

**K982532 PARIETEX**Jan 20, 1999  
184 days to decisionK982532 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k982532/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Jul 20, 1998
Decision date	Jan 20, 1999
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cogent</b>
Location	Washington, Dc, DC, US
Contact	HOWARD M HOLSTEIN
Website	<a href="http://www.cogentco.com/">http://www.cogentco.com/</a>
510(k) history	1 submissions · 1 cleared · 1999-1999

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982532/> 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 July 4, 2026