

K810530 PERI-PATCH GLUE MOLDMar 24, 1981
26 days to decisionK810530 · Product code: **KGZ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k810530/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter (KGZ)
Date received	Feb 26, 1981
Decision date	Mar 24, 1981
Days to decision	26 days
Third-party review	No

APPLICANT

Company	Quinton, Inc.
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k810530/> 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 20, 2026