FDA 510(k) Premarket Notification

K893959 P.S.A. FEMORAL HIP PROSTHESIS

CLEARED

Jul 25, 1989
54 days to decision

K893959 · Product code: **KWY** · Orthopedic
Source: https://www.510kdatabase.net/k893959/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Hemi-, Femoral, Metal/polymer, Cemented Or Uncemented (KWY)
Date received	Jun 1, 1989
Decision date	Jul 25, 1989
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Protek, Inc.**
Location	Indianapolis, IN, US
Contact	KENNETH EPLING
510(k) history	25 submissions · 20 cleared · 1985-1989