

Global Clinical Data Platform CHOLERA CASE REPORT FORM (CRF) MODULE 2

INTRODUCTION

The CRF is designed to collect data obtained direct from patient examination and interview, and from review of hospital or clinical notes of people with suspected, probable or confirmed cholera.

The CRF captures data from patients being managed as inpatients in cholera treatment centre (CTC), cholera treatment unit (CTU) or dedicated inpatient ward. Data may be collected prospectively or retrospectively. The data collection period is defined as the period from hospital admission, or first visit, to discharge from care, transfer or death.

This CRF has three modules:

Module 1: To be completed on the first day of presentation or admission to the CTC,

CTU or ward.

Module 2: Daily form: to be completed daily on inpatient days.

Module 3: To be completed at hospital discharge, transfer or death.

GENERAL GUIDANCE

Participant identification numbers consist of a site code and a participant number.

Please e-mail the data management team at <u>globalclinicaldataplatform@who.int</u> and they will provide instructions for data entry and will assign you a 5-digit site code at that time.



IV Potassium

| | PARTICIPANT ID I | - 11 | - 11 | - 11 | 11 | l l | - 11 | - 11 | 11 | - 1 |
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| I. CASE IDENTIFICATION and TIM | E OF INTER | M REVIEW | | | | | | |
|--|------------------------------------|--|---------------------------|---------------------|----------------------|---------------------|--|--|
| ID number: [|] | Site/facility nan | ne: [| | |] | | |
| Date of admission: (dd/mm/yyyy) [][] /[][] / 20 [][] | | Date of this review: (dd/mm/yyyy) [][] /[] / 20 [][] | | | | | | |
| II. VITAL SIGNS AT INTERIM REVI | | | | | | | | |
| Temperature (°C): [][] . [] | Heart rat | e (bpm) | | | / rate (/min) | • • • | | |
| BP (mmHg): [][] (systolic) | O ₂ satura | ation room air (% | 6): [][][_ | _] Level of co | nsciousness | s: A / V / P / U | | |
| [][] (diastolic) | on □Rod on □CP | | Oxygen ther Ventilator | rapy Pain score | Pain score [][] /10 | | | |
| Capillary refill ≥ 3 sec? ☐ Yes ☐ No | Weight (I | kg): [][][_ |].[_] | Oedema | | □ Yes □ No | | |
| Absent or weak pulse ☐ Yes ☐ No | | 0, 1 1 1 | | Location: | | + _++ _+++ | | |
| Sunken eyes ☐ Yes ☐ N | lo Drinking | ☐ Able ☐ N | lot able | Mobility: | | | | |
| Slow skin pinch (> 2 secs) ☐ Yes ☐ N | | ☐ Able ☐ N | | ☐ Indepen | dent | | | |
| Passed urine in past 12 hours ☐ Yes ☐ No | | □PO □N | G | ☐ Walks w | ith help | | | |
| | ☐ Breas | stfeeding | | ☐ Unable | ☐ Unable to mobilize | | | |
| Episodes of vomiting in last 24 hours: [][Episodes of diarrhoea in last 24 hours: [][|] Clinical] □ None, □ Mode □ Sever | rate | sessment | | | | | |
| III. CLINICAL DETAILS AT INTERIM REVIEW | | | | | | | | |
| Symptoms (at time of review) | | | | | | | | |
| Confusion/irritability ☐ Yes ☐ No | | | orexia | ☐ Yes | □ No | ☐ Unknown | | |
| Shortness of breath ☐ Yes ☐ No | | | usea | ☐ Yes | □ No | ☐ Unknown | | |
| Weakness ☐ Yes ☐ No | | | rrhoea | □ Yes | □ No | ☐ Unknown | | |
| Myalgia ☐ Yes ☐ No | | miting | □ Yes | □ No | ☐ Unknown | | | |
| Abdominal pain ☐ Yes ☐ No | | any | □ Yes | □ No | □ Unknown | | | |
| | _ | cramps | □ Yes | □ No | □ Unknown | | | |
| | | Thi | rst | ☐ Yes | □ No | ☐ Unknown | | |
| IV. DAILY FLUID BALANCE (time | of daily total | s: | _hrs) | | | | | |
| DAILY INPUT | | | DAILY OU | JTPUT | | | | |
| Total oral intake (PO, NG) in 24 hrs | n | nL/24 hours | Total vom | itus in 24 hrs | |][_][_] mL/24 hours | | |
| Total IV intake in 24 hrs | | mL/24 hours | Total urine | e output in 24 hrs | |][] mL/24 hours | | |
| | | | | l output in 24 hrs1 | |][] mL/24 hours | | |
| TOTAL IN | | mL/24 hours | TOTAL O | UT | [_][|][] mL/24 hours | | |
| V. CURRENT TREATMENT | | | | | | | | |
| ORS □ Yes | □ No | | | Zinc | ☐ Yes | □ No | | |
| F75 □ Yes □ No | | | | Doxycycline | ☐ Yes | □ No | | |
| F100/Ready to use therapeutic Food | ☐ Yes | □ No | | Other antibiotic | ☐ Yes | □ No | | |
| IV fluids (tick all that apply) ☐ Ringer's I☐ 5% Dextrose | % w/v (Normal) s | saline | Furosemide | ☐ Yes | □ No | | | |
| IV Glucose (50%) ☐ Yes | □ No | | | | | | | |

□ No

☐ Yes

¹ If stool and urine are measured together, e.g. bucket, then enter the total volume as "stool".



| PARTICIPANT ID I II II II II II II II II | ıl |
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|--|----|

VI. LABORATORY TEST RESULTS

| Test | Collection date (dd/mm/yyyy) | Result |
|---------------------------------|------------------------------|--|
| Haemoglobin (Hb) | [_][_] /[_][_] / 20 [_][_] | [_][_] . [_] g/dL |
| Haematocrit | [_][_] /[_][_] / 20 [_][_] | [_][_].[_] % |
| Platelets | [_][_] /[_][_] / 20 [_][_] | [_][_][_] x10 ⁹ /L |
| Sodium | [_][_] /[_][_] / 20 [_][_] | [][] . [] mmol/L |
| Potassium (K+) | [_][_] /[_][_] / 20 [_][_] | [][] . [] mmol/L |
| Urea | [_][_] /[_][_] / 20 [_][_] | [_][_] . [_] mmol/L OR [][_][_] mg/dL |
| Creatinine | [_][_] /[_][_] / 20 [_][_] | [_][_] . [_] mg/dL OR [_][_][_] µmol/L |
| Chloride | [_][_] /[_][_] / 20 [_][_] | [][] . [] mmol/L |
| Bicarbonate (HCO ₃) | [_][_] /[_][_] / 20 [_][_] | [][] . [] mmol/L |
| Lactate | [_][_] /[_][_] / 20 [_][_] | [][] . [] mmol/L |
| Magnesium | [_][_] /[_][_] / 20 [_][_] | [_][_] . [_] mmol/L |