FDA 510(k) Premarket Notification

K982532 PARIETEX

CLEARED

Jan 20, 1999
184 days to decision

K982532 · Product code: **FTL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k982532/

SUBMISSION DETAILS

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

Company	**Cogent**
Location	Washington, Dc, DC, US
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510(k) history	1 submissions · 1 cleared · 1999-1999