

K073095 TUMARK PROFESSIONAL, MODEL 271560Mar 19, 2008
139 days to decisionK073095 · Product code: **NEU** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k073095/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Radiographic, Implantable (NEU)
Date received	Nov 1, 2007
Decision date	Mar 19, 2008
Days to decision	139 days
Third-party review	No
Summary / Statement	Summary
Other names	MR- TUMARK PROFESSIONAL, MODEL 601560

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/k073095/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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