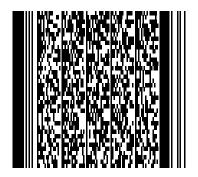
R&D Retic-I for DxH 800 HEMATOLOGY CONTROLS

CONTROL



Assay Values and Expected Ranges

QCP Data Months: December, January



153112250, 164212250, 175312250



2026-02-05

		LEVEL 1 LOT	153112250	LEVEL 2 LOT	164212250	LEVEL 3 LOT	175312250
		Mean	Range	Mean	Range	Mean	Range
Beckman Coulter® UniCel® DxH™900/800/690T/600	Retic %	1.0	± 1.0	6.2	± 2.5	11.0 ±	3.8
RBC*	RBC × 10 ⁶ /µL	3.55	NA	3.55	NA	3.50	NA

^{*}Use this RBC value to calculate the absolute reticulocyte number.

INTENDED USE

R&D Retic-I for DxH 800 is a control designed to monitor values obtained on automated reticulocyte counting methods. Please refer to the assay table for specific instrument models.

SUMMARY AND PRINCIPLE

It is an established laboratory practice to use a stable control to monitor the performance of diagnostic tests. This control is composed of stable materials that provide a means of monitoring the performance of reticulocyte counting methods. It is sampled in the same manner as a patient specimen.





REAGENTS

R&D Retic-I for DxH 800 I is an in vitro diagnostic reagent composed of human erythrocytes suspended in a plasmalike fluid with preservatives.



PRECAUTION

R&D Retic-I for DxH 800 is intended for in vitro diagnostic use only by trained personnel.



WARNING:

POTENTIALLY BIOHAZARDOUS MATERIAL. For in vitro diagnostic use. Each human donor/unit used in the preparation of this product has been tested by a FDA licensed method/test and found to be negative or nonreactive for the presence of HBsAg, Anti-HCV, NAT testing for HIV-1, HCV (RNA) and HIV-1/2.

Each unit is also negative by a serological test for Syphilis (RPR or STS). Because no test method can offer complete assurance that infectious agents are absent, this material should be handled as potentially infectious. When handling or disposing of vials follow precautions for patient specimens as specified in the OSHA Bloodborne Pathogen Rule (29 CFR Part 1910.1030) or other equivalent biosafety procedures.



STABILITY AND STORAGE

Store R&D Retic-I for DxH 800 upright at 2 - 8° C (35 -46 F) when not in use. Protect vials/ tubes from overheating and freezing. Unopened vials/tubes are stable until the expiration date. Opened vials/tubes are stable for at least 14 days provided they are handled properly.

INDICATIONS OF DETERIORATION

After mixing, product should be similar in appearance to fresh whole blood. In unmixed vials/tubes, the supernatant may appear pink; this is normal and does not indicate deterioration. Other discoloration, very dark red supernatant or unacceptable results may indicate deterioration. Do not use the product if deterioration is suspected.

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INSTRUCTIONS FOR USE

- Remove vials/tubes from the refrigerator and allow to warm to room temperature (15 - 30°C or 59 - 86°F) for 15 minutes before mixing.
- 2. To mix, hold a vial/tube horizontally between the palms of the hands. Do not pre-mix on a mechanical mixer.
 - a) Roll the vial/tube back and forth for 20 30 seconds; occasionally invert the vial/tube. Mix vigorously, but do not shake.
 - b) Continue to mix in this manner until the red cells are completely suspended. Vials/tubes stored for a long time may require extra mixing.
 - c) Gently invert the vial/tube 8 10 times immediately before sampling.
- 3. Analyze the sample as instructed in the Quality Control section of the Operator's Manual for your instrument.
- 4. After sampling:
 - a) If tube has been open for sampling, clean residual material from the cap and tube rim with a lint-free tissue. Replace the cap tightly.
 - b) Return tubes to refrigerator within 30 minutes of use.

QC FILE SETUP PROCEDURE FOR BECKMAN COULTER DXH 900/800/690T/600:

2 DIMENSIONAL BARCODE SETUP:

- At the workstation, select Menu, Set Up, and then Quality Control. Select New Control From Bar Code button. A dialog box appears that reads "Waiting for 2D bar code to be scanned from assay sheet..."
- Scan the 2D barcode using the hand held scanner. The scanned lot numbers will appear. Click on Select All and OK.
- The assay information will be transmitted to the QC Retic files. Level 1 values will load into the Retic Level 1 file. Level 2 values will load into the Retic Level 2 file. Level 3 values will load into the Retic Level 3 file.

EXPECTED RESULTS

Verify that the lot number on the vial/tube matches the lot number on the table of assay values. Assay values are determined on well-maintained, properly calibrated instruments using the instrument manufacturer's recommended reagents. Reagent differences, maintenance, operating technique, and calibration may contribute to interlaboratory variation.

PERFORMANCE CHARACTERISTICS

Assigned values are presented as a Mean and Range. The Mean is derived from replicate testing on instruments operated and maintained according to the manufacturer's instructions. The Range is an estimate of variation between laboratories and also takes into account inherent imprecision of the method and expected biological variability of the control material.

Assay values on a new lot of control should be confirmed before the new lot is put into routine use. Test the new lot when the instrument is in good working order and quality control results on the old lot are acceptable. The laboratory's recovered mean should be within the assay range.

For greater control sensitivity each laboratory should establish its own mean and acceptable range and periodically reevaluate the mean. The laboratory range may include values outside of the assay range. The user may establish assay values not listed on the Assay Sheet, if the control is suitable for the method.

LIMITATIONS

The performance of this product is assured only if it is properly stored and used as described in this insert. Incomplete mixing of a vial/tube prior to use invalidates both the sample withdrawn and any remaining material in the vial/tube.

TECHNICAL ASSISTANCE AND CUSTOMER SERVICE

For assistance in resolving control recovery problems, please call Technical Service at (800) 523-3395. For additional information on R&D Systems, Inc. hematology controls and calibrators, or to place an order, call Customer Service at (800) 428-4246.

QUALITY CONTROL PROGRAM

For information on CBC-Monitor, our Inter-Laboratory Quality Control Program, call (800) 523-3395 ext. 4435.

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