

K091142 TUTOPATCH, TUTOMESH, MODELS 68350, 68351, 68352, 68353, 68354, 68355, 68356, 68357, 68358Aug 31, 2009
133 days to decisionK091142 · Product code: **FTM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k091142/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical (FTM)
Date received	Apr 20, 2009
Decision date	Aug 31, 2009
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Rti Biologics, Inc.
Location	Alachua, FL, US
Contact	TRAVIS AROLA
510(k) history	3 submissions · 3 cleared · 2008-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091142/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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