

K072933 SPI CONTACT PLATFORM O 4.0 MMNov 15, 2007
30 days to decisionK072933 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k072933/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Oct 16, 2007
Decision date	Nov 15, 2007
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Thommen Medical AG
Location	San Diego, CA, US
Contact	FLOYD G LARSON
510(k) history	26 submissions · 26 cleared · 2002-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072933/> 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 June 26, 2026