

**K110594 GF-210R MULTI-GAS MODULE FOR NOHON KOHDEN
BEDSIDE MONITORS**Oct 28, 2011
240 days to decisionK110594 · Product code: **CCK** · Anesthesiology
Source: <https://www.510kdatabase.net/k110594/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Mar 2, 2011
Decision date	Oct 28, 2011
Days to decision	240 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nihon Kohden Corp.
Location	Foothill, CA, US
Contact	STEVE GEERDES
510(k) history	9 submissions · 9 cleared · 2010-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k110594/> 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 May 24, 2026