FDA 510(k) Premarket Notification

K925351 SYNTHES(U.S.A.) ANTERIOR SPINAL PLATE

CLEARED

Nov 5, 1993
379 days to decision

K925351 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k925351/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Oct 22, 1992
Decision date	Nov 5, 1993
Days to decision	379 days
Third-party review	No

APPLICANT

Company	**Synthes (Usa)**
Location	Mchenry, IL, US
Contact	DIANE C TIERNAN
510(k) history	411 submissions · 394 cleared · 1977-2015