

BACTERIAL PATHOGENICITY

## Haemolysin produced by *Vibrio cholerae* non-O1 is not enterotoxigenic

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Of 28 isolates of *Vibrio cholerae* non-O1 (10 from diarrhoeal patients and 18 from environmental sources) examined for haemolytic activity and its correlation, if any, with enterotoxigenic activity, 24 showed haemolysis. The four non-haemolytic isolates showed haemolysis after consecutive passages through rabbit ileal loops (RILs). The titres of haemolytic activity were 4–64 HU/ml irrespective of their source. Eight (28.5%) of the non-O1 isolates caused fluid accumulation; six (25%) were haemolytic and two (50%) non-haemolytic. The remaining isolates showed enterotoxigenic activity after one-to-three consecutive passages through RILs irrespective of their haemolytic character and source. Environmental isolates caused significantly more fluid accumulation than the diarrhoeal isolates. All these isolates reverted to their original non-toxigenic character on repeated subculture or on storage in the laboratory, but continued to show haemolytic activity. The results of the present study indicate that *V. cholerae* non-O1 strains are potentially enterotoxigenic independent of their haemolytic character and source, and enterotoxin, not haemolysin, is the factor most likely to be responsible for their enterotoxigenic activity.

### Introduction

*Vibrio cholerae* non-O1 strains have been implicated in sporadic and localised outbreaks of cholera-like diarrhoea [1, 2], which is sometimes accompanied by fever, and blood and mucus in the stool. Apart from their diarrhoeagenic potential, these organisms have also been implicated in a few cases of extra-intestinal infections, such as wound infection, septicaemia and cellulitis [3–5]. A large number of *V. cholerae* non-O1 strains produce cholera-like toxins, although some strains do not [2, 6]. In addition to cholera toxin (CT), several other extracellular products such as the new cholera toxin [7], heat stable enterotoxin (NAG-ST), a thermostable direct haemolysin, shiga-like toxin and haemagglutinin have been reported to play a role in the disease process [8–14]. More recent reports suggest that the haemolysin produced by *V. cholerae* non-O1 is identical to *V. cholerae* biotype El Tor haemolysin [15, 16]. It has also been suggested that the El Tor-like haemolysin of *V. cholerae* non-O1 can cause fluid accumulation mixed with mucous and blood in rabbit ileal loops (RIL) [17]. Alm *et al.* [18] indicated that the haemolysin may be enterotoxigenic because fluid accumulation in RILs was not observed when the whole determinant was deleted from the chromosome of a CT gene-negative but diarrhoea-

producing strain of El Tor biotype. However, earlier observations made in this laboratory indicate that El Tor-like haemolysin is not responsible for fluid outpouring in RILs because non-haemolytic strains of *V. cholerae* biotype El Tor caused fluid accumulation [19]. These reports caused confusion about the nature of the enterotoxin in *V. cholerae* non-O1. Therefore, 28 diarrhoeal and environmental isolates of *V. cholerae* non-O1 were examined for haemolytic and enterotoxigenic activities, with particular interest in any change in enterotoxigenicity after passage through RILs and any correlation with haemolytic activity.

### Materials and methods

#### Bacterial strains

Twenty-eight strains of *V. cholerae* non-O1 isolated from diarrhoeal stools (10) and from environmental sources, such as the River Ganga (15) and sewage (3) were included in the study. The strains were identified by the method recommended by WHO [20]. The strains were maintained in peptone agar stab cultures at room temperature and did not undergo more than three subcultures before the experiments.

#### Detection of haemolysis

Preliminary testing for haemolysis by *V. cholerae* non-O1 strains was performed by inoculating 4–5-h growths of each organism in Brain Heart Infusion Broth (BHIB,

Difco) on to sheep blood 5% v/v agar. After incubation for 24 h at 37°C, the blood agar plates were examined for haemolysis around the colonies.

#### Preparation of culture filtrates for haemolysin production

Culture filtrates (CFs) of *V. cholerae* non-O1 strains shown to be haemolytic were prepared by the method of Richardson *et al.* [21]. Briefly, 10 ml of BHIB contained in a 50-ml conical flask were inoculated with five or six colonies from an overnight culture on blood agar plates. The flasks were incubated at 37°C in a shaking water bath for 16–18 h with 80–120 oscillations/min. The cultures were centrifuged at 22 000 *g* for 20 min at 4°C, supernates were filtered through a membrane filter (Millipore, 0.22 µm) and stored at 4°C. These CFs were used for haemolysin assay.

#### Preparation of culture filtrates for enterotoxin production

CFs of *V. cholerae* non-O1 strains that gave positive ileal loop reactions were prepared in AKI medium by the method described above. The AKI medium (Bacto-peptone 1.5% w/v, yeast extract 0.4%, NaCl 0.5% and NaHCO<sub>3</sub> 0.3%) was prepared as described by Iwanaga *et al.* [22]. The sodium bicarbonate was filter sterilised separately and mixed with the other autoclaved ingredients. Freshly prepared medium was always used; the pH of this medium was *c.* 7.4 without adjustment.

#### Titration of haemolysin

The production of haemolysin by each strain of *V. cholerae* non-O1 was confirmed by the method of Smith [23] as modified by Rennie and Arbuthnott [24] and as described by Singh and Sanyal [25]. Briefly, sheep erythrocytes (SRBC) were washed three times in isotonic saline and a 2% suspension was prepared in 0.04 M phosphate-buffered saline (PBS, pH 7.4). Haemolytic activity was determined by mixing 0.5 ml of two-fold serial dilutions of CF of each strain with an equal volume of SRBC 2% suspension. After incubation at 37°C in a water-bath for 2 h and standing at 4°C

for 12 h, the lysed portion was diluted four-fold with sterile normal saline and the optical density was measured in a colorimeter. Standardisation of erythrocyte suspension was by lysis of 0.5 ml of SRBC 2% with a few crystals of saponin. An optical density of 0.5 at 540 nm was considered to indicate a standardised SRBC 2%. The negative control was 0.5 ml of saline instead of CF. One haemolytic unit (HU) was defined as the amount of CF that caused 50% haemolysis under experimental conditions.

#### Ileal loop test

Live cells and CFs of all the strains of *V. cholerae* non-O1 were tested for enterotoxin production in adult albino rabbits by the method of De and Chatterjee [26]. Briefly, bacteria grown in peptone water for 3 h were diluted 10-fold in the same medium and 1 ml (containing 10<sup>5</sup>–10<sup>6</sup> cfu) was inoculated into the RIL. A peptone water culture of toxigenic strain 569B of *V. cholerae* and unseeded peptone water served as positive and negative controls, respectively. CFs (1 ml, prepared in AKI medium) were also tested in the same way. Each test was done in two rabbits. Rabbits were killed after 8 h.

#### Passage through RIL

Strains of *V. cholerae* non-O1 that caused little or no fluid accumulation in the initial tests were passaged through RILs according to the method of Sanyal *et al.* [27, 28]. Briefly, each strain was cultured aseptically from RIL on nutrient agar and incubated overnight; five or six colonies were inoculated into peptone water and incubated for 3 h; and 1 ml of diluted culture was inoculated again into a RIL. The process was continued until a positive response was obtained.

## Results

Twenty-four of the 28 isolates of *V. cholerae* non-O1 tested produced haemolysis on sheep blood 5% agar plates (Table 1). CFs of these isolates also caused lysis of sheep erythrocytes in 2% suspension. The remaining

**Table 1.** Haemolytic activity of *V. cholerae* non-O1 strains before and after passage through RILs\*

Source and haemolytic character	Before passage		After passage	
	Number tested	Haemolysin production (HU/ml)	Number tested	Haemolysin production (HU/ml)
Diarrhoeal				
Haemolytic	8	8–64	0	...
Non-haemolytic	2	ND	2	8–16
Environmental				
Haemolytic	16	8–64	0	...
Non-haemolytic	2	ND	2	8–32

ND, not detected.

\*Non-haemolytic strains tested only after animal passage.

four isolates became haemolytic after one-to-three consecutive passages through RILs. The titres of haemolytic activity were 4–64 HU/ml. No difference in the titres of haemolytic activity was observed in relation to the source of the strains. However, there was strain-to-strain variation in the titres of haemolysin produced (Table 1).

In the initial tests with *V. cholerae* non-O1 in RILs, live cells of eight of the 28 isolates caused fluid accumulation; six of them were haemolytic and two were non-haemolytic. The remaining haemolytic strains did not cause any fluid outpouring. CFs of the RIL-positive strains caused a similar secretory response. With organisms from both sources, there were strain variations in the volume of fluid accumulation, and also variation between individual loops inoculated with the same strain (Table 2).

Environmental *V. cholerae* non-O1 strains caused significantly (Students *t* test,  $p < 0.05$ ) more fluid accumulation than the diarrhoeal isolates regardless of their haemolytic character (Fig. 1).

After one-to-three consecutive passages through RILs, 18 (75%) of the 24 haemolytic and two (50%) of the four non-haemolytic *V. cholerae* strains that caused little or no accumulation of fluid in the initial experiments did so (Table 2).

Two non-haemolytic and 13 haemolytic strains of *V. cholerae* non-O1 tested by consecutive passages in RILs, caused marked increase in fluid accumulation after each passage (Table 3). The titres of haemolysin produced also increased after each passage. However, on repeated subcultures or on storage in the laboratory, all these strains reverted to their original haemolytic but non-toxigenic character. The three non-toxigenic isolates of environmental origin that caused no fluid accumulation even after four consecutive passages through RILs, showed lysis of sheep erythrocytes 2% suspension (Table 3).

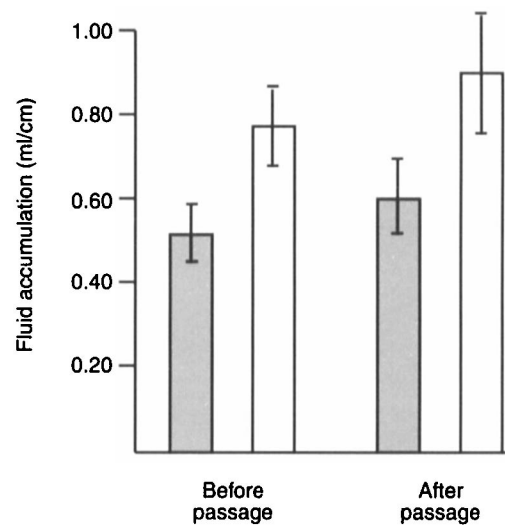


Fig. 1. Enterotoxicity of diarrhoeal (▨) and environmental (□) isolates of *V. cholerae* non-O1 strains before and after passage through RILs. The bars on the columns indicate the range of fluid accumulation caused by different strains.

### Discussion

The majority of strains of *V. cholerae* non-O1 showed haemolysis on sheep blood 5% agar plates in the initial tests. However, a few of the non-haemolytic strains became haemolytic after one-to-three consecutive passages through RILs. Moreover, the titres of haemolysin in CFs also increased after consecutive passages. This phenomenon may be explained by the genetic evidence that the sequences homologous to El Tor haemolysin gene are present in both haemolytic and non-haemolytic strains of *V. cholerae* non-O1 but are probably not expressed in non-haemolytic isolates [29, 30].

Several workers suggested that El Tor-like haemolysin produced by *V. cholerae* non-O1 strains may be responsible for their enterotoxic activity [15–17].

Table 2. Haemolytic activity and enterotoxicity of *V. cholerae* non-O1 strains

Source and haemolytic character	Number of strains tested	Before passage		After passage*	
		Number showing fluid accumulation	Fluid accumulation (ml/cm of RIL)	Number showing fluid accumulation	Fluid accumulation (ml/cm of RIL)
Diarrhoeal					
Haemolytic	8	2	0.46–0.6	6	0.42–0.8
Non-haemolytic	2	1	0.44–0.6	1	ND
Environmental					
Haemolytic	16	4	0.60–1.0	9	0.66–1.2
Non-haemolytic	2	1	0.60–0.8	1	ND
Positive control†	1	1	0.90–1.2		1.0–1.4
Negative control‡	0	0	0		0

ND, not determined.

\*These strains were tested only after animal passage.

†Peptone water culture of *V. cholerae* 569B inoculated into RIL of each animal.

‡Peptone water.

**Table 3.** Influence of RIL passage on haemolytic character, fluid accumulation and haemolysis titre of *V. cholerae* non-O1 strains

Strain no.	Haemolytic character	Fluid accumulation (ml/cm) and haemolysis titre (HU/ml) after passage no.									
		0		1		2		3		4	
		ml/cm	HU/ml	ml/cm	HU/ml	ml/cm	HU/ml	ml/cm	HU/ml	ml/cm	HU/ml
D-12	Haemolytic	*ND	64	0.46	†64	0.60	NT	NT	NT	NT	NT
D-13	Haemolytic	*ND	8	0.60	†16	0.72	NT	NT	NT	NT	NT
D-12475 B	Haemolytic	*ND	32	ND	32	0.58	†64	NT	NT	NT	NT
D-13094 RAN	Haemolytic	*0.7	†16	0.86	64	NT	NT	NT	NT	NT	NT
D-12158 RAN	Non-haemolytic	0.44	†ND	*0.68	8	0.76	32	NT	NT	NT	NT
E-RP-86	Haemolytic	*ND	8	ND	16	0.80	†64	0.90	NT	NT	NT
E-DSM-86	Haemolytic	*0.8	†64	1.0	128	1.2	256	1.4	NT	NT	NT
E-AS-88	Haemolytic	*ND	64	ND	64	ND	128	0.68	†256	NT	NT
E-SH-88	Haemolytic	*ND	32	0.66	†64	0.90	128	1.1	128	NT	NT
E-TG2-86	Haemolytic	*ND	16	ND	32	ND	32	0.70	†64	0.90	NT
E-HG-86	Haemolytic	*ND	64	ND	64	ND	128	0.70	†256	0.9	NT
E-CH-86	Haemolytic	*ND	32	ND	64	ND	NT	ND	NT	ND	NT
E-HGS-86	Haemolytic	*ND	8	ND	16	ND	32	ND	NT	ND	NT
E-SHS-86	Haemolytic	*ND	16	ND	32	ND	32	ND	64	ND	NT
E-TG1-86	Non-haemolytic	ND	ND	*ND	16	0.50	†32	0.66	64	NT	NT

ND, not detected; NT, not tested; D, diarrhoeal; E, environmental.

\*Change to haemolysis detected; †enterotoxin produced.

However, the observation that more than two-thirds of the haemolytic strains were unable to produce a secretory response in the initial tests indicates that haemolysis may not be responsible for enterotoxic activity. This observation is further strengthened by the fact that three haemolytic strains did not cause accumulation of fluid even after four consecutive passages, although they showed enhanced haemolytic activity. Moreover, the observation that 50% of the non-haemolytic strains showed enterotoxic activity also supports this conclusion. All these data suggest that the secretory response without any blood and mucous observed in RILs is most probably due to the enterotoxin(s) produced by the organism.

The environmental strains of *V. cholerae* non-O1 caused significantly more fluid accumulation than the diarrhoeal isolates. A probable explanation of this difference in the degree of enterotoxicity may be the observation, made in this laboratory that, that compared to diarrhoeal isolates, strains of environmental origin produced quantitatively more constitutive chitinase. Some strains even elaborated inducible chitinase and both these properties correlated with greater enterotoxicity (unpublished data). Similar observations were also made in relation to *Aeromonas* spp. of clinical and environmental origin [31].

The strains of *V. cholerae* non-O1 that failed to produce enterotoxin in the initial tests caused fluid accumulation after one-to-three consecutive passages through RILs, suggesting that almost all the strains of *V. cholerae* non-O1 were potentially enterotoxigenic. Similar observations on the effect of passage through the gut of a susceptible host were made in earlier studies with *V. cholerae* non-O1 [2, 32] and

also with many other organisms, such as *V. cholerae* O1 [27, 28], *V. fluvialis* [33], *V. mimicus* [27, 34], *Aeromonas* spp. [35–38], *Plesiomonas shigelloides* [39] and *Klebsiella pneumoniae* [40]. Furthermore, it is also known that the so-called hypertoxigenic strain 569B of *V. cholerae* O1 also requires passage through the rabbit gut for enhancement of toxin production. A mechanism of repression and derepression of the toxin gene may account for this phenomenon. It is possible that the toxigenicity of the fresh isolates in this study might have decreased during one-to-three subcultures *in vitro* before the ileal loop tests. It appears that expression of toxin genes in a bacterial strain is a means of adaptation to a particular micro-environment as occurs in *V. cholerae* [41].

The observation that the majority of strains were non-toxigenic but haemolytic in the initial tests indicates that there is no correlation between these two properties. Furthermore, the three haemolytic strains of *V. cholerae* non-O1 that failed to cause accumulation of fluid after four consecutive passages through RILs also confirms this conclusion.

The present study indicates that strains of *V. cholerae* non-O1, irrespective of their haemolytic character and sources, are potentially enterotoxigenic and that enterotoxin, not haemolysin, is the factor most likely to be responsible for their enterotoxic activity. Passage through the gut of a susceptible host may control the expression of the gene responsible for haemolysin and enterotoxin production.

The study was supported in part by the Council of Scientific and Industrial Research, New Delhi as a fellowship to D.V.S. The authors are grateful to Mr D. Venu Gopal for secretarial assistance and Ms Anjali Tikoo for technical assistance during the study.

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