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ETHICAL REVIEW COMMITTEE, ICODR, B.

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Date OCTOBER 03, 1989

incipal Investigator Dr. D. Mahalanabis Trai	NOVEMBER 19, 1989
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inic-based cohort# study. ()	Continuation with change
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Source of Population: 5.	ollowing (If Not Applicable waite NA)
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(a) Nature and purposes of	PAROL to review, the following information
	should be included in the abstract cummany.
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SECTION I

ICDDR,B LIBRARY DHAKA 1212

Title

Prognostic and risk factors for prolongation of acute diarrhoea: a clinic-based cohort study

P. Investigator

Dilip Mahalanabis

Co-PI's:

Md. Shahadat Hossain,CSD Abu Faruque, CSD Saul Tzipori, LSD Hamidur Rahman, LSD Mr. Mujibur Rahaman, LSD

Budget

US\$ 114,000

Starting Date:

December 1989

Ending date:

June, 1991

Signature of Division Head:

Stabilli

Dr. D. Mahalanabis

Clinical Sciences Division

Date: October 03, 1989

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Abstract Summary:

- 1. Prognostic factors that reliably identify a child in whom an acute diarrhocal episode is likely to become persistent will be sought and used as an indicator of the need for early, appropriate treatment.
- 2. Risk factors that influence the duration of a diarrhoeal episode or determine severe nutritional impact from persistent diarrhoea will be determined; risk factors that can be modified by specific community or facility-based interventions will be of particular interest.

Review Committees:

Research	h Review Commit	tee:	· · · · · · · · · · · · · · · · · · ·	
	Review Committe	7		
Director	· · · · · · · · · · · · · · · · · · ·	••••••	• • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •

OBJECTIVES AND RATIONALE

State clearly the objectives of the project and discuss briefly their revelance to the control of diarrhocal discusses. Summarize, briefly, the present statum of acjentific knowledge relevant to the proposed project (important references should be cited). Indicate why the project is feasible within the estimated time and with the resources requested.

<u>Objective</u>

1. Prognostic factors that reliably identify a child in whom an acute diarrhoeal episode is likely to become persistent will be sought and used as an indicator of the need for early, appropriate treatment. 2. Risk factors that influence the duration of a diarrhoeal episode or determine severe nutritional impact from persistent diarrhoea will be determined; risk factors that can be modified by specific community or facility based interventions will be of particular

Possible risk/prognostic factors for persistent diarrhoea

- a) Hostfactors:
 - young age, especialy <12 months
 - malnutrition
 - impaired cell-mediated immunity
- b) Previous Infections:
 - recent acute diarrhoea
 - previous persistent diarrhoea
 - recent viral illness, particularly measles
- c) Pre-illness feeding:
 - Recent introduction of animal milk
- d) Microbial isolates during acute phase:
 - enteroadherent E.coli.
 - Shigella
 - more than one pathogen
- e) Treatment/diet used during acute phase:
 - antiparasitic drugs, antiperistaltic drugs
 - lack of breastfeeding, full strength cow-milk based formula feeding
 - type of ORS used (e.g. rice-based ORS vs glucose/sucrose-based ORS) /

Risk/Prognostic factors of major interest

a) Type of ORS: It was shown that rice-based ORS reduces diarrhoeal stool and duration of acute diarrhoea compared to glucose-based ORS. A field study

(USE ADDITIONAL PAGES IF NECESSARY - MAXIMUM 4 PAGES)

(Bari et al)showed that children treated with rice-ORS had shorter average duration of diarrhoea, smaller number of children with persistent diarrhoea (as defined by a diarrhoea duration of >14 days) and better weight gain compared with the group treated with glucose ORS. However, serious methodological problems with the study design made the conclusions tentative at best. This factor will be evaluated by direct comparison of the two groups for duration of diarrhoea, weight gain and mid-arm circumference. If rice ORS can be shown to reduce the risk of developing persistent diarrhoea, the finding will have important implications for its prevention.

- b) Cell-mediated immunity: An earlier study at ICDDR, B and a recent ongoing study in Lima indicate that the risk of developing persistent diarrhoea can be predicted from the capacity of children to produce normal, delayed-type hypersensitivity reactions to standard skin test antigens. Children with impaired skin test responses were more likely to develop persistent diarrhoea on followup than were children with normal responses. This effect persisted after controlling for both age and initial nutritional status. Impaired skin test reactivity in these children were often transient. Its relation with factors like micronutrient status (zinc, iron, folate, B₁₂) and anaemia, previous illness (e.g. acute respiratory infection, measles), nutritional status, type of pathogens associated with acute illness, etc., is of interest.
- c) Recent acute diarrhoca and previous persistent diarrhoca: In Guatemala and India the risk of developing persistent diarrhoca increased two- to four-fold during the two months following an episode of acute diarrhoca. Whether this association reflects the damage inflicted on the gut mucosa during the earlier episode or other alteration in host defenses that in some way predispose to persistent diarrhoca is not clear; nor is it clear if the risk is related to the specific etiology of the preceding acute episode. Two studies in Guatemala have shown that children suffering from one documented episode of persistent diarrhoca have a three- to six-fold increased risk of developing at least one additional episode during the same year. Furthermore, in northeastern Brazil nearly half of all diarrhocal days documented during the 30 months of active surveillance were experienced by children who suffered at least one episode of persistent diarrhoca.
 - d) Etiological agents: The association of EPEC and EAEC with persistent diarrhoea is of particular interest and requires further investigation. The entero-adherent strains, some of which are also of EPEC scrotypes, are characterised by their capacity to adhere to the intestinal mucosal brush border and to cells in tissue culture. At least 3 patterns of adhesion are recognised: localised adhesion (LA), diffuse adhesion (DA), and autoaggressive adhesion (AA). LA E.coli have been definitely associated with persistent diarrhoea. In India, for example, $\Lambda\Lambda$ E.coli were isolated from 2% of healthy controls, 9% of children with acute diarrhoea and 26% of children with persistent diarrhoea under 3 years of age.
 - e) Pre-illness feeding practices: The possibility that pre-illness feeding

practices, especially breast-feeding may affect the risk of developing persistent diarrrhoea has been studied in India and Peru. According to preliminary results of these studies, there is no evidence that the risk of developing persistent diarrhoea is related to preillness feeding practices, i.e. exclusive breast-feeding, supplemented breast-feeding, and use of breast-milk substitutes. These results are surprising given the marked protective effect of breast-feeding on the incidence and severity of acute diarrhoea, especially in infants, and further acudy of this topic is needed. The study in India has shown, however, that the incidence of persistent diarrhoea does increase two- to three-fold during the frist month following the introduction of animal milk. could be several reasons for this (singly or in combination) such as, reduced intake of protective factors in breast-milk, contamination of animal milk proteins and often related intolerance to lactose. Λ recent study indicated that use of unmodified cow's milk in infants may be associated with prolongation of acute diarrhoea.

f) Etiological agents: Most of the bacteria and parasites that are known to cause acute diarrhoea have also been isolated from patients with persistent diarrhoea. Whether certain pathogens are especially able to cause persistent diarrhoea has however not been extensively studied. Those enteropathogens that are isolated with about equal frequency from episodes of acute and persistent diarrhoea include Shigella, non-typhoid Salmonella, entereotoxigenic E.coli, C. jejuni, A. hydrophila and less ferquently G. Lamblia, Y. Enterocolitica, C. difficile, E. histolytica. Those that are isolated with greater frequency from episodes of persistent diarrhoea include enteroadherent E.coli (EAEC), enteropathogenic E.coli and cryptosporidium. Cryptosporidium and possibly Shigella may be particularly important pathogens associated with persistent diarrhoea in children with pre-existing malnutrition. The mechanisms by which these agents cause persistent diarrhoea probably related to their capacity to adhere to or invade the bowel mucosa.

Background

Intestinal Microflora and Persistent Diarrhoea

Most enteropathogens that acuse acute diarrhoea have also been associated with persistent diarrhoea, notable exceptions being vibrios and viruses, especially rotavirus. Those organisms that are isolated with about equal frequency from episodes of acute and persistent diarrhoea include non-typhoid Salmonella, enterotoxigenic E.Coli, Campylobacter Jejuni, and aeromonas hydrophila. Their continued presence in episodes of persistent diarrhoea probably reflects an impaired ability of the host to terminate infection. Organisms isolated with greater frequency for episodes of persistent diarrhoea include Shigella, enteroadherent E.Coli (EAEC) enteropathogenic E.Coli

and Cryptosporidium (Black, Penny - Unpublished; Bhan et al - Unpublished).

Epidemiological studies at ICDDR,B suggest (Black et al) that 7% of episodes of acute diarrhoea associated with enterotoxigenic E.Coli persisted for longer than 3 weeks compared with 3% of episodes of rotavirus diarrhoea.

The association of EPEC and EAEC with persistent diarrhoca is of particular interest and requires further investigation. Enteroadherent strains, nome of which are also of EPEC serotypes are characterised by their capacity to adhere to cells in tissue culture (e.g. HEP-2 cells). E.Coli with localised and autoaggregative types of adhesion are associated with persistent diarrhoca. The latter type which usually does not belong to EPEC serotypes may play an important role in persistent diarrhoea. In India, for example, they were isolated from 2% of healthy controls, 9% of children with acute diarrhoea and 26% of children (under age 3 years) with persistent diarrhoea (Bhan et al.)—Unpublished).

In most studies, less than half the children with persistent diarrhoea have recognised enteric pathogens in their faeces. However, a few studies such as the one in Peru (Penny et al) have shown that patients with persistent diarrhoea have increased number of aerobic and anaerobic fecal bacteria in the small bowel in comparison to findings in healthy controls from developed countries in whom the upper small bowel only contains very small numbers of respiratory-type commensal bacteria. However, studies in children with acute diarrhoea and in locally recruited healthy controls in Lima showed similar results as those in persistent diarrhoea patients, raising doubts as to their significance in relation to the pathogenesis of persistent diarrhoea.

Background

Diarrhoeagenic E.coli and persistent diarrhoea.

Bray and Beaban suggested in 1945 that certain members of E.coli were associated with infectious diarrhoea. However, advances in our understanding of basic mechanisms, made only in the last decade, have finally established this charismatic group of micro organisms as a cause of severeal diarrhoeal syndromes. Enterotoxigenic, enteropathogenic and enteroadherent E.coli are associated with acute watery diarrhoea and persistent diarrhoea, enteroinvasive E.coli with dysentery and enterohaemorrhagic E.coli with haemorrhagic colitis.

Enterotoxigenic E.coli in persistent diarrhoea

Epidemiological studies (Black et al) have shown a link between episodes of enterotoxigenic <u>E.coli</u> (and <u>Shigella</u>) associated diarrhoea and subsequent poor growth in children. It was also shown that 7% of episodes of diarrhoea associated with enterotoxigenic E.coli persistent no longer that 3 weeks (16% of episodes of Shigellosis) compared with 3% of episodes

Enteropathogenic E.coli

The term enteropathogenic E.coli was coined in the 1950's to refer to E.coli of O serogroups including 026, 055, 086, 0111, 0119, 0125 and 0128 that were isolated more frequently from children with diarrhoca compared with controls. They do not produce CFA's, LT or ST neither do they show Shigella like invasiveness of enteroinvasive E.coli. Examination of electron micrographs of jejunal and colonic biopsies from infants with persistent diarrhoea associated with enteropathogenic E.coli gave insights into mechanisms of pathogenecity. At sites of mucosal attachment of enteropathogenic E.coli the brush border was destroyed (effaced) and the E.coli was closely attached to cup like projections (pedestals) of bare plasma membrane. The bacterial determinants of this pathognomic process have been studied recently using HeP2 and HeLA cells in tissue culture. Typically enteropathogenic E.coli induce acute self limiting watery distributes associated with fever and vomiting in infants. In some the illness may

Enteroadherent E.coli

The HeP2 adhesion assay has been exploited to identify strains of E.coli which do not produce LT or ST and whose serotypes may differ from enteropathogenic E.col: Such strains were first identified in travellers diarrhoea. Unpublished cpidemiological data (MK Bhan et al) as stated earlier suggests a role for enteroadherent E.coli in acute and specially persistent diarrhoea. A cross sectional age cohort of 580 children below 3 years of age residing in a village in North India were seen at weekly intervals for 12 months. Faecal samples were available for 346 of 565 diarrhoeal episodes. Seventy of which were persistent. Enteroadherent E.coli were isolated from 2.4% of controls, 9.2% of patients with acute diarrhoea and 26% of children with persistent diarrhoea. The majority had watery or loose stools but 11.6% had bloody stools,

raising the possibility of relationships between some strains of enteroadherent and enterohaemorrhagic E.coli.

EXPERIMENTAL DESIGN AND METHODOLOGY

Include all relevant details on experimental design, methodology and statistical methods as well as approximate schedule for each part of the proposed plan of work. This plan and schedule should indicate clearly the logical progression of the work towards the objectives of the project. Please refer to the guidelines.

Study design

- the study will be conducted at the Clinical Research Centre, Dhaka.
- a random sample (cohort 1) of children aged 3 to 35 months attending the treatment centre with acute watery diarrhoea of 5 days or less will be included; the sampling period will cover 12 consecutive months. A second random sample (cohort 2) of children aged 3 months to 35 months with acut. bloody diarrhoea of 5 days or less will also be included in the study.
- patients in cohort 1 and 2 will be assigned randomly to either rice ORS or glucose ORS groups. The group in glucose ORS will be treated with glucose ORS; the rest of the treatment will not be interfered with. Rice ORS is routinely used at the Clinical Research Centre and the group assigned to rice ORS group will receive the usual treatment;
- mothers will be interviewed to fill up a pretested questionnaire; admissiphysical findings will be recorded; stool/RS will be cultured for pathogen. (E.coli and Klebsiella strains will be saved); stool microscopy will be carried out for cryptosporidium, G. lamblia, and E. histolytica; duration of hospital stay and condition at discharge recorded.
- A followup examination will be made (and a questionnaire administered) after 7 and 15 days of onset of illness (parents will be compensated for travel cost and wage loss).
- blood test for total leucocytes and lymphocyte countron, admission.
- followup at 1 month from the last examination to record if a second attac of diarrhoea occurred and relate to preceding episode and its ethology.
- patients will receive a skin test with multiple antigens (Multitest IMC -Institut Merieux); those negative for tuberculin will receive a BCG vaccination on discharge from hospital and a tuberculin test after 6 weeks of receiving BCG.
- E.coli/klebsiella strains will be tested for adhesion to tissue culture cells and pathogenecity in 'ritard model' of rabbits.
- random sample of the patients in target age group for 12 consecutive months will be enrolled;

Sample size

Diarrhoea ceased by day 7 (Bari et al): For nondysenteric diarrhoea glucose ORS group rice oRS group

80% 95%

Sample size to detect this difference is

(USE ADDITIONAL PAGES IF NECESSARY - MAXIMUM 6 PAGES)

$$2 \times \frac{80 \times 20 + 90 \times 5}{15^2} \times 10.5 = 194 \text{ say } 200$$

with 10% drop out from followup the sample size will be 220.

- 2. Cellmediated Immunity (CMI)
- a) let us assume that 70% of acute diarrhoea patients have a positive response to skin test with multiple antigens and in patients with prolonged diarrhoea only 30% are positive. To detect this difference the sample size (90% power and (= .05).

$$2 \times \frac{70 \times 30 + 30 \times 70}{40^2} \times 10.5 = 28$$

+ 10% dropout = 31

Assuming 10% of patients will have prolonged diarrhoea we need $280\,$ patients in the cohort.

- b) we assume 90% of the patients will be negative for tuberculin on 'Multitest'. After BCG innoculation we expect a conversion rate of 90%. If we assume that the patients with prolonged diarrhoea are relatively anergic and that 50% only become positive for tuberculin after BCG then the sample size should be
- $2 \times \frac{90 \times 10 + 50 \times 50}{40^2} \times 10.5 = 34, \text{ say } 40.$

add 10% for tuberculin positive = 44 add 20% for drop outs = 54 patients only.

we assume 10% of the patients will have prolonged diarrhoea. Therefore, the size of the cohort will be 540.

3. Incidence of adherent E.coli (Bhan's data)

Let us assume 20% of acute nondysenteric diarrhoea become persistent. From Bhan's study 2% of nondiarrhoea controls, 9% of acute diarrhoea and 26% of persistent diarrhoea in children under 3 years had enteroadherent E.coli in stool culture. Sample size required to detect this degree of difference between acute and persistent diarrhoea.

No. of persistent diarrhoea patients required:

= $\frac{26 \times (100-26) + 9 \times (100-9)}{(26-9)^2} \times 10.5 = 95$, say 100.

We assume 12% of patients to have prolonged diarrhoea; therefore the size of the cohort should be 835±10% for drop outs = 920

Data Analysis

- 1. Type of ORS and diarrhoea duration: Group (cohort 1) with acute watery diarrhoea will be used for this anlaysis. Data will initially be analysed by direct comparison of the two groups using diarrhoea duration as a continuous variable (t-test, equivalent nonparametric test and multiple regression). The groups will be evaluated for comparability. Finally the groups will be compared for the incidence of persistent diarrhoea (diarrhoea persisting beyond 14 days) as a dichotomous variable (X test and logistic regression).
- 2. Impaired cell mediated immunity: two types of outcome variables will be evaluated, summary response to 7 antigens for skin test (multitest CMI Institute Merieux, France) and response to BCG (conversion tested by tuberculin test after 6 weeks). Summary skin test response will be compared with diarrhoea duration by appropriate regression analysis. Response to BCG (dichotomous variable) will be evaluated by multiple logistic regression.
- 3. EAEC: Its presence and type (of adhesion) will be analysed against diarrhoea duration. Multivariate analysis will be used to control for confounders (e.g. age, nutritional status, breastfeeding, immune status etc.).
- 4. Recent acute diarrhoea and previous persistent diarrhoea: These will be fitted into a multiple regression model using diarrhoea duration as a dependent variable. Occurrence of persistent diarrhoea will be analysed for the presence or absence of a recent attack of acute diarrhoea and a past attack of persistent diarrhoea by logistic regression.

Procedures

- 1. A systematic sampling technique will be used. Patients will be selected from a register of children aged 3 to 35 months entering the hospital (at triage area) with diarrhoea (or dysentery) of 5 days or less.
- 2. Two cohorts will be selected; a systematic sample of acute watery diarrhoea will form cohort 1 and a systematic sample of dysentery syndrome will form cohort 2.
- 3. A questionnaire form and a physical examination form will be filled. A skin test will be administered and planned laboratory tests (Stool/rectal swab on admission, blood count Nb, TC,DC after adequate hydration, skin test with multitest on admission) will be performed. Skin test will be read after 48 hours and tuberculin test will be repeated after 6 weeks.

- 4. Patients in cohort 1 and 2 will be randomised into either rice ORS group or into the glucose ORS group. The rice ORS group will receive routine treatment in the hospital and on discharge will be provided with rice ORS (precooked) packets for home use. Other patients will receive standard glucose ORS in place of rice ORS and the rest of the treatment will be the same as for other patients in the hospital; these patients on discharge will be supplied with glucose ORS packets for home use. Home treatment advice will be standardised and will be the same for both groups.
- 5. Patients will be asked to return on day 7 and day 15 of illness. If they do not return, they will be visited at home and a questionnaire and physical examination form will be filled in.
- 6. Patients will be asked to return on day 45 of illness for repeat questionnaire and physicial examination.

Test for Cell Mediated Immunity (CMI) (Skin test for delayed type hypersensitivity)

A skin test for evaluation of cell-mediated immunity developed by Institut Merieux will be used (Multitest CMI) $^{\rm R}$. This test is carried out by a multiple head applicator loaded with several antigens. The disposable unit has 8 heads and 9 times on each head linked by the support and loaded with 7 different antigens and a glycerine control. The following antigens are used - tetanus, diphtheria, streptococcus (group C), tuberculin, candida (ablicans), trichophyton (mentagrophytos) and proteus (mirabilis). In addition one head contains glycerine control. Each multitest CMI head numbered from 1 to 8 carries a drop of antigen in a glycerine solution at 70% weight for volume. The test will be applied on the forearms of the infants under study; because there is insufficient space for all the 8 heads to be applied on the same arm in small infants four heads in a row will be applied on one arm and the other four heads in a row will be applied on the other arm. Reading will be carried out after 48 hours. Positivity is defined only by induration. If a reaction is oval then two diameters are averaged. Diameter of inducation is measured on each antigen site. A reaction is positive if the average diameter expressed in milimeter is equal to or greater than 2 milimeter. of less than 2 mm is considered to be 0 and therefore negative. score will be used i.e. the score is the sum of average diameters of the positive reactions. A compound score is the sum of average diameters and the number of antigens with a positove response. Tuberculin test will be carried out by a skin test by multipuncture for the detection of, tul-aculin Monotest^R produced by Institut Merieux will be used for this purpose. In this test multipuncture applicator is loaded with 300,000 international unit per ml of PPD (purified protein derivative). Readings of reaction will be read after 48 hours to make it comparable with the multitest.

Microbiology (stool samples)

For routine isolation of pathogens, methods described in "Manual for laboratory investigations of acute enteric infections" WHO CDD/83.3 will generally be followed. Rotavirus antigen will be detected by ELISA.

Four strains of each E.coli and a single strain of Klebsjella isolated will be stored on both Dorsett egg media and frozen in trypticase say broth at -70° C to attempt maximum preservation of plasmids and virulence factors.

E.coli: all of the four strains isolated will be tested for serogroup by commercial grouping sera. Hemolytic ability will be noted after growth on blood agar.

Secretory toxins LT and ST will be identified by Y1 adrenal cell assay and infant mouse assay. Adherent <u>E.coli</u> will be identified by examination in tissue culture (HeLa and HeP2 cells) adherence assay, looking for localised, diffuse or aggregative adherence. The presence of cytotoxin will be noted; a cell-free supernatant of broth culture will be inoculated into HeLa cell monolayers and toxic effects determined after 48 hours.

Any <u>Klebsiella</u> isolated from small bowel will be examined in tissue culture for adherence. The effects of cell free supernatants will also be examined on tissue culture (CHO or Yl adrenal) cells for evidence of secretory toxins.

RESEARCH TRAINING OPPORTUNITIES OFFERED BY THE PROJECT

Describe how performance of this project will provide training for persons who work on the project. Be specific with regard to the persons who will acquire new or improved skills, their position in the institution, their role in the project, the skills which will be acquired or improved, and how this will enhance institutional capability for future research on diarrhoeal diseases. In addition, specify whether it will be necessary to train some workers in special areas prior to commencement of the project.

Two investigators (Drs. Md. Shahdat Hossain and Abu Faruque) will receive training in study design, conduct, analysis and reporting.

(USE ADDITIONAL PAGES IF NECESSARY)

PROGNOSTIC AND RISK FACTORS FOR PROLONGATION OF ACUTE DIARRHOEA: A CLINIC-BASED COHORT STUDY

Patient's name	•
Father's name	:
Detailed address and location	:
	<u></u>
	·
Name of the interviewer	•
Date	•
Time	:

MEDICAL HISTORY FORM

PART A

Variable name	Column	Code	
1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2	, 1	·/	
2.Randomization	2	/	
WHO-ORS=1, Rice-ORS=2			
3.Study No. 4.Date of attendance	3-6 7-12	//	// · '///
5. Time of attendance (h/min)	13-14	//_	//
6. How old is your child (months)?	15-16	//	
7. What is the sex of your child? Male=1, Female=2	1.7	/	·
8. What was the date you have observed that your child having diarrhoea?	18	//_	//
9. For how many hours the child has been suffering from diarrhoea?	19-23	//_	//
10. For how many hours the child has been suffering from vomiting? (Not applicable=99999)	24-28	//_	//
11. Has your child had any sickness in last one month (before onset of present diarrhea)?			
None [Yes=1, No=2]	29	/	
Cough [Yes=1, No=2]	30	/	
Fever [Yes=1, No=2]	31	/	•

Cough & fever	32	/
[Yes=1, No=2] Fever, sneezing and running	33	· /
[Yes=1, No=2] Frequent cough, fever and respiration	34	/
[Yes=1, No=2] Mumps [Yes=1, No=2]	35	/
Measles	36	/
[Yes=1, No=2] Ear discharge [Yes=1, No=2]	37	
Scables ([Yes=1, No=2]	38	/
Conjunctivitis [Yes=1, No=2]	39	/
Boils/skin infection [Yes=1, No=2]	40	/
Angular stomatitis [Yes=1, No=2]	41	/
Other [Yes=1, No=2]	42	/
12.Had your child had any diarrhoea in last one month (other than the present one)? [Yes=1, No=2]	43	/
13.If yes, what was the nature of the stool at the onset? Watery/loose=1, Bloody or bloody mucoid=2, Mucoid=3 Not applicable=9		/
14.Did the child receive any outside treatment at home for the new episode of diarrhoea ? [Yes=1, No=2]	45	/
15.If yes:		
antibacterial drugs [Yes=1, No=2 not applicable=9]	46	/
name of the drugs		
anthelmentic drugs [Yes=1, No=2 not applicable=9]	47	/
name of the drugs	•	

antiprtozoal drugs [Yes=1, No=2 not applicable=9]	48	/
name of the drugs		
antiperistaltic drugs [Yes=1, No=2 not applicable=9]	49	/
name of the drugs		
other drugs [Yes=1, No=2 not applicable=9]	50	/
name of the drugs		
16.Did the child get any .ORS for the present episode of diarrhoea ? [Yes=1, No=2]	51	/
17.If yes, what were the ingredients? Prepackaged=1 Home-made=2	. 52	/

PHYSICAL FINDINGS

PHYSIC	AL FINDINGS	
PART B		• .
18.Dehydration none=1, mild=2, moderate3, severe=4	53	/
19. Signs of vitamin A deficiency		
Normal [Yes=1, No=2]	54	/
Night blindness (XN) [Yes=1, No=2]	55	/
Conjunctival xerosis (X1A) [Yes=1, No=2]	56	/
Bitot's spot (X1B) [Yes=1, NO=2]	57	/
Corneal ulcer <1/3 (X3A) [Yes=1, NO=2]	. 58	/
Corneal ulcer >1/3 (X3B) [Yes=1, No=2]	59	/
Xerophthalmia scar (XS) [Yes=1, No=2]	60	/
Xerophthalmia fundus (XF) [Yes=1, No=2]	61	
20.Otitis media [Yes=1, No=2]	62	/
21.Throat normal	63	/
[Yes=1, No=2] inflammed/tonsillitis [Yes=1, No=2]	64	/
22.Lungs clear	65	/
[Yes=1, No=2] rhonchi [Yes=1, No=2]	66	/
crepitations [Yes=1, No=2]	67	/
23.Admission weight (kg)	68	///
24.Length (cm)	69	//

70

25.Mid arm circumference (cm)

1

26.Skin fold thickness (mm)		
Triceps	71-73	//
Subscapular	74-76	//
27.Rectal temperature (c)	77-79	//
28.Duration of stay in hospital	80-82	//
29.Outcome	83	/
Discharged with diarrhoea=1 Discharged after recovery=2' Absconded=3 DAMA=4 Referred=5 Died=6 Other=7		·. ,
30. Treatment received		, *
ORS [Yes=1, No=2]	84	
ORS + IV [Yes=1, No=2]	85	/
Anibiotics [Yes=1, No=2]	86	/ `
Other drugs [Yes=1, No=2]	87 -	/
Other [Yes=1, No=2]	88	/
31. Time of starting of ORS at hospital	89-92	///
32.Type of ORS:	93	/
Glucose=1, Rice-ORS=2		
INTAKE OUTPUT CHART		
Stool volume (ml)		
A. (8 hourly)// at		dt
B. (8 hourly)// at		dt
C. (8 hourly)/// at		dt

Urine volume (ml)	•
A. (8 hourly)/_/_/ at	dl
B. (8 hourly)// at	dt
C. (8 hourly)//_ at	dl
34. Total at 24 Hours	98–101///
ORS intake (ml)	
A. Total volume given (ml)	
Total volume leftover (ml)	,
Total volume taken (ml)	· ·
B. Total volume given (ml)	
Total volume leftover (ml)	
Total volume taken (ml)	
C. Total volume given (ml)	
Total volume leftover (ml)	·
Total volume taken (ml)	
A. (8 hourly)/_/_/ at	dt
B. (8 hourly)// at	dl
C. (8 hourly)// at	dt
35. Total at 24 Hours	102-105///
36.Frequency of vomiting for:	
1st 8 hours	106-107//
2nd 8 hours	108-109//
3rd 8 hours	110-111//

37. Frequency of feed for:

Not applicable=9

	• •			
	1st 8 hours			
	BM	112-	-113	//
	FM	. 114-	-115/	·/
	Water	116-	-117/	/ <u>·</u> _/
	Semi-solid	118-	-119/	
	Other	120-	-121/	/
	2nd 8 hours		·	,
	BM	122-	-123	<u>'</u> ;
	FM	124-	-125/	/
÷	Water	126-	-124/	<u></u>
	Semi-solid	128-	-129/	/
	Other	130-	-131/	<u>'</u> / .
	3rd 8 hours	<i>'</i> .		
	BM	132-	·133/	
	FM	134-	-135/	/
•	Water	136-	.137/	/
	Semi-solid	138-	-139/	/
	Other Other	140-	-141/	/
	EDULED IV THERAPY applicable=99999)		·	
38.We	ight before IV	. 142-	-145/	
be	hydration status fore IV Moderate=1 Moderate-severe=2 Severe=3	146	/	

€.

*

•	-	
40.Weight at 24 hrs	147-148	//
41.Weight at 48 hrs	149-152	//
42.Duration of stay in hospital	153-157	////
43.Outcome	158	/
Discharged with diarrhoea=1 Discharged after recovery=2 Absconded=3 DAMA=4 Referred=5 Died=6 Other=7		
44.Treatment received		
ORS [Yes=1, No=2]	159	/
ORS + IV [Yes=1, No=2]	160	<u></u> / .
Anibiotics [Yes=1, No=2]	161	/
Other drugs [Yes=1, No=2]	162	/
Other [Yes=1, No=2]	163	
45.Weight at discharge	. 164–167	///
46.Date of discharge	168-173	/////

SOCIO-ECONOMIC-DEMOGRAPHIC HISTORY

	47. What is your (mother) age? (year)	174-175	
	48.What class did you reach in educational institute? No education=00	176-177	· · ·// .
	49.What class did your husband reach in educational institute? No education=00	178–179	//
	50.What is the primary occupation of your husband? Farmer=01, Day labourer=02, Rickshaw/pushcart puller=03, Taxi/bus/truck driver=04,	180-181	//
	Mill worker=05, Non-executive=06, Office executive=07, Petty business=08, Big business=09, Overseas employment=10, Boatman=11, Other=88		• •
	51. How much does your family earn on an average per month? (Taka)	182-186	////
	52.Number of sleeping room for the family	187	
!	53.Floor of the house Cemented=1, Non-cemented=2	188	/
!	54.Re ID No.	189-192	//

FEEDING PRACTICES

Variable name	Column	Code
1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2	1	/
2.Randomization WHO-ORS=1, Rice-ORS=2	2	
3.Study No.	3–6	///
A. DURING ILLNESS		
4. What did you feed your child after onset of diarrhoea?		
Breast-milk	7	/ .
[Yes=1, No=2]	* -	
Formula milk [Yes=1, No=2]	8	/
Cow's milk	9.	<i>i</i>
[Yes=1, No=2]		
Semi-solid	10	/
[Yes=1, No=2]	1 1	,
Solid	11	/
[Yes=1, No=2] Other	12	1
[Yes=1, No=2]	12	/
[165-1, 110-2]		
5. How often did you feed your child during previous 24 hour period ? Not applicable=99		
breast-milk	13-14	/
unbranded powder milk	15-16	//
Commercial baby food	17-18	//
cow's milk	19-20	//
Goat's milk	21-22	//
Buffalo's milk	23-24	//
Rice gruel	25-26	/ /

Barley water	27-28	/
semi-solid '	29-30	/
solid [Yes=1, No=2]	31-32	/
other	33-34	/
B.PRIOR TO DIARRHEAL ILLNESS	•	
6.If breast-fed is that exclusive=1 partial=2 not applicable=9	- 35	/
7.If partially breast-fed, how many weeks before this episode did you start formula milk [not applicable=99]		
Unbranded powder milk	36-37	/
Commercial baby food	38-39	/
Cow's milk	40-41	/
Goat's milk	42-43	/
Buffalo's milk	44-45	/
Barley water	46-47	/
Rice gruel	48-49	/
Other	50-51	/
8.If formula fed, what was the quality of milk before illness	52	/
diluted=1, Undiluted=2, not applicable=9		
9.If formula-fed what did you give your child before diarrhoeal illness: [Not applicable=9]		
Unbranded powder milk + rice suii	53	/

Unbranded poweder milk + wheat suji [Yes=1, No=2]	54	/
Commercial baby food + rice suji [Yes=1, No=2]	55	/
Commercial baby food + wheat suji [Yes=1, No=2]	56	· , , , , , , , , , , , , , , , , , , ,
Cow's milk + rice suji [Yes=1, No=2]	57	/
Cow's milk + wheat suji [Yes=1, No=2]	58	/
Goat's milk + rice suji [Yes=1, No=2]	59	
Goat's milk + wheat suji [Yes=1, No=2]	60 🟎	/
Buffalo's milk + rice suji [Yes=1, No=2]	61	/
Buffalo's milk + wheat suji [Yes=1, No=2]	62	
C. DISTANT PAST	•	
10.Did your child receive breast milk after birth ? [Yes=1, No=2, Don't know=3]	63	/
11.What did you do with the first milk of the breast (colostrum) ?	64	
Thrown away=1, Given to the child=2, Don't know=3 [Yes=1, No=2]		
12.Re ID NO.	65-68	//

LABORATORY DATA

Variable name	Column	Code
1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2	1	/
2.Randomization WHO-ORS=1, Rice-ORS=2	2	<u></u>
3.Study No.	3-6	///
STOOL MICROSCOPICAL EXAMINATION		
4.Date of specimen received	7–12	////
5.Color yellow=1, brown=2, green=3,	13	
<pre>greenish=4, pale yellow=5, rice watery=6, creamy=7, other=8, not done=9</pre>	.	
6.Stool consistency liquid=1, watery=2, mucoid=3, bloody=4, bloody-mucoid=5, loose=6, soft=7, other=8, not done=9	14	
7.Blood none=0, trace=1, moderate=2,	15	/
heavy=3 8.Mucus	16	/
(same as above) 9.Worm	17	/
[Yes=1, No=2] 10.pH	18	/
[Acidic=1, Alkaline=2] 11.RBC none=1, 1-10=2, 11-20=3,	19	/
21-50=4, 51+=5 12.Pus cells less 10=2, 11-20=3, 21-50=4, 51+=5	20	
13.Macrophage none=2, 1-5=3, 6-10=4, 11+=5	21	/
14.Neutral fat none=2, few=3, mod=4, many=5	22	<u>/</u>

none=2, few=3, mod=4, many=5 16.Giardia	15. Yeast	23	/
many=5 16.Giardia none=0, cyst=1, trophozoite=2, cyst + trophozoite=3 17.E.H. none=0, cyst=1, trophozoite=2, cyst + trophozoite=3 18.Ascaris none=0, few=1, mod=2, many=3 19.Trichuris none=0, few=1, mod=2, many=3 20.Hookworm none=0, few=1, mod=2, many=3 21.S.Stercoralis none=0, few=1, mod=2, many=3 22.Trichomonas hominis none=0, few=1, mod=2, many=3 22.Trichomonas hominis none=0, few=1, mod=2, many=3 23.Other (Yes=1, No=2) 24.Cryptosporidium (Yes=1, No=2) STCOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae (Yes=1, No=2) 28.Vibrio cholerae serotype Inaba=1, Ogwa=2 29.Vibrio cholerae serotype Inaba=1, Ogwa=2 30.Other vibrios (Yes=1, No=2) 31.Other vibrios (Yes=1, No=2) 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, ACT, other=8, not applicable=9 31.EFEC (Yes=1, No=2) 31.ETEC (Yes=1, No=2) 31.ETEC	none=2, few=3, mod=4,		 '
16. Giardia			
none=0, cyst=1, trophozoite=2, cyst + trophozoite=3 17.E.H. none=0, cyst=1, trophozoite=2, cyst + trophozoite=3 18.Ascaris none=0, few=1, mod=2, many=3 19.Trichuris 27	*	24	/
cyst + trophozoite=3 17.E.H. none=0, cyst=1, trophozoite=2, cyst + trophozoite=3 18.Ascaris none=0, few=1, mod=2, many=3 19.Trichuris none=0, few=1, mod=2, many=3 20.Hookworm none=0, few=1, mod=2, many=3 21.S.Stercoralis none=0, few=1, mod=2, many=3 22.Trichomones hominis none=0, few=1, mod=2, many=3 22.Trichomones hominis none=0, few=1, mod=2, many=3 23.Other [Yes=1, No=2] 24.Cryptosporidium [Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 33-38 26.Vibrio cholerae [Yes=1, No=2] 27.Vibrio cholerae biotype E1 Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Oga=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC7, other=8, not applicable=9 37.EFEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3			
17.E.H.			·
none=0, cyst=1, trophozoite=2, cyst + trophozoite=3 18.Ascaris		25	/
Cyst + trophozoite=3 18.Ascaris		23	
18. Ascaris none=0, few=1, mod=2, many=3 19.Trichuris 27			
none=0, few=1, mod=2, many=3 19.Trichuris		oc	,
many=3 19.Trichuris		20	/
19.Trichuris		×	
none=0, few=1, mod=2, many=3 20.Hookworm none=0, few=1, mod=2, many=3 21.S.Stercoralis none=0, few=1, mod=2, many=3 22.Trichomomas hominis none=0, few=1, mod=2, many=3 22.Trichomomas hominis none=0, few=1, mod=2, many=3 23.Other [Yes=1, No=2] 24.Cryptosporidium [Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogava=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3			
many=3 20.Hookworm none=0, few=1, mod=2, many=3 21.S.Stercoralis none=0, few=1, mod=2, many=3 22.Trichomonas hominis none=0, few=1, mod=2, many=3 23.Other		27	/
20. Hookworm none=0, few=1, mod=2, many=3 21. S. Stercoralis none=0, few=1, mod=2, many=3 22. Trichomonas hominis none=0, few=1, mod=2, many=3 22. Trichomonas hominis none=0, few=1, mod=2, many=3 23. Other [Yes=1, No=2] 24. Cryptosporidium [Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION 25. Date of specimen sent 26. Vibrio cholerae [Yes=1, No=2] 28. Vibrio cholerae biotype El Tor=1, Classical=2 29. Vibrio cholerae serotype Inaba=1, Ogava=2 30. Other vibrios [Yes=1, No=2] 31. Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32. EFEC [Yes=1, No=2] 33. ETEC [Yes=1, No=2] 34. ETEC type ST=1, LT=2, LT/ST=3			
none=0, few=1, mod=2, many=3 21.S.Stercoralis none=0, few=1, mod=2, many=3 22.Trichomonas hominis none=0, few=1, mod=2, many=3 23.Other [Yes=1, No=2] 24.Cryptosporidium [Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3			
none=0, few=1, mod=2, many=3 21.S.Stercoralis none=0, few=1, mod=2, many=3 22.Trichomonas hominis none=0, few=1, mod=2, many=3 23.Other [Yes=1, No=2] 24.Cryptosporidium [Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VVH=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3		28	/ -
many=3 21.S.Stercoralis	none=0, $few=1$, $mod=2$,	•	 '
none=0, few=1, mod=2, many=3 22.Trichomonas hominis			
none=0, few=1, mod=2, many=3 22.Trichomonas hominis	21.S.Stercoralis	29	/
Many=3 22.Trichomomas hominis 30	none=0, few=1, mod=2,		<u> </u>
22.Trichomonas hominis			
none=0, few=1, mod=2, many=3 23.Other		30	/
Many=3 23.Other		•	
23.Other [Yes=1, No=2] 24.Cryptosporidium [Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 33-38/// 26.Vibrio cholerae		-	
[Yes=1, No=2] 24.Cryptosporidium [Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VP=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3		91	,
24.Cryptosporidium		31	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 33-38		00	,
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent		32	<u></u> / .
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3	[Yes=1, No=2]	•	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3	•	,	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3	CHOOL DACETTAL COLORS BURNEY,		
26.Vibrio cholerae	STOOL BACTERIOLOGICAL EXAMINATION		
26.Vibrio cholerae	05.00		
[Yes=1, No=2] 28.Vibrio cholerae biotype	25.Date of specimen sent	33-38	///
[Yes=1, No=2] 28.Vibrio cholerae biotype			
28. Vibrio cholerae biotype El Tor=1, Classical=2 29. Vibrio cholerae serotype Inaba=1, Ogawa=2 30. Other vibrios [Yes=1, No=2] 31. Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32. EPEC [Yes=1, No=2] 33. ETEC [Yes=1, No=2] 34. ETEC type ST=1, LT=2, LT/ST=3		39	/
El Tor=1, Classical=2 29.Vibrio cholerae serotype	[Yes=1, No=2]		
29. Vibrio cholerae serotype	28. Vibrio cholerae biotype	40	/
29. Vibrio cholerae serotype	El Tor=1, Classical=2		,
Inaba=1, Ogawa=2 30.Other vibrios		41	/
30.Other vibrios			 '
[Yes=1, No=2] 31.Other vibrios type		42	/
31.Other vibrios type			·
VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC		43	/
AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC		10	
not applicable=9 32.EPEC	AS-5 AH-6 AC-7 Olbon-9		
32.EPEC 44/ [Yes=1, No=2] 33.ETEC 45/ [Yes=1, No=2] 34.ETEC type 46/ ST=1, LT=2, LT/ST=3			
[Yes=1, No=2] 33.ETEC	• •	4.4	,
33.ETEC	•	44	/
[Yes=1, No=2] 34.ETEC type 46/ ST=1, LT=2, LT/ST=3		40	,
34.ETEC type 46/ ST=1, LT=2, LT/ST=3		40	/
ST=1, LT=2, LT/ST=3		40	,
		46	/
35.EIEC 47/		a m a	
	35.EIEC	47	/

36.EAEC	48	/
[Yes=1, No=2] 37.EAEC type	49	/
LA=1, DA=2, AA=3		
38.EHEC	50	/
[Yes=1, No=2]		
39.Klebsiella	51	/
[Yes=1, No=2]		
40.Klebsiella type	52	/
LA=1, DA=2, AA=3		
41.Rotavirus	53	/
[Yes=1, No=2]		
42.Campylobacter	54	/
[Yes=1, No=2]	•	
43.Campylobacter type	55	/
jejuni=1, coli=2, other=3		
44.Salmonellae	56	/.
[Yes=1, No=2]		
45.Salmonellae type	57	/
salm. typhi=1		
salm. other=2		•
46.Shigellae	58	/
[Yes=1, No=2]		
47.Shigellae type	59 🕶	/
sh. dyst I=1, sh. flex=2,		
sh. boydii=3, sh. sonnei=4,		
sh. dyst 2=5,		
sh. dyst (3-10)=6		, ,
sh. dyst (3-10)=6 48.Sensitivity pattern of:	60-61	//
48. Sensitivity pattern of:	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14,	60-61`	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19,	60-61	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99	60-61 `	//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49. Sensitive to: Tetra		//
48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2]		//
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49. Sensitive to: Tetra	62	/
48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2] Ampi	62	
48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2] Ampi [Sen=1, Res=2]	62 63 64	/
48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2] Ampi [Sen=1, Res=2] TMP-SMX	62 63	/
<pre>48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2] Ampi [Sen=1, Res=2] TMP-SMX [Sen=1, Res=2]</pre>	62 63 64 65	/
<pre>48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2] Ampi [Sen=1, Res=2] TMP-SMX [Sen=1, Res=2] Furox</pre>	62 63 64	/
<pre>48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2] Ampi [Sen=1, Res=2] TMP-SMX [Sen=1, Res=2] Furox [Sen=1, No=2]</pre>	62 63 64 65 66	/
<pre>48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to: Tetra [Sen=1, Res=2] Ampi [Sen=1, Res=2] TMP-SMX [Sen=1, Res=2] Furox [Sen=1, No=2] Chlor</pre>	62 63 64 65	/

[Sen=1, Res=2]

Nalidixic acid	68	/
[Sen=1, Res=2] Sulpha	69	/
[Sen=1, Res=2] Pivmecillinum	70	/
[Sen=1, Res=2]		
Other [Sen=1, Res=2]	71	
50.Sensitivity pattern of:	72-73	//
VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99		
51.Sensitive to :		,
Tetra	74 ~	
[Sen=1, Res=2] Ampi	75	/
[Sen=1, Res=2]	¹ 76	/
[Sen=1, Res=2]		/
Furox [Sen=1, No=2]	77	
Chlor	78	/
[Sen=1, Res=2] Genta	79	/
[Sen=1, Res=2]	22	,
Nalidixic acid [Sen=1, Res=2]	80	/
Sulpha	81	/
[Sen=1, Res=2] Pivmecillinum	82	/
[Sen=1, Res=2] Other	83	1
[Sen=1, Res=2]	00	
BLOOD EXAMINATION		
52.Date of specimen sent	84-89	//////
53.Total WBC		y y m m u u
% Polys	90-91	//
% Bands	92-93	//

% Monocyte	96-97	//
% Eosinophil	98-99	//
% Basophil	100-101	//
54. % HCT	102-103	//
55. Total Protein	104-106	//
56. Specific gravity	107-111	//
MULTITEST CMI REACTION RESULTS		
57.TETANUS diameter (a) nun diameter (b) nun average nun positivity [Yes=1, No=2]	112-113 114-115 116-117 118	// // /
58.DIPHTHERIA	n -	·
diameter (a) mm diameter (b) mm average num positivity [Yes=1, No=2]	119-120 121-122 123-124 125	
59.STREPTOCOCCUS		
diameter (a) nun diameter (b) nun average nun positivity [Yes=1, No=2]	126-127 128-129 130-131 132	// / /
60.TUBERCULIN		
diameter (a) nun diameter (b) nun average nun positivity [Yes=1, No=2]	133-134 135-136 137-138 139	
61.CONTROL		
diameter (a) num diameter (b) num average num positivity {Yes=1,_No=2}	140-141 142-143 144-145 146	

62.CANDIDINE		
diameter (a) mm diameter (b) mm average mm positivity [Yes=1, No=2]	147-148 149-150 151-152 153	// // /
63.TRICOPHYTON		·
diameter (a) min diameter (b) min average min positivity [Yes=1, No=2]	154-155 156-157 158-159 160	// / /
64.PROTEUS		
diameter (a) mm diameter (b) mm average mm positivity [Yes=1, No=2]	161–162 163–164 165–166 167	
65.Re ID No.	168-171	/ / / /

FIRST FOLLOW-UP VISIT (Day 8 of illness)

PART A: MEDICAL HISTORY

1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2	1	/
2.Randomization WHO-ORS=1, Rice-ORS=2	2	
<pre>3.Study No. 4.If recovered, what was the duration of illness (days)? not applicable=99</pre>	3-6 7-8	///
5.If the child has diarrhoea, is the episode: Still continuing=1, New episode=2, Not applicable=9	9	/
6.If new episode, what is the duration of diarrhoea (hrs) not applicable=99999	10-14	///
7.If new episode, what is the duration of vomiting (hrs) not applicable=99999	15 - 19 -	////
8.if new episode, what is the character of stool? watery/loose=1, bloody or bloody-mucoid=2, mucoid=3, not applicable=9	20	/
9.Did your child take ORS that were given at the time of discharge? [Yes=1, No=2]	21	
10.Did the child receive any outside treatment at home after discharge? [Yes=1, No=2]	22	/

	•	
11.If yes:	,	•
antibacterial drugs [Yes=1, No=2 not applicable=9]	23	/
name of the drugs		
anthelmentic drugs [Yes=1, No=2 not applicable=9]	24	/
name of the drugs		
antiprotozoal drugs [Yes=1, No=2 not applicable=9]	25	/
name of the drugs		
antiperistaltic drugs [Yes=1, No=2 not applicable=9]	26	/
name of the drugs	-	
other drugs [Yes=1, No=2 not applicable=9]	27	/
name of the drugs		
12.Did the child receive any outside treatment at home for the new episode? [Yes=1, No=2]	25	/
13.If yes: antibacterial drugs [Yes=1, No=2 not applicable=9]	26	/
name of the drugs		
anthelmentic drugs [Yes=1, No=2 not applicable=9]	27	/
name of the drugs		

antiprotozoal drugs [Yes=1, No=2 not applicable=9]	. 28	
name of the drugs		
antiperistaltic drugs [Yes=1, No=2 not applicable=9] name of the drugs	29	/ ,
the drugs		
other drugs [Yes=1, No=2 not applicable=9]	30	/
name of the drugs		
PART B: PHYSICAL FINDINGS		
14. Dehydration	31	/
none=1, mild=2, moderate=3, severe=4 not applicable=9		
15.Rectal temperature	32-34	
16. Signs of vitamin A deficiency		
Normal [Yes=1, No=2]	, 35	
Night blindness (XN) [Yes=1, No=2]	36	/
Conjunctival xerosis (X1A) [Yes=1, No=2]	37	/
Bitot's spot (X1B) [Yes=1, NO=2]	38	/
Corneal ulcer <1/3 (X3A) [Yes=1, NO=2]	39	/
Corneal ulcer >1/3 (X3B) {Yes=1, No=2}	40	/
Xerophthalmia scar (XS) [Yes=1, No=2]	41 .	/
Xerophthalmia fundus (XF) [Yes=1, No=2]	42	·/
17.Otitis media		_
	43	/

IYes=1. No=2

18. Throat		
normal [Yes=1, No=2]	44	
inflammed/tonsillitis [Yes=1, No=2]	45	/
19.Lungs	·	
clear [Yes=1, No=2]	46	/
rhonchi [Yes=1, No=2]	47	<i>'</i>
crepitations [Yes=1, No=2]	48	/
20.Weight (kg)	49-52	
21.Length (cm)	53-55	_/_/
22. Mid arm circumference (cm)	56-58	
23.Skin fold thickness (mm)		
Triceps	59-61 ⊷	//
Subscapular ,	62-64	/ / /

·	ea)	•
1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2		/
2.Randomization WHO-ORS=1, Rice-ORS=2	2	/
3.Study No.	3-6	///
STOOL MICROSCOPICAL EXAMINATION		
4.Date of specimen received	7-12	///////
5.Color yellow=1, brown=2, green=3, greenish=4, pale yellow=5, rice watery=6, creamy=7, other=8, not done=9	13	
6.Stool consistency liquid=1, watery=2, mucoid=3, bloody=4, bloody-mucoid=5, loose=6, soft=7, other=8, not done=9	14 •	
7.Blood none=0, trace=1, moderate=2, heavy=3	15	/
8.Mucus (same as above)	16	/
9.Worm [Yes=1, No=2]	17	/
10.pH [Acidic=1, Alkaline=2]	18	/
11.RBC none=1, 1-10=2, 11-20=3, 21-50=4, 51+=5	1,9	/
12.Pus cells less 10=2, 11-20=3, 21-50=4, 51+=5	20	
13.Macrophage none=2, 1-5=3, 6-10=4, 11+=5	21	
14. Neutral fat none=2, few=3, mod=4, many=5	22	/
15.Yeast none=2, few=3, mod=4, many=5	23	

16.Giardia		-
	24	/
none=0, cyst=1, trophozoite=2, cyst + trophozoite=3	•	
17.E.H.		
none=0, eyst=1, trophozoite=2,	25	/
∪yau ⊤ Urophozoi1p-3		-
18. Ascaris	26	
none=0, few=1, mod=2,	. 20	/
many=3	•	•
19.Trichuris	27	,
none=0, few=1, mod=2,		/
many=3 20.Hookworm		
none=0, few=1, mod=2,	28	<i>J</i>
many=3		
21.S.Stercoralis		
none=0, few=1, mod=2,	29	/
many=3		,
22.Trichomonas hominis	30	
none=0, few=1, mod=2.	30	/
many=3		
23.0ther	31	,
[Yes=1, No=2]	01	/
24.Cryptosporidium	32 •	
[Yes=1, No=2]		 /
•		
STOOL BACTERIOLOGICAL EXAMINATION		
25.Date of specimen sent	33-38	
	00 00	/////
26. Vibrio cholerae	39	,
[Yes=1, No=2]		/
28. Vibrio cholerae biotype	40	1
El Tor=1, Classical=2		/
29.Vibrio cholerae serotype Inaba=1, Ogawa=2	41	/
30.Other vibrios		 '
[Yes=1, No=2]	42	/
31.0ther vibrios type	43	
VP=1, $VF=2$, $VM=3$, $PS=4$.	40	/
AS=5, $AH=6$, $AC=7$, other=8.		
not appircable=9		
32.EPEC	44	/
[Yes=1, No=2] 33.ETEC		/
[Yes=1, No=2]	45	/
34.ETEC Lype		
ST=1, LT=2, LT/ST=3	46	/
35.EIEC 2, 27, 27, 27, 27, 27, 27, 27, 27, 27,	A.17	
[Yes=1, No=2]	47	/
36.EAEC	48	,
[Yes=1, No=2]	1 U	/
37.EAEC type		

38.EHEC		
	50	/
[Yes=1, No=2] 39.Klebsiella		/
[Yes=1, No=2]	51	/
40.Klebsiella type		 -
LA=1, $DA=2$, $AA=3$	52	/
41.Rotavirus	5 0	
[Yes=1, No=2]	53	/
42. Campylobacter	54	
[Yes=1, No=2]	94	/
43.Campylobacter type	55	,
jejuni=1, coli=2, other=3		/
44.5almonellae	56	,
[Yes=1, No=2]	00	/
45.Salmonellae type	57	,
salm. typhi=1	01	/
salm. other=2		
46.Shigellae	58	,
[Yes=1, No=2]		/
47.Shigellae type	59	,
sh. dyst I=1, sh. flex=2,	•	/
sh. boydii=3, sh. sonnei=4,		
sh. dyst 2=5,		
sh. dyst (3-10)=6	4-	
48. Sensitivity pattern of:	60-61	/ /
VC 01-01 VD 00 17 00		/
VC 01=01, VP=02, VF=03, VM=04 PS=05		
VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10,		
Salm. typhi=11, Salm. other=12	•	
Sh. dyst I=13, Sh. flex=14,	,	
Sh. sonnei=15, Sh. boydii=16,		
Sh. dyst 2=17,		
Sh. dyst (3-10)=18, Other=19,		
Not done=88, Not applicable=99		
and do, not applicable=99		•
49. Sensitive to:		
1		
Tetra	62	
[Sen=1, Res=2]	02	/
Ampi	63	. ,
[Sen=1, Res=2]		/
TMP-SMX	64	/
[Sen=1, Res=2] Furox		/
	65	,
[Sen=1, No=2] Chlor	•	
[Sen=1, Res=2]	66	1
Genta		 *,
[Sen=1, Res=2]	67	_ /
Nalidixie acid	60	<u>-</u> -
[Sen=1, Res=2]	68	/
Sulpha	69	
[Sen=1, Res=2]	ua	/

		·
Pivmecillinum	70	/
[Sen=1, Res=2] Other		
[Sen=1, Res=2]	71	/
50. Sensitivity pattern of:	72-73	//
VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99	•	
51. Sensitive to:		
Tetra [Sen=1, Res=2]	74	/
Ampi [Sen=1, Res=2]	75	/
TMP-SMX [Sen=1, Res=2]	76 👡	
Furox	77	
[Sen=1, No=2]	11 -	/
Chlor	70	,
	78	/
[Sen=1, Res=2]	5 0	
Genta	79	/
[Sen=1, Res=2] Nalidixic acid	00	,
[Sen=1, Res=2]	80	/
Sulpha	81	,
[Sen=1, Res=2]	01	•
Pivmecillinum	82	/
[Sen=1, Res=2]	02	***************************************
Other	83	/
[Sen=1, Res=2]	00	<u> </u>
BLOOD EXAMINATION		
52.Date of specimen sent	84-89	Y Y M M D D
53. Total WBC		Y Y M M D D
% Polys	90-91	//
% Bands	92-93	//
% Lymohocyte	94-95	//
% Monocyte	96-97	//

% Basophil	100-101	//
54. % HCT	102-103	//
55. Total Protein	104-106	///
56. Specific gravity	107-111	///
57 Re ID No.	112-115	/ / / /

SECOND FOLLOW-UP VISIT (Day 16 of illness)

[Yes=1, No=2]

PART A: MEDICAL HISTORY 1 1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2 2 2. Randomization WHO-ORS=1, Rice-ORS=2 3-6 3.Study No. 7-8 4. If recovered, what was the duration of illness (days)? not applicable=99 5.If the child has diarrhoea, is the episode: Still continuing=1, New episode=2, Not applicable=9 10 - 146.If new episode, what is the duration of diarrhoea (hrs) not applicable=99999 15 - 197.If new episode, what is the duration of vomiting (hrs) not applicable=99999 20 8.if new episode, what is the character of stool? watery/loose=1, bloody or bloody-mucoid=2. mucoid=3, not applicable=9 21 9.Did your child take ORS that were given at the time of discharge? [Yes=1, No=2] 22 10.Did the child receive any outside treatment at home after discharge?

11.If yes:		
antibacterial drugs [Yes=1, No=2 not applicable=9]	.23	/
name of the drugs	-	
anthelmentic drugs [Yes=1, No=2 not applicable=9]	24 .	/
name of the drugs	•	
antiprotozoal drugs [Yes=1, No=2 not applicable=9]	25	/
name of the drugs	-	
antiperistaltic drugs [Yes=1, No=2 not applicable=9]	26	/
name of the drugs	_	
other drugs [Yes=1, No=2 not applicable=9]	27	/
name of the drugs	-	
12.Did the child receive any outside treatment at home for the new episode ? [Yes=1, No=2]	25	/
13.If yes: antibacterial drugs [Yes=1, No=2 not applicable=9]	26	/
name of the drugs		
anthelmentic drugs [Yes=1, No=2 not applicable=9]	27	/
name of the drugs		

antiprotozoal drugs [Yes=1, No=2 not applicable=9]	28	/
name of the drugs		•
antiperistaltic drugs [Yes=1, No=2 . not applicable=9]	29	
name of the drugs		
other drugs [Yes=1, No=2 not applicable=9] name of the drugs	30 .	
PART B: PHYSICAL FINDINGS	8-	·
14. Dehydration	31	/
none=1, mild=2, moderate=3, severe=4 not applicable=9		
15.Rectal temperature	32-34	//
16. Signs of vitamin A deficiency		
Normal [Yes=1, No=2]	35	/
Night blindness (XN) [Yes=1, No=2]	36	
Conjunctival xerosis (X1A) [Yes=1, No=2]	37	/
Bitol's spot (X1B) [Yes=1, NO=2]	38	/
Corneal ulcer <1/3 (X3A) [Yes=1, NO=2]	39	/
Corneal ulcer >1/3 (X3B) [Yes=1, No=2]	40	/
Xerophthalmia scar (XS)	41	/
[Yes=1, No=2] Xerophthalmia fundus (XF) [Yes=1, No=2]	42	
17.Otitis media [Yes=1, No=2]	43	/

18.Throat		
normal	44	/
[Yes=1, $No=2$]		
inflammed/tonsillitis	45	/
[Yes=1, No=2]		
19.Lungs		
clear	46	/
[Yes=1, No=2]		
rhonchi	47	/
[Yes=1, No=2]		
crepitations	48	/
[Yes=1, No=2]		
20.Weight (kg)	19-52	///
21.Length (cm)	53-55	//
22.Mid arm circumference (cm)	56-58	//
23.Skin fold thickness (mm)		
Triceps	59-61-	//
Subscapular	62-64	/ / /

<u> many=5</u>

1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2	1	/
2.Randomization WHO-ORS=1, Rice-ORS=2	2	/
3.Study No.	3-6	///
STOOL MICROSCOPICAL EXAMINATION		
4.Date of specimen received	7-12	/////
5.Color yellow=1, brown=2, green=3, greenish=4, pale yellow=5, rice watery=6, creamy=7, other=8, not done=9	13	/
6.Stool consistency liquid=1, watery=2, mucoid=3, bloody=4, bloody-mucoid=5, loose=6, soft=7, other=8, not done=9	14 •	/
7.Blood none=0, trace=1, moderate=2, heavy=3	15	
8. Mucus (same as above)	16	/
9.Worm	17	/
[Yes=1, No=2]		 -
10.pH		
-	18	
[Acidic=1, Alkaline=2] 11.RBC	18 19	/ /
[Acidic=1, Alkaline=2] 11.RBC none=1, 1-10=2, 11-20=3,		/ /
[Acidic=1, Alkaline=2] 11.RBC none=1, 1-10=2, 11-20=3, 21-50=4, 51+=5 12.Pus cells less 10=2, 11-20=3,		/ /
[Acidic=1, Alkaline=2] 11.RBC none=1, 1-10=2, 11-20=3, 21-50=4, 51+=5 12.Pus cells less 10=2, 11-20=3, 21-50=4, 51+=5 13.Macrophage none=2, 1-5=3, 6-10=4,	19	//
[Acidic=1, Alkaline=2] 11.RBC none=1, 1-10=2, 11-20=3, 21-50=4, 51+=5 12.Pus cells less 10=2, 11-20=3, 21-50=4, 51+=5 13.Macrophage	19	/////

16.Giardia	24	,		
none=0, cyst=1, trophozoite=2,	~ .	/	•	
cyst + trophozoite=3				
17.E.H.	25 .	/	•	
none=0, cyst=1, trophozoite=2,		<u></u> ′		
cyst + trophozoite=3	,			
18. Ascaris	26	/		
none=0, few=1, mod=2,		—·		
many=3				
19.Trichuris	27	/		
none=0, few=1, mod=2,				
many=3 20.Hookworm				
	28	/		
none=0, few=1, mod=2,				
, many=3 21.S.Stercoralis			•	
	29	/		
none=0, few=1, mod=2,				
many=3			,	
22. Trichomonas hominis	30	/		
none=0, few=1, mod=2, many=3				
23.0ther	0.1			
[Yes=1, No=2]	31	/		
24.Cryptosporidium	20	ý	•	
	32 👞	/		
[Yes=1, No=2] STOOL BACTERIOLOGICAL EXAMINATION		_		
STOOL BACTERIOLOGICAL EXAMINATION	33-38	//	'//	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent		//	'//	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae	33 -38 	//	'///	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2]	39	//	'//	/
STOOL BACTERIOLOGICAL EXAMINATION 25. Date of specimen sent 26. Vibrio cholerae [Yes=1, No=2] 28. Vibrio cholerae biotype		//	'//	/
5.Date of specimen sent 6.Vibrio cholerae [Yes=1, No=2] 8.Vibrio cholerae biotype El Tor=1, Classical=2	39 40	//	'///	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2	39	// / /	'///	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2	39 40 41	// / /	'/// ,	/
5.Date of specimen sent 6.Vibrio cholerae [Yes=1, No=2] 8.Vibrio cholerae biotype El Tor=1, Classical=2 9.Vibrio cholerae serotype Inaba=1, Ogawa=2 0.Other vibrios [Yes=1, No=2]	39 40	// / /	'///	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 0.Other vibrios [Yes=1, No=2] 1.Other vibrios type	39 40 41 42	// / /	'///	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4,	39 40 41	// / /	'/// '//// '	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 1.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8,	39 40 41 42	// / / /	'/// '/	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 1.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9	39 40 41 42	// / / /	'// ,	/
ETCOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 1.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 2.EPEC	39 40 41 42	// / / /	'/// -	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 11.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 2.EPEC [Yes=1, No=2]	39 40 41 42 43	// / / /	'/// -	/
6.Vibrio cholerae [Yes=1, No=2] 8.Vibrio cholerae biotype El Tor=1, Classical=2 9.Vibrio cholerae serotype Inaba=1, Ogawa=2 0.Other vibrios [Yes=1, No=2] 1.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 2.EPEC [Yes=1, No=2] 3.ETEC	39 40 41 42 43	// / / /	'/// '	
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 20.Other vibrios [Yes=1, No=2] 1.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 2.EPEC [Yes=1, No=2] 3.ETEC [Yes=1, No=2]	39 40 41 42 43	//////	'/// -	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 3.Other vibrios [Yes=1, No=2] 1.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 2.EPEC [Yes=1, No=2] 3.ETEC [Yes=1, No=2] 4.ETEC type	39 40 41 42 43	/////	'// -	/
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 1.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 2.EPEC [Yes=1, No=2] 3.ETEC [Yes=1, No=2] 4.ETEC type ST=1, LT=2, LT/ST=3	39 40 41 42 43 44 45 46	//////		
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 44.ETEC type ST=1, LT=2, LT/ST=3 5.EIEC	39 40 41 42 43	///////	'/// -	,
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 44.ETEC type ST=1, LT=2, LT/ST=3	39 40 41 42 43 44 45 46 47	///////	'/// -	
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 34.ETEC [Yes=1, No=2] 44.ETEC type ST=1, LT=2, LT/ST=3 5.EIEC [Yes=1, No=2]	39 40 41 42 43 44 45 46	///////	'/// - -	/

38.EHEC [Yes=1, No=2] 39.Klebsiella [Yes=1, No=2] 40.Klebsiella type LA=1, DA=2, AA=3 41.Rotavirus [Yes=1, No=2] 42.Campylobacter [Yes=1, No=2] 43.Campylobacter type jejuni=1, coli=2, other=3 44.Salmonellae [Yes=1, No=2] 45.Salmonellae type salm. typhi=1 salm. other=2 46.Shigellae [Yes=1, No=2] 47.Shigellae type sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst 2	_/.
39.Klebsiella [Yes=1, No=2] 40.Klebsiella type	_/.
[Yes=1, No=2] 40.Klebsiella type	/
40.Klebsiella type	/
IA=1, DA=2, AA=3 41.Rotavirus	/
41. Rotavirus 53	/
[Yes=1, No=2] 42.Campylobacter [Yes=1, No=2] 43.Campylobacter type	/
42.Campylobacter [Yes=1, No=2] 43.Campylobacter type	/
[Yes=1, No=2] 43.Campylobacter type	/
43.Campylobacter type	/
jejuni=1, coli=2, other=3 44.Salmonellae [Yes=1, No=2] 45.Salmonellae type salm. typhi=1 salm. other=2 46.Shigellae [Yes=1, No=2] 47.Shigellae type sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	/
44.Salmonellae [Yes=1, No=2] 45.Salmonellae type 57 salm. typhi=1 salm. other=2 46.Shigellae 58 [Yes=1, No=2] 47.Shigellae type 59 sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: 60-61 VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
[Yes=1, No=2] 45.Salmonellae type	_/
45.Salmonellae type 57 salm. typhi=1 salm. other=2 46.Shigellae 58 [Yes=1, No=2] 47.Shigellae type 59 sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: 60-61 VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
salm. typhi=1 salm. other=2 46.Shigellae [Yes=1, No=2] 47.Shigellae type sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
salm. other=2 46.Shigellae [Yes=1, No=2] 47.Shigellae type sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
46.Shigellae [Yes=1, No=2] 47.Shigellae type sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
[Yes=1, No=2] 47.Shigellae type sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
[Yes=1, No=2] 47.Shigellae type	_/
sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
sh. dyst I=1, sh. flex=2, sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
sh. boydii=3, sh. sonnei=4, sh. dyst 2=5, sh. dyst (3-10)=6 48. Sensitivity pattern of: 60-61/ VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
sh. dyst 2=5, sh. dyst (3-10)=6 48.Sensitivity pattern of: 60-61/_ VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
48. Sensitivity pattern of: VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	_/
VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	/
VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	
VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	
AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	
Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	
Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16.	
Sh. sonnei=15, Sh. boydii=16.	
Sh. dyst 2=17,	
w/20 B-1/4	
Sh. dyst (3-10)=18, Other=19,	
Not done=88, Not applicable=99	
Not done=00; Not applicable=99	
49. Sensitive to:	
Tetra 62	
[Sen=1, Res=2]	
Ampi 63	
[Sen=1, Res=2]	
TMD CMC	
[Sen=1, Res=2].	
The same of the sa	
[Sen=1, No=2]	
C1. 1	
(Sen=1, Res=2)	
Court	
[Sen=1, Res=2]	
37. 7 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
[Sen=1, Res=2]	
01.1	
Surpna 69/	
(DOII-14 IVCS-6)	

Pivmecillinum	70	
[Sen=1, Res=2]	70	/·
Other	71	,
[Sen=1, Res=2]	•	/
50 Sensitivity and		
50. Sensitivity pattern of:	72-73	//
VC 01=01, VP=02, VF=03,		•
VM=04, PS=05, AH=06, AS=07,		
AC=08, ETEC=09, Campy=10,		
Salm. typhi=11, Salm. other=12	7	
Sh. dyst I=13, Sh. flex=14,	۵,	
Sh. sonnei=15, Sh. boydii=16,		
Sh. dyst 2=17,		
Sh. dyst (3-10)=18, Other=19.		
Not done=88, Not applicable=99)	
51. Sensitive to :		
or sensitive to :		
Tetra	, ,,	
[Sen=1, Res=2]	74	/
Ampi	75	
[Sen=1, Res=2]	75 .	/
TMP-SMX	76	
[Sen=1, Res=2]	70 m.	
Furox	7 7	,
[Sen=1, No=2]		/
Chlor	78	
[Sen=1, Res=2] Genta		·
[Sen=1, Res=2]	79	/
Nalidixic acid	2.0	
[Sen=1, Res=2]	80	/
Sulpha	0.1	
[Sen=1, Res=2]	81	/
Pivmecillinum	82	,
[Sen=1, Res=2]		/
Other	83	/
[Sen=1, Res=2]		
BLOOD EXAMINATION		•
22000 INDITION TON		
52. Date of specimen sent	0.1.00	
	84 -89	<u>Y</u> <u>Y</u> <u>M</u> <u>M</u> <u>D</u> <u>D</u>
53. Total WBC		Y Y M M D D
% Polys	90-91	
. 9 Parata		
% Bands	92-93	//
% Lymohocyte	04.05	
	94-95	//
% Monocyte	96-97	, ,
·	JU-31	//

% Basophil	100-101	//
54. % HCT	102-103	//
55. Total Protein	104-106	//
56. Specific gravity	107-111	///
57.Re ID No.	112-115	/_ / / /

THIRD FOLLOW-UP VISIT (Day 32 of illness)

[Yes=1, No=2]

PART A: MEDICAL HISTORY 1. Cohort No. 1 Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2 2.Randomization 2 WHO-ORS=1. Rice-ORS=2 3.Study No. 3 - 64.If recovered, what was the 7 - 8duration of illness (days)? not applicable=99 5.If the child has diarrhoea, is the episode: Still continuing=1, New episode=2, Not applicable=9 6.If new episode, what is the 10 - 14duration of diarrhoea (hrs) not applicable=99999 7.If new episode, what is the 15 - 19duration of vomiting (hrs) not applicable=99999 8.if new episode, what is the 20 character of stool? watery/loose=1, bloody or bloody-mucoid=2, mucoid=3, not applicable=9 9.Did your child take ORS 21 that were given at the time of discharge? [Yes=1, No=2] 10.Did the child receive any 22 outside treatment at home after discharge?

11.If yes:		
antibacterial dru [Yes=1, No=2 not applicable=9]	gs 23	·
name of the drugs		
anthelmentic drugs [Yes=1, No=2 not applicable=9]	s 24	/
name of the drugs		
antiprotozoal drug [Yes=1, No=2 not applicable=9]	ર્લ ક 25	/
name of the drugs		
antiperistaltic dr [Yes=1, No=2 not applicable=9]	rugs 26	/
name of the drugs		
other drugs [Yes=1, No=2 not applicable=9]	27	/
name of the drugs		
12.Did the child receive outside treatment at 1 for the new episode? [Yes=1, No=2]	any 25 home	/
13.If yes: antibacterial drugs [Yes=1, No=2 not applicable=9]	s 26	/
name of the drugs		
anthelmentic drugs [Yes=1, No=2 not applicable=9]	27	/

name of the drugs _

antiprotozoal drugs [Yes=1, No=2 not applicable=9]	28	/
name of the drugs	_	
antiperistaltic drugs [Yes=1, No=2 not applicable=9]	29	
name of the drugs		
other drugs [Yes=1, No=2 not applicable=9]	30	·/
name of the drugs		
PART B: PHYSICAL FINDINGS		
14. Dehydration	31	/
none=1, mild=2, moderate=3, severe=4 not applicable=9		
15.Rectal temperature	32-34	//
16.Signs of vitamin A deficiency		
Normal [Yes=1, No=2]	35	/
Night blindness (XN) [Yes=1, No=2]	36	/
Conjunctival xerosis (X1A) [Yes=1, No=2]	37	/
Bitot's spot (X1B)	38	/
[Yes=1, NO=2] Corneal ulcer <1/3 (X3A)	39	/
[Yes=1, NO=2] Corneal ulcer >1/3 (X3B)	40	
(Yes=1, No=2) Xerophthalmia scar (XS)	41	/
[Yes=1, No=2] Xerophthalmia fundus (XF) [Yes=1, No=2]	42	/
17.Otitis media	43	/

18.Throat		
normal	44	/
[Yes=1, No=2]		
inflammed/tonsillitis	45	/
[Yes=1, No=2]		
19.Lungs		
clear	46	/
[Yes=1, No=2]		
rhonchi	47	/
[Yes=1, No=2]		
crepitations	48	/
[Yes=1, No=2]		
20.Weight (kg)	49-52	///
21.Length (cm)	53-55	///
22.Mid arm circumference (cm)	56-58	/ / /
•		
23.Skin fold thickness (mm)		
Triceps	59-64	//
Subscapular	62-64	/ / /

1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2	1	/
2.Randomization WHO-ORS=1, Rice-ORS=2	2	
3.Study No.	3-6	///
STOOL MICROSCOPICAL EXAMINATION		
4.Date of specimen received	7-12	////
5.Color yellow=1, brown=2, green=3, greenish=4, pale yellow=5, rice watery=6, creamy=7, other=8, not done=9	13	/
6.Stool consistency liquid=1, watery=2, mucoid=3, bloody=4, bloody-mucoid=5, loose=6, soft=7, other=8, not done=9	14 •	/
7.Blood none=0, trace=1, moderate=2, heavy=3	15	/
8. Mucus	16	/
(same as above) 9.Worm	17	/
[Yes=1, No=2]		
10.pH [Acidic=1, Alkaline=2]	18	
11.RBC	19	/
none=1, 1-10=2, 11-20=3, 21-50=4, 51+=5		
12.Pus cells less 10=2, 11-20=3, 21-50=4, 51+=5	20	
13.Macrophage none=2, 1-5=3, 6-10=4,	21	/
11+=5 14.Neutral fat none=2, few=3, mod=4, many=5	22	/
15.Yeast none=2, few=3, mod=4, many=5	23	

•		
16.Giardia	24	/
none=0, cyst=1, trophozoite=2,		······································
cyst + trophozoite=3		
17.E.H.	25	/
none=0, cyst=1, trophozoite=2,		· <u>-</u> '
cyst + trophozoite=3		
18. Ascaris	26	/
none= 0 , few= 1 , mod= 2 ,		·
many=3		
19.Trichuris	27	/
none= 0 , few= 1 , mod= 2 ,		
many=3		
20.Hookworm	28	/
none=0, few=1, mod=2,		
many=3		
21.S.Stercoralis	29	/
none= 0 , few= 1 , mod= 2 ,	1	
many=3		
22.Trichomonas hominis	30	/
none=0, $\Gamma ew=1$, $mod=2$,		
many=3		,
23.Other	31	/
[Yes=1, No=2]		•
24.Cryptosporidium	32 📥	/
[Yes=1, No=2]		
•		
CTOOL DACTED TOLOCTOR THANKS AND ON		
STOOL BACTERIOLOGICAL EXAMINATION		
25.Date of specimen sent	33-38	
20.Date of specimen sent	33 - 36	////
26. Vibrio cholerae	39	,
[Yes=1, No=2]		'
28. Vibrio cholerae biotype	40	/
El Tor=1, Classical=2	••	·/
29. Vibrio cholerae serotype	41	/
Inaba=1, Ogawa=2		
30.Other vibrios	42	/ .
[Yes=1, No=2]		
31.Other vibrios type	43	/
VP=1, VF=2, VM=3, PS=4,		
AS=5, AH=6, AC=7, other=8,		
not applicable=9		-
32.EPEC	44	/
$\{Yes=1, No=2\}$		
33.ETEC	45	/
[Yes=1, No=2]		
34.ETEC type	46	/
ST=1, LT=2, LT/ST=3	4.5	
35.EIEC	47	/
[Yes=1, No=2]		
36.EAEC	10	
\	48	/
[Yes=1, No=2] 37.EAEC Lype	48 49	/

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	51 52 53 54 55 56 57 58 59

•		
Pivmecillinum	70	
[Sen=1, Res=2] Olher	71	,
[Sen=1, Res=2]	11	/
50.Sensitivity pattern of:	72-73	answered series
VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99		
51. Sensitive to:		•
Tetra	74	
[Sen=1, Res=2] .Ampi	75	/
[Sen=1, Res=2]	10	/
TMP-SMX	76	
[Sen=1, Res=2] Furox	77	/
[Sen=1, No=2]	* *	 ′
Chlor	78	/
[Sen=1, Res=2] Genta	79	,
[Sen=1, Res=2]		/
Nalidixic acid	80	
[Sen=1, Res=2]	0.1	,
Sulpha [Sen=1, Res=2]	81	/
Pivmecillinum	82	/
[Sen=1, Res=2] Olher	83	,
[Sen=1, Res=2]	03	/
BLOOD EXAMINATION		
52.Date of specimen sent	84-89	//////
53. Total WBC		
% Polys	90-91	//
% Bands	92-93	//
% Lymohocyte	94-95	//
% Monocyte	96-97	//

% Basophil	100-101	/
54. % HCT	102-103	//
55. Total , Protein	104-106	//
56. Specific gravity	107-111	/////
57.Re ID No.	112-115	///

FOURTH FOLLOW-UP VISIT (Day 48 of illness)

PART A: MEDICAL HISTORY 1.Cohort No. 1 Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2 2.Randomization 2 WHO-ORS=1. Rice-ORS=2 3.Study No. 3 - 64. If recovered, what was the 7-8 duration of illness (days)? not applicable=99 5.If the child has diarrhoea, 9 is the episode: Still continuing=1, New episode=2, Not applicable=9 6. If new episode, what is the 10 - 14duration of diarrhoea (hrs) not applicable=99999 7. If new episode, what is the 15 - 19/___/__/ duration of vomiting (hrs) not applicable=99999 8.if new episode, what is the 20 character of stool? watery/loose=1, bloody or bloody-mucoid=2, mucoid=3, not applicable=9 % 9.Did your child take ORS 21 that were given at the time of discharge? [Yes=1, No=2] 10.Did the child receive any 22 outside treatment at home after discharge? [Yes=1, No=2]

11.If yes:

antibacterial drugs [Yes=1, No=2 not applicable=9]	23	/
name of the drugs		
anthelmentic drugs [Yes=1, No=2 not applicable=9]	. 24	/
name of the drugs	_	
antiprotozoal drugs [Yes=1, No=2 not applicable=9]	25	/
name of the drugs		
antiperistaltic drugs [Yes=1, No=2 not applicable=9]	26 •-	
name of the drugs		
other drugs [Yes=1, No=2 not applicable=9]	27	<u> </u>
name of the drugs	_	
12.Did the child receive any outside treatment at home for the new episode ? [Yes=1, No=2]	· · 25	/
13.If yes:		
antibacterial drugs [Yes=1, No=2 not applicable=9]	26	/
name of the drugs	-	
anthelmentic drugs [Yes=1, No=2 not applicable=9]	27	. —_/
name of the drugs	-	

antiprotozoal drugs [Yes=1, No=2 not applicable=9]	28	/
name of the drugs		
antiperistaltic drugs [Yes=1, No=2 not applicable=9]	29	/
name of the drugs	<u> </u>	
other drugs [Yes=1, No=2 not applicable=9]	30	/
name of the drugs	_	
PART B: PHYSICAL FINDINGS	a -	
14. Dehydration	31	/
none=1, mild=2, moderate=3, severe=4 not applicable=9		
15.Rectal temperature	32-34	//
16.Signs of vitamin A deficiency		
Normal [Yes=1, No=2]	35	
Night blindness (XN) [Yes=1, No=2]	36	/
Conjunctival xerosis (X1A) [Yes=1, No=2]	37	/
Bitot's spot (X1B) [Yes=1, NO=2]	38	/
Corneal ulcer <1/3 (X3A) [Yes=1, NO=2]	39	/
Corneal ulcer >1/3 (X3B) [Yes=1, No=2]	40	/
Xerophthalmia scar (XS) [Yes=1, No=2]	41	/
Xerophthalmia fundus (XF) [Yes=1, No=2]	42	/
17.0titis media [Yes=1, No=2]	43	/

18.Throat		
normal	44	·/
[Yes=1, No=2]		
inflammed/tonsillitis	45	/
[Yes=1, No=2]		
19.Lungs		
clear	. 46	/
[Yes=1, No=2]	4.77	,
rhonchi [Yes=1, No=2]	47	/
crepitations	48	/ ·
[Yes=1, No=2]		** ** ** *
20 Matala (ba)	40.50	, , , ,
20.Weight (kg)	49-52	///
21.Length (cm)	53-55	//
22.Mid arm circumference (cm)	56-58	//
23.Skin fold thickness (mm)		
Triceps	59-6.1	//
Subscapular	62-64	//

1.Cohort No. Cohort I (watery diarrhoea)=1 Cohort II (bloody diarrhoea)=2	1	
2.Randomization WHO-ORS=1, Rice-ORS=2	2	
3.Study No.	3-6	//
STOOL MICROSCOPICAL EXAMINATION		·
4.Date of specimen received	7-12	////
5.Color yellow=1, brown=2, green=3, greenish=4, pale yellow=5, rice watery=6, creamy=7, other=8, not done=9	13	
6.Stool consistency liquid=1, watery=2, mucoid=3, bloody=4, bloody-mucoid=5, loose=6, soft=7, other=8, not done=9	14 •	
7.Blood none=0, trace=1, moderate=2, heavy=3	15	/
8.Mucus	16	
(same as above) 9.Worm	17	/
[Yes=1, No=2]		· · · · · · · · · · · · · · · · · · ·
10.pH [Acidic=1, Alkaline=2]	18	/
11.RBC	19	/
none=1, 1-10=2, 11-20=3, 21-50=4, 51+=5		
12.Pus cells less 10=2, 11-20=3, 21-50=4, 51+=5	20	
13. Macrophage	21	/
none=2, 1-5=3, 6-10=4, 11+=5		•
14.Neutral fat none=2, few=3, mod=4, many=5	22	/
15. Yeast	23	/
none=2, few=3, mod=4,		

16.Giardia none=0, cyst=1, trophozoite=2,	24	/
<pre>cyst + trophozoite=3 17.E.H. none=0, cyst=1, trophozoite=2,</pre>	25	·/
cyst + trophozoite=3		
18.Ascaris none=0, few=1, mod=2, many=3	26	
19.Trichuris	27	/
none=0, few=1, mod=2, many=3		,
20. Hookworm	28	,
none=0, few=1, mod=2,	20	/
many=3 21.S.Stercoralis	29	,
none=0, few=1, mod=2,	29	/
many=3		
22.Trichomonas hominis none=0, few=1, mod=2,	30	/
many=3		,
23.Other	31	/
[Yes=1, No=2]	~~	 '
24.Cryptosporidium [Yes=1, No=2]	32 -	/
STOOL BACTERIOLOGICAL EXAMINATION		4
STOOL BACTERIOLOGICAL EXAMINATION 25.Date of specimen sent	33-38	///
25.Date of specimen sent	33-38 39	
25.Date of specimen sent		
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2	39	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype	39	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2	39 40 41	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios	39 40	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4,	39 40 41	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8,	39 40 41 42	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9	39 40 41 42 43	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC	39 40 41 42	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2]	39 40 41 42 43	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC	39 40 41 42 43	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type	39 40 41 42 43	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3 35.EIEC	39 40 41 42 43	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3 35.EIEC [Yes=1, No=21] 36.EAEC	39 40 41 42 43 . 44 45	
25.Date of specimen sent 26.Vibrio cholerae [Yes=1, No=2] 28.Vibrio cholerae biotype El Tor=1, Classical=2 29.Vibrio cholerae serotype Inaba=1, Ogawa=2 30.Other vibrios [Yes=1, No=2] 31.Other vibrios type VP=1, VF=2, VM=3, PS=4, AS=5, AH=6, AC=7, other=8, not applicable=9 32.EPEC [Yes=1, No=2] 33.ETEC [Yes=1, No=2] 34.ETEC type ST=1, LT=2, LT/ST=3 35.EIEC [Yes=1, No=2]	39 40 41 42 43 . 44 45 46 47	

38.EHEC	50 .	/
[Yes=1, No=2]		
39.Klebsiella	51	/
[Yes=1, No=2]	•	•
40.Klebsiella type	52	/
LA=1, DA=2, AA=3 41.Rotavirus	53	,
[Yes=1, No=2]	55	<u>—</u> ./
42.Campylobacter	54	,
[Yes=1, No=2]	•	/
43.Campylobacter type	55	/
jejuni=1, coli=2, other=3		
44.Salmonellae	56	/
[Yes=1, No=2]		
45.Salmonellae type	57	/
salm. typhi=1		
salm. other=2	50	
46.Shigellae [Yes=1, No=2]	58	/
47.Shigellae type	59	,
sh. dyst I=1, sh. flex=2,	00	/
sh. boydii=3, sh. sonnei=4,		
sh. dyst 2=5.		
sh. dyst (3-10)=6	••	
48.Sensitivity pattern of:	60-61	/
VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99 49.Sensitive to:		
,		
Tetra	62	/
[Sen=1, Res=2]		
Ampi	63	/
[Sen=1, Res=2] TMP-SMX	64	,
[Sen=1, Res=2]	04	/
Furox	65	·/
[Sen=1, No=2]		
Chlor	66	/
[Sen=1, Res=2]		
Genta	67	/
[Sen=1, Res=2]	CO	,
Nalidixic acid [Sen=1, Res=2]	68	/
Sulpha	69	1
[Sen=1, Res=2]		<u></u> ′

		•
Pivmecillinum [Sen=1, Res=2]	70	·/
Other	71	,
[Sen=1, Res=2]	1.1	/
[
50. Sensitivity pattern of:	72-73	//
VC 01=01, VP=02, VF=03, VM=04, PS=05, AH=06, AS=07, AC=08, ETEC=09, Campy=10, Salm. typhi=11, Salm. other=12, Sh. dyst I=13, Sh. flex=14, Sh. sonnei=15, Sh. boydii=16, Sh. dyst 2=17, Sh. dyst (3-10)=18, Other=19, Not done=88, Not applicable=99		
51. Sensitive to:		
Tetra [Sen=1, Res=2]	74	/
Ampi	75	/
[Sen=1, Res=2]		
TMP-SMX	76 📥	/
[Sen=1, Res=2]		
Furox	77	/
[Sen=1, No=2] Chlor	T.O.	,
[Sen=1, Res=2]	78	/
Genta	79	,
[Sen=1, Res=2]	13	/
Nalidixic acid	80	/
[Sen=1, Res=2]	•	
Sulpha	81	/
[Sen=1, Res=2]		
Pivmecillinum	82 .	/
[Sen=1, Res=2]		
Other [Sen=1, Res=2]	83	/
(Ben-1, Nes-2)		
BLOOD EXAMINATION		
52.Date of specimen sent	84-89	///////
53.Total WBC		Y Y M M D D
% Polys	90-91	
% Bands	92-93	//
% Lymohocyte	94-95	//
% Monocyte	96-97	//

% Basophil	100-101	//
54. % HCT	102-103	//
55. Total Protein	104-106	//
56. Specific gravity	107-111	
57.Re ID No.	112-115	/ / / /