

K810661 CATHDAMPMar 27, 1981
16 days to decisionK810661 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k810661/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026