FDA 510(k) Premarket Notification

K800502 ADDITIONAL COMPONENTS KRONNER COMPRES

CLEARED

Apr 2, 1980
29 days to decision

K800502 · Product code: **HRS** · Orthopedic
Source: https://www.510kdatabase.net/k800502/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Mar 4, 1980
Decision date	Apr 2, 1980
Days to decision	29 days
Third-party review	No

APPLICANT

Company	**Richard F. Kronner, M.D.**
Location	Walker, MI, US
510(k) history	3 submissions · 3 cleared · 1977-1980