

**K770403 CATHETER-THRU-NEEDLE DEVICE**Mar 16, 1977  
13 days to decisionK770403 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k770403/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                                       |
| Submission type       | Traditional  |
| Device classification | Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ) |
| Date received         | Mar 3, 1977  |
| Decision date         | Mar 16, 1977   |
| Days to decision      | 13 days  |
| Third-party review    | No   |

**APPLICANT**

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|----------------|---|
| Company        | <b>Abbott Laboratories</b>                                |
| Location       | Abbott Park, IL, US                                       |
| Website        | <a href="http://www.abbott.com">http://www.abbott.com</a> |
| 510(k) history | 883 submissions · 868 cleared · 1976-2026                 |

Abbott Laboratories is an American multinational medical devices and health care company headquartered in Abbott Park, Illinois. The company operates in over 160 countries and produces pharmaceuticals, diagnostics, nutritional products, and medical devices. Abbott maintains a substantial FDA 510(k) regulatory record with FDA 510(k) cleared devices from total submissions since 1976. The company's cleared devices span chemistry, microbiology, hematology, immunology, and toxicology categories. The latest clearance in 2025 reflects continued regulatory activity and product de...

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