FDA 510(k) Premarket Notification

K041888 CURVED PEEK TETRIS SPINAL IMPLANT

CLEARED

Aug 10, 2004
29 days to decision

K041888 · Product code: **MQP** · Orthopedic
Source: https://www.510kdatabase.net/k041888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jul 12, 2004
Decision date	Aug 10, 2004
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Signus Medizintechnik GmbH**
Location	Minneapolis, MN, US
Contact	TRACY L GRAY
510(k) history	23 submissions · 22 cleared · 1999-2025