

K914362 TOP CERVICAL SPINE PLATES & TITANIUM BONE SCREWSApr 29, 1992
211 days to decisionK914362 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914362/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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