FDA 510(k) Premarket Notification

CLEARED

K153571 Cavilon Advanced High Endurance Skin Protectant

Aug 23, 2016
252 days to decision

K153571 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k153571/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Dec 15, 2015
Decision date	Aug 23, 2016
Days to decision	252 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**3M Healthcare**
Location	St. Paul, MN, US
Contact	MARIA RUIZ
510(k) history	21 submissions · 20 cleared · 1991-2020

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k153571/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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