

K880152 PAR(TM) BRAND DRUG OF ABUSE CONTROLSMay 31, 1988
139 days to decisionK880152 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k880152/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Jan 13, 1988
Decision date	May 31, 1988
Days to decision	139 days
Third-party review	No

APPLICANT

Company	Medical Analysis Systems, Inc.
Location	Mchenry, IL, US
Contact	DONNA L ANDERSON
510(k) history	107 submissions · 107 cleared · 1978-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880152/>; 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 14, 2026