

K823331 PC-204 TUNNELORJan 26, 1983
79 days to decisionK823331 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k823331/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Nov 8, 1982
Decision date	Jan 26, 1983
Days to decision	79 days
Third-party review	No

APPLICANT

Company	Medigroup
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1983-1983

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823331/> 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 June 30, 2026