FDA 510(k) Premarket Notification

K810297 DATASCOPE PERCOR=DL DUAL-LUMEN INTRA

CLEARED

May 6, 1981
90 days to decision

K810297 · Product code: **DSP** · Cardiovascular
Source: https://www.510kdatabase.net/k810297/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Balloon, Intra-aortic And Control (DSP)
Date received	Feb 5, 1981
Decision date	May 6, 1981
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Datascope Corp.**
Location	Mchenry, IL, US
510(k) history	136 submissions · 135 cleared · 1976-2019