

K041888 CURVED PEEK TETRIS SPINAL IMPLANTAug 10, 2004
29 days to decisionK041888 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k041888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026