

K800823 QUICK FILLApr 29, 1980
15 days to decisionK800823 · Product code: **DXT** · Cardiovascular
Source: <https://www.510kdatabase.net/k800823/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Apr 14, 1980
Decision date	Apr 29, 1980
Days to decision	15 days
Third-party review	No

APPLICANT

Company	Medrad, Inc.
Location	Mchenry, IL, US
Website	http://www.medrad.com/
510(k) history	105 submissions · 105 cleared · 1976-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k800823/>; 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 17, 2026