

K873647 VENTRESCREEN CAPILLARY BLOOD COLLECTION TUBEOct 20, 1987
41 days to decisionK873647 · Product code: **GIO** · Hematology
Source: <https://www.510kdatabase.net/k873647/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Collection, Capillary Blood (GIO)
Date received	Sep 9, 1987
Decision date	Oct 20, 1987
Days to decision	41 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k873647/>; 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