FDA 510(k) Premarket Notification

K022645 CURAD

CLEARED

Nov 5, 2002
89 days to decision

K022645 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k022645/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Aug 8, 2002
Decision date	Nov 5, 2002
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Beiersdorf AG**
Location	North Attleboro, MA, US
Contact	DANIEL J DILLON
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510(k) history	3 submissions · 3 cleared · 2002-2004