FDA 510(k) Premarket Notification

K760744 TAVERNETTI-TENNANT

CLEARED

Oct 7, 1976
7 days to decision

K760744 · Product code: **HRY** · Orthopedic
Source: https://www.510kdatabase.net/k760744/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Sep 30, 1976
Decision date	Oct 7, 1976
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**Cutter Laboratories, Inc.**
Location	Mchenry, IL, US
Website	https://www.bayer.com
510(k) history	39 submissions · 39 cleared · 1976-1986