

K091246 ILED 3 MOBILE (I300M), TRULIGHT 5500 MOBILE (T500M), TRULIGHT 5300 MOBILE (T300M)Sep 28, 2009
154 days to decisionK091246 · Product code: **FSS** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k091246/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Light, Surgical, Floor Standing (FSS)
Date received	Apr 27, 2009
Decision date	Sep 28, 2009
Days to decision	154 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Trumpf Medizin Systeme GmbH + Co. KG
Location	Charleston, SC, US
Contact	LINDSEY HENGEL
510(k) history	1 submissions · 1 cleared · 2009-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091246/> 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 22, 2026