FDA 510(k) Premarket Notification

K894472 CROSS ROUND & SQUARE SPINAL RODS & DIST. SPINAL

CLEARED

Oct 10, 1989
84 days to decision

K894472 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k894472/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Jul 18, 1989
Decision date	Oct 10, 1989
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**Buckman Co., Inc.**
Location	Mchenry, IL, US
Contact	BROMM, R.N.
510(k) history	111 submissions · 104 cleared · 1983-1998