FDA 510(k) Premarket Notification

K802924 KONTRON PERCUTAEOUS DOUBLE-LUMEN

CLEARED

Mar 11, 1981
113 days to decision

K802924 · Product code: **DSP** · Cardiovascular
Source: https://www.510kdatabase.net/k802924/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Balloon, Intra-aortic And Control (DSP)
Date received	Nov 18, 1980
Decision date	Mar 11, 1981
Days to decision	113 days
Third-party review	No

APPLICANT

Company	**Avco Medical Products**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1981