

**K960138 SAS SALMONELLA 0 FACTOR 8 ANTISERUM AND/OR  
SAS SALMONELLA 0 GROUP C2 FACTOR 9 ANTISERUM**Mar 6, 1996  
55 days to decisionK960138 · Product code: **GRM** · Microbiology  
Source: <https://www.510kdatabase.net/k960138/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, All Groups, Salmonella Spp. (GRM)
Date received	Jan 11, 1996
Decision date	Mar 6, 1996
Days to decision	55 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Sa Scientific, Inc.</b>
Location	San Antonio, TX, US
Contact	HARBI SHADFAN
510(k) history	199 submissions · 199 cleared · 1993-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960138/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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