

K050311 ATRIUM MEDICAL CORPORATION C-QUR MESHMar 31, 2006
416 days to decisionK050311 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k050311/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Feb 8, 2005
Decision date	Mar 31, 2006
Days to decision	416 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Atrium Medical Corp.
Location	Mchenry, IL, US
Contact	JOSEPH P DE PAOLO
510(k) history	63 submissions · 61 cleared · 1981-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k050311/> 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 May 25, 2026