

**K960435 PHOENIX CRX VALVE**Aug 27, 1996  
209 days to decisionK960435 · Product code: **JXG** · Neurology  
Source: <https://www.510kdatabase.net/k960435/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Central Nervous System And Components (JXG)
Date received	Jan 31, 1996
Decision date	Aug 27, 1996
Days to decision	209 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Phoenix Biomedical Corp.</b>
Location	Valley Forge, PA, US
Contact	HORACE J WILMER
510(k) history	12 submissions · 11 cleared · 1996-2003

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960435/> 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 June 30, 2026