

K790411 AUTO TRAUMA KITJun 1, 1979
95 days to decisionK790411 · Product code: **KME** · General Hospital
Source: <https://www.510kdatabase.net/k790411/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bedding, Disposable, Medical (KME)
Date received	Feb 26, 1979
Decision date	Jun 1, 1979
Days to decision	95 days
Third-party review	No

APPLICANT

Company	Marshall Electronics, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1978-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k790411/> 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 June 30, 2026