FDA 510(k) Premarket Notification

K840144 YAG LASER 8000 GASTROINTESTINAL BLEE

CLEARED

May 18, 1984
126 days to decision

K840144 · Product code: **FCG** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k840144/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Biopsy Needle (FCG)
Date received	Jan 13, 1984
Decision date	May 18, 1984
Days to decision	126 days
Third-party review	No

APPLICANT

Company	**Cooper Medical Corp.**
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984