

K082388 MINICAP IMMUNOTYPING, MODEL: 2300Apr 14, 2009
238 days to decisionK082388 · Product code: **CFF** · Immunology
Source: <https://www.510kdatabase.net/k082388/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Immuno-electrophoretic, Immunoglobulins, (g, A, M) (CFF)
Date received	Aug 19, 2008
Decision date	Apr 14, 2009
Days to decision	238 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Sebia
Location	Chelsea, MI, US
Contact	KAREN ANDERSON
Website	http://www.sebia.com/
510(k) history	32 submissions · 32 cleared · 1995-2024

Sebia is a global specialized in vitro diagnostic (IVD) player providing powerful diagnostic tools for chronic and metabolic diseases. The company operates with a manufacturing facility in Chelsea, US, and serves laboratories worldwide with instruments, tests, and software solutions. Sebia has received FDA 510(k) clearances from total submissions since 1995, with no denied submissions on record. The company specializes in immunology devices, including capillary electrophoresis and immunofixation technologies. Latest clearance in 2024 confirms active regulatory engagement....
