

K822631 ARGYLE DUO-TUBESep 9, 1982
9 days to decisionK822631 · Product code: **KNT** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k822631/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Aug 31, 1982
Decision date	Sep 9, 1982
Days to decision	9 days
Third-party review	No

APPLICANT

Company	Sherwood Medical Co.
Location	Mchenry, IL, US
510(k) history	191 submissions · 177 cleared · 1976-1998

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k822631/> 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 14, 2026