FDA 510(k) Premarket Notification

K902320 BAUER FLEXI-TEMNO BIOPSY NEEDLE

CLEARED

Oct 1, 1990
131 days to decision

K902320 · Product code: **FCG** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k902320/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Biopsy Needle (FCG)
Date received	May 23, 1990
Decision date	Oct 1, 1990
Days to decision	131 days
Third-party review	No

APPLICANT

Company	**Proact, Ltd.**
Location	State College, PA, US
Contact	JOHN N ZINK
510(k) history	5 submissions · 5 cleared · 1990-1994