FDA 510(k) Premarket Notification

K073095 TUMARK PROFESSIONAL, MODEL 271560

CLEARED

Mar 19, 2008
139 days to decision

K073095 · Product code: **NEU** · General & Plastic Surgery
Source: 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