FDA 510(k) Premarket Notification

K013820 FINN CHAMBER (R)

CLEARED

Jan 30, 2002
75 days to decision

K013820 · Product code: **KXF** · General Hospital
Source: https://www.510kdatabase.net/k013820/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Applicator, Absorbent Tipped, Non-sterile (KXF)
Date received	Nov 16, 2001
Decision date	Jan 30, 2002
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Epitest Ltd. OY**
Location	Petaluma, CA, US
Contact	CLAWSON BOWMAN
510(k) history	1 submissions · 1 cleared · 2002-2002