

**K113311 TIO2MESH TM**Jul 17, 2012  
251 days to decisionK113311 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k113311/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Nov 9, 2011
Decision date	Jul 17, 2012
Days to decision	251 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biocer Entwicklungs GmbH</b>
Location	Austin, TX, US
Contact	CHERYL FISHER
510(k) history	1 submissions · 1 cleared · 2012-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k113311/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 14, 2026