FDA 510(k) Premarket Notification

K802462 SYNTHES SPINAL IMPLANT SYSTEM

CLEARED

Dec 16, 1980
69 days to decision

K802462 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k802462/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Oct 8, 1980
Decision date	Dec 16, 1980
Days to decision	69 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k802462/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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