

K013820 FINN CHAMBER (R)Jan 30, 2002
75 days to decisionK013820 · 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

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