



Vitamin A Laboratory - External Quality Assurance Program (VITAL-EQA)
Round 35 / Iron Indicators / CRP

Form Approved
OMB No. 0920-xxxx
Exp. Date xx/xx/20xx

INSTRUCTIONS

- E-mail this completed worksheet to: **vitaminalab@cdc.gov**
- Round deadline: **December 18, 2020**
- All cells highlighted **red** are required or have invalid data that needs to be revised. Click on cell to see data requirements.
- Run these samples in the same way as you would routine patient samples.
- Run in singlicate over a period of two consecutive days (Day 1 and Day 2).
- Record all results with three significant figures (i.e., 105, 10.5, 1.05, 0.105).
- Retain the vials in your ultra-cold freezer for at least two weeks after data submission in case there is a question about the ID code or results.

Laboratory ID

Received Date (MM/DD/YY) Res Stripper arrived cold No Stripper did not arrive cold

Assay Type

Instrument (Manufacturer/Model)
Kit Information (if used) (Name/Manufacturer)

Calibration Range Low High mg/L

LOD mg/L

Run	Assay Date	Sample ID	Result (mg/L)	< LOD	Note
1			1	<input type="checkbox"/>	
			1	<input type="checkbox"/>	
			1	<input type="checkbox"/>	
2			1	<input type="checkbox"/>	
			1	<input type="checkbox"/>	
			1	<input type="checkbox"/>	

Laboratory Notes

OFFICIAL USE ONLY