

K801352 HEMORRHOIDAL LIGATION INSTR. SYSTEMJul 8, 1980
29 days to decisionK801352 · Product code: **FHN** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k801352/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ligator, Hemorrhoidal (FHN)
Date received	Jun 9, 1980
Decision date	Jul 8, 1980
Days to decision	29 days
Third-party review	No

APPLICANT

Company	Depuy, Inc.
Location	Mchenry, IL, US
510(k) history	303 submissions · 239 cleared · 1976-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k801352/> 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