Amendment Issue Date: August 26, 2008

Proposal Due Date/Time: November 6, 2008, at 3:00 P.M., local time
[Unchanged]

Issued By: Karen M. Gamble
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Offerors must acknowledge receipt of this Amendment on each copy of the proposal submitted. Failure to receive your acknowledgment of this Amendment may result in the rejection of your proposal.

The hour and date specified for receipt of proposals HAS NOT been extended.

The following corrections/insertions have been included in the above-referenced solicitation:

- 1. Pg. 25: Article I.3., Additional Contract Clauses, paragraph a., has been revised to insert the following FAR Clause:**

19. FAR Clause 52.230-2, Cost Accounting Standards (April 1998).

Subsequent clauses under subparagraph a., are renumbered accordingly.

2. Pg. 31: The title of Amendment 4 is revised to change the word "Meetings" to "Agreements."

3. Attachment 5, Reporting Requirements and Deliverables, has been modified to add the following:

- Paragraph a.7) on page 3, Report on Select Agents or Toxins and/or Highly Pathogenic Agents.
- Paragraph b.1) on page 3., Source Code and Object Code (the previous paragraphs 1) and 2) are renumbered accordingly).
- Article F.2., paragraph b. table on pages 5-7.

a. Technical Progress Reports

7) Report on Select Agents or Toxins and/or Highly Pathogenic Agents

For work involving the possession, use, or transfer of a *Select Agent or Toxin* and/or a *Highly Pathogenic Agent*, the following information shall also be included in each Semiannual Progress Report:

- a) Any changes in the use of the *Select Agent or Toxin* and/or a *Highly Pathogenic Agent*, that have resulted in a change in the required biocontainment level, and any resultant change in location, if applicable, as determined by the IBC or equivalent body or institutional biosafety official.
- b) If work with a new or additional *Select Agent or Toxin* and/or a Highly Pathogenic Agent will be conducted in the upcoming reporting period, provide:
 1. A list of each new or additional *Select Agent or Toxin* and/or a *Highly Pathogenic Agent* that will be studied;
 2. A description of the work that will be done with each new or additional *Select Agent or Toxin* and/or a *Highly Pathogenic Agent*;
 3. The title and location for each biocontainment resource/facility, including the name of the organization that operates the facility, and the biocontainment level at which the work will be conducted, with documentation of approval by your IBC or equivalent body or institutional biosafety official. It must be noted if the work is being done in a new location.

If the IBC or equivalent body or institutional biosafety official has determined, for example, by conducting a risk assessment, that the work that has been performed or is planned to be performed under this contract may be conducted at a biocontainment safety level that is lower than BSL3, a statement to that effect shall be included in each Semiannual Progress Report.

If no work involving a *Select Agent or Toxin* and/or a *Highly Pathogenic Agent* has been performed or is planned to be performed under this contract, a statement to that effect shall be included in each Semiannual Progress Report.

b. Other Reports and Deliverables:

1) Source Code and Object Code

Unless otherwise specified herein, the Contractor shall deliver to the Government, upon the expiration date of the contract, all source code and object code developed, modified, and/or enhanced under this contract.

4. ARTICLE F – DELIVERIES, paragraph b. Other Reports and Deliverables (Delivery Schedule), is amended to add the following Item 25:

b. Other Reports and Deliverables (Delivery Schedule)

Item	Deliverables	SOW Reference	Recipient	Delivery Schedule
25.	Source Code and Object Code	Article C.2., paragraph b.1)	Project Officer	On/before the completion date of the contract.