FDA 510(k) Premarket Notification

K910823 PERCUDISC(TM) SYSTEM

CLEARED

Dec 30, 1991
307 days to decision

K910823 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k910823/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 26, 1991
Decision date	Dec 30, 1991
Days to decision	307 days
Third-party review	No

APPLICANT

Company	**Percudisc, Inc.**
Location	Mountain View, CA, US
Contact	CHARLES L ROSE
510(k) history	2 submissions · 1 cleared · 1991-1992