FDA 510(k) Premarket Notification

K873647 VENTRESCREEN CAPILLARY BLOOD COLLECTION TUBE

CLEARED

Oct 20, 1987
41 days to decision

K873647 · 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