FDA 510(k) Premarket Notification

K820897 VENTRE/SPEC IGE CONTROLS

CLEARED

May 5, 1982
36 days to decision

K820897 · 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