FDA 510(k) Premarket Notification

K980382 NON-WOVEN COMPRESS

CLEARED

Mar 9, 1998
35 days to decision

K980382 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k980382/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Feb 2, 1998
Decision date	Mar 9, 1998
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Convatec, A Division of E.R. Squibb & Sons**
Location	Walker, MI, US
Contact	ADRIENNE MCNALLY
510(k) history	81 submissions · 68 cleared · 1982-2002