Women, Infants, and Children (WIC) Competent Professional Authority (CPA) Training Program

Laboratory Procedure Manual



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	Chapter 1. Introduction
Purpose	This manual is designed to be used as a reference tool to help WIC staff safely and accurately perform hemoglobin tests. Use the hyperlinks in the Table of Contents above to quickly locate the information needed.
Anemia and Hemoglobin Testing	The Nevada WIC Program conducts hemoglobin testing on its participants as a screening tool for anemia, which is a condition resulting from low levels of healthy red blood cells. Its symptoms include poor appetite, tiredness, weakness, developmental delays and learning problems. Iron deficiency anemia is the most common type of anemia and represents a significant global health concern. Even though the hemoglobin testing performed within Nevada WIC clinics is not able to be used to diagnose anemia, let alone the specific type of anemia, the obtained hemoglobin values can still be used for participant education and community healthcare referrals.
Types of Hemoglobin Testing	The Nevada WIC Program utilizes two different types of hemoglobin tests. The Masimo Pronto is a non-invasive hemoglobin testing device that is required to be used on WIC participants aged 2 years and older. Capillary sampling is the other, more invasive type of hemoglobin testing completed within the Nevada WIC program, and is only designed to be used on participants under the age of 2, but can be used on participants over the age of 2 if no hemoglobin value is able to be obtained with the Masimo Pronto. Capillary sampling involves taking a small sample of a participant's blood and measuring the amount of hemoglobin it contains using a hemoglobin analyzer. We'll discuss both types of hemoglobin tests in future chapters of this manual.
Individual and Local Agency Hematology Training Requirements	 In order for CPAs to complete unsupervised hemoglobin blood tests within Nevada WIC clinics, it's mandatory that they: Complete the Hemoglobin LMS course, and pass the corresponding post test Complete the Hemoglobin Guidebook, and discuss their answers with their local agency trainer Complete the required Hematology observations as indicated in the New Employee Training Plan In addition to these individual requirements, each local agency must also designate one or more trainers to provide hematology training and observations with staff. Designated trainers must be trained and re-certified at least once every three years by attending Anthropometric / Hematology Trainings provided by the Nevada WIC State office. Local agencies must also receive and maintain a Laboratory Certification in order to security hematology is a security of the security of the security in tests.

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Chapter 2. Universal / Standard Precautions

Universal
 Precautions
 The umbrella term "Universal Precautions" refers to approaches to minimize
 occupational exposure to bloodborne pathogens. A key component of taking
 Universal Precautions is to treat the blood and bodily fluid of others as if it contained
 a bloodborne pathogen such as Human Immuno-deficiency Virus (HIV), Hepatitis B
 Virus (HBV), or Hepatitis C Virus (HCV), regardless if the person has a known or
 suspected bloodborne pathogen.

There are 2 types of universal precautions: "Standard Precautions" and "Transmission-Based Precautions". Standard Precautions are a basic set of precautions that need to be taken whenever coming into potential contact with blood or other bodily fluids. Transmission-Based Precautions are steps that should be taken in addition to "Standard Precautions" if the person you're coming into contact with has a known or suspected condition that can be spread through the air (e.g. chickenpox), through droplets from coughs or sneezes (e.g. influenza) or by coming into contact with the skin (e.g. herpes simplex virus).

There are four main components of Standard Precautions that are required to be followed in all Nevada clinics: Hand Hygiene, Personal Protective Equipment, Sharps Safety, and Cleaning and Disinfection. We'll discuss each of these topics in the following sections of this chapter.

Transmission-Based Precautions will not be discussed in this document. Speak with your supervisor if you ever believe that additional Transmission-Based Precautions are necessary in order to protect your health and safety.

Standard Precautions (Hand Hygiene) The Centers for Disease Control and Prevention (CDC) defines Hand Hygiene as "Cleaning your hands using either handwashing (washing hands with soap and water), antiseptic hand wash, antiseptic hand rub (i.e. alcohol-based hand sanitizer including foam or gel), or surgical hand antisepsis." In the WIC program, there are two methods generally used for hand hygiene: alcohol-based hand sanitizer and washing hands with soap and water. The table below describes examples of when each is recommended.

Source: CDC

When to Use Alcohol-Based Hand Sanitizer	When to Wash with Soap and Water
Immediately before touching a patient	When hands are visibly soiled
Before moving from work on a soiled body site to a clean body site on the same patient	After caring for a person with known or suspected infectious diarrhea
After touching a patient or the patient's immediate environment	After known or suspected exposure to spores (e.g. B. anthracis, C difficile outbreaks)
After contact with blood, body fluids, or contaminated surfaces	
Immediately after glove removal	

Alcohol-Based Hand Sanitizer vs Soap and Water

Source: CDC

How to Use Alcohol-Based Hand Sanitizer:

Apply the amount of product recommended by the manufacturer in your hands, and continuously rub the product over all parts of your hands until your hands feel dry (approximately 20 seconds).

How to Wash Hands with Soap and Water:

Wet your hands first. Apply the amount of product recommended by the manufacturer in your hands, and vigorously rub it over all parts of your hands for at least 20 seconds. Rinse your hands with water and dry them with disposable towels. Use a disposable towel to turn off the water.

Source: CDC

Standard	Wearing personal protection equipment (PPE) is essential in order to help
Precautions	protect your safety, and the safety of others while working in the lab and must
(Personal	be worn whenever it can be reasonably assumed that contact with blood or
Protection	other potentially infectious materials may occur. Gloves are by far the most
Equipment)	common form of PPE used in WIC clinics, so they are the only form of PPE
	discussed in this manual. However, you must wear additional forms of PPE (e.g.
	gloves, masks, goggles, face shields etc.) if required by your local agency's
	policies. Gloves provide an additional layer of protection, helping to stop the
	possible spread of infection from the wearer and others. However, wearing
	gloves should never replace hand hygiene. At WIC, gloves should be worn in the
	following situations:

- When starting to perform a capillary sampling test, perform hand hygiene and cover any breaks in the skin with bandages before applying gloves.
- Change gloves in between capillary sampling tests, even if performing the tests on members of the same family.
- Immediately change gloves and perform hand hygiene if gloves ever become damaged while performing a capillary sampling test.
- If gloves become visibly soiled with blood or other body fluids, carefully remove gloves, and perform hand hygiene after completing the capillary sampling test.

To remove gloves, take the following steps:

- 1. Using a gloved hand, pinch the palm area of the other gloved hand and peel off the first glove.
- 2. Using the gloved hand only, roll-up the removed glove into a ball, and hold it in the palm of the gloved hand.
- 3. Using your ungloved hand, slide a finger under your gloved hand and peel off the second glove over the first rolled-up glove.
- 4. Discard both gloves in the waste container designated by your local agency.

Source: CDC

Standard Precautions (Sharps Safety) Retractable lancets are used during capillary sampling tests. Even though retractable lancets contain safety features (e.g. retractable needles) that help to prevent the user from accidental needle sticks, it's still important to take safety precautions. These include:



- Focusing on the task at hand
- Keeping fingers away from the opening of the lancet
- Immediately disposing of used lancets and microcuvettes in a "sharps" container.

There are a variety of styles, and all are clearly marked with a biohazard symbol (see figure to the left). The container must be rigid, puncture-resistant, leak-proof, and disposable with a locking lid.

When this container is filled to the acceptable level, it must be properly disposed of as biohazardous waste.

Standard Precautions (Cleaning and Disinfection) There are different receptacles for trash that is generated in the lab.

- Lancets and microcuvettes must be disposed of in sharps containers.
- Throw away trash that is <u>saturated with blood</u> in a red, plastic biohazard bag. All waste that is <u>saturated and dripping with blood</u> must be, sterilized, incinerated or, chemically disinfected prior to mixing and disposing with ordinary waste. Find out from your supervisor how to handle biohazardous waste since it must be decontaminated before it can be disposed of in a landfill.
- Waste, such as lint-free tissue, alcohol preps, gloves, bandages & wrappers, that contain blood but are not <u>dripping</u>, can be discarded in a regular trash bag if there are no means for biohazard waste disposal.

It's important that all lab work surfaces are cleaned thoroughly at least once a day, but also must be cleaned immediately if they are visibly soiled. Lab work surfaces must be cleaned using either a 10% bleach solution or a disinfectant approved for use in your local agency.

- To prepare a 10% bleach solution, mix 1-part household bleach with 9 parts tap water. Store at room temperature in an opaque plastic bottle labeled "10% Bleach."
- The date of preparation and the expiration date should be clearly marked on the outside. The expiration date is seven calendar days after the date the solution was prepared.

Note: Keep the sharps container, biohazard bag, lab trash, and all cleaning products out of the reach of children.

Exposure Control Plan

A written exposure control plan must be kept in an easily accessible location at each WIC clinic. The plan in your clinic will include a copy of your local agencies' policies and procedures for employee safety and a procedure for reporting accidents. Speak with your supervisor to ensure that you know where to locate the exposure control plan in your clinic and familiarize yourself with its contents.

- If blood touches your skin or hair, wash the area with soap and water immediately.
- If blood enters your eye(s), flush them with water immediately.
- If you are accidentally stuck by a used lancet, have blood touch your skin/hair, have blood enter your eye(s), or if you have any reason to believe that you may have been exposed to an infectious substance, notify your supervisor immediately to discuss seeing a licensed healthcare provider to be tested for infectious diseases such as HIV, HBV, and HCV.

Chapter 3. Capillary Sampling Equipment

Lancets	Lancets are small, single-use devices that are designed to pierce the skin, allowing a blood sample to be collected with a microcuvette. There are various lancets available depending on incision depth and intended puncture location (e.g. finger or heel). Check with local agency trainers / supervisors to ensure that the correct lancets are used based on differing situations. In addition, refer to the chart below for lancet requirements for each WIC participant category.				
		Infants certified at < 9 months old	Infants certified from 9 to 11 months	Children (1 to 5 years)	Women (Pregnant / Postpartum / Breastfeeding)
	Finger Puncture Depth	N/A	1.5 mm or less	1.5 mm or less	2.4 mm or less
	Heel Puncture Depth	N/A	2.0 mm or less*	N/A	N/A
	*May only b	e performed on inf	ants who have not yet	begun to learn to wa	
Microcuvettes	Microcuv which the hemoglot	ettes are man w were desigr pin analyzer(s)	ufactured to be ned. Therefore, c used in your loc	used only in the only use the mic cal agency clinic	hemoglobin analyzer for rocuvettes designed for the
HemoCue 201+ Microcuvettes	HemoCue work insid yellow co	201 Microcu de of the Hem lor at the tip c	vettes contain a oCue 201+ whic of each microcuv	special reagent h is noticeable o vette (see pictur	designed to due to its re).
Hb 201 HemoCue	Once the vial of HemoCue 201+ Microcuvettes has been opened, they are able to be used up to <u>90 days</u> afterwards. Upon opening a vial for the first time, write the date the vial was opened, as well as the date that the microcuvettes must be used by on the side of the vial.				
HemoCue 201+ Microcuvettes	In addition, always use microcuvettes by the expiration date written on the side of the each vile, regardless of whether the vial has been opened or not.		LOT 6110751 2008 11 08		
HemoCue 301 Microcuvettes	Unlike Ho Microcuv are comp	emoCue 201 vettes do not pletely clear (+ Microcuvette contain any re (see picture).	es, HemoCue 3 eagent, so thei	01 r tips



HemoCue 301 Microcuvettes Once the vial of HemoCue 301 Microcuvettes has been opened, they are able to be used up to <u>24 months</u> afterwards. Upon opening a vial for the first time, write the date the vial was opened, as well as the date that the microcuvettes must be used by on the side of the vial.

In addition, always use microcuvettes by the expiration date written on the side of the each vile, regardless of whether the vial has been opened or not.



Hemoglobin	The purpose of hemoglobin analyzers is to measure the amount of hemoglobin
Analyzers	contained in a blood sample. The Nevada WIC Program utilizes 2 different types of
	Hemoglobin Analyzers: HemoCue 201+, and HemoCue 301.



The HemoCue 201+ hemoglobin analyzer is a portable instrument with a sliding cuvette holder and display screen. It can operate on AC power (AC adapter included) or 4 AA batteries.

HemoCue 201+Hemoglobin
MeasurementTo obtain a reading, turn the device on by pressing the power button, and rotate
the cuvette holder into the loading position. Place a filled microcuvette in the
cuvette holder and rotate the cuvette holder into the measuring position. The
measurement takes between 40 and 240 seconds to display, and the hemoglobin
value is expressed as grams per deciliter or g/dl. After reading the sample, the
value will remain displayed on the screen as long as the cuvette holder is in the
measuring position. Once the cuvette holder is rotated back to the loading position
the previously recorded value will be erased.

HemoCue 201+ Cuvette Holder Cleaning	Clean cuvette holder after each day of use by following these steps:				
	 Turn off the analyzer. Pull the cuvette holder out to the loading position. 				
	 While pressing the catch carefully pull out the cuvette holder. 				
	3. Remove the cuvette holder from the analyzer. It will come off the stainless-steel pin it rotates on.				
	 4. Clean the cuvette holder with alcohol (20-70%) or mild detergent. 				
	Wait 15 minutes to air dry before replacing the cuvette holder (making sure that it's locked in place by the catch) and using the HemoCue 201+.				
HemoCue 201+ Optical Parts Cleaning	The optronic unit inside the HemoCue 201+ (see picture) only needs to be cleaned when an error code appears on the display (e.g. 901 or 902), or if the hemoglobin values obtained are higher or lower than anticipated.				
	To clean the optronic unit, first follow steps 1-3 as outlined above for cleaning the HemoCue 201+ cuvette holder.				
	 4. Push a HemoCue Cleaner, or a lint free cotton swab, moistened with alcohol (20-70% without additives) or water, inside the device as far as possible. Move it from side to side 5 to 10 times. Do not attempt to use an alcohol prep pad to clean inside the device. 				

	 Push and pull the HemoCue Cleaner or cotton swab in and out of the device opening (on the left side where the optronic unit is located). 		
	6. Remove the HemoCue Cleaner or cotton swab from the device. If the HemoCue Cleaner or cotton swab is stained, repeat steps 4 and 5.		
	Wait 15 minutes before replacing the cuvette holder and using the HemoCue 201+.		
HemoCue 201 Supply Ordering	Local agencies are responsible for procuring all related supplies for capillary sampling.		
	HemoCue 201 hemoglobin analyzers, and microcuvettes will be purchased by local agencies directly from HemoCue. Please place your order with:		
	Sales Support Email: <u>insidesales@hemocue.com</u> Phone: 800-881-1611		
	HemoCue Microcuvette Ordering Information		
	Part Number 111716 - Box of 200 (4 vials of 50 each) HemoCue 201+ Microcuvettes		
	For agencies that need smaller quantities: Part Number 111715 – 100 individually wrapped HemoCue Microcuvettes		
HemoCue 201+ Troubleshooting	When the analyzer is turned on, it will perform a self-test to verify the performance of the optronic unit. If the self-test fails, or if experiencing any other issue, troubleshooting may be required to discover and resolve the problem.		
	For troubleshooting information regarding the HemoCue 201+, review the HemoCue 201+ Troubleshooting Guide (Appendix B).		
	If experiencing problems that are not able to resolved by taking the steps outlined in the troubleshooting guide, speak with your supervisor or call HemoCue Technical Support at 1-800-426-7256 or email: <u>technicalsupport@hemocue.com</u> .		
HemoCue 201+ User Manual	For additional information regarding the HemoCue 201+, review the HemoCue 201+ User Manual, which can be viewed by visiting the following link:		
	https://www.hemocue.us/wp-content/uploads/2020/07/Manual_Glu_201.pdf		

1

HemoCue 301 Hemoglobin Measurement



HemoCue 301+ Cuvette Holder Cleaning

HemoCue 301

Optical Parts

Cleaning

The HemoCue 301 hemoglobin analyzer operates in much the same way as the HemoCue 201+ hemoglobin analyzer. However, instead of measurements taking 40 to 240 seconds to display, once the filled microcuvette has been placed in the cuvette holder and rotated into the measuring position, the HemoCue 301 will display the measured hemoglobin value within 3 seconds.

 Turn off the analyzer. While the cuvette holder is still in the loading position, press the catch using a

pointed object.

Clean cuvette holder after each day of use by following these steps:

- 2. While still pressing the catch, carefully pull out the cuvette holder.
- 3. Clean the cuvette holder with alcohol (20-70%) or mild detergent.
- 4. Wait 15 minutes to air dry before replacing the cuvette holder (making sure that it's locked in place by the catch) and using the HemoCue 301.

E01, E02, E08, E10-30), or if the hemoglobin values obtained are

higher or lower than anticipated.



2

3



To clean the optronic unit, first follow steps 1-2 as outlined above for cleaning the HemoCue 301 cuvette holder.

- Push a HemoCue Cleaner, or a lint free cotton swab, moistened with alcohol (20-70% without additives) or water, inside the device as far as possible. Move it from side to side 5 to 10 times. Do not attempt to use an alcohol prep pad to clean inside the device.
- 4. Push and pull the HemoCue Cleaner or cotton swab in and out of the device opening (of the left side where the optronic unit is located).
- 5. Remove the HemoCue Cleaner or cotton swab from the device. If the HemoCue Cleaner or cotton swab is stained, repeat steps 3 and 4.
- 6. Wait 15 minutes to air dry before replacing the cuvette holder (making sure that it's locked in place by the catch) and using the HemoCue 301.



Part Number 111809 - Box of 200 (4 vials of 50 each) HemoCue 301+ Microcuvettes





When the analyzer is turned on, it will perform a self-test to verify the performance of the optronic unit. If the self-test fails, or if experiencing any other issued, troubleshooting may be required to discover and resolve the problem.
For troubleshooting information regarding the HemoCue 301, review the HemoCue Hb 301 HemoCue Troubleshooting Guide (Appendix C).
If experiencing problems that are not able to resolved by taking the steps outlined in the troubleshooting guide, speak with your supervisor or call HemoCue Technical Support at 1-800-426-7256 or email: <u>technicalsupport@hemocue.com</u> .
For additional information regarding the HemoCue 301, review the HemoCue 301
User Manual, which can be viewed by visiting the following link:
https://www.hemocue.us/wp-content/uploads/2020/08/HB-301_Operating- Manual_US.pdf

Chapter 4. How to Perform a Capillary Sampling Hemoglobin Test

Participant Consent	Prior to any lab testing, have the authorized representative provide consent to perform hemoglobin test. If testing is being done by third party, provide authorized representative with WIC Hemoglobin test referral slip and record 'Results Deferred' which will give a 90-day window for getting results back from participant(s).			
Select Work Area	Select a work area to complete the capillary sampling test.			
	An ideal work area:			
	• Is clean			
	 Ensures participant and staff safety 			
	 Has a surface which is smooth, free of cracks, and washable 			
	Ensures participant privacy			
	 Is away from noise and confusion 			
	Has a chair and table			
Fxplain	Explain the procedure to the Authorized Representative in simple terms			
Procedure				
	Example:			
	"I am going to make a little poke in your child's finger to get a few drops of blood to			
	put into this little container. Then I am going to put it into this machine to find out			
	how much hemoglobin it has in it."			
	Reassure them that you will make it as quick and painless as possible, but also be			
	that it may burt a little, but only for a moment			
	that it may null a little, but only for a moment.			
Perform Hand	Perform Hand Hygiene (either by washing hands or using an alcohol-based hand			
Hygiene and	sanitizer) before putting on unused gloves.			
Put on Gloves				
Assemble	Assemble all necessary supplies in order to perform the capillary sampling test.			
Supplies				
	Hemoglobin Analyzer			
	Alcohol prep pads			
	Sterile lancet Lint free tissues (conitizing wines or Course rade			
	 Lint-free tissues/sanitizing wipes of Gauze pads Microsuvette (remove 1 at a time & receal container) 			
	 Bandages (not for children under age 2) 			
	 Sharps container 			
	Biohazard bag			
	 10% bleach solution or EPA-registered disinfectant 			

Daily Steps for Performing Hemoglobin Tests Continued

Choose Site	The ring and middle fingers are the primary testing sites for all WIC participants for whom capillary sampling is indicated. Avoid using fingers with rings on them. The heel may only be used for infants 9-11 months of age who have not yet began to learn to walk.
Position Participant	When performing a fingerstick, position the participant comfortably seated with their arm extended, (but lower than their heart), with their palm facing up.
	When performing a heel stick, position the infant laying comfortably on their back.
Warm the Site (If Necessary)	The site should not be cold, blue, swollen or calloused. If cold, warm the site by holding it in your hands, rubbing the site for a minute, or by have the Authorized Representative wash the site vigorously with warm running water and soap.
Cleanse the Site	 Cleanse the site thoroughly with an alcohol pad (unless the Authorized Representative already washed the site with warm water and soap). Wipe the site with a tissue or lint-free wipe. <u>Be sure skin is dry.</u> Do not touch the prepared site after cleaning.
	<u>Note</u> : Pooled alcohol at the puncture site will dilute and hemolyze the blood, giving a lower reading, if the skin surface is not dried completely.

Hold the Site



When performing a fingerstick, apply gentle pressure near the participants first finger joint.

When performing a heel stick, grasp the heel between your thumb and forefinger with your other fingers underneath the infant's calf. Apply a small amount of pressure to flex the foot back.

Puncture



If performing a fingerstick, puncture off the centerline of the participant's fingertip using a lancet. (See figure to the left.)

If performing a heel stick, puncture only on the medial or lateral side of the bottom surface of the heel using a lancet. (See figure to the left.)

Note: If the lancet is blade-shaped, it should be placed perpendicularly to the whorls of the fingerprint/footprint, so the blood is more easily collected into the microcuvette.

Fill the Microcuvette	To ensure accuracy, wipe away the first two to three drops of blood. This will stimulate spontaneous blood flow, resulting in a better sample. If necessary, press gently again with thumb and forefinger until another drop of blood appears. Avoid squeezing or milking the site as this can dilute the blood and gives a false low reading.
	All drops should be large enough, so they "sit" on top of the heel or finger like a bead. Ensure that the drop of blood is big enough to fill the entire microcuvette tip .
	Touch the tip of the microcuvette, pointing downward, into the middle of the blood drop so the microcuvette touches the skin. Allow the microcuvette to fill in one step. The microcuvette will fill itself automatically. Never "top off" the microcuvette if it doesn't fill in the first attempt.
	If the microcuvette does not fill completely on the first attempt, or if air bubbles are visible, discard the microcuvette, wipe the puncture site and allow a new, larger bead of blood to form before collecting blood into the new microcuvette.
Prepare the Cuvette for Testing	Wipe off the flat outside surfaces of the microcuvette using gauze or a lint-free wipe. Avoid wiping the open slit of the microcuvette.
Measuring Hemoglobin Value	Measure the hemoglobin value in the microcuvette by following the manufacturer's directions for the hemoglobin analyzer used in your local agency's clinic. (See Chapter 3.) Immediately record the value obtained, so that it can be entered into the WISH Assessment, Blood Panel.

Seal and Bandage Site	Place dry gauze or lint-free tissue over the puncture site and apply gentle pressure until the wound is clotted. Elevating the hand or foot above the level of the heart will help to stop the blood flow. Apply the bandage. <u>Note</u> : Do not use bandages on the finger of a child less than two years of age to prevent potential ingestion and choking.
Cleanse Surface	If any blood spills on the hemoglobin analyzer, work surfaces, or your skin, clean with a 10% bleach solution or disinfectant spray immediately.
Disposal of Supplies	Wastebasket: Throw away any paper wrappers, alcohol preps, gauze, lint-free tissues, gloves and other supplies which are <u>not saturated</u> and dripping with blood in a wastebasket.
	Biohazard bag: Throw away any supplies that are saturated and dripping with blood in the red biohazard bag.
	Sharps container: Throw away all lancets and used microcuvettes in the sharps container.
Remove Gloves and Perform Hand Hygiene	Remove and discard gloves after disposing the other supplies. If your gloves are contaminated with blood, place them in the biohazard bag with the other supplies. Perform hand hygiene after removing your gloves.
Perform Another Capillary Sampling Test (If Necessary)	If performing another capillary sampling test, begin by putting on a new pair of gloves, and assembling new supplies needed for another test.

Chapter 5. Masimo Pronto Equipment



The Masimo Pronto measures total hemoglobin concentration of the blood via a sensor attached to the machine that emits multiple light waves through the capillary bed of the fingertip. Changes in the light absorption can be used to measure the functional arterial oxygen saturation of the blood, from which the hemoglobin concentration can be calculated.

Masimo Pronto Pulse CO-Oximeter



Displays parameter/measurement numeric hemoglobin values once a spot check test is complete.

Parameter/ Measurement Numeric Display



Flashes with patient's pulse reading (BPM) during spot check test period

Pulse Indicator



Incrementally illuminates upward after a SpHb spot check has been initiated. This indicates progress towards completion of a SpHb spot check. A fully illuminated spot check progress indicator indicates a completed spot check.

Spot Check Progress Indicator



Displays parameter/measurement label data once a spot check test is complete.

Battery Level Indicator

0

Low SIQ

Low SIQ Indicator

Sensor Use Indicator Battery charge level is indicated by four LED indicators. All four indicators will be lit when the batteries are full, with fewer indicators being lit as the batteries lose their charge.

The Low SIQ Indicator illuminates when the signal is too low to complete the test, indicating that it will be necessary to restart the test.

The Sensor Use Indicator illuminates to display the approximate number of uses remaining for the attached sensor. The bottom LED will turn red when the remaining uses for the connected sensor are low. The approximate number of sensor uses remaining is displayed upon power up (if a sensor is attached) and when a sensor is connected.

Pronto Controls



CO-005128



Powers the instrument on or off. Press the button once to power on, press and hold the Button for 2 seconds to power off.



SpHb Button



SpHb Button - Press to initiate total hemoglobin (SpHb) spot check information on display or to display a Total Hemoglobin (SpHb) spot check test. If configured, oxygen saturation (SpO2), perfusion index (PI) and pulse rate (PR) are automatically displayed. When in the configuration menu, pressing this button will confirm a menu setting and navigate to the next menu option.

Use the Up and Down arrow buttons to scroll between parameter or measurement spot check results. When in the configuration menu, use the Up and Down arrow buttons to scroll through menu setting options.

Perfusion Index



Perfusion Index (PI) **Perfusion Index (PI)** is a numeric indication of the pulse strength at the measurement site. The PI numerical value on the Masimo Pronto ranges between 0.02% and 20%. To successfully conduct a test the "PI" perfusion index must measure 1.0 or greater

- PI varies between monitoring sites and from patient to patient, as physiologic conditions vary
- Lower values indicate lower perfusion meaning decreased blood flow
- Several possible causes (e.g. cold hands, Vasoconstriction (narrowing of the blood vessels), Hypovolemia (decreased blood volume), Peripheral Vascular Disease, etc.)

4-6% of the population have subclinical issues that will prevent a good reading • (carboxyhemoglobin, other irregularities) Monitor the trend of the PI for changes in physiologic conditions Battery Compartment located in back panel • Holds 4 "AA" alkaline or rechargeable batteries Operates up to 8 hours (usually 200-250 tests per battery replacement) The Pronto cannot be plugged in, it only uses battery power Battery Compartment **Disabling Pulse** Pulse rate and oxygen saturation are features of the Masimo Pronto that are not Rate and utilized by the WIC program. Disabling these features is optional but will help to Oxygen prolong battery life. To disable them take the following steps. Saturation (Optional) 1. Press Up and Down arrow buttons simultaneously for 5 seconds 2. Press the Sp/Hb button repeatedly until "O2" (Oxygen Saturation) appears on the display. 3. Press either the up or down arrow to turn "off" oxygen saturation 4. Press SpHb to confirm your choice 5. "Pr" (Pulse Rate) will appear next in the display 6. Press either the up or down arrow to turn "off" pulse rate. 7. Press SpHb to confirm your choice

Patient Cable

One-foot patient cable that connects to rainbow DCI and rainbow DCIP digit sensors. To connect the Patient Cable, plug the red end of the cable into the port on top of the Masimo Pronto.

Adult-sized sensor that can be connected to the RC1 Patient Cable when performing



tests on adults. Includes a slender digit gauge (see slender digit gauge description below).

Rainbow DCI Digit Sensor



Rainbow DCIP Digit Sensor Pediatric-sized sensor that can be connected to the RC1 Patient Cable when performing tests on children or women with slender fingers. The finger of a child must be long enough and wide enough reach the end of the sensor. If the child's finger isn't long enough, the hemoglobin value will need to be obtained via a capillary sampling test.



Masimo ProntoFor Masimo Pronto troubleshooting, review the Masimo Pronto TroubleshootingTroubleshooting(Appendix D).									
	If experiencing problems that are not able to resolved by taking the steps outlined in the troubleshooting guide, speak with your supervisor or call Masimo technical support at 1-800-326-4890 or email: <u>TechSupport-US@masimo.com</u> .								
Masimo Pronto User Manual	For all other information regarding the Masimo Pronto, Review the Masimo Pronto User Manual, which can be viewed by visiting the following link:								
	pronto-manual.pdf								

Chapter 6. How to Perform a Hemoglobin Test Using the Masimo Pronto

Participant Consent	Prior to any hematology testing (including the completion of a hemoglobin blood test using the Masimo Pronto), have the Authorized Representative consent to test(s). Make sure to provide information on different test options.								
Choose a Location	The selected location for using the Pronto device should be a comfortable area, free of excessive noise or distractions, such as a private office space. It is not recommended to conduct hemoglobin tests with the Pronto device in a common lab area where other measurements or capillary sampling tests are being completed. The sights and sounds of other participants who may be stressed or upset by lab procedures may make it more difficult to keep participants relaxed and motionless while using the Pronto.								
Explain Procedure	 Explain the procedure to the Authorized Representative in simple terms. Example: "I am going to place a sensor on your child's finger to find out how much hemoglobin is in his/her blood. Your child will need to sit completely still for a couple of minutes in order to complete the test, but don't worry, it's completely painless." 								
Perform Hand Hygiene	Perform Hand Hygiene (either by washing hands or using an alcohol-based hand sanitizer).								
Position Participant	Position the participant comfortably seated with their arm extended, (but lower than their heart), with their palm facing down.								
Selecting Child or Adult Sensor	 Measure the participant's finger size (diameter) at the cuticle, using the sensor size gauge, to determine the correct sensor size. Slide the Slender Digit Gauge circle on the digit as shown in Figure A. If the gauge circle stops at any point of the nail bed before the cuticle, the sensor can be used on that digit. If the gauge slides past the cuticle as shown in Figure B, the digit is too slender for this sensor. Select a different digit or use a pediatric / slender digit sensor on the patient (see instructions below for disconnecting and connecting sensors). Remove the gauge from the digit before sensor application Fig.B 								

Disconnect and Connect Sensor (If necessary)	 To disconnect the sensor, lift the protective cover (labeled 3) to gain access to the sensor connector (labeled 1). Pull firmly on the sensor connector to remove from the patient cable (labeled 2). To avoid damage, pull on the sensor connector, not the cable. 							
	 To connect the sensor, properly orient the sensor connector and insert the sensor connector completely into the patient cable connector. 							
	Completely close the protective cover.							
Sensor Site Selection	 Remove anything from participant's arm that can impede blood flow to the sensor site, such as restrictive garments, accessories, purses, backpacks, watches, and jewelry. 							
	 Select the participants testing finger in the following priority: 							
	 Ring or middle finger The thumb may be used instead for children at least 2 years old with very small fingers. 							
	Note: Do not use pinky or index fingers. Do not use anatomically abnormal fingers (e.g. clubbed, deviated, etc.).							
	• You can use the Pronto on sites with nail polish and/or acrylic nails. Some types of nail polish may interfere with the test and prevent a reading, but it is advised to attempt tests with nail polish. If the Pronto displays a result from a site with nail polish, the result is within the specifications for accuracy required by FDA. If a reading fails, it does not reduce the number of tests left on the sensor.							

Cleanse the Site Cleanse the site thoroughly with an alcohol pad.

Sensor Placement	 Place the participant's selected finger inside of the sensor. Make sure fingertip is inserted all the way and touching the finger stop inside the sensor (allowing long fingernails to extend beyond the finger stop). Examine the finger while placed in the sensor to ensure the emitter and detector are directly aligned on top of each other and there is no gap between the sensor and fingertip. See figures C and D for proper alignment. 								
	Cable Position: Run the cable from sensor over the top of the hand without kinks or twists so the cable does not pull on the sensor. (See figure E) Kinks and twists in the cable may increase the time needed to complete tests and cause tests to fail more often.								
Light Shielding	Cover the sensor with a hand (either your hand or a caregivers) or cloth to shield the sensor from excessive light. Excessive light may increase the time needed to complete tests and cause tests to fail more often.								
Starting a Hemoglobin Test using the Masimo Pronto	 Press the power button to activate the Pronto. a. The Pronto will run a Self-Test. b. The Pronto will display the number of sensor tests remaining on the sensor. (New sensors will only function for a specific number of tests (200, 400, or 1000). c. When the Pronto reads SEN OFF it is ready for testing. d. Scrolling zeros appear to indicate sensor initialization. e. Dashes appear to indicate testing has begun. f. The letters "Pl" (perfusion index) will automatically display after start up in the lower window display. The upper window display will show numerical reading of the Pl. g. Pulse indicator light will flash with each heartbeat. h. It will take 1 to 3 minutes for the Pronto to acquire and display a hemoglobin measurement. i. The Spot Check Progress indicator incrementally illuminates from bottom to top and an <u>audible tone</u> will sound when the hemoglobin measurement is ready to display. 								
Display Hemoglobin Test Result	Press SpHb button when ready to view hemoglobin results (hemoglobin data will available for 5 minutes after the test has completed by depressing either the up or down arrow).								
Record Hemoglobin Test Result	Immediately record the value obtained, so that it can be entered into the WISH Blood Panel.								

Remove Sensor	Remove the sensor from the participant's finger.
Turn off Masimo Pronto	Turn off the Masimo Pronto by holding down the Power Button for 2 seconds. (It will also turn off automatically if not used for more than 1 minute.

Chapter 7: Hemoglobin Value Results

Different Hemoglobin Values	The CDC has established hemoglobin cutoff values below which someone is considered at risk for anemia (see Appendix A). Hemoglobin cut-off values vary, depending upon participant smoking status, level of elevation, and category. There are different tables based on elevation level. From there, each table is broken down by smoking status, and category (pregnant [by trimester], postpartum, children and infants). One of these tables should be posted in your lab for you to evaluate your participant's test result. After obtaining hemoglobin values (via the Masimo Pronto or capillary sampling) and entering them into the WISH system, there are three possible courses of action depending on the risk assigned, and the hemoglobin value obtained. Each of these three courses of action will be reviewed in this chapter.
Normal Hemoglobin Value	If the hemoglobin value obtained is above the "Low Hemoglobin" range based on the CDC cutoff values for hemoglobin (Appendix A), the participant's hemoglobin is considered normal, so no risk code will be generated when the value is entered into WISH, and no further action is necessary.
"Low Hemoglobin" Value	If the hemoglobin value obtained is within the "Low Hemoglobin" range based on the CDC cutoff values for hemoglobin (Appendix A), risk code 201 will be generated when the value is entered into WISH. A referral to RD/Nutritionist is required if just below cutoff value. However, a referral to a healthcare provider may be provided to determine the cause of the low hemoglobin and to discuss possible treatment options. Since iron deficiency anemia is by far the most common cause of low hemoglobin values, it may also be useful to attempt to determine the potential cause and possibly discuss tips to increase dietary iron intake.
"Very Low Hemoglobin " Value	A very low hemoglobin value (see Appendix A) is a serious medical concern and is life- threatening. If a very low hemoglobin value is obtained, have a second hemoglobin test completed (preferably by another staff member), using a finger on the participant's other hand. The higher of the two hemoglobin values shall be entered into the WISH computer system. Local Agencies shall also establish a referral plan (in conjunction with county/ agency Health Program Officers), and train staff to ensure that all participants with confirmed very low hemoglobin values are referred for an immediate medical evaluation, either with their primary care provider or at an emergency medical center. All referrals shall be documented in WISH. It's also required to refer participants with very low hemoglobin values to a medium risk nutritionist or high-risk nutritionist since it is a medium risk code (201.1).

Cutoff Values from Hig	h Risk Manual Hct Table
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		Category									
Table 201 – A		1 ^{et} Trimester	2 nd Trimester	3 rd Trimester	Nonpreg 12 - < 15 yrs	Nonpreg 15 - < 18 yrs	Nonpreg ≥18 yrs	Infants 0 - < 6 mo	Infants 6 - < 12 mo	Child 1 - < 2 yrs	Child 2 - < 5 yrs
					Hct <						Hct <
	Nonsmokers	33.0	32.0	33.0	35.7	35.9	35.7		33.0	32.9	33.0
No altitude	Up to < 1 pack/day	34.0	33.0	34.0	36.7	36.9	36.7				
adjustment	1 - 2 packs/day	34.5	33.5	34.5	37.2	37.4	37.2				
	> 2 packs/day	35.0	34.0	35.0	37.7	37.9	37.7				
	Nonsmokers	33.5	32.5	33.5	36.2	36.4	36.2		33.5	33.4	33.5
	Up to < 1 pack/day	34.5	33.5	34.5	37.2	37.4	37.2				
3000-3999 ft	1 - 2 packs/day	35.0	34.5	35.0	37.7	37.9	37.7				
	> 2 packs/day	35.5	34.5	35.5	38.2	38.4	38.2				
	Nonsmokers	34.0	33.0	34.0	36.7	36.9	36.7		34.0	33.9	34.0
4000 4000 5	Up to < 1 pack/day	35.0	34.0	35.0	37.7	37.9	37.7				
4000- 4999 ft	1 - 2 packs/day	35.5	34.5	35.5	38.2	38.4	38.2				
	> 2 packs/day	36.0	35.0	36.0	38.7	38.9	38.7				
	Nonsmokers	34.5	33.5	34.5	37.2	37.4	37.2		34.5	34.4	34.5
E000 E000 #	Up to < 1 pack/day	35.5	34.5	35.5	38.2	38.4	38.2				
5000-555511	1 - 2 packs/day	36.0	35.0	36.0	38.7	38.9	38.7				
	> 2 packs/day	36.5	35.5	36.5	39.2	39.4	39.2				
	Nonsmokers	35.0	34.0	35.0	37.7	37.9	37.7		35.0	34.9	35.0
6000 6999 fr	Up to < 1 pack/day	36.0	35.0	36.0	38.7	38.9	38.7				
0000-0555 ft	1 - 2 packs/day	36.5	35.5	36.5	39.2	39.4	39.2				
	> 2 packs/day	37.0	36.0	37.0	39.7	39.9	39.7				

Table 201 – B		Category									
		1 [#] Trimester	2 nd Trimester	3 rd Trimester	Nonpreg 12 - < 15 yrs	Nonpreg 15 - < 18 yrs	Nonpreg ≥18 yrs	Infants 0 - < 6 mo	Infants 6 - < 12 mo	Child 1 - < 2 yrs	Child 2 - < 5 yrs
											Hgb <
	Nonsmokers	11.0	10.5	11.0	11.8	12.0	12.0		11.0	11.0	11.1
No altitude	Up to < 1 pack/day	11.3	10.8	11.3	12.1	12.3	12.3				
adjustment	1- 2 packs/day	11.5	11.0	11.5	12.3	12.5	12.5				
	> 2 packs/day	11.7	11.2	11.7	12.5	12.7	12.7				
	Nonsmokers	11.2	10.7	11.2	12.0	12.2	12.2		11.2	11.2	11.3
	Up to <1 pack/day	11.5	11.0	11.5	12.3	12.5	12.5				
3000- 3999 ft	1-2 packs/day	11.7	11.2	11.7	12.5	12.7	12.7				
	> 2 packs/day	11.9	11.4	11.9	12.7	12.9	12.9				
	Nonsmokers	11.3	10.8	11.3	12.1	12.3	12.3		11.3	11.3	11.4
	Up to <1 pack/day	11.6	11.1	11.6	12.4	12.6	12.6				
4000- 4999 ft	1- 2 packs/day	11.8	11.3	11.8	12.6	12.8	12.8				
	> 2 packs/day	12.0	11.5	12.0	12.8	13.0	13.0				
	Nonsmokers	11.5	11.0	11.5	12.3	12.5	12.5		11.5	11.5	11.6
E000 E000 #	Up to < 1 pack/day	11.8	11.3	11.8	12.6	12.8	12.8				
5000-5555 ft	1- 2 packs/day	12.0	11.5	12.0	12.8	13.0	13.0				
	> 2 packs/day	12.2	11.7	12.2	13.0	13.2	13.2				
	Nonsmokers	11.7	11.2	11.7	12.5	12.7	12.7		11.7	11.7	11.8
C000 C000 D	Up to <1 pack/day	12.0	11.5	12.0	12.8	13.0	13.0				
6000-6555 ft	1-2 packs/day	12.2	11.7	12.2	13.0	13.2	13.2				
	> 2 packs/day	12.4	11.9	12.4	13.2	13.4	13.4				

Cutoff Values from High Risk Manual Hgb Table

Cutoff values for Nevada at <3000 – 3999 feet

AREA AND	HCT/HGB	BLOOD VALUES FOR HEMOGLOBIN AND HEMATOCRIT								
ELEVATION	NS AND S	Women			Infants	Children				
		Pregnant w	veeks gestation:		Nonpregnant			1		
Less than 3000 feet	Hemoglobin g%	0-13	14-29	30-40	12-14 yrs	15-17 yrs	≥18 yrs	6-24 mo	2-5 yrs	
Henderson Las Vegas	Nonsmoker Smoker	11.0	10.5	11.0	11.8	12.0	12.0	11.0		
Pahrump	<1 pack/day	11.3	10.8	11.3	12.1	12.3	12.3		11.1	
Mesquite	1-2 packs/day ≥2 packs/day	11.5 11.7	11.0 11.2	11.5 11.7	12.3 12.5	12.5 12.7	12.5 12.7			
	Hematocrit %									
	Nonsmoker	33.0	32.0	33.0	35.7	35.9	35.7	22.0		
	<1 nack/day	34.0	33.0	34.0	36.7	36.9	36.7	52.9	33.0	
	1-2 packs/dav	34.5	33.5	34.5	37.2	37.4	37.2		0010	
	≥2 packs/day	35.0	34.0	35.0	37.7	37.9	37.7			
3000-3999 feet	Hemoglobin g%									
Alamo	Nonsmoker	11.2	10.7	11.2	12.0	12.2	12.2	11.2	11.3	
Beatty	Smoker									
Fallon	<1 pack/day	11.5	11.0	11.5	12.3	12.5	12.5			
Lovelock	1-2 packs/day	11.7	11.2	11.7	12.5	12.7	12.7			
Stillwater	≥2 packs/day	11.9	11.4	11.9	12.7	12.9	12.9			
	Hematocrit %									
	Nonsmoker	33.5	32.5	33.5	36.2	36.4	36.2	33.4	33.5	
	Smoker	34 5	33.5	34 5	37.2	37 /	37.2			
	1-2nacks/dav	34.5	33.5	34.5	37.2	37.4	37.2			
	>2 nacks/day	35.5	34.5	35.5	38.2	38.4	38.2			
	Le paulo day		••							

Cutoff values for Nevada at 4,000 - 5,999 feet

AREA AND	HCT/HGB	BLOOD V							
ELEVATION	NS AND S	Women		Infants	Children				
	Pregnant	weeks ges	tation:	Nonpre	gnant				
4000-4999 feet	Hemoglobin g%								
Battle Mountain Babbit	Nonsmoker Smoker	11.3	10.8	11.3	12.1	12.3	12.3	11.3	11.4
Caliente Carlin	<1 pack/day 1-2 packs/day	11.6 11.8	11.1 11.3	11.6 11.8	12.4 12.6	12.6 12.8	12.6 12.8		
Carson City Dayton	≥2 packs/day	12.0	11.5	12.0	12.8	13.0	13.0		
Dresslerville Dyer	Hematocrit %								
Fernley Gabbs	Nonsmoker Smoker	34.0	33.0	34.0	36.7	36.9	36.7	33.9	34.0
Gardnerville Hawthorne	<1 pack/day 1-2 packs/day	35.0 35.5	34.0 34.5	35.0 35.5	37.7 38.2	37.9 38.4	37.7 38.2		
May McDermitt Mina Minden	≥2 packs/day	30.0	35.0	30.0	38.7	38.9	38.7		
Orvada Panaca Reno									
Schurz Silver Springs Wendover Winnemucca Yerington									
5000 -5999 feet	Hemoglobin g%	0-13	14-29	30-40	12-14 yrs	15-17 yrs	≥18 yrs	6-24 mo	2-5 yrs
Duckwater	Nonsmoker	11.5	11.0	11.5	12.3	12.5	12.5	11.5	11.6
Goldfield Jackpot Lund Mountain City	<pre>Smoker <1 pack/day 1-2 packs/day ≥2 packs/day</pre>	11.8 12.0 12.2	11.3 11.5 11.7	11.8 12.0 12.2	12.6 12.8 13.0	12.8 13.0 13.2	12.8 13.0 13.2		
Owyhee Topaz	Hematocrit %								
Wells	Nonsmoker	34.5	33.5	34.5	37.2	37.4	37.2	34.4	34.5
	Smoker <1 pack/day 1-2 packs/day ≥2 packs/day	35.5 36.0 36.5	34.5 35.0 35.5	35.5 36.0 36.5	38.2 38.7 39.2	38.4 38.9 39.4	38.2 38.7 39.2		

Cutoff values for Nevada at 6,000 - 6,999 feet

AREA AND ELEVATION	ID HCT/HGB BLOOD VALUES FOR HEMOGLOBIN AND HEMATOCRIT								
		Women							Children
		Pregnant v	veeks gestation:		Nonpregnant			-	
6000 6000 foot	Homoglobin a%			[[[
0000-0333 1661	nemoglobin g //								
Austin	Nonsmoker	11.7	11.2	11.7	12.5	12.7	12.7	11.7	11.8
Beowawe	Smoker								
Ely	<1 pack/day	12.0	11.5	12.0	12.8	13.0	13.0		
Eureka	1-2 packs/day	12.2	11.7	12.2	13.0	13.2	13.2		
McGill Pioche	≥2 packs/day	12.4	11.9	12.4	13.4	13.4	13.4		
Round Mountain Ruth	Hematocrit %								
Stateline	Nonsmoker	35.0	34.0	35.0	37.7	37.9	37.7	34.9	35.0
Tonopah	Smoker								
Virginia City	<1 pack/day	36.0	35.0	36.0	38.7	38.9	38.7		
Zepher Cove	1-2packs/day	36.5	35.5	36.5	39.2	39.4	39.2		
	≥2 packs/day	37.0	36.0	37.0	39.7	39.9	39.7		

Us Troubleshooting Guide If you are unable to resolve the problem by following this Troubleshooting Guide, please call your HemoCue distributor or HemoCue America. LED = Light Emitting Diode. Note! Do not open the cover of the analyzer. The warranty is voided if the analyzer has been opened.

Symptom	Explanation	Action
The analyzer shows "ERROR" and a digit code 900–930.	May be an occasional fault.	Turn off the analyzer and turn it on again after 30 seconds. Take a new microcuvette and repeat the measurement. If the problem continues, see specific error code below.
900	No stable endpoint found within the time range. 1. The microcuvette is faulty. 2. The circuit board is out of order.	 Check expiration date for the microcuvettes. Take a new microcuvette and repeat the measurement. The analyzer needs service. Call your distributor.
901	Light intensity for the compensating LED is too low. 1. The optronic unit is dirty. 2. The optronic unit is out of order.	 Clean the optronic unit, as described in the Maintenance section. The analyzer needs service. Call your distributor.
902	Light intensity for the measuring LED is too low. 1. The optronic unit is dirty. 2. The optronic unit is out of order.	 Clean the optronic unit, as described in the Maintenance section. The analyzer needs service. Call your distributor.
903	1. The optronic unit is out of order.	1. The analyzer needs service. Call your distributor.
905	1. Light intensity for one of the LED's is too high.	1. The analyzer needs service. Call your distributor.
906	1. Unstable blank value. The analyzer might be cold.	 Turn off the analyzer and allow it to reach room temperature. If the problem continues, the analyzer needs service. Call your distributor.
907	1. The battery power is too low.	 The batteries need to be replaced. Turn off the analyzer and replace the batteries, 4 type AA. Use the power adapter.
908	The absorbance is too high. 1. Light blocking item in the cuvette holder.	 Check that the analyzer and microcuvettes are used according to the HemoCue Glucose 201 operating manual and instruction for use. The analyzer needs service. Call your distributor.
910	Seen during all phases of use, and is a fatal error. A read or write operation to the EEPROM did not succeed. 1. The EEPROM memory is out of order.	 Turn off the analyzer and turn it on again after 30 seconds and retry testing. The analyzer needs service. Call your distributor.
911	 Seen during start up and is a fatal error. The analyzer cannot detect a valid EEPROM memory configuration. 	 1a. Turn off the analyzer and turn it on again after 30 seconds and retry testing. 1b. The analyzer needs service. Call your distributor.
913	Seen during start up and is a fatal error. Self test of RAM memory failed. 1. The electronics are out of order.	1. The analyzer needs service. Call your distributor.
925	Seen at start up and is a fatal error. Calibration checksum is not valid. 1. The analyzer needs calibration.	1. The analyzer needs service. Call your distributor.
929	1. Communication error during internal hardware test.	1. The analyzer needs service. Call your distributor.
930	1. The electronic "SELFTEST" failed.	1a. Turn off the analyzer and allow it to reach room temperature.1b. If the problem continues, the analyzer needs service. Call your distributor.
ннн	1. Measured value exceeds 444 mg/dL (24.6 mmol/L).	 Check the expiration date of the microcuvettes. Remeasure sample with fresh microcuvette.
No characters on the display.	 The analyzer is not receiving power. If on battery power, the batteries need to be replaced. The display is out of order. 	 Check that the power adapter is connected to the power source. Check that the power adapter is securely connected to the analyzer. Check that the adapter wire is not damaged. Turn off the analyzer and replace the batteries, 4 type AA. The analyzer needs service. Call your distributor.
The display gives erroneous characters.	 The display is out of order. The microprocessor is out of order. 	 The analyzer needs service. Call your distributor. The analyzer needs service. Call your distributor.

Symptom	Explanation	Action
The display shows " 💶 "	 The batteries need to be replaced. If using power adapter, the adapter or circuit board is out of order. 	 Turn the analyzer off and replace the batteries, 4 type AA. Check that the power adaptor is properly connected and working. The analyzer needs service. Call your distributor.
The display does not switch from "SELFTEST" to "READY" or from "READY" to " " (measuring).	 The magnet in the cuvette holder may be missing. The magnetic sensor is out of order. 	 The analyzer needs service. Call your distributor. The analyzer needs service. Call your distributor.
Measurements on control materials out of range – either too HIGH or too LOW.	 The microcuvettes are beyond their expiration date, damaged or have been improperly stored. The optical eye of the microcuvette is contaminated. The control has not been mixed well and/or is not at room temperature. Air bubbles are present in the microcuvette. The optronic unit is dirty. The control is not suitable for use with the HemoCue Glucose 201 system. The controls are beyond their expiration dates or have been improperly stored. 	 Check the expiration date and the storage conditions of the microcuvettes. Remeasure the sample with a new microcuvette. Make sure that the control is mixed well and at room temperature. Check the microcuvette for air bubbles. Remeasure the sample with a new microcuvette. Clean the optronic unit, as described in the Maintenance section. Only use controls intended for the HemoCue Glucose 201 system recommended by HemoCue. The analyzer needs service. Call your distributor. Check the expiration date and the storage conditions of the control. Take a new microcuvette and repeat the measurement
Measurements on patient samples are higher or lower than anticipated.	 The microcuvettes are beyond their expiration date, damaged or have been improperly stored. The optical eye of the microcuvette is contaminated. Air bubbles are present in the microcuvette. The optronic unit is dirty. The calibration of the analyzer has been changed. 	 Check the expiration date and the storage conditions of the microcuvettes. Remeasure the sample with a new microcuvette. Check the microcuvette for air bubbles. Remeasure the sample with a new microcuvette. Clean the optronic unit, as described in the Maintenance section. The analyzer needs service. Call your distributor.

Appendix C – HemoCue 301 Troubleshooting Guide

Troubleshooting

If problem remains after recommended actions below are taken, contact the HemoCue distributor or HemoCue AB. The analyzer should be cleaned prior to service, see section *Maintenance* for instructions. If any spare parts are lost or damaged, see section *Components*. Do not open the cover of the analyzer. Warranty is void if analyzer has been opened.

* Follow relevant section in this operating manual.

Error Code	Explanation	Action
Error code is displayed.	May be a temporary fault or faulty electronics/optical parts.	Turn analyzer off. Turn on after 30 seconds. If the problem continues, see specific error code below or the analyzer needs service.
E00	 Faulty microcuvette. Internal error in analyzer. 	 Check expiry date. Fill a new microcuvette and perform measurement. * Clean optical parts. *
E01-E02	 Dirty optical parts. Analyzer too hot/cold. 	 Clean optical parts. * Turn analyzer off, allow to reach operating temperature before use.
E03	1. Analyzer exposed to direct light.	1. Avoid direct light exposure.
E05, E06	1. Analyzer too hot/cold or exposed to direct light.	 Turn analyzer off, allow to reach operating temperature before use. Avoid direct light exposure.
E07	1. Battery power too low.	1a. Batteries need to be replaced. * 1b. Use power adapter.
E08	 The absorbance is too high. Faulty microcuvette or sample. Dirty or blocked optical parts. 	1a and 1b. Ensure analyzer and microcuvettes are used according to operating manual and relevant package insert. 1a and 1c. Clean optical parts. *
E10-E30	1. Internal error in analyzer.	1. Clean optical parts. *

Troubleshooting

Measuring Deviations	Explanation	Action
ННН	1. Result exceeds measuring range.	1a. Fill a new microcuvette and perform measurement. * 1b. Confirm result with laboratory method.
No characters displayed	 Analyzer not receiving power. Faulty display. 	 Ensure power adapter is properly connected and cable not damaged. Batteries need to be replaced. * Analyzer needs service.
Erroneous characters displayed	1. Faulty display/electronics.	1. Analyzer needs service.
The display shows "FIR"	This function is for manufacturing use only.	 Replace all cables and/or batteries, and restart. Analyzer needs service.
is displayed	 Low battery. Faulty power adapter. 	 Batteries need to be replaced. * Ensure power adapter is properly connected and cable not damaged.
$\overline{\mathbb{Z}}$ and "Hb" continuously displayed	1. Faulty cuvette holder/analyzer.	1. Analyzer needs service.
Unexpected patient or control results	Patient sample and control solution 1. Microcuvettes expired, damaged or improperly stored. 2. Microcuvette contaminated. 3. Dirty optical parts. 4. Air bubbles in filled microcuvette. 5. Faulty optical parts. 6. Incorrect sampling technique. 7. Measurement started later than 40 seconds after filling the microcuvette. Control solution 8. Control solution not compatible. 9. Inproper handling of control solution. 10. Control solution expired or improperly stored.	Patient sample and control solution 1. Check expiry date and storage conditions. 2. Fill a new microcuvette and perform measurement. * 3. Clean optical parts. * 4. Fill a new microcuvette and perform measurement. * 5. The analyzer needs service. 6. Fill a new microcuvette and perform measurement. * 7. Start measurement no later than 40 seconds after filling the microcuvette. Control solution 8. Only use control solutions recommended by HemoCue. 9. Always follow instructions for use for control solution. 10. Check expiry date and storage conditions.

Appendix D – Masimo Pronto Troubleshooting Guide

Masimo Pronto Troubleshooting Guide					
Problem	Possible Cause(s)	Recommendation(s)			
Difficulty or No SpHb Reading	Coiled / Twisted Sensor Cable	Uncoil / Untwist sensor cable			
(Including Low SIQ)	Soiled Sensor	Clean sensor with soap and water solution			
	Participant wearing metallic, acrylic, or dark fingernail polish	Move sensor to another finger or obtain Hgb value via capillary sampling test			
	Inappropriate Sensor/Finger Selection	Verify that appropriate finger and sensor are chosen			
	Excessive Motion	Minimize or eliminate motion at the measurement site			
	Low Perfusion Index (less than 1.0)	Warm Hands / Choose Another Finger on Other Hand			
	Excessive Ambient Light	Shield the sensor from excessive light			
Unit does not Power On	Low Battery	Check / replace batteries			
Sensor Fails to Initialize (Circulating LEDs)	Incorrectly connected sensor cable	Disconnect and reconnect sensor cable from the patient cable.			
Continuous Speaker Tone	Internal Failure	Return Masimo Pronto for Service			
Buttons don't work when pressed	Internal Failure	Return Masimo Pronto for Service			