FDA 510(k) Premarket Notification

K890170 SCISSORS

CLEARED

Feb 3, 1989
25 days to decision

K890170 · Product code: **LRW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k890170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scissors, General, Surgical (LRW)
Date received	Jan 9, 1989
Decision date	Feb 3, 1989
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Dhason Brothers, Inc.**
Location	Hyattsville, MD, US
Contact	ROBERT DHASON
510(k) history	14 submissions · 14 cleared · 1989-1989