FDA 510(k) Premarket Notification

K780470 H.P. MEDICAL GRADE SILICONE TUBING

CLEARED

Apr 27, 1978
36 days to decision

K780470 · Product code: **KDL** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k780470/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Perfusion, Kidney, Disposable (KDL)
Date received	Mar 22, 1978
Decision date	Apr 27, 1978
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Dow Corning Corp. Healthcare Industries Materials**
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1976-1985