

K110110 ATRIUM CENTRILFX MESHFeb 15, 2011
32 days to decisionK110110 · Product code: **FTL** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k110110/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Jan 14, 2011
Decision date	Feb 15, 2011
Days to decision	32 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k110110/> 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 26, 2026