

**K811559 VENTRE/SEP IGE RIA TEST SET**Jul 10, 1981  
38 days to decisionK811559 · Product code: **DGC** · Chemistry  
Source: <https://www.510kdatabase.net/k811559/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ige, Antigen, Antiserum, Control (DGC)
Date received	Jun 2, 1981
Decision date	Jul 10, 1981
Days to decision	38 days
Third-party review	No

**APPLICANT**

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Company	<b>Ventrex Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	82 submissions · 82 cleared · 1979-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k811559/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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