Non-gastrointestinal *Bacillus cereus* infections: an analysis of exotoxin production by strains isolated over a two-year period

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SUMMARY Isolates of *Bacillus cereus* from 118 cases, and two maternity unit outbreaks, of non-gastrointestinal infection were grouped on the basis of their estimated probable involvement in the infections from which they were isolated: (i) high probability—48 strains; (ii) intermediate—16 strains; (iii) low—7 strains; (iv) very low ("irrelevant")—49 strains.

Rabbit skin test, haemolysin and phospholipase assays were used to determine exotoxin activities of strains within each group.

The results suggest a significant relation between the virulence of an isolate as reflected in the degree to which it appeared responsible for the signs and symptoms of an infection, and its toxigenicity in the skin test. This is attributed to the ability of *B cereus* strains to synthesise, in varying degrees, a necrotic enterotoxin, possibly in conjunction with the primary haemolysin (cereolysin).

The cases analysed in this study support the contention that *B cereus*, when isolated from an infection, may not be an inconsequential contaminant and should not be too readily dismissed as such.

The number of reports of infections caused by *Bacillus cereus* continues to increase.¹⁻⁴ They comprise a wide range of clinical conditions, often severe in nature and occasionally fatal. It has been proposed² that an exotoxin, studied initially in relation to diarrhoeal-syndrome *B cereus* food poisoning,^{5,6} is a principal determinant of virulence in non-gastrointestinal *B cereus* infections. This toxin has been partially purified and characterised,⁷ and shown to exhibit the properties of a true enterotoxin.⁸

Injected intradermally into rabbits the toxin produces increased vascular permeability and necrosis to a degree proportional to its concentration. However this test is not entirely specific for the enterotoxin: the *B cereus* haemolysin, cereolysin, can also alter capillary permeability and cause similar dermal necrosis. In order to assess the contributory roles of these two exotoxins to the signs and symptoms of an infection, both the skin test and

haemolytic activities of a clinical isolate must be determined.

Almost all *B cereus* strains elaborate an extracellular phospholipase-C (responsible for the diagnostic lecithinase reaction). Early beliefs that this metabolite might have a pathogenic role, possibly analogous to that of *Clostridium perfringens* α -toxin (which also has lecithinase activity), are now in doubt.³

The results of a previous analysis of the exotoxin activities of isolates from 21 human cases of *B cereus* non-gastrointestinal infection and three cases of bovine mastitis indicated that only the vascular permeability reaction/necrotic toxin appeared to be related to the severity of the infection.² The substantially larger number of isolates now being referred to this laboratory has enabled this more extensive investigation into the relation between exotoxin production and virulence to be undertaken.

Cases and methods

TYPES OF INFECTION FROM WHICH CULTURES WERE RECEIVED

Isolates of *B. cereus* were received from a total of 136 cases encompassing a broad spectrum of infections: traumatic (postoperative or casualty) wounds and drain sites—74 cases (three deaths) including 10 described as necrotic or gangrenous; panophthalmitis—11 cases; bacteraemia/septicaemia—7 cases (excluding those placed within other categories such as wound or central nervous system infections with bacteraemia); abscesses and pustules—7 cases; peritonitis or ascites—7 cases; respiratory tract infections—6 cases (three deaths); burns—3 cases (one death); central nervous system infections—3 cases (two deaths); “sticky” eyes in infants—3 cases; infected umbilici in infants—3 cases; ulcers—2 cases; cot death—1 case. Two maternity unit outbreaks^{9,10} accounted for a further 6 and 3 cases respectively: six of the affected infants had infected umbilical areas and three had upper respiratory tract infections.

ASSESSMENT OF RELEVANCE

Bacillus spp are commonly encountered in clinical laboratory cultures and tend to be regarded as environmental contaminants. It was therefore necessary to assess the relevance of the *B. cereus* isolates in the cases listed above. Specific details considered important to the assessment (Table 1) were sought from those clinicians and microbiologists directly involved in each case. Greater significance was attached to multiple isolations, particularly from blood cultures. Where *B. cereus* was not the only isolate recovered, attempts were also made to determine the proportions and types of other organisms present.

Fig. 1 illustrates how an assessment of relevance was made. At one extreme (upper left) a scanty *B. cereus* isolation in mixed culture from a patient showing no clinical evidence of infection was considered to be “irrelevant”. At the opposite extreme (lower right) isolation of a heavy growth of the

Fig. 1 Interrelation of principal factors used to assess the relevance of *Bacillus cereus* to an infection.

organism in pure culture from an obviously infected site, with the patient in a toxic state, was regarded as being “relevant”.

Many isolates, as expected, fell between these two extremes. So, for the purposes of analysing the importance of exotoxin production to the clinical condition apparent, strains were placed into one of four categories of probable relevance.

Sufficient information was available on 105 of the 136 cases to permit classification in this way. Serological typing and consistent toxin profiles indicated that the two outbreaks of non-gastrointestinal *B. cereus* infection could be represented by just two strains.

A further 21 cases of human infection previously reported² were re-examined under the new criteria of assessment: the information on 13 of these was judged adequate to permit their inclusion in this study. The analysis was therefore carried out on a total of 120 strains.

ASSAYS OF SKIN TEST, HAEMOLYTIC AND PHOSPHOLIPOLYTIC ACTIVITIES

The methods used were essentially as described previously.^{2,7} For toxin production, 18 h nutrient broth cultures of the test strains were used to inoculate 10 ml volumes of brain heart infusion broth (Difco) + 0.1% wt/vol glucose in 250 ml Erlenmeyer flasks. These were incubated with shaking (200 cycles/min) for 6–7 h at 36°C, pH control being by addition of 0.1 ml 1 N NaOH to each culture at 3–3.5 h and again at 4–4.5 h. Bacterial cells were removed by centrifugation and filtration through 0.45 µm membrane filters (Millipore Ltd, London).

Table 1 Criteria used to assess the relevance of *Bacillus cereus* to an infection

1	Isolation of <i>B. cereus</i>	—extent present (+ to +++) —in pure/mixed culture —on one/more than one occasion
2	Isolation of other organisms—identities and types (if known)	—relative proportions (%)
3	Nature and condition of infection site	
4	Patient response (for example: pyrexia, toxic reaction)	
5	Factors predisposing to infection	
6	Relevant chemotherapy	

For the rabbit skin test assay, 0.05 ml of each cell-free culture filtrate (CCF) was injected intradermally in duplicate into the depilated backs of two adult rabbits. At 3 h postinoculation, 4 ml of 2% wt/vol Evans Blue (Sigma, London) was injected intravenously; measurements of zones of blueing and necrosis were made after a further hour. Necrotic activity, when present, was rechecked at 24 h. In previous studies,^{2,3,7} skin test readings were placed into five categories ranging from very weak (category 1) to very strong (category 5) toxin production. However, it has since proved more practical to use just three groups, designated "weak" (formerly categories 1 and 2), "intermediate" (formerly category 3) and "strong" (formerly categories 4 and 5).

Haemolysin titrations were carried out in WHO plates using twofold serial dilutions of CCF incubated for 2 h at 36°C with 0.5% vol/vol washed rabbit erythrocytes suspended in 0.01 M phosphate buffered saline. Test strains were divided into four groups on the basis of their reciprocal haemolysin titres: <10, 20–80, 160–640, and >640.

Phospholipase activities were determined as the mean diameters of zones of opacity produced (after 20 h at 36°C) in a 10% vol/vol egg-yolk emulsion agar around duplicate 5 mm wells containing 50 µl CCF. Strains were allocated to one of three categories according to their phospholipolytic activity: <10 mm, 10–20 mm, and >20 mm.

Results

RELEVANCE OF ISOLATES

The 120 strains included in this study were grouped according to their probable relevance to the infections from which they were isolated, as follows: (i) high probability—48 strains; (ii) intermediate—16 strains; (iii) low—7 strains; (iv) very low (of no clinical significance)—49 strains.

With reference to the 10 patients who died it was considered that *B. cereus* infection was a contributory factor in the deaths of seven (intermediate probability of relevance in six; low probability in one), and that in a further two cases (including the cot death) the organism probably did not have any role. The significance of the *B. cereus* isolation in the remaining case could not be assessed.

EXOTOXIN PRODUCTION VERSUS RELEVANCE

The histograms (Figs. 2, 3 and 4) show the relations found between the rabbit skin test, haemolytic and phospholipolytic activities of the strains and the estimated relevance to their respective infections. For further comparison, the results of tests on environmental isolates accumulated over some years

were also included. These strains were isolated from dairy products, meat and poultry, hospital environments, animal feedstuffs and other miscellaneous sources.

In Fig. 2 a marked bias is apparent towards strong skin test activity among the strains assessed as having some degree of clinical relevance, particularly those in the "high" or "intermediate" probability categories. In contrast, strains assessed as being of "very low" relevance show a bias towards weak skin test activity. Furthermore, an approximately normal distribution of toxigenicity was found among the 68 environmental isolates.

In Fig. 3, haemolysin production by the same clinical isolates is analysed. Haemolysin titrations were only carried out on 27 of the environmental isolates. As with the skin test activity, there is a notable bias towards high haemolysin titres among those isolates considered to have some degree of clinical relevance, although the contrast between the "relevant" and the "irrelevant" strains is not as marked as in Fig. 2. Nonetheless, haemolysin production by the "irrelevant" strains lacked any bias towards high titres and had a distribution comparable with that of the environmental isolates.

From a similar analysis of phospholipase production (Fig. 4) a bias towards higher enzyme activity

Fig. 2 Skin test toxigenicity of (a) 120 clinical isolates of *Bacillus cereus* and (b) 68 environmental isolates.

Fig. 3 Haemolysin production by (a) 120 clinical isolates of *Bacillus cereus* and (b) 27 environmental isolates.

was shown equally by strains of *B. cereus* from all sources.

Without supplying absolute proof, the results presented in Fig. 2 indicate a correlation between the

Fig. 4 Phospholipase production by (a) 120 clinical isolates of *Bacillus cereus* and (b) 27 environmental isolates.

degree of reaction obtained with the rabbit skin test and the reported manifestation of symptoms.

Support for this conclusion seems to lie in the contrasting absence of any apparent association between phospholipase production and assessed relevance of a strain to its associated infection (Fig. 4); there is no obvious difference between strains deemed to have caused or contributed to symptoms, those considered irrelevant and the environmental strains.

Conclusions with respect to haemolysin production (Fig. 3) are less easy to make. A bias towards high titres is an apparent characteristic of those strains judged relevant to some degree. This bias is essentially absent among the "irrelevant" strains. However, a contrasting bias towards low skin test activity by the latter was not apparent either, and distribution curves for the "irrelevant" and environmental strains were both similar. Possible interpretations of the haemolysin results are discussed below.

ANTIMICROBIAL SENSITIVITIES

The results of in vitro antimicrobial sensitivity tests, carried out in all instances by disc diffusion, were supplied with 68 of the clinical isolates. These data are summarised in Table 2.

Discussion

The cases analysed in this study in which *B. cereus* was considered either contributory or causative, substantially enhances the increasing evidence that this bacterium should be viewed more seriously as a potential pathogen and not be too readily dismissed as a saprophyte.

Underlying disease or debilitation did not feature greatly in the histories obtained on these cases: seven patients were under treatment for malignancies, one was on steroid therapy and one a diabetic. Apart from two premature infants, the remainder of

Table 2 *In vitro* antimicrobial sensitivities of 68 strains of *Bacillus cereus* isolated from non-gastrointestinal infections

Antimicrobial		No of strains		
		Sensitive	Intermediate	Resistant
Cefuroxime	30 µg	0	1	24
Cephalexin	25 µg	1	37	2
Chloramphenicol	10 µg	27	0	0
Erythromycin	5 µg	59	7	0
Gentamicin	10 µg	67	1	0
Methicillin	10 µg	8	1	17
Penicillin G	1 unit	1	4	61
Tetracycline	10 µg	49	7	9
Trimethoprim	2.5 µg	0	0	26
Sulphonamide	50 µg	23	1	2

the patients appeared to have been in good health prior to the event (accident, burns, surgery etc.) which led to the *B. cereus* infection.

As a clinical entity, the proportion of cases of panophthalmitis stands out noticeably. The importance of *B. cereus* as a cause of panophthalmitis has been discussed elsewhere.¹¹

The seven deaths associated with *B. cereus* infections included four cases of respiratory tract involvement, two cases (one a premature neonate) of central nervous system involvement and one (also a premature neonate) involving both the respiratory tract and central nervous system. *B. cereus* is known to produce a "lethal toxin" the nature of which is becoming clearer,³ and the lethal potential of this organism has been observed in both naturally¹² and experimentally^{13,14} infected animals.

Age and predisposition to infection did not appear related. In the two maternity unit outbreaks, the infections in the first⁹ were mild; in the second,¹⁰ only three of 28 infants from which the common serotype was isolated exhibited clinical signs, and from these *Staphylococcus aureus* was also isolated. Of the remaining cases in which the *B. cereus* isolations were considered relevant, a further two were infants and one was recorded as elderly.

With reference to possible modes of *B. cereus* pathogenesis, the results support previous contentions^{2,3,7,14} that there is a significant relation between the skin test toxigenicity of a strain and the extent of clinical manifestation of the infection.

The nature of the skin test reaction has been reviewed extensively.³ The response in the skin of rabbits or guinea pigs to intradermal injection of crude *B. cereus* culture filtrates is attributable to a combination of two distinct factors: (i) a partially characterised necrotic enterotoxin of a molecular weight of approximately 50 000 daltons and (ii) the primary *B. cereus* haemolysin, cereolysin. Their individual roles can, as yet, only be assessed indirectly by testing separately the haemolytic activity of the culture filtrate as was done here. In its purified form the enterotoxin has little or no inherent in vitro haemolytic activity,⁷ though this is something of a paradox since it possesses strongly necrotic and cytotoxic properties. Furthermore, certain strains at least, are known to produce a secondary (minor) haemolysin. For these reasons in vitro haemolysis tests as used here are not entirely satisfactory and alternative immunological assays are awaited.

The finding of a similar, though less marked, correlation between the haemolytic activity of an isolate and severity of infection suggests that haemolysin production is also involved in *B. cereus* pathogenicity. However some qualification is necessary since cereolysin activity has been shown to be neutralised

in vitro by normal serum.³ The manner and extent, if any, by which this occurs in vivo is not yet known.

The results also indicate that phospholipase-C is probably not of direct pathological significance *per se* in *B. cereus* infections. There is some evidence³ to suggest that phospholipase-C might play an auxiliary pathogenic role by breaking-down host-cell membrane phospholipids exposed by the action of other enzymes or toxins. The contribution of "secondary virulence factors" would not necessarily be apparent in a study of this kind.

An incidental observation made during this investigation was the distribution of serotypes among the *B. cereus* isolates: 13.5% of strains were serotype 20, a further 25.6% fell into 18 different serogroups, and 60.9% were not typable. The reason for the prevalence of type 20 among human isolates is not clear, but a similar phenomenon has been noted in investigations of *B. cereus* emetic-syndrome food poisoning, where 70% of isolates were type 1,¹⁵ and *B. cereus*-induced bovine mastitis, where strains of serotype 12 were predominant.¹² The unusual degree of heat resistance exhibited by type 1 spores¹⁶ may account for the frequent association of this serotype with food poisoning, but no analogous explanation has yet been found for the disproportionate representation of types 20 and 12 in human and veterinary isolations.

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