

**K890898 MODEL 611101 CONTRAST INJECTION DEVICE**Apr 12, 1989  
49 days to decisionK890898 · Product code: **DXT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k890898/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Feb 22, 1989
Decision date	Apr 12, 1989
Days to decision	49 days
Third-party review	No

**APPLICANT**

---

Company	<b>Medtronic Versaflex, Inc.</b>
Location	San Diego, CA, US
Contact	TIMOTHY J JOHNSON
510(k) history	2 submissions · 2 cleared · 1989-1989

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890898/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026