

**K962903 PRESTIGE MEDICAL SERIES 2100 CLINICAL
AUTOCLAVE**Mar 3, 1998
586 days to decisionK962903 · Product code: **FLE** · General Hospital
Source: <https://www.510kdatabase.net/k962903/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilizer, Steam (FLE)
Date received	Jul 25, 1996
Decision date	Mar 3, 1998
Days to decision	586 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Prestige Medical Corp.
Location	Washington, DC, US
Contact	MARK A HELLER
510(k) history	1 submissions · 1 cleared · 1998-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962903/> 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 July 5, 2026