

K931688 SEMI-TUBULAR PLATEApr 25, 1994
385 days to decisionK931688 · Product code: **HRS** · Orthopedic
Source: <https://www.510kdatabase.net/k931688/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Apr 5, 1993
Decision date	Apr 25, 1994
Days to decision	385 days
Third-party review	No

APPLICANT

Company	Onyx Medical Corp.
Location	Memphis, TN, US
Contact	LARAIN B GILMORE
510(k) history	47 submissions · 22 cleared · 1990-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931688/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026