

K120935 SECURACATHApr 27, 2012
30 days to decisionK120935 · Product code: **OKC** · General Hospital
Source: <https://www.510kdatabase.net/k120935/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implanted Subcutaneous Securement Catheter (OKC)
Date received	Mar 28, 2012
Decision date	Apr 27, 2012
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interrad Medical, Inc.
Location	Plymouth, MN, US
Contact	Sew-Wah Tay
510(k) history	7 submissions · 7 cleared · 2008-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120935/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026