

K820978 VENTRE/SPEC ALLERGEN CONTROL-INHALANTJun 11, 1982
66 days to decisionK820978 · Product code: **DGC** · Microbiology
Source: <https://www.510kdatabase.net/k820978/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ige, Antigen, Antiserum, Control (DGC)
Date received	Apr 6, 1982
Decision date	Jun 11, 1982
Days to decision	66 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k820978/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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