FDA 510(k) Premarket Notification

K900903 KOPANS HOOKWIRE LESION LOCALIZATION NEEDLE

CLEARED

May 3, 1990
65 days to decision

K900903 · Product code: **GDF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k900903/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Needle, Surgical (GDF)
Date received	Feb 27, 1990
Decision date	May 3, 1990
Days to decision	65 days
Third-party review	No

APPLICANT

Company	**Milex Products, Inc.**
Location	Mchenry, IL, US
Contact	ROBERT SHAW
510(k) history	8 submissions · 8 cleared · 1976-2000