FDA 510(k) Premarket Notification

K913632 BATEMAN UPF-III BIPOLAR ENDOPROSTHESIS SYSTEM

CLEARED

Nov 13, 1991
90 days to decision

K913632 · Product code: **KWY** · Orthopedic
Source: https://www.510kdatabase.net/k913632/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Hemi-, Femoral, Metal/polymer, Cemented Or Uncemented (KWY)
Date received	Aug 15, 1991
Decision date	Nov 13, 1991
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Kirschner Medical Corp.**
Location	Mchenry, IL, US
Contact	SAM SON
510(k) history	76 submissions · 57 cleared · 1983-1995