FDA 510(k) Premarket Notification

K931688 SEMI-TUBULAR PLATE

CLEARED

Apr 25, 1994
385 days to decision

K931688 · 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	LARAINE B GILMORE
510(k) history	47 submissions · 22 cleared · 1990-1994