

K000921 BAYER ADVIA IMS SYSTEMMay 18, 2000
57 days to decisionK000921 · Product code: **DER** · Chemistry
Source: <https://www.510kdatabase.net/k000921/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alpha-1-lipoprotein, Antigen, Antiserum, Control (DER)
Date received	Mar 22, 2000
Decision date	May 18, 2000
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary
Other names	200 + 3

APPLICANT

Company	Bayer Corp.
Location	Elkhart, IN, US
Contact	FREDRICK CLERIE
510(k) history	96 submissions · 96 cleared · 1989-2003

Bayer Corp. is the American subsidiary of Bayer AG, headquartered in Whippany, New Jersey. The company operates 40 fully consolidated subsidiaries across 19 states. Bayer Corp. received FDA 510(k) clearances from total submissions, with no denied submissions on record. The company's regulatory activity spans from 1989 to 2003, with a primary focus on chemistry devices and immunology assays. Notable cleared devices include the ASCENSIA BREEZE BLOOD GLUCOSE METER, CLINITEST PREGNANCY TEST, and the ADVIA CENTAUR immunoassay system. This represents a historical regulatory rec...
