FDA 510(k) Premarket Notification

K063202 PRODERMA LIQUID BANDAGE

CLEARED

May 2, 2007
194 days to decision

K063202 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k063202/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Oct 20, 2006
Decision date	May 2, 2007
Days to decision	194 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Procurement Technology Systems, LLC**
Location	Greer, SC, US
Contact	WILLIAM J GRISWOLD
510(k) history	1 submissions · 1 cleared · 2007-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k063202/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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