FDA 510(k) Premarket Notification

K812640 URE SIL EMBOLECTOMY/THROMBECTOMY CATH

CLEARED

Oct 2, 1981
16 days to decision

K812640 · Product code: **DXE** · Cardiovascular
Source: https://www.510kdatabase.net/k812640/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Sep 16, 1981
Decision date	Oct 2, 1981
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Uresil Corp.**
Location	Mchenry, IL, US
510(k) history	45 submissions · 44 cleared · 1981-2001