



SPECIAL 510(k) SUBSTANTIAL EQUIVALENCE DETERMINATION DECISION SUMMARY

I Background Information:

A 510(k) Number

K203790

B Applicant

Instrumentation Laboratory Co.

C Proprietary and Established Names

GEM Premier 5000

D Regulatory Information

Product Code(s)	Classification	Regulation Section	Panel
CHL	Class II	21 CFR 862.1120 - Blood Gases (pCO ₂ , pO ₂) And Blood pH Test System	CH - Clinical Chemistry
JGS	Class II	21 CFR 862.1665 - Sodium test system	CH - Clinical Chemistry
CEM	Class II	21 CFR 862.1600 - Potassium test system	CH - Clinical Chemistry
CGZ	Class II	21 CFR 862.1170 - Chloride test system	CH - Clinical Chemistry
JFP	Class II	21 CFR 862.1145 - Calcium test system	CH - Clinical Chemistry
CGA	Class II	21 CFR 862.1345 - Glucose test system	CH - Clinical Chemistry
KHP	Class I	21 CFR 862.1450 - Lactic acid test system	CH - Clinical Chemistry
MQM	Class I, reserved	21 CFR 862.1113 - Bilirubin (total and unbound) in the neonate test system	CH - Clinical Chemistry
GKF	Class II	21 CFR 864.5600 - Automated hematocrit instrument	HE - Hematology

Product Code(s)	Classification	Regulation Section	Panel
GHS	Class II	21 CFR 864.7425 - Carboxyhemoglobin assay	HE - Hematology
GKR	Class II	21 CFR 864.5620 - Automated hemoglobin system	HE - Hematology
GLY	Class II	21 CFR 864.7500 - Whole blood hemoglobin assays	HE - Hematology

II Review Summary:

This 510(k) submission contains information/data on modifications made to the submitter's own Class II device requiring 510(k). The following items are present and acceptable.

1. The name and 510(k) number of the SUBMITTER'S previously cleared device, the GEM Premier 5000, K173403, K160415, K160402, and K160225.
2. Submitter's statement that the **INDICATIONS FOR USE/INTENDED USE** of the modified device as described in its labeling **HAS NOT CHANGED** along with the proposed labeling which includes instructions for use, package labeling, and, if available, advertisements or promotional materials (labeling changes are permitted as long as they do not affect the intended use).
3. A description of the device **MODIFICATION(S)**, including clearly labeled diagrams, engineering drawings, photographs, user's and/or service manuals in sufficient detail to demonstrate that the **FUNDAMENTAL SCIENTIFIC TECHNOLOGY** of the modified device **has not changed. This change was for the upgrade of the operating system from Fedora 17 Linux to WindRiver LTS 18 Linux for the instrument.**
4. Comparison Information (i.e., similarities and differences) to the submitter's legally marketed predicate device including, labeling, intended use, and physical characteristics.
5. A Design Control Activities Summary which includes:
 - a) Identification of Risk Analysis method(s) used to assess the impact of the modification on the device and its components, and the results of the analysis.
 - b) Based on the Risk Analysis, an identification of the verification and/or validation activities required, including methods or tests used and acceptance criteria to be applied.

The labeling for this modified subject device has been reviewed to verify that the indication/intended use for the device is unaffected by the modification. In addition, the submitter's description of the particular modification(s) and the comparative information between the modified and unmodified devices demonstrate that the fundamental scientific technology has not changed. The submitter has provided the design control information as specified in The New 510(k) Paradigm and on this basis, I recommend the device be determined substantially equivalent to the previously cleared (or their preamendment) device.