

K820247 EXTERNAL VENTRICULAR DRAINAGE SYS.Feb 18, 1982
20 days to decisionK820247 · Product code: **JXG** · Neurology
Source: <https://www.510kdatabase.net/k820247/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Central Nervous System And Components (JXG)
Date received	Jan 29, 1982
Decision date	Feb 18, 1982
Days to decision	20 days
Third-party review	No

APPLICANT

Company	American Heyer Schulte
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1984

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820247/> 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 21, 2026