SALMONELLA RUBISLAW WITH THREE "NORMAL" FLAGELLAR ANTIGENS

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Edwards, McWhorter, and Douglas (J. Bacteriol. **84**:95, 1962) and McWhorter and Edwards (J. Bacteriol. **85**:1440, 1963) reviewed the subject of diphasic Salmonella serotypes that possess three or more recognizable flagellar components. These investigators noted that in certain instances complex forms possessed well-known and recognized Salmonella flagellar antigens, such as in S. salinatis $(4,12:d,e,h:d,e,n,z_{15})$, whereas in others the microorganisms contained more recently characterized antigens, such as z_{43} or z_{45} , as a major component of both phases. The culture (2209-63) described here was an unusual example of the first of the two above-mentioned categories.

Culture 2209-63 was isolated from a stool specimen of a male child, aged 2 months, who was ill with gastroenteritis and was hospitalized in Louisiana. The biochemical reactions given by the culture were similar to those given by members of the genus Salmonella, and the strain was a member of subgenus I of Kauffmann (Acta Pathol. Microbiol. Scand. 49:293, 1960; 58:109, 1963). However, it differed from the standard strain of S. rubislaw in that it utilized l-tartrate but failed to utilize i-tartrate and mucate.

The strain was a member of Salmonella O antigen group 11, was agglutinated to the titer of an O antiserum prepared with S. aberdeen (O11), and in absorption tests removed all agglutinin from that antiserum. When received, the culture

Table 1. Agglutination reactions of the flagellar antigens of Salmonella rubislaw 2209-63

Antigen	Antiserum		
	S. typhi (d)	S. rubis- law phase 1 (r)	S. abortus equi (e,n,x)
2209-63, phase 1*	6,400	100	<100
2209-63, phase 2*	3,200	< 100	3,200
2209-63, r phase†	< 100	6,400	< 100
2209-63, e, n, x phase†	< 100	400	3,200

^{*} From single colonies from original culture.

was flocculated to the titer of H antiserum d derived from S. typhi, but not by diagnostic dilutions of other Salmonella H antisera. When numerous single colonies from platings were examined, it was found that about 85% agglutinated in diagnostic dilutions of d antiserum alone, whereas the remainder reacted strongly in d and e, n, x antisera (phases 1 and 2, respectively, Table 1). Five colonies that were agglutinated by d antiserum alone were placed in semisolid medium containing d antiserum. Four of these yielded forms that were agglutinated only by r antiserum, and one colony gave rise to a form that was flocculated only by e, n, x antiserum. When similarly treated, five colonies, which originally were agglutinated by both d and e, n, xantisera, yielded e, n, x forms and a sixth yielded a r form. Single-colony isolations from more than 20 cultures (including the 11 colonies mentioned above) that had been passed through d antiserum were placed in semisolid medium that contained both r and e, n, x antisera and passed serially through five transfers. In every instance, these cultures were immobilized and the d form was not recovered.

The r and e,n,x phases (Table 1) of culture 2209-63 were identical with those of S. rubislaw phase 1 (r) and S. abortus equi (S. abortivoequina) (e,n,x), respectively, as demonstrated by appropriate agglutinin-absorption tests. The d antigen of 2209-63 reduced the titer of S. typhi d antiserum from 1:12,800 to 1:400 in absorption tests; hence, it was not identical with the d antigen of S. typhi.

Thus, the antigenic composition of culture 2209-63 was characterized as 11:d,r:d,e,n,x. Since it was possible to derive a culture indistinguishable from S. rubislaw (11:r:e,n,x) from culture 2209-63, it was regarded as a complex form of that serotype, and completely comparable with complex forms of S. montgomery (Edwards, Kauffmann, and Huey, Acta Pathol. Microbiol. Scand. 41:517, 1957) and S. salinatis (Edwards and Bruner, J. Bacteriol. 44:289, 1942).

[†] The r and e, n, x phases of culture 2209-63 were obtained by passage of phase 1 and phase 2, respectively, through d antiserum.