

K812999 PET POLYETHYLENE TORQUE CATHETERNov 16, 1981
21 days to decisionK812999 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k812999/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Oct 26, 1981
Decision date	Nov 16, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Universal Medical Instrument Corp.
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1977-1996

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k812999/> 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 30, 2026