FDA 510(k) Premarket Notification

K801352 HEMORRHOIDAL LIGATION INSTR. SYSTEM

CLEARED

Jul 8, 1980
29 days to decision

K801352 · Product code: **FHN** · Gastroenterology & Urology
Source: 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