

**K811964 VISITEC NEEDLE RANGE (CARDIOVASCULAR)**Jul 20, 1981  
10 days to decisionK811964 · Product code: **DRC** · CardiovascularSource: <https://www.510kdatabase.net/k811964/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Trocar (DRC)
Date received	Jul 10, 1981
Decision date	Jul 20, 1981
Days to decision	10 days
Third-party review	No

**APPLICANT**

---

Company	<b>Visitec Co.</b>
Location	Walker, MI, US
510(k) history	49 submissions · 49 cleared · 1979-1995

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k811964/>; 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 23, 2026