

K800502 ADDITIONAL COMPONENTS KRONNER COMPRESApr 2, 1980
29 days to decisionK800502 · 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

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Device record: <https://www.510kdatabase.net/k800502/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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