

K980092 SMITH & NEPHEW DYONICS INTELIJET REUSABLE CANNULASMar 9, 1998
59 days to decisionK980092 · Product code: **FRN** · General Hospital
Source: <https://www.510kdatabase.net/k980092/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Jan 9, 1998
Decision date	Mar 9, 1998
Days to decision	59 days
Third-party review	No

APPLICANT

Company	Smith & Nephew, Inc.
Location	Mchenry, IL, US
Contact	DEBORAH J CONNOR
Website	http://www.smith-nephew.com/
510(k) history	530 submissions · 517 cleared · 1980-2026

Smith & Nephew, Inc. is a medical technology company focused on repair, regeneration, and replacement of soft and hard tissues. The company operates with a manufacturing facility in McHenry, US. Smith & Nephew has established a significant regulatory track record with the FDA. The company has received FDA 510(k) clearances from total submissions since 1980. Orthopedic devices represent the dominant category, accounting for 71% of submissions. The company remains active, with the latest clearance in 2025. Recent cleared devices reflect a strong focus on orthopedic surgical...

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k980092/> 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 15, 2026