FDA 510(k) Premarket Notification

K812970 S/P CAPILLARY TUBES, HEPARINIZED (3A)

CLEARED

Nov 6, 1981
15 days to decision

K812970 · Product code: **GIO** · Hematology
Source: https://www.510kdatabase.net/k812970/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Collection, Capillary Blood (GIO)
Date received	Oct 22, 1981
Decision date	Nov 6, 1981
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**American Dade**
Location	Mchenry, IL, US
510(k) history	149 submissions · 149 cleared · 1980-1987