

BC-BF

HEMATOLOGY CONTROLS

CONTROL

Assay Values and Expected Ranges

LOT BFI1124A

QCP Data Months: **November, December**



2025-01-10

Analyzer	Parameter	LEVEL 1		LEVEL 2		LEVEL 3	
		LOT	BFI1124A-1	LOT	BFI1124A-2	LOT	BFI1124A-3
		Mean	Range	Mean	Range	Mean	Range
Mindray BC-6800,BC-6600	WBC-BF 10 ⁹ /L	0.070	± 0.035	0.239	± 0.191	0.988	± 0.198
	RBC-BF 10 ¹² /L	0.024	± 0.007	0.072	± 0.014	0.470	± 0.033
	TC-BF # 10 ⁹ /L	0.070	± 0.035	0.239	± 0.191	0.988	± 0.198
	PMN %	72.5	± 12.0	70.6	± 8.0	86.9	± 6.0
	PMN # 10 ⁹ /L	0.051	± 0.038	0.169	± 0.169	0.859	± 0.243
	MN %	27.5	± 12.0	29.4	± 8.0	13.1	± 6.0
	MN # 10 ⁹ /L	0.019	± 0.019	0.070	± 0.070	0.129	± 0.098
Mindray BC-6000, BC-6100, BC-6200, BC-6000Plus, BC-6100Plus	WBC-BF 10 ⁹ /L	0.074	± 0.037	0.247	± 0.198	1.007	± 0.201
	RBC-BF 10 ¹² /L	0.023	± 0.007	0.069	± 0.014	0.456	± 0.032
	TC-BF # 10 ⁹ /L	0.074	± 0.037	0.247	± 0.198	1.008	± 0.202
	PMN %	70.2	± 12.0	71.3	± 8.0	86.9	± 6.0
	PMN # 10 ⁹ /L	0.052	± 0.039	0.176	± 0.176	0.875	± 0.247
	MN %	29.8	± 12.0	28.7	± 8.0	13.1	± 6.0
	MN # 10 ⁹ /L	0.022	± 0.022	0.071	± 0.071	0.132	± 0.099
Mindray BC-6600Plus, BC-6700Plus, BC-6800Plus	WBC-BF 10 ⁹ /L	0.075	± 0.038	0.259	± 0.207	1.056	± 0.211
	RBC-BF 10 ¹² /L	0.023	± 0.007	0.073	± 0.015	0.474	± 0.033
	TC-BF # 10 ⁹ /L	0.075	± 0.038	0.261	± 0.209	1.057	± 0.211
	PMN %	71.3	± 12.0	71.2	± 8.0	86.8	± 6.0
	PMN # 10 ⁹ /L	0.053	± 0.041	0.184	± 0.184	0.917	± 0.259
	MN%	28.7	± 12.0	28.8	± 8.0	13.2	± 6.0
	MN # 10 ⁹ /L	0.022	± 0.022	0.075	± 0.075	0.139	± 0.104
Mindray BC-700[R], BC-720[R],BC-760[R], BC-780[R],BC-700 [R] CS, BC-760[R] CS,BC-700[B], BC-760[B]	WBC-BF 10 ⁹ /L	0.071	± 0.036	0.244	± 0.195	0.997	± 0.199
	RBC-BF 10 ¹² /L	0.022	± 0.007	0.067	± 0.013	0.431	± 0.030
	TC-BF # 10 ⁹ /L	0.071	± 0.036	0.245	± 0.196	0.998	± 0.200
	PMN %	69.0	± 12.0	69.9	± 8.0	86.4	± 6.0
	PMN # 10 ⁹ /L	0.049	± 0.038	0.171	± 0.171	0.861	± 0.244
	MN%	31.0	± 12.0	30.1	± 8.0	13.6	± 6.0
	MN # 10 ⁹ /L	0.022	± 0.022	0.073	± 0.073	0.136	± 0.098
Mindray BC-7500[B] CRP, BC-7500 [R] CRP, BC-7500 [N] CRP, BC-7500 [NR] CRP	WBC-BF 10 ⁹ /L	0.067	± 0.034	0.232	± 0.186	0.958	± 0.192
	RBC-BF 10 ¹² /L	0.023	± 0.007	0.073	± 0.015	0.469	± 0.033
	TC-BF # 10 ⁹ /L	0.067	± 0.034	0.233	± 0.186	0.959	± 0.192
	PMN %	70.1	± 12.0	71.5	± 8.0	86.9	± 6.0
	PMN # 10 ⁹ /L	0.047	± 0.036	0.166	± 0.166	0.833	± 0.235
	MN%	29.9	± 12.0	28.5	± 8.0	13.1	± 6.0
	MN # 10 ⁹ /L	0.020	± 0.020	0.066	± 0.066	0.125	± 0.095

BC-BF

HEMATOLOGY CONTROLS CONTROL

INTENDED USE

The BC-BF Control is an assayed hematology control intended to monitor the reliability of hematology instruments that quantitatively measure red and white blood cell counts in cerebrospinal fluids, serous fluids, and synovial fluids.

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 which provide a means of monitoring the performance of hematology blood cell counters. It is sampled in the same manner as a patient specimen.

REAGENTS

BC-BF Control is an in vitro diagnostic reagent composed of human erythrocytes and bovine leukocytes suspended in a fluid with preservatives.



PRECAUTION

BC-BF is intended for in vitro diagnostic use only by trained personnel.



WARNING:

POTENTIAL BIOHAZARDOUS MATERIAL For in vitro diagnostic use. Each human donor/unit used in the preparation R&D Systems' products has been tested, and yielded non-reactive / negative results, according to FDA guidelines as contained in 21 CFR 610.40(a)(b).

Specifically, a sample from each donation used has been tested by FDA-licensed tests and found nonreactive / negative for:

1. Antibodies to human immunodeficiency virus (anti-HIV 1,2), hepatitis C virus (anti-HCV), and antibodies to Trypanosoma cruzi (T cruzi, the causative agent of Chagas disease), and nonreactive for hepatitis B surface antigen (HBsAg).

2. Nucleic acid tests (NAT) for HCV ribonucleic acid (RNA), HIV-1 RNA, HBV deoxyribonucleic acid (DNA) and West Nile virus (WNV) RNA

3. Serologic test for syphilis

Because no test method can offer complete assurance that infectious agents are absent, 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 BC-BF upright at 2 -8°C (35 - 46°F) when not in use. Protect tubes from overheating and freezing. Unopened tubes are stable through the expiration date. Opened tubes are stable 30 days, provided they are handled properly.

INDICATIONS OF DETERIORATION

After mixing, product should be similar in appearance to diluted whole blood. In unopened tubes, the supernatant may appear cloudy and reddish; 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

Note: Begin with a system rinse to reduce carryover. It is critical that the background counts be low prior to running body fluid controls. Run controls from lowest to highest concentration to reduce carryover

- Remove tubes from the refrigerator and allow to warm to room temperature (15 to 30°C or 59 to 86°F) for 15 minutes before mixing.
- To mix, hold a tube horizontally between the palms of the hands. **Do not pre-mix on a mechanical mixer.**
 - Roll the tube back and forth for 20 - 30 seconds; occasionally invert the tube. Mix vigorously, but do not shake.
 - Continue to mix in this manner until the red cells are completely suspended. Tubes stored for a long time may require extra mixing.
 - Gently invert the tube 8 - 10 times immediately before sampling.
- Analyze the sample as instructed in the Operator's Manual for your instrument.
- After sampling:
 - 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.
 - Return tubes to refrigerator within 30 minutes of use.

EXPECTED RESULTS

Verify that the lot number on the 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 inter-laboratory 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.

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 tube prior to use invalidates both the sample withdrawn and any remaining material in the tube.

Assay values and expected ranges given are intended only as guidelines and each laboratory should perform their own test system validation and establish tolerance limits.

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TECHNICAL ASSISTANCE AND CUSTOMER SERVICE

For additional information on Mindray hematology controls and calibrators, to place an order or for assistance in resolving control recovery problems, please email your local Mindray representative at service@mindray.com.

