FDA 510(k) Premarket Notification

K790835 DYNAJECT

CLEARED

Jun 22, 1979
53 days to decision

K790835 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k790835/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Apr 30, 1979
Decision date	Jun 22, 1979
Days to decision	53 days
Third-party review	No

APPLICANT

Company	**Atomic Products Corp.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1979-1992

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k790835/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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