

K893959 P.S.A. FEMORAL HIP PROSTHESISJul 25, 1989
54 days to decisionK893959 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893959/>; 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