

**GEM<sup>®</sup> PREMIER<sup>™</sup> 3500**  
with iQM<sup>®</sup>

# Test assured in Critical Care diagnostics.



The most advanced Critical Care testing solutions. Anywhere, everywhere.

 **Instrumentation  
Laboratory**  
A Werfen Company



# Introducing the GEM Premier 3500 system

Fast, reliable whole blood testing for your lab and hospital.

Building on unprecedented simplicity, flexibility and reliability of the GEM Premier 3000, the GEM Premier 3500 offers new capabilities in an enhanced system, adaptable to the needs—and volume—of your hospital and lab.

- **Simple.** Maintenance-free, multi-use, disposable GEM PAKs and intuitive touchscreen menus are very easy to use.
- **Flexible.** Customized cartridge configurations and a broad test menu meet the needs of any location and any testing capacity, cost-effectively.
- **iQM.** IL's proprietary Intelligent Quality Management provides continuous, real-time quality assurance for the most accurate results, every time.
- **Total Connectivity.** GEMweb® software allows information management and real-time communication throughout the hospital.





GEM PAKs contain all components required for patient testing, are replaced every 21 days, and require no refrigeration.

## More versatility and flexibility for faster, easier, more efficient Critical Care testing.

### Self-Contained GEM PAK Cartridges

- Non-refrigerated disposable GEM PAKs include all components for patient testing, and are maintenance-free

### Intuitive Touchscreen

- Basic operation learned in minutes—simply press 'Go!' and present sample
- Easy-to-use, touchscreen displays and clear, concise menus simplify selection and customization of parameters and viewing of results

### GEMweb Connectivity, Enhanced with HL-7

- Allows wireless communication to LIS or HIS
- Patient and quality results can be viewed remotely from any networked PC

### Enhanced Features

- Larger sampling area with LED light facilitates sampling
- Barcode scanner allows rapid data input

### iQM

- Monitors all testing processes and components while providing continuous error detection, correction and documentation 24 hours a day, 7 days a week

### Complete Test Menu

- Customized cartridges include blood gases, electrolytes, metabolites, and hematocrit, with optional CO-Oximetry<sup>†</sup> and coagulation<sup>††</sup> modules

### Multiple Cartridge Configurations

Analyte Menu	Tests/PAK	Onboard Use-Life (weeks)
BG, <sup>*</sup> Hct	75	4
	150	3
	300	3
	450	3
	600	2
BG, Lytes, <sup>**</sup> Hct	75	4
	150	3
	300	3
	450	3
	600	2
BG, Lytes, Glu, Lac, Hct	75	3
	150	3
	300	3
	450	3
	600	2

\* BG = pH,  $p\text{CO}_2$ ,  $p\text{O}_2$

\*\* Lytes =  $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Ca}^{++}$

† GEM OPL™ portable CO-Oximeter

†† GEM PCL™ Plus portable coagulation analyzer



## iQM—patented, real-time, automated, continuous quality assurance system—helps to ensure optimal test results for enhanced patient care.

### Consistent Quality Assurance

- Active, continuous, real-time quality processing—even during use
- Reduces error detection time when compared to traditional (auto or manual) QC
- Ensures optimal quality control protocol at all times, regardless of time of day or level of operator training
- Generates reports for regulatory compliance
- Continuously monitors and checks all critical components in real time
  - Sensors
  - Process Control Solutions
  - Pattern Recognition software
  - Process stability

### Improved Patient Care

- Automatically and continuously monitors, detects and corrects potential errors
- Assesses functionality and initiates and documents corrective action
- Prevents reporting of results when instrument tolerance limits are exceeded
- Helps ensure the quality and accuracy of each patient result

### Reduces Error Detection Time from Hours to Minutes<sup>1,2</sup>

	pH	pO <sub>2</sub>	pCO <sub>2</sub>	Na <sup>+</sup>	K <sup>+</sup>	Ca <sup>++</sup>	Glu	Lac	Hct
iQM*	3 min	3 min	3 min	17 min	3 min	3 min	11 min	6 min	3 min
Traditional/ Auto QC	≥ 8 hr	≥ 8 hr	≥ 8 hr	≥ 8 hr	≥ 8 hr	≥ 8 hr	≥ 8 hr	≥ 8 hr	≥ 8 hr

\* Represents average time to error detection during sample processing.  
Statistical presentation of an average error detection time with 95% confidence.



# “iQM: A new standard for the future of QC.”

– James O. Westgard, PhD\*

**A published study, analyzing more than 10,000 patient samples, confirms iQM is not only valid in the research environment, but is also proven in the *clinical* setting.<sup>1</sup>**

## Study Details

Conducted to clinically validate the performance claims of iQM, as reported by Westgard *et al.*<sup>2</sup>

- 10,550 patient samples
- Four major teaching institutions
- Compared iQM-measured QC values to traditional QC results
- Calculated the average error detection time for each measured analyte

## Conclusions

Study results were published in the peer-reviewed, laboratory reference journal, *Clinica Chimica Acta*, as follows:

*“The findings from our study confirm that (a) iQM precision in a clinical setting is comparable to that found in previous studies done in a research setting, (b) the improved precision of control material in iQM is likely because the internal control fluids are sealed and not susceptible to exposure from handling, and (c) the system detects and often corrects errors in specific samples that might not be reported by traditional analytical systems...iQM provides quality control results comparable to or better than those obtained with traditional QC methods running on the GEM or other benchtop analyzers...Furthermore, the error detection capabilities that function on every sample provide an additional safeguard against reporting erroneous results due to clots or interferences.”<sup>1</sup>*

## FDA-Cleared Intended Use Statement:<sup>†</sup>

iQM is an active quality process control program designed to provide continuous monitoring of the analytical process with real-time, automatic error detection, automatic correction of the system and automatic documentation of all corrective actions, replacing the use of traditional external quality controls.<sup>††</sup>

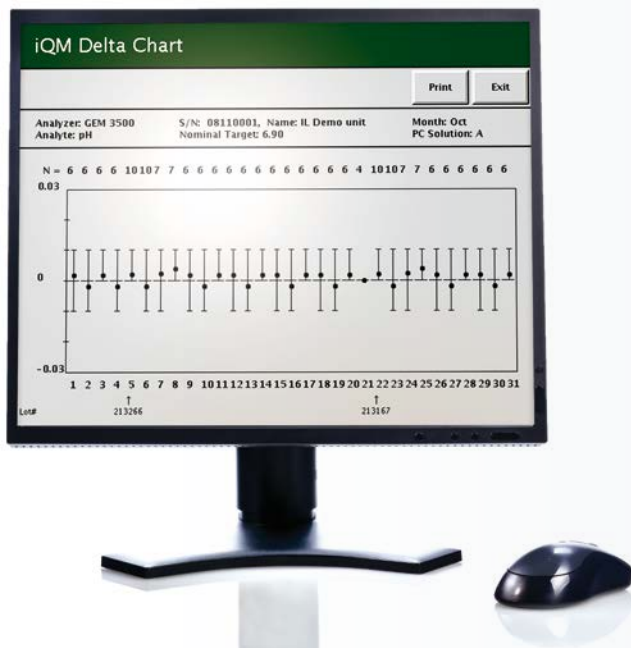
<sup>†</sup> Cleared for GEM Premier 3000 and GEM Premier 4000.

<sup>††</sup> To ensure that a total quality management system is adhered to, you should follow local, state and federal regulatory guidelines.

1. Toffaletti JG, McDonnell EH, Ramanathan LV, Tolnai J, Templin R, Pompa L. Validation of a quality assessment for a blood gas and electrolyte testing. *Clinica Chimica Acta*, 382 (2007) 65–70.

2. Westgard JO, Fallon KD, Mansouri S. Validation of iQM Active Process Control Technology. *Point of Care, The Journal of Near-Patient Testing and Technology*, 2003: Vol. 2, No. 1.

\* Professor, Pathology and Laboratory Medicine, University of Wisconsin, and developer of ‘Westgard Rules’.



**GEMweb facilitates information exchange and remote management:** iQM Delta Charts display detection and correction of errors to monitor quality and accuracy—even remotely.

## **GEMweb connectivity software—integrated information management for complete control throughout the hospital system. *Now wireless.***

- New wireless configuration option provides mobility and greater flexibility for data management
- Remotely view status for all networked analyzers, in real time
- Search and view patient and quality results on all networked analyzers from any networked PC
- Request patient demographic information from the HIS/LIS
- Connects to HIS/LIS via ASTM or HL-7 protocol

Communicate wirelessly to LIS or HIS  
with GEMweb connectivity software



## Technical Specifications

### Dimensions and Weight

#### Analyzer

H: 17.5 in, W: 13 in, D: 11.8 in, Wt: 31.2 lbs

#### PAK

H: 6 in, W: 8.5 in, D: 3 in, Wt: 4.2 lbs

### Sample Volume

135µL	BG,* Hct cartridges
135µL	BG, Lytes,** Hct cartridges
145µL	BG, Lytes, Glu, Lac, Hct cartridges (capillary mode)
150µL	BG, Lytes, Glu, Lac, Hct cartridges

\* BG = pH,  $p\text{CO}_2$ ,  $p\text{O}_2$

\*\* Lytes = Na+, K+ and Ca++

### Sample Type

Heparinized whole blood (sodium or lithium only)

### Time to Results

All tests: 85 seconds from sample introduction

### Measurement Methodology

Amperometric:	$p\text{O}_2$ , Glu, Lac
Potentiometric:	pH, $p\text{CO}_2$ , Na+, K+, Ca++
Conductivity:	Hct

### Power Requirements

Universal power input, 100–240 VAC, 50/60 Hz  
60-minute power interrupt allows transport without power.

### Temperature Control

Electrode Chamber maintained at 37°C nominal

### Data Output Port

3 RS-232 Serial I/O Ports, 1 Parallel Printer Port,  
1 Ethernet Port, 4 USB Ports

### Product Safety

Complies with IEC 610101, IEC 61326, ISTA,  
and ASTM 999.

### Interface Protocols

ASTM or HL-7 data transmission to a Laboratory,  
Hospital or third-party Information System via  
a wired or wireless connection.

### Measured Analytes<sup>†</sup>

Analyte	Displayed Ranges	Resolution
pH	6.80 to 7.80	0.01
$p\text{CO}_2$	5 to 115 mmHg <sup>††</sup>	1 mmHg
$p\text{O}_2$	0 to 760 mmHg	1 mmHg
Na <sup>+</sup>	100 to 200 mmol/L	1 mmol/L
K <sup>+</sup>	0.1 to 20.0 mmol/L	0.1 mmol/L
Ca <sup>++</sup>	0.10 to 5.00 mmol/L	0.01 mmol/L
Glu	5 to 500 mg/dL	1 mg/dL
Lac	0.2 to 15.0 mmol/L	0.1 mmol/L
Hct	15 to 65%	1%

<sup>†</sup> See Operator's Manual for complete validated ranges,  
specifications and performance characteristics

<sup>††</sup>  $p\text{CO}_2$  trending to 150 mmHg available

### Derived (calculated) Parameters

Derived Analytes	Displayed Ranges	Resolution
$\text{HCO}_3^-$	3.0 to 60.0 mmol/L	0.1 mmol/L
$\text{HCO}_3^-$ std	3.0 to 60.0 mmol/L	0.1 mmol/L
$\text{TCO}_2$	3.0 to 60.0 mmol/L	0.1 mmol/L
BE(B) ( <i>in vitro</i> )	–30.0 to 30.0 mmol/L	0.1 mmol/L
BE(ecf) ( <i>in vivo</i> )	–30.0 to 30.0 mmol/L	0.1 mmol/L
$\text{SO}_2\text{c}$	0 to 100%	1%
Ca <sup>++</sup> (7.4)	0.10 to 5.00 mmol/L	0.01 mmol/L

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