

K822536 MARK VSep 28, 1982
35 days to decisionK822536 · Product code: **DXT** · Cardiovascular
Source: <https://www.510kdatabase.net/k822536/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Injector And Syringe, Angiographic (DXT) |
| Date received | Aug 24, 1982 |
| Decision date | Sep 28, 1982 |
| Days to decision | 35 days |
| Third-party review | No |

APPLICANT

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|----------------|---|
| 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/k822536/>; 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