Product Data Sheet
49 cu. ft. Blood Bank Refrigerator, High Performance, FDA listed Class II medical device

NSBR492WSW/0

Product Description

Corepoint™ Scientific Blood Bank Refrigerators are designed in accordance with FDA listed Class II medical devices. In addition, blood bank refrigerators also conform to the requirements set forth by AABB for the refrigerated cold storage of blood-based products.

Backed by optimal temperature control and EPA SNAP compliant refrigerants, these high-performance units protect blood, prevent waste, and allow for peak delivery. Corepoint™ Scientific blood bank refrigerators utilize smart controllers and feature a full array of alarms, LED interior lighting, stainless steel interiors, sliding drawers and probe access port.

General Description and Application

Description	Double solid door Blood Bank refrigerator
Operational environment	Indoor use only, +18°C to +26°C (+65°F to +78°F), <70% RH
Storage capacity	49 cu. ft. gross volume, up to 832 blood bags with optional 8th drawer per door
Door	Two swing solid doors, self-closing, stay position at 100° open, right and left hinged to open from the middle, non-reversible, magnetic sealed gasket, keyed lock
Drawers	7 drawers standard per door (14 drawers total) with option for 8th drawer per door (16 drawers total), 304 SS drawers, 65 lb. capacity each
Mounting and Installation	4 swivel casters, front casters locking
Interior lighting	Shielded, switched LED lighting, full coverage, balanced spectrum
External probe access	Rear wall port (3/4") dia.
Insulation	Cabinet is foamed-in-place with EPA compliant high density urethane foam
Exterior materials	White powder coated steel
Access control	Key lock
General warranty	Two (2) years parts and labor warranty, excluding calibrations
Compressor warranty	Seven (7) years compressor warranty
Product Weight	636 lbs. (14 drawers)
Shipping Weight	755 lbs. (14 drawers)
Rated Amperage	4.5 Amps
Power Plug/Power Cord	Hospital grade, NEMA 5-15, 9 ft nominal (2.7 m)
Facility Electrical Requirement	110-120V AC: 15 A (minimum)
Agency Listing and Certification	FDA listed Class II medical device, 21CFR part 820 compliant, ETL, CETL Listed (certified to UL471 standard, hydrocarbon refrigerant safety)
Optional Accessories	Additional 8th drawer per door, Chart paper, Mounting anchor, Upper solid ballast, Lower glycerol bottle kit (factory install only), IQ/OQ/PQ

Refrigeration System

Compressor	Hermetic, variable speed (VSC). Rated speed range: 1300-4000 rpm
Refrigerant	EPA SNAP compliant, R600a, Isobutane
Condenser	Anti-fouling tube and grid design, ultra-quite multi-speed fan
Evaporator	Fin and tube design, high efficiency fan
Defrost	Cycle optimized, zero energy

Performance

Uniformity[1] (Cabinet air)	+/- 0.4°C
Stability[2] (Cabinet air)	+/- 0.5°C
Maximum temperature variation (Cabinet air)	+/- 0.7°C
Stability[2] (Simulator ballast)	+/- 0.1°C
Stability[2] (Simulator bag)	+/- 0.1°C
Temperature Rise after 8 sec Door Openings	Temperature did not exceed 4.5°C at any probe
Recovery after 3 min Door Opening	All probes under 6.2°C throughout opening
Energy Consumption	1.47 KWh/day[3]
Average Heat Rejection	3.27 KWh/day (320 BTU/h)[3]
Noise Pressure Level (dBA)	38 or less installed

Controller, Configuration, Alarms and Monitoring

Controller technology	Proportional Integral Derivative (PID) microprocessor with LCD display
Battery Backup	24V high capacity battery, controller, all alarms active, temperature monitoring DAQ and event logging active on battery backup
Digital Communication	RS-485 (MODBUS)
Chart Recorder	n/a
Temperature setpoint range	1°C to 10°C
Display probe	Calibrated, stainless steel
External alarm connection	State switching remote alarm contacts
Alarms	Visual and audible indicators, Power failure, Temperature sensor failure, Battery voltage monitor and replacement, High / Low temperature, Door ajar
Simulator ballast	Upper probe: 4 oz. (120 ml) bottle, 50% glycerol mixture. Lower probe: Solid thermal media

Performance data acquired at 22°C ambient, 4°C nominal set point in an empty cabinet with drawers using validation ballast probes, during stabilized steady state operation and a DAQ sampling rate of one measurement every 10 seconds

1 - Uniformity is defined as the maximum variance in temperature across all probes at any point in time over the testing period

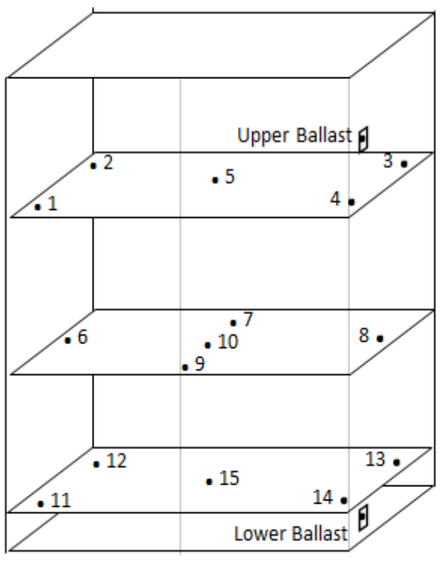
2 - Stability is defined as the maximum variance in temperature experienced by any single probe over the testing period

3 - Data per Energy Star test results or equivalent testing and calculation. Heat rejection based on daily averages, not continuous operation. Performance exceeds Energy Star requirements.

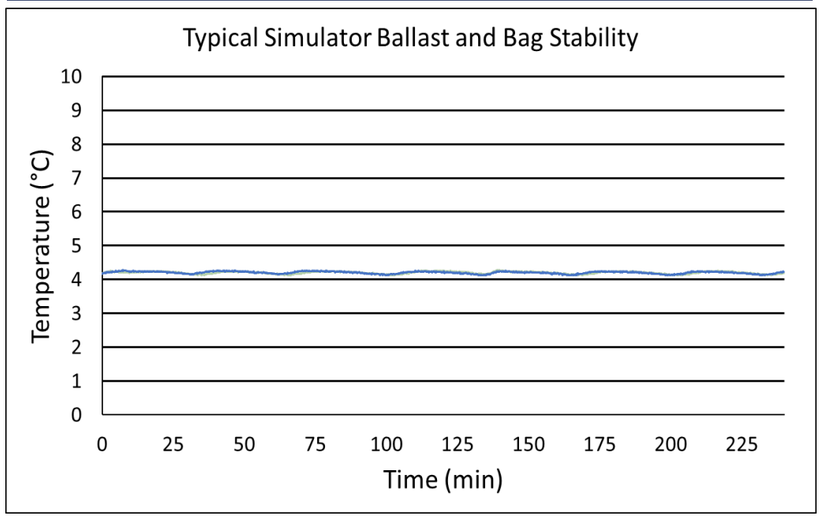
Certifications

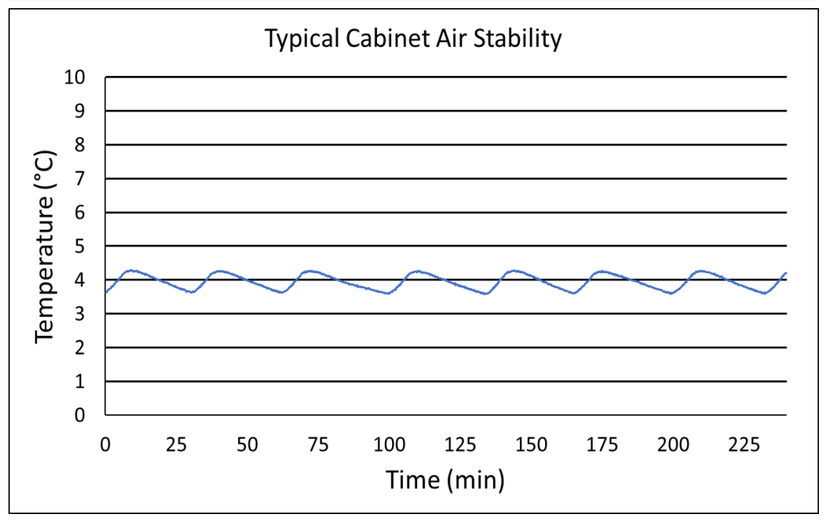
Temperature Probes

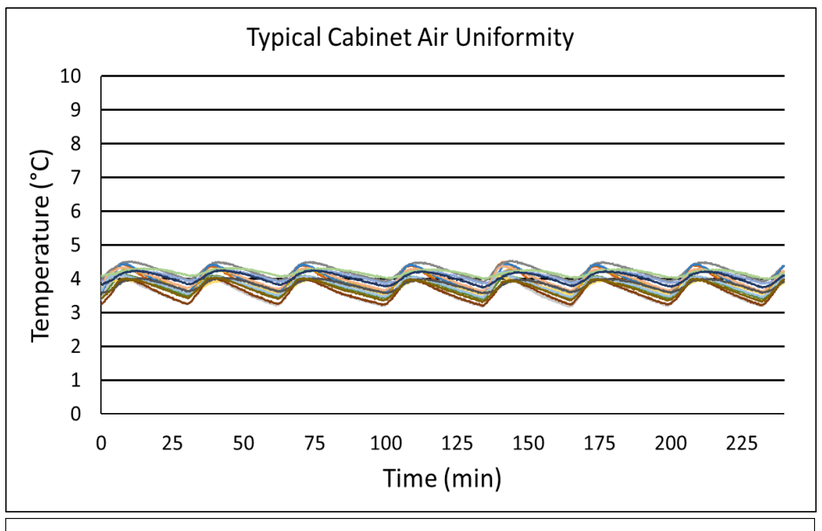
Probe	Ave	Min	Max
1	3.9	3.4	4.5
2	4.2	3.9	4.5
3	3.9	3.6	4.3
4	4.0	3.6	4.4
5	3.8	3.4	4.1
6	4.1	3.8	4.3
7	4.0	3.6	4.4
8	3.6	3.2	4.2
9	3.8	3.5	4.1
10	3.8	3.4	4.2
11	4.2	4.0	4.3
12	4.0	3.7	4.3
13	3.6	3.2	4.0
14	3.8	3.6	4.0
15	3.7	3.3	4.0
Bal	4.2	4.1	4.3
Bag	4.2	4.1	4.3

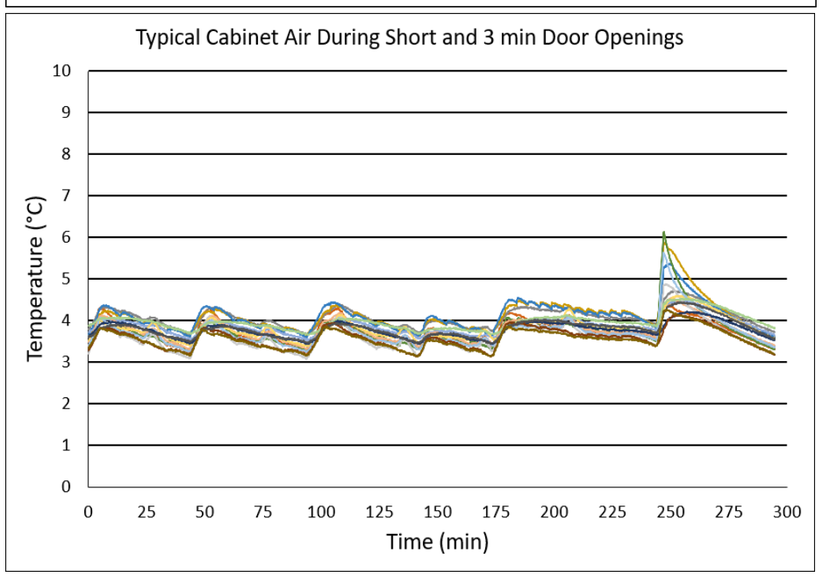


Temperature Charts









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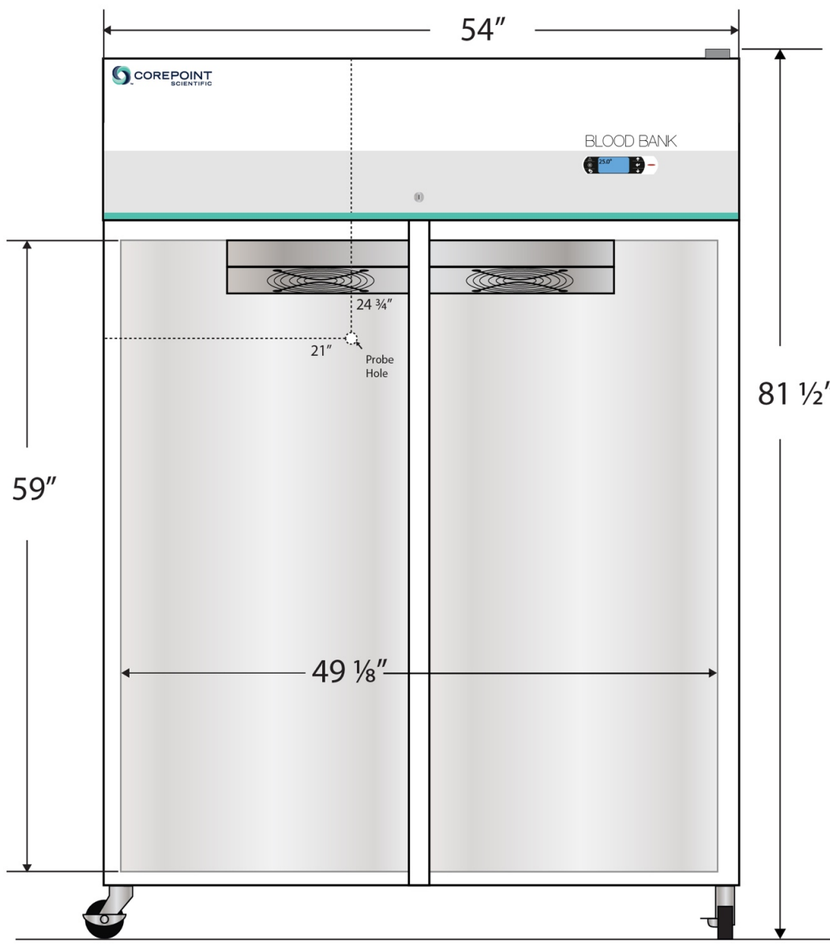
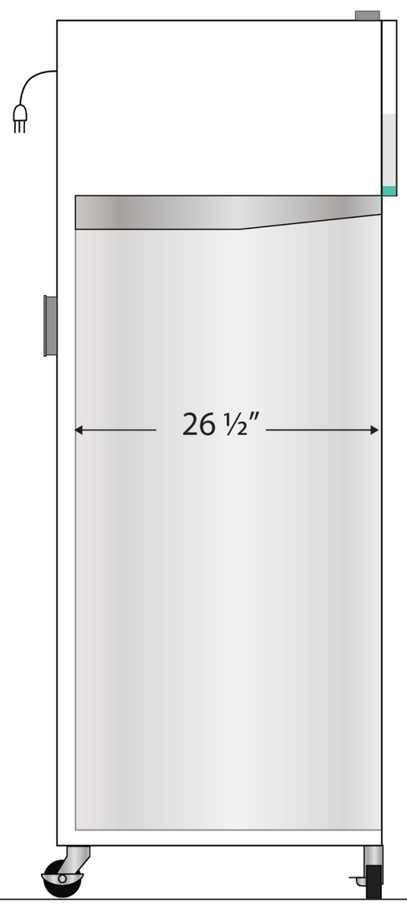
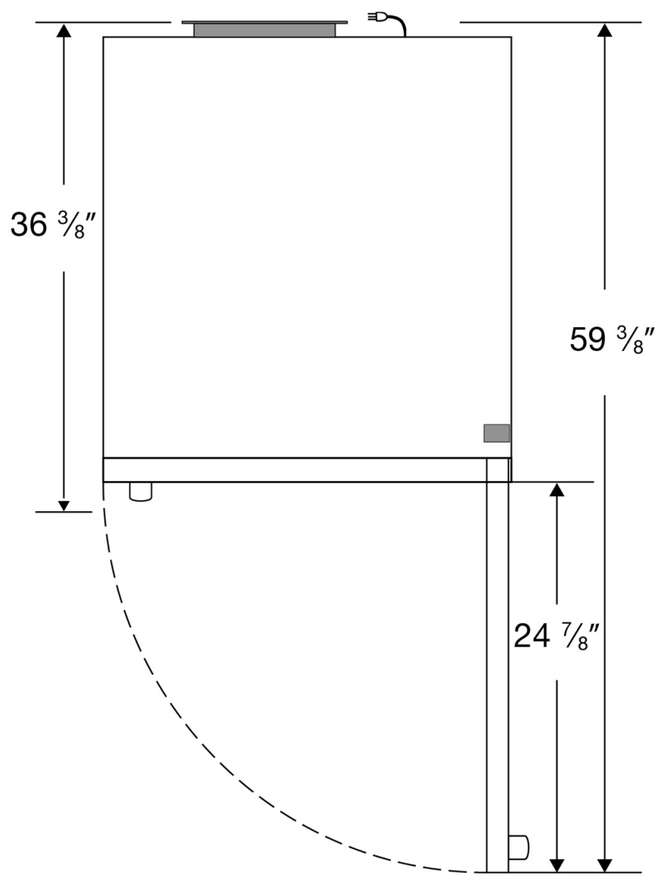
Images

Dimensions

	Width	Depth	Height	Door Swing	Total open Depth
Exterior	54"	36 3/8'"	81 1/2"	24 7/8"	59 3/8"
Interior	49 1/8"	26 1/2"	59"		







Contact

Customer Service	800-648-4041 Option 3	customerservice@horizonscientific.com
Technical Service	800-648-4041 Option 5, Parts Option 4	technicalservice@horizonscientific.com

HS-MKT-DOC-0008_Rev 1