FDA 510(k) Premarket Notification

CLEARED

K830149 IL VIDEO 22,21 & II ATOMIC SPECTROPHOTO-

Feb 28, 1983
42 days to decision

K830149 · Product code: **DWD** · Chemistry
Source: https://www.510kdatabase.net/k830149/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suction Control, Intracardiac, Cardiopulmonary Bypass (DWD)
Date received	Jan 17, 1983
Decision date	Feb 28, 1983
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Instrumentation Laboratory CO**
Location	Mchenry, IL, US
510(k) history	321 submissions · 320 cleared · 1976-2023