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Abstract

Objective—To define the benefit of rice oral rehydration salts solution in relation to the glucose based World Health Organisation oral rehydration salts solution for treating and preventing dehydration in patients with severe dehydrating diarrhoea.

Design—Meta-analysis using data from 13 available randomised trials that compared these two formulations.

Subjects—The studies compared 1367 patients with cholera, severe cholera-like diarrhoea, or acute non-cholera diarrhoea. 668 received the standard WHO solution and 699 the rice based solution.

Intervention—Each trial report was reviewed to determine patient eligibility, the number of patients who were randomised and the number of these included from analysis, details of the randomisation procedure, and the precise timing of the outcome

cholera diarrhoea and should be more precisely defined before its practical value can be judged.

Introduction

Oral rehydration therapy with the glucose and electrolyte solution recommended by the World Health Organisation and Unicef is the preferred method for treating children with dehydration due to diarrhoea, provided that they are able to drink and do not have signs of shock.¹ Although the solution is both safe and effective (D Mahalanabis, unpublished WHO document), it has important limitations: it neither reduces the rate of stool loss nor shortens the duration of illness.^{2,5} Mothers often do not understand the relation between diarrhoea and dehydration, and their primary concern, shared by many health workers, is to see the diarrhoea stop. This probably accounts for

first 24 hours; weighted estimates of the difference in mean stool output between treatments.

Results—The rice solution significantly reduced the rate of stool output during the first 24 hours by 36% (95% confidence interval 28 to 44%) in adults with cholera and by 32% (19 to 45%) in children with cholera. The rate of stool loss in infants and children with acute non-cholera diarrhoea was reduced by only 18% (6 to 30%).

Conclusions—The benefit of rice oral rehydration salts solution for patients with cholera is sufficiently great to warrant its use in such patients. The benefit is considerably smaller for children with acute, non-

instead of, or in addition to, oral rehydration salts solution (WHO diarrhoeal diseases control programme, seventh programme report, 1988-89, 1990).

If a packaged oral rehydration salts formulation could be developed that not only had the positive features of the WHO formulation, including low cost and safety and stability during prolonged storage, but also substantially reduced the duration of diarrhoea or the rate of stool loss, it would have considerable advantages. In particular, it could be promoted as having a real antidiarrhoeal effect. This should improve its acceptance and use by both health workers and mothers, especially if its benefits were sufficiently great



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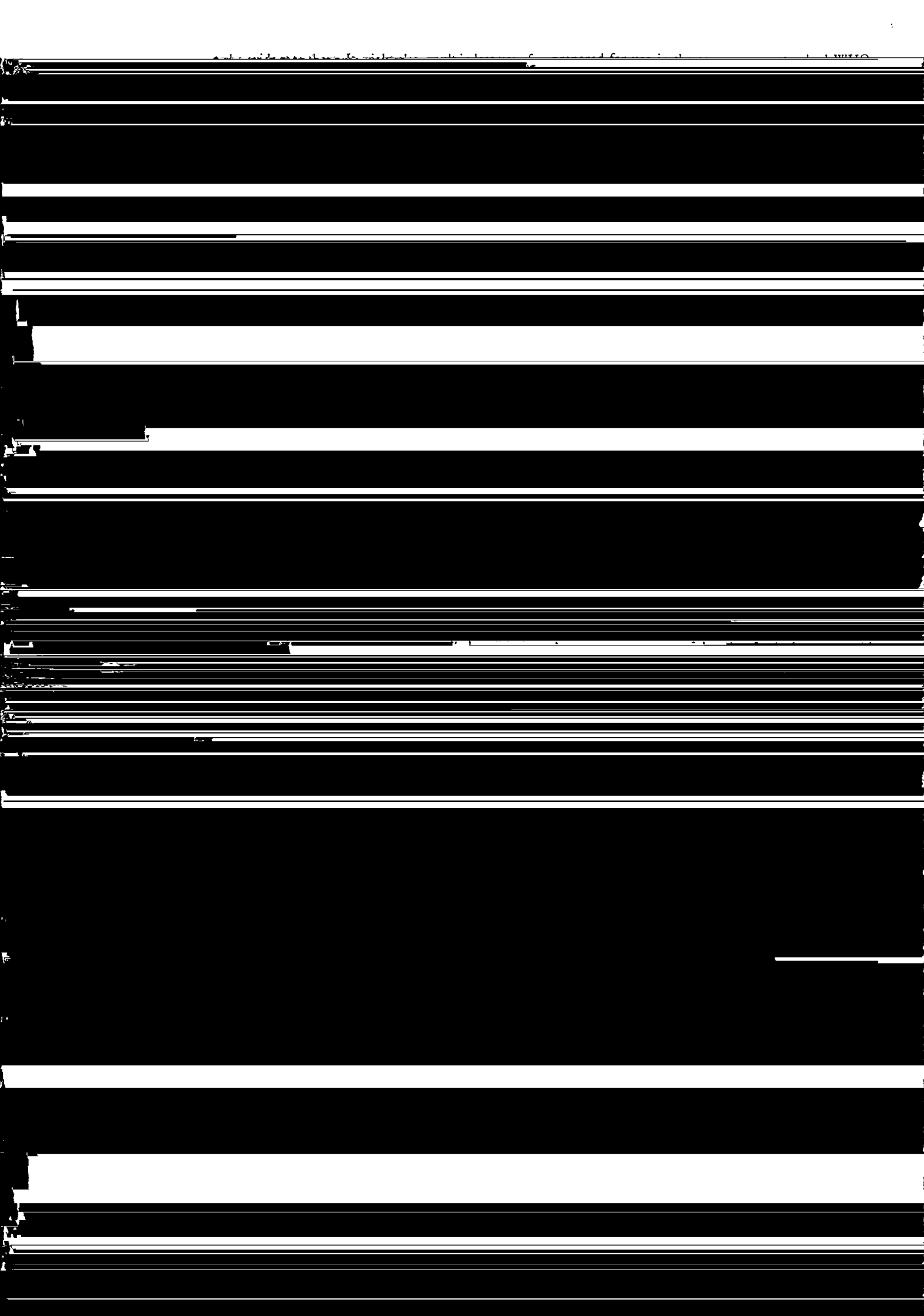


TABLE II—Stool output in first 24 hours in adults with diarrhoea associated with cholera or cholera-like illness

Comparison	No analysed taking WHO/rice solution	Treatment with WHO solution			Mean reduction (variance) in stool output during treatment with rice solution (g or ml/kg)	Study weight
		Mean (SD) stool output (g or ml/kg) in first 24 h	Mean stool output/SD	Mean stool output/mean intake of solution		
Moechtar <i>et al</i> (1) 6-24 hours	83/81	133 (92)	1.4	0.63	39 (148)	0.45
Moechtar <i>et al</i> (2) 6-24 hours	12/14	106 (55)	1.9	0.63	48 (407)	0.16
Alam <i>et al</i> (1)	47/46	391 (157)	2.5	0.75	164 (829)	0.08
Alam <i>et al</i> (2)	42/47	366 (174)	2.1	0.81	144 (1068)	0.06
Molla <i>et al</i> (1) ^a	72/85	159 (109)	1.5	0.64	44 (266)	0.25

TABLE III—Stool output in first 24 hours in children with diarrhoea associated with cholera or cholera-like illness

Comparison	No analysed taking WHO/rice solution	Treatment with WHO solution			Mean reduction (variance) in stool output during treatment with rice solution (g or ml/kg)	Study weight
		Mean (SD) stool output (g or ml/kg) in first 24 h	Mean stool output/SD	Mean stool output/mean intake of solution		
Molla <i>et al</i> (2) ^a	101/84	204 (140)	1.5	0.54	49 (362)	0.42
Molla <i>et al</i> (3) ^a	42/37	343 (151)	2.3	1.49	181 (628)	Zero
Molla <i>et al</i> (4) ^a	25/27	210 (158)	1.3	0.69	105 (1206)	0.12
Alam <i>et al</i> (3) ^{a*}	19/20	290 (190)	1.5	0.98	160 (2500)	0.06
Alam <i>et al</i> (4) ^{a*}	7/6	90 (75)	1.2	0.57	-40 (700)	0.21
Patra <i>et al</i> ^a	24/24	166 (114)	1.5	0.62	69 (794)	0.19

^aData not reported quantitatively; values in this table are approximated from graphic presentation.

TABLE IV—Stool output in first 24 hours in children with diarrhoea not associated with cholera

Comparison	No analysed taking WHO/rice solution	Treatment with WHO solution			Mean reduction (variance) in stool output during treatment with rice solution (g or ml/kg)	Study weight
		Mean (SD) stool output (g or ml/kg) in first 24 h	Mean stool output/SD	Mean stool output/mean intake of solution		
Guiraldes <i>et al</i>	48/49	126 (64)	2.0	0.30	14 (219)	0.21
Kenya <i>et al</i> ^a	49/50	103 (31)	3.3	0.48	3 (35)	Zero
Durta <i>et al</i> ^a	33/37	103 (55)	1.9	0.60	17 (145)	0.32
Bhan <i>et al</i> ^a	33/31	77 (58)	1.3	0.49	10 (161)	0.29
El Moughi <i>et al</i> ^a	26/25	245 (129) [*]	1.9	0.72	82 (1115)	0.04
Mohan <i>et al</i> ^a	23/23	110 (69)	1.6		22 (340)	0.14

^{*}The reported SD (25.3) was very low in relation to the large observed difference in stool output; we assumed that the reported SD values were actually SEs and revised the SDs accordingly.

immediately before treatment with oral rehydration salts solution began—that is, after the completion of any intravenous treatment for severe dehydration. However, in no trial was it stated when patients with severe dehydration were randomised and outcome measurements initiated—that is, before or after initial intravenous rehydration. Thus it was unclear whether the first 24 hour measurement of stool output began when intravenous rehydration was started or when oral rehydration salts solution was first given, as should have been the case.

In one study patients were randomised irrespective of age, but were stratified into arbitrary age groups during analysis.¹ Ideally, such stratification should have been part of the randomisation plan. Stratification during analysis was also done in two other studies (Moechtar *et al*),⁹ but this was based on aetiology and so was unavoidable.

Exclusion from analysis

Pragmatic analysis according to intention to treat requires that all randomised patients continue to be monitored and that their data be included in the analysis. Nevertheless, in seven trials (Guiraldes *et al*)^{6, 8, 9, 12-15} 1-15% of randomised patients were excluded from the analysis (table I), either because they were considered to be "treatment failures" (usually because additional intravenous treatment was required) or because they had been randomised in error. In two trials that used a permuted block¹⁰ or factorial design (Alam *et al*) it seems that some patients were randomised but not reported on, as the numbers specified in the different treatment groups differed appreciably. The reasons for these differences were not stated.

Analysis and internal consistency of outcome data

Whereas all studies reported stool output and oral rehydration salts solution intake during the first 24 hours, few reported total stool output until diarrhoea stopped, and only seven studies reported the duration of diarrhoea. Our analysis therefore focused largely on stool output during the first 24 hours. The following results for the first 24 hours are reported: mean (standard deviation) stool output (in g or ml/kg body weight) for patients randomised to WHO oral rehydration salts solution; the ratio of mean stool output to its standard deviation; the ratio of mean stool output to mean intake of WHO oral rehydration salts solution; and the mean reduction in stool output (in g or ml/kg) for patients given rice oral rehydration salts solution compared with those given WHO oral rehydration salts solution, and the variance of that value.

Tables II, III, and IV show the mean (SD) stool output (in g or ml/kg) during the first 24 hours for patients in each study who were randomised to receive WHO oral rehydration salts solution. Whether the data were for adults with cholera or with cholera-like diarrhoea (severe dehydrating diarrhoea, clinically resembling that associated with cholera but from which *Vibrio cholerae* 01 was not isolated) (table II), children with cholera or cholera-like diarrhoea (table III), or children with only acute non-cholera diarrhoea (table IV), the ratios of mean to standard deviation for stool output were roughly constant, averaging 1.6 and ranging (with one exception) from 1.2 to 2.5. This regularity indicates the need for logarithmic transformation; however, no study reported logarithmically transformed data or performed calculations on that scale. This finding also provides a criterion for judging the internal consistency of key outcome data. By this

criterion one trial seems to be atypical with a ratio of 3.3,¹⁴ twice the mean value reported in other studies. Even more extreme was the ratio of 10 from the data reported in another study.¹⁵ We suspected confusion between standard error and standard deviation in this study, and therefore table IV shows what we believe to

similar to those in the other studies.

Tables II, III, and IV also show a second measure by which to assess the internal consistency of trial data—namely, the ratio of mean stool output to mean intake of WHO oral rehydration solution. Mean stool output averages about two thirds of mean oral rehydration salts solution intake. By this criterion one trial seems to be atypical,¹⁰ the mean stool output being almost 50% greater than the mean intake of oral rehydration salts solution (table III). In the analyses that follow, two studies^{10,14} have been excluded (that is, zero weighted) for the reasons mentioned above.

SUBSTANTIVE RESULTS

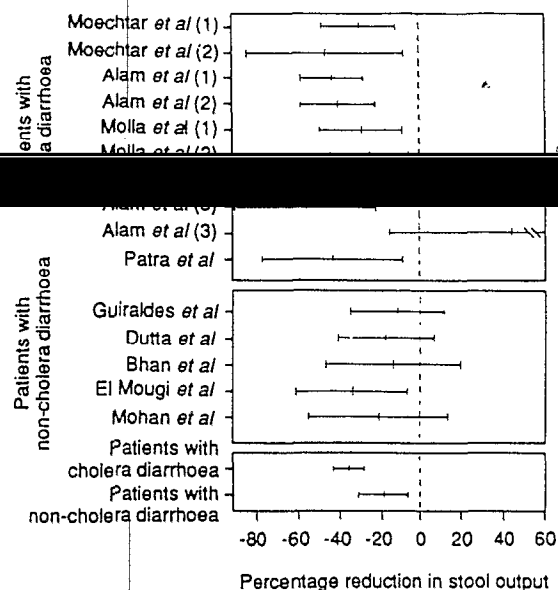
The results of the analysis of stool output and intake of oral rehydration salts solutions during the first 24 hours have been grouped into three sets according to the patient's age and aetiology of diarrhoea, as shown in tables II, III, and IV. For the duration of diarrhoea all data have been combined, but comparisons in patients with suspected cholera who received tetracycline before diarrhoea stopped have been zero

Adults with cholera or cholera-like diarrhoea—Table II shows the weights assigned to each of the five comparisons in this set. By using these weights the estimated mean stool output for patients given WHO oral rehydration salts solution was 170 ml/kg. For patients given rice oral rehydration salts solution this was reduced by a mean of 58 ml/kg (36%, 95% confidence interval 28 to 44%).

Children with cholera or cholera-like diarrhoea—Table III shows the weights assigned to each of the five comparisons in this set. One study¹⁵ was zero weighted for reasons described above. With those weights the estimated mean stool output for patients given the WHO oral solution was 178 ml/kg. For patients given the rice solution this was reduced by a mean of 48 ml/kg (32%, 19% to 45%).

Children with non-cholera diarrhoea—In this set of six comparisons, one study¹⁴ was zero weighted for reasons described above (table IV). By using the weights calculated for the other five comparisons the estimated mean stool output for patients given the WHO solution was 107 ml/kg. For patients given the rice solution this was reduced by a mean of 18 ml/kg (18%; 6% to 30%). It is noteworthy that the estimated mean percentage reduction in stool output associated with the rice solution in the zero weighted study is outside the 95% confidence interval derived from the other five studies.

Overall reduction in stool output—The figure presents the percentage reduction in mean stool output (with



Mean percentage reduction in 24 hour stool output in individual studies of adults and children with cholera or cholera-like diarrhoea and children with non-cholera diarrhoea given rice oral rehydration salts solution. Pooled (weighted) estimates of percentage reduction in mean stool output (95% confidence interval) for each group of studies are shown in shaded box.

95% confidence intervals) for patients treated with the

this overview, as well as the pooled (weighted) estimates of the percentage reduction in mean stool output for patients with cholera (adults and children) and without cholera (details of these calculations are not shown). The effect of the rice solution on stool output was significantly less in children with non-cholera diarrhoea than in children and adults with cholera or cholera-like diarrhoea (95% confidence interval 3% to 31% for the difference in percentage reduction in stool output in patients with cholera or cholera-like diarrhoea v patients with non-cholera diarrhoea).

Duration of diarrhoea—Data from six comparisons, including both adults and children with cholera (who had not received tetracycline before diarrhoea stopped) and acute non-cholera diarrhoea were considered for this analysis (table V). The estimated mean duration in patients given the WHO oral rehydration solution was 68 hours. For those given the rice solution the duration was reduced by a mean of eight hours (12%; 5% to

indicating a modest but significant reduction in the duration of diarrhoea.

Discussion

Irrespective of their age, patients with cholera who were given rice oral rehydration salts solution had substantially lower rates of stool loss than those who were given WHO oral rehydration salts solution. Stool volume was reduced by a mean of 48-58 ml/kg during

presumably reflects the fact that a greater amount of glucose (and amino acids) is released when rice powder is fully digested than is present in the WHO solution. Assuming that glucose facilitated absorption of sodium proceeds on an equimolar basis, 50-80 g/l of rice powder would release sufficient glucose and amino acids to promote the absorption of all the sodium (and water) in the rehydration solution and, in addition, reabsorption of at least part of the sodium (and water) secreted into the bowel as part of the diarrhoeal process, thus diminishing stool output.¹⁸ In contrast, the WHO solution contains only enough glucose (20 g/l) to promote the absorption of the sodium and water in

TABLE II. Duration of diarrhoea

Comparison	No analysed taking WHO/rice solution	Mean (SD) No of patients	Mean duration/SD	(variance) duration of diarrhoea with rice solution	Study weight
Adults with cholera					
Moechtar et al (1)	83/81	39 (11)	3.5	2 (3)	Zero
Moechtar et al (2)	12/14	36 (7)	5.1	7 (13)	Zero
Alam et al (1)	47/46	86 (22)	3.9	9 (26)	0.23
Alam et al (2)	42/47	85 (20)	4.2	4 (22)	0.27
Children with cholera					
Alam et al (3)*	24/24	90 (43)	2.1	12 (109)	0.05
Patra et al*	24/24	43 (22)	2.0	13 (36)	0.17
Children without cholera					
Kenya et al ¹⁴	49/50	46 (9)	5.1	4 (4)	Zero
Dutta et al ¹⁴	33/37	79 (37)	2.1	10 (67)	0.09
El Mougi et al ¹⁴	26/25	34 (12)	2.8	6 (31)	0.19

the solution, thus leaving the rate of stool loss essentially unaffected.¹⁹ The lower osmolarity of the rice solution (about 200 mmol/l *v* about 310 mmol/l) would also enhance the intestinal absorption of water, but not of sodium.²⁰

In contrast with stool output in cholera patients, that in children with acute non-cholera diarrhoea was

one litre packet about three times the cost of the standard packet of WHO oral rehydration salt packet. On the other hand, if the effect on total stool output is appreciably greater, owing to a concurrent shortening of illness, a change in oral rehydration salts formulation might be justified. This requires further study. In the meantime the current data show that rice oral rehy-