

K904212 HIGH-PRESSURE CONNECTOR TUBEDec 3, 1990
82 days to decisionK904212 · Product code: **DXT** · CardiovascularSource: <https://www.510kdatabase.net/k904212/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Sep 12, 1990
Decision date	Dec 3, 1990
Days to decision	82 days
Third-party review	No

APPLICANT

Company	Mallinckrodt Medical
Location	St Louis, MO, US
Contact	DAVID E BROWN
510(k) history	43 submissions · 42 cleared · 1990-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k904212/>; 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