FDA 510(k) Premarket Notification

K810661 CATHDAMP

CLEARED

Mar 27, 1981
16 days to decision

K810661 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k810661/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Mar 11, 1981
Decision date	Mar 27, 1981
Days to decision	16 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	**Medline Industries, Inc.**
Location	Mchenry, IL, US
510(k) history	238 submissions · 234 cleared · 1977-2024