FDA 510(k) Premarket Notification

K905069 IODOSORB GEL

CLEARED

Jan 31, 1991
83 days to decision

K905069 · Product code: **KOZ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k905069/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Beads, Hydrophilic, For Wound Exudate Absorption (KOZ)
Date received	Nov 9, 1990
Decision date	Jan 31, 1991
Days to decision	83 days
Third-party review	No

APPLICANT

Company	**Perstorp AB C/O Robert Joel Slomoff**
Location	Potomac, MD, US
Contact	SLOMOFF
510(k) history	2 submissions · 1 cleared · 1987-1991