

GTFCC Laboratory Reporting Form for Cholera Suspected Case

The laboratory is to complete this form and send a copy to the relevant health authorities and requesting clinician.

The type of results reported in this form are based on the methods recommended by the GTFCC and that match the contents of the WHO cholera laboratory kits. Other tests and results may be reported.

For more information on testing for cholera refer to GTFCC Job Aids [Rapid Diagnostic Test \(RDT\) for cholera detection](#), [Isolation and Presumptive Identification of *Vibrio cholerae* O1/O139 from fecal specimens](#), [Antimicrobial Susceptibility Testing for Treatment and Control of Cholera](#).

Report made by

Name/Address of laboratory (or stamp)

Name of laboratory director/contact person:

Phone: E-mail:

Signature:

Patient and specimen information

Patient full name: **Patient ID:** **Sex:** Male Female

Age: ___ Years/___ Months/___ Days or date of birth DD MM YYYY ___/___/___

Date of onset of illness: DD MM YYYY ___/___/___ **Specimen ID:**

Date that sample was collected: DD MM YYYY ___/___/___

Date and time of receipt at laboratory: DD MM YYYY Hour Minute ___/___/___ ___:___

Specimen condition for testing: Adequate Not adequate, specify

Laboratory results

RDT
Performed in laboratory: <input type="checkbox"/> No <input type="checkbox"/> Yes, specify: <input type="checkbox"/> Enriched RDT / <input type="checkbox"/> Direct RDT
Name of kit used:
Date test performed: <small>DD MM YYYY</small> ___/___/___
Result: <input type="checkbox"/> Reactive O1 <input type="checkbox"/> Reactive O139 <input type="checkbox"/> Reactive O1 and O139 <input type="checkbox"/> Non-reactive <input type="checkbox"/> Invalid

Oxidase test
Performed: <input type="checkbox"/> Yes <input type="checkbox"/> No
Date test performed: <small>DD MM YYYY</small> ___/___/___
Result: <input type="checkbox"/> Positive <input type="checkbox"/> Negative

Culture
<input type="checkbox"/> on TCBS: Directly from sample: <input type="checkbox"/> Yes <input type="checkbox"/> No After enrichment in APW: <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> on Non Selective Agar (NSA): Directly from sample: <input type="checkbox"/> Yes <input type="checkbox"/> No After enrichment in APW: <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Others, specify:
Date test performed: <small>DD MM YYYY</small> ___/___/___
Results: <input type="checkbox"/> Growth on TCBS, specify color and aspect of colonies:
<input type="checkbox"/> Growth on NSA

Seroagglutination test

Performed: Yes No

DD MM YYYY

Date test performed: ____ / ____ / ____

Results:

Self-agglutination in saline: Yes No

Serogroup identification:

Positive O1 Positive O139 Negative

Serotype identification (for O1):

Positive Inaba Positive Ogawa

Polymerase Chain Reaction test

Performed: Yes No

Use of commercial kit: Yes No / Name:

or in-house assay used: Yes No

DD MM YYYY

Date test performed: ____ / ____ / ____

Species confirmation, *V. cholerae* Target:

Positive Negative Indeterminate

Serogroup O1 Target:

Positive Negative Indeterminate

Serogroup O139 Target:

Positive Negative Indeterminate

Toxin detection, Target *ctxA*:

Positive Negative Indeterminate

Others, target:

Positive Negative Indeterminate

Antimicrobial Susceptibility Testing

Performed: Yes No

Method: Agar Disk Diffusion Method
 Minimum Inhibitory Concentration test-strips
 Other, specify

DD MM YYYY

Date test performed: ____ / ____ / ____

Erythromycin (EM): Not tested Susceptible
 Intermediate Resistant

Pefloxacin (PEF): Not tested Susceptible
 Intermediate Resistant

Tetracycline (TE): Not tested Susceptible
 Intermediate Resistant

Doxycycline (DOX): Not tested Susceptible
 Intermediate Resistant

Azithromycin (AZ): Not tested Susceptible
 Intermediate Resistant

Ciprofloxacin (CIP): Not tested Susceptible
 Intermediate Resistant

Others:

Susceptible Intermediate Resistant

Other tests performed (e.g. lysis by phages, string test etc)

Specify:

DD MM YYYY

Date test performed: ____ / ____ / ____

Results:

Summary of results

Vibrio cholerae: Yes No Serogroup: O1 O139 Toxicogenicity: Positive Negative

Serotype (if known): Biotype (if known):

Alternative diagnosis or coinfections:

Pathogen identified	Diagnostic method used (eg. Culture, PCR...)

Additional comments (including information regarding susceptibility/resistance to antimicrobials, or awaiting further results):

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DD MM YYYY

Date laboratory results are sent back to referring health facility: ____ / ____ / ____

DD MM YYYY

Date laboratory results are sent to health authorities: ____ / ____ / ____