FDA 510(k) Premarket Notification

K833561 LIGHT REFLECTION RHEOGRAPHY

CLEARED

Jan 8, 1985
454 days to decision

K833561 · Product code: **JOM** · Cardiovascular
Source: https://www.510kdatabase.net/k833561/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Photoelectric, Pneumatic Or Hydraulic (JOM)
Date received	Oct 12, 1983
Decision date	Jan 8, 1985
Days to decision	454 days
Third-party review	No

APPLICANT

Company	**Instru-Med, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1985-1985