FDA 510(k) Premarket Notification

**CLEARED**

# K091142 TUTOPATCH, TUTOMESH, MODELS 68350, 68351, 68352, 68353, 68354, 68355, 68356, 68357, 68358

Aug 31, 2009
133 days to decision

K091142 · Product code: **FTM** · General & Plastic Surgery
Source: 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 |