An epidemic of gastroenteritis due to an uncommon variant of *Escherichia coli* 0.128

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**Synopsis** A small epidemic of gastroenteritis in a children’s hospital is described, and also the biochemical characteristics of the causative organism. Stress is laid on the fact that failure to recognize that epidemic strains of *Escherichia coli* may be late lactose fermentors can lead to errors or delay in diagnosis.

The role of certain enteropathic (epidemic) strains of *Escherichia coli* in the aetiology of infantile gastroenteritis is now well established. The subject has recently been reviewed by Neter (1959, 1960) and by Rogers and Taylor (1961). Adam (1923) suggested that these strains might be recognized by their biochemical properties and was able to isolate them from the faeces of children suffering from gastroenteritis. Bray (1945), who investigated 44 cases of this condition, used serological methods for the identification of the strains which he isolated. From 42 cases he isolated the same serotype which he named *E. coli* var *neapolitanum*. This organism was later classified serologically by Kauffmann (1947) as *E. coli* 0.111. Bray noted that 34 of the strains which he isolated fermented lactose whereas eight did not. The eight non-lactose fermenting strains were agglutinated to the end titre by an antiserum prepared against a standard lactose-fermenting strain. Later the same serological type was isolated by numerous workers, including Taylor, Powell, and Wright (1949), who studied eight outbreaks of gastroenteritis in children, involving a total of 116 cases. They isolated 178 epidemic *E. coli* strains, all of which fermented lactose within 24 hours.

The identification of epidemic *E. coli* strains in a busy laboratory is difficult as reliance on the criterion of colonial morphology is unreliable (Charter and Taylor, 1952). Occasionally a strain appears with some peculiar biological characteristic and is not identified by routine laboratory investigation. An epidemic caused by such a strain is reported here.

**Clinical Aspects**

In July and August 1961 there was an outbreak of gastroenteritis in the Children’s Hospital, Sheffield. Eight children under the age of 5 months were involved. All had been patients in the same ward. As soon as a clinical diagnosis of gastroenteritis was made, the child was transferred to an isolation unit in another part of the hospital.

The original source of the organism was never conclusively established but it is reasonable to surmise that it was introduced by Case 1. This child had been admitted to another hospital with diarrhoea and vomiting. Stool culture on several occasions failed to show any pathogen and he was transferred to a cubicle attached to the main ward on 21 July. Cases 2 and 3 were twins aged 3 months who were admitted on 23 July to a cubicle opposite for adjustment of their feeding régime. They were discharged home on 26 July and readmitted to the isolation unit on 2 August. Case 2 had severe diarrhoea and vomiting for five days before readmission and Case 3 had a milder illness with loose stools for one day. Case 4 was admitted to the main ward on 9 August with pyloric stenosis which was treated surgically and he was discharged home on 14 August. Four days later he was readmitted to a cubicle in the isolation unit in a moribund state. He had had diarrhoea and vomiting for 24 hours, and had been vaguely unwell for three days before that. Cases 5 to 8 were mildly ill. These children developed diarrhoea on 15 August, 19 August, 20 August, and 21 August respectively. Cases 5, 6, and 7 were children in the main ward. Case 8 had been admitted to the cubicle formerly occupied by Case 1.

The severity of the illness varied markedly from patient to patient; Cases 1, 2, and 4 were very severely ill, requiring parenteral fluid replacement. At the other extreme Case 7 passed only one or two loose stools. The other four cases

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were also mild. All were treated with Neomycin by mouth. The clinical response was excellent and all eight recovered.

Bacteriology

Rectal swabs taken from each of the eight patients were plated on MacConkey agar and desoxycholate citrate agar, and were inoculated into selenite F broth. The latter was incubated at 37°C. for 12 hours and then subcultured on desoxycholate citrate agar. All plate cultures were examined after 24 hours' aerobic incubation at 37°C. and again after 48 hours. 'Pale' colonies (non-lactose fermenting) were also cultured and subcultured on MacConkey agar, and Dulcitol. Mannitol, rhamnose, sorbitol, xylose, and trehalose; acid and gas after three to four days in lactose, sucrose, raffinose, and salicin (one strain, from Case 4, fermented lactose in four hours); and acid and gas after eight to 10 days in dulcitol. No strain fermented adonitol. All strains produced indole and gave a positive methyl red reaction. Ammonium citrate was not utilized and the Voges-Proskauer test was negative.

Discussion

The colonial appearance of epidemic E. coli serotypes after 24 or 48 hours' incubation on MacConkey or desoxycholate citrate agar is indistinguishable from that of the normal bowel commensal species. It has therefore become customary to search for the pathogenic varieties by screening lactose-fermenting (pink) colonies by a slide agglutination test with polyvalent epidemic coli 0 antiserum. The bacteriological findings in this outbreak emphasize that epidemic E. coli strains may give pale, non-lactose fermenting colonies on a MacConkey plate and ferment lactose so slowly that they may be designated non-lactose fermenters in a routine laboratory. Failure to appreciate the possible significance of these pale colonies may result in serious delay in diagnosis.

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References

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