FDA 510(k) Premarket Notification

K960908 SRRS PRESSED FIT FEMORAL REVISION COMPONENT

CLEARED

Aug 8, 1996
156 days to decision

K960908 · Product code: **LWJ** · Orthopedic
Source: https://www.510kdatabase.net/k960908/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Uncemented (LWJ)
Date received	Mar 5, 1996
Decision date	Aug 8, 1996
Days to decision	156 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Stelkast Company**
Location	Pittsburgh, PA, US
Contact	DONALD A STEVENS
510(k) history	40 submissions · 37 cleared · 1994-2013