

K760744 TAVERNETTI-TENNANTOct 7, 1976
7 days to decisionK760744 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k760744/>; 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 May 22, 2026