FDA 510(k) Premarket Notification

K880152 PAR(TM) BRAND DRUG OF ABUSE CONTROLS

CLEARED

May 31, 1988
139 days to decision

K880152 · 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