

**K113744 QUILL PDO KNOTLESS TISSUE-CLOSURE DEVICE
(POLYDIOXANONE)**Feb 1, 2012
43 days to decisionK113744 · Product code: **NEW** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k113744/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Surgical, Absorbable, Polydioxanone (NEW)
Date received	Dec 20, 2011
Decision date	Feb 1, 2012
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Angiotech
Location	Herndon, VA, US
Contact	KIRSTEN STOWELL
510(k) history	7 submissions · 6 cleared · 2007-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k113744/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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