FDA 510(k) Premarket Notification

K914362 TOP CERVICAL SPINE PLATES & TITANIUM BONE SCREWS

CLEARED

Apr 29, 1992
211 days to decision

K914362 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k914362/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Oct 1, 1991
Decision date	Apr 29, 1992
Days to decision	211 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Acorn Group, Inc.**
Location	Walnut Creek, CA, US
Contact	JAMES P.STOUT
510(k) history	4 submissions · 4 cleared · 1990-1992