FDA 510(k) Premarket Notification

K982439 PROXIDERM, MODELS TN 460, TN 90 460, BK 460

CLEARED

Sep 8, 1998
56 days to decision

K982439 · Product code: **MKY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k982439/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Skin Closure (MKY)
Date received	Jul 14, 1998
Decision date	Sep 8, 1998
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Progressive Surgical Products, Inc.**
Location	Westbury, NY, US
Contact	ROBERT ODDSEN
510(k) history	3 submissions · 3 cleared · 1994-1998