

K820897 VENTRE/SPEC IGE CONTROLSMay 5, 1982
36 days to decisionK820897 · Product code: **DGC** · Immunology
Source: <https://www.510kdatabase.net/k820897/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ige, Antigen, Antiserum, Control (DGC)
Date received	Mar 30, 1982
Decision date	May 5, 1982
Days to decision	36 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820897/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026