

**K963080 TESTOSTERONE ASSAY FOR THE TECHNICON  
IMMUNO 1 SYSTEM (IN-VITRO DIAGNOSTIC SYSTEM)**Sep 27, 1996  
50 days to decisionK963080 · Product code: **CDZ** · Chemistry  
Source: <https://www.510kdatabase.net/k963080/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Testosterones And Dihydrotestosterone (CDZ)
Date received	Aug 8, 1996
Decision date	Sep 27, 1996
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bayer Corp.</b>
Location	Elkhart, IN, US
Contact	GABRIEL J MURACA, JR.
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...

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Device record: <https://www.510kdatabase.net/k963080/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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