

**K902320 BAUER FLEXI-TEMNO BIOPSY NEEDLE**Oct 1, 1990  
131 days to decisionK902320 · Product code: **FCG** · Gastroenterology & UrologySource: <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	<b>Proact, Ltd.</b>
Location	State College, PA, US
Contact	JOHN N ZINK
510(k) history	5 submissions · 5 cleared · 1990-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902320/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 2, 2026