

**K093064 TUMARK PROFESSIONAL, MODEL 271560, MR-
TUMARK PROFESSIONAL, MODEL 601560**Feb 17, 2010
140 days to decisionK093064 · Product code: **NEU** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k093064/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Radiographic, Implantable (NEU)
Date received	Sep 30, 2009
Decision date	Feb 17, 2010
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Somatex Medical Technologies GmbH
Location	Cambridge, MA, US
Contact	SUSANNE RAAB
510(k) history	7 submissions · 7 cleared · 2008-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k093064/>; 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 27, 2026