

K151386 C-QUR, C-QUR FX, C-QUR TacShield, C-QUR V-Patch, C-QUR CentriFX, C-QUR Mosaic

Oct 22, 2015
149 days to decision

K151386 · Product code: FTL · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k151386/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	May 26, 2015
Decision date	Oct 22, 2015
Days to decision	149 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Atrium Medical Corporation
Location	Hudson, NH, US
Contact	Timothy J Talcott
510(k) history	6 submissions · 6 cleared · 2011-2024

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k151386/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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