ABL90 Flex Plus – OPERATOR Training and Competency Record



On completion of the training session the participant will be able to demonstrate competence in the use of the ABL90 Flex Plus blood gas analyser.

Name (PRINT):	Dept:	

1. Overview

Participants have been provided with an overview of the analyser including:				
Printer and how to change paper	Touch screen, Barcode scanner, Power/battery indicator			
Sensor cassette and solution pack	Inlet			
Analyser software: Ready mode, parameter bar, analyser status (traffic lights), consumable status, date, time, AQURE connectivity, data logs, log on/off				

2. Pre-analytical Guidance

Participants have been informed regarding the importance of the pre-analytical steps involved to ensure a good quality sample, including:				
Use of an appropriate pre-heparinised syringe or capillary: slows down the clotting process				
Patient must be in a stable respiratory condition for 5 mins prior to sampling or wait 20 mins following change of ventilation				
Appropriate patient and sample identification: prevent mix up, loss of samples, misdiagnosis and treatment				
Visually inspect sample and expel air/bubbles. Cap syringe to avoid contact with room air (affect O_2 and CO_2 values)				
Prevention of haemolysis: positive bias to K ⁺ and negative bias to Na ⁺ and Ca ²⁺				
Analyse sample immediately and dispose of sample correctly				
A-Line sampling – withdraw adequate flush solution from the arterial line prior to sampling to avoid dilution of blood sample				
Adequate mixing of sample: reduces risk of clotting and ensures accurate Hb results. Collect correct volume of blood – minimum/maximum fill volume				
Ensure capillary is full				
How to use and benefit from end caps, mixing wire and clot catcher				
Introduce clean end of capillary				

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3. Analyser Use

Participants have been provided with a demonstration of how to analyse a sample and provided with the following information:
Logging on and off the analyser
Operator passwords/barcodes and importance of NOT SHARING
Results affected by pre-analytical errors
Analyser has been validated for whole blood samples
Interference of test results – refer to IFU
Limitation of use: only analyse human whole blood, operators must be trained, interpret results in relevant clinical context, refer to IFU for full details
Who to contact if the analyser is not working
Automated calibration and quality control
Correct procedure for cleaning and decontamination
Importance of reporting analyser adverse incidents

4. Competency Check

The trainer has observed the participant competently demonstrate the following:			
Identify the analyser is ready to use	Yes	No	
Log on/off	Yes	No	
Introduce sample in a safe and appropriate manner	Yes	No	
Input patient information	Yes	No	
View results (screen/printed), understands error messages, ranges and critical limits		No	
Retrieve previous patient results	Yes	No	

Operator	Trainer	
Name:	Name:	
Signature:	Signature:	
Date:	Date:	