

K882329 L-F COILED EXTENSION TUBINGSep 2, 1988
88 days to decisionK882329 · Product code: **DXT** · Cardiovascular
Source: <https://www.510kdatabase.net/k882329/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Jun 6, 1988
Decision date	Sep 2, 1988
Days to decision	88 days
Third-party review	No

APPLICANT

Company	Mallinckrodt Group, Inc.
Location	McHenry, IL, US
Contact	THOMAS F DONOHUE
Website	https://www.mallinckrodt.com
510(k) history	27 submissions · 27 cleared · 1976-1995

Mallinckrodt Group, Inc. is a pharmaceutical and medical device company based in McHenry, US. Now part of a restructured corporate family following a 2025 merger and subsequent spin-off, the company's regulatory history reflects its historical role in medical device manufacturing. Mallinckrodt received FDA 510(k) clearances from total submissions, with no denied submissions on record. The company's cleared devices span radiology, cardiovascular, gastroenterology, and surgical specialties, with clearances issued between 1976 and 1995. This regulatory record is now historic...
