

**K942381 LOW PRESSURE INJECTION TUBING, COILED LINE,
EXTENSION TUBING**Mar 17, 1995
302 days to decisionK942381 · Product code: **DXT** · Cardiovascular
Source: <https://www.510kdatabase.net/k942381/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	May 19, 1994
Decision date	Mar 17, 1995
Days to decision	302 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Apogee, Inc.
Location	Raleigh, NC, US
Contact	DIANE N PEPER
510(k) history	2 submissions · 2 cleared · 1995-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942381/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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