PH-BSI-NSF-UCBI-0404

Product Data Sheet
Undercounter 4.6 cu. ft. Built-In Vaccine Refrigerator -
Certified to NSF/ANSI 456 Standard for Vaccine Storage

Product Description

These built-in undercounter refrigerators are designed in accordance with the NSF/ANSI 456 Standard for Vaccine Storage. Units protect pharmaceuticals at optimal temperatures, preventing waste and allowing for peak delivery.

The solid door refrigerators utilize microprocessor controllers and feature temperature alarms, remote alarm contacts, and probe access ports with included probes. Vaccine Storage Refrigerators utilize HFC-free refrigerant for environmental health and energy efficiency.

Certifications

*-one or more of these certifications may apply to this unit.

General Description and Application

Description	Single Solid Door Pharmacy/Vaccine Undercounter Refrigerator Built-In
Operational environment	Indoor use only, +18°C to +26°C (+65°F to +78°F), <70% RH
Storage capacity	4.6 cu. ft. gross volume
Door	One swing solid door, self-closing, right hinged, non-reversible, magnetic sealed gasket, keyed lock
Shelves	Three shelves (two adjustable/one fixed) with guard rail on back
Mounting	Low profile roller wheels and leveling legs
Interior lighting	N/A
Airflow management	Forced Air technology, patent pending
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	Pyxis®, Omnicell® and AcuDose RX® compatible
General warranty	Two (2) years parts and labor warranty, excluding display probe calibration
Compressor warranty	Five (5) years compressor warranty
Product Weight	100 lbs.
Shipping Weight	140 lbs.
Rated Amperage	1.74 Amps
Power Plug/Power Cord	NEMA 5-15 plug, 8 to 10 ft typical, conforms to UL471 requirements, Vaccine storage power cord warning label
Facility Electrical Requirement	110-120V AC: 15 A (minimum)
Agency Listing and Certification	Certified in accordance with the NSF/ANSI 456 Standard for Vaccine Storage. UL, C-UL, ETL, C-ETL listed (either single or dual agency listings) and certified to UL471 standard, hydrocarbon refrigerant safety.
Included Accessories	Temperature monitor device (TMD) complies with the current CDC guidelines, with 3 years certification of calibration, “buffered” probe in the product simulated solution, min/max memory, field installable, and visual & audible temp alarm Pharmacy refrigerator/freezer toolkit and temperature logs

Refrigeration System

Compressor	Hermetic, high performance
Refrigerant	EPA SNAP compliant, R600a, Isobutane
Condenser	Hybrid fin and tube with low noise fan
Evaporator	Plate wall
Defrost	Cycle optimized, zero energy

Performance

Uniformity[1] (Cabinet air)	+/- 0.8°C
Stability[2] (Cabinet air)	+/- 1.2°C
Maximum temperature variation (Cabinet air)	+/- 1.4°C
Temperature rise after 8 sec door openings	Temperature did not exceed 6.4°C at any probe for all required NSF/ANSI 456 testing protocols[3]
Recovery after 3 min door opening	All probes recover to under 8°C within 4.8 min.
Energy consumption	1.15 KWh/day[4]
Average heat rejection	1.57 KWh/day (224 BTU/h)[4]
Noise pressure level (dBA)	43 or less installed
Pull down time to nominal operating temp	35 min

Controller, Configuration, Alarms and Monitoring

Controller technology	Parametric, microprocessor, LED display with 0.1°C resolution
Temperature setpoint range	1°C to 10°C (Setpoint must remain unaltered from the factory setting to remain compliant with NSF/ANSI 456 Standard for Vaccine Storage requirements)
Display probe	Calibrated, stainless steel
External alarm connection	State switching remote alarm contacts Visual and audible indicators
Alarms	High / Low temperature, compliant with alarm requirements defined in the NSF/ANSI 456 Standard for Vaccine Storage
Simulator ballast	Glass bead thermal media

Performance data acquired at 22°C ambient, using NSF/ANSI 456 compliant validation ballast probes, empty chamber, 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

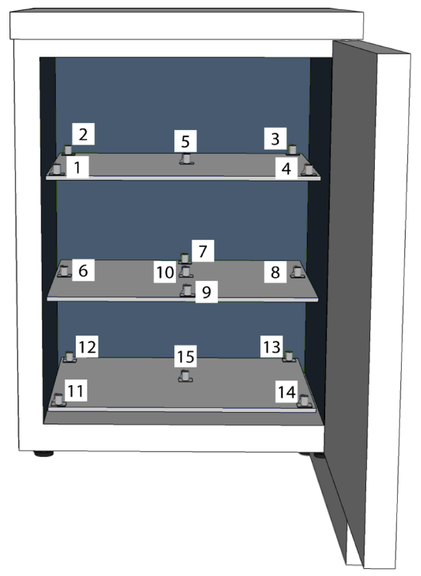
2 - Stability is defined as the maximum variance in temperature experienced by any single probe over the testing period

3 - Temperature performance for all loaded and unloaded door opening protocols, all alarm, controller and probe requirements as defined in the NSF/ANSI 456 standard for vaccine storage

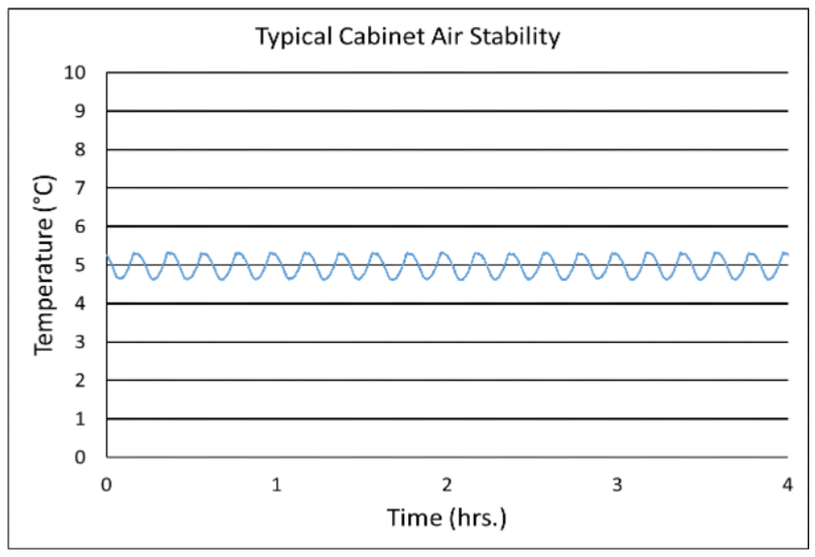
4 - Data per Energy Star test results or equivalent testing and calculation. Heat rejection based on daily averages, not continuous operation. Performance exceeds Energy Star requirements.

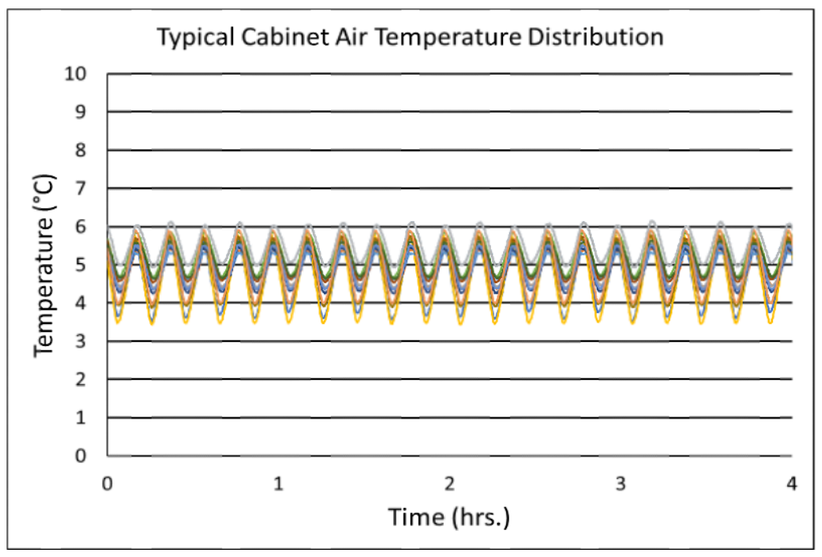
Temperature Probes

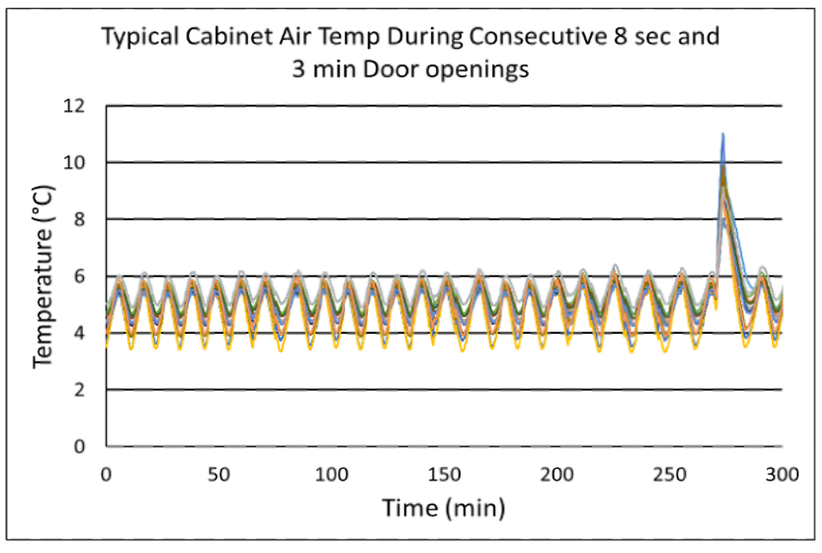
Probe	Ave	Min	Max
1	4.6	3.5	5.8
2	4.9	4.3	5.4
3	5.0	4.4	5.6
4	4.6	3.4	5.8
5	5.0	4.6	5.3
6	5.3	4.7	5.9
7	4.8	4.2	5.5
8	5.1	4.5	5.8
9	4.8	3.9	5.8
10	4.8	3.9	5.8
11	5.5	4.9	6.2
12	5.1	4.6	5.6
13	4.9	4.3	5.5
14	4.9	4.0	5.9
15	5.5	4.9	6.2



Temperature Charts







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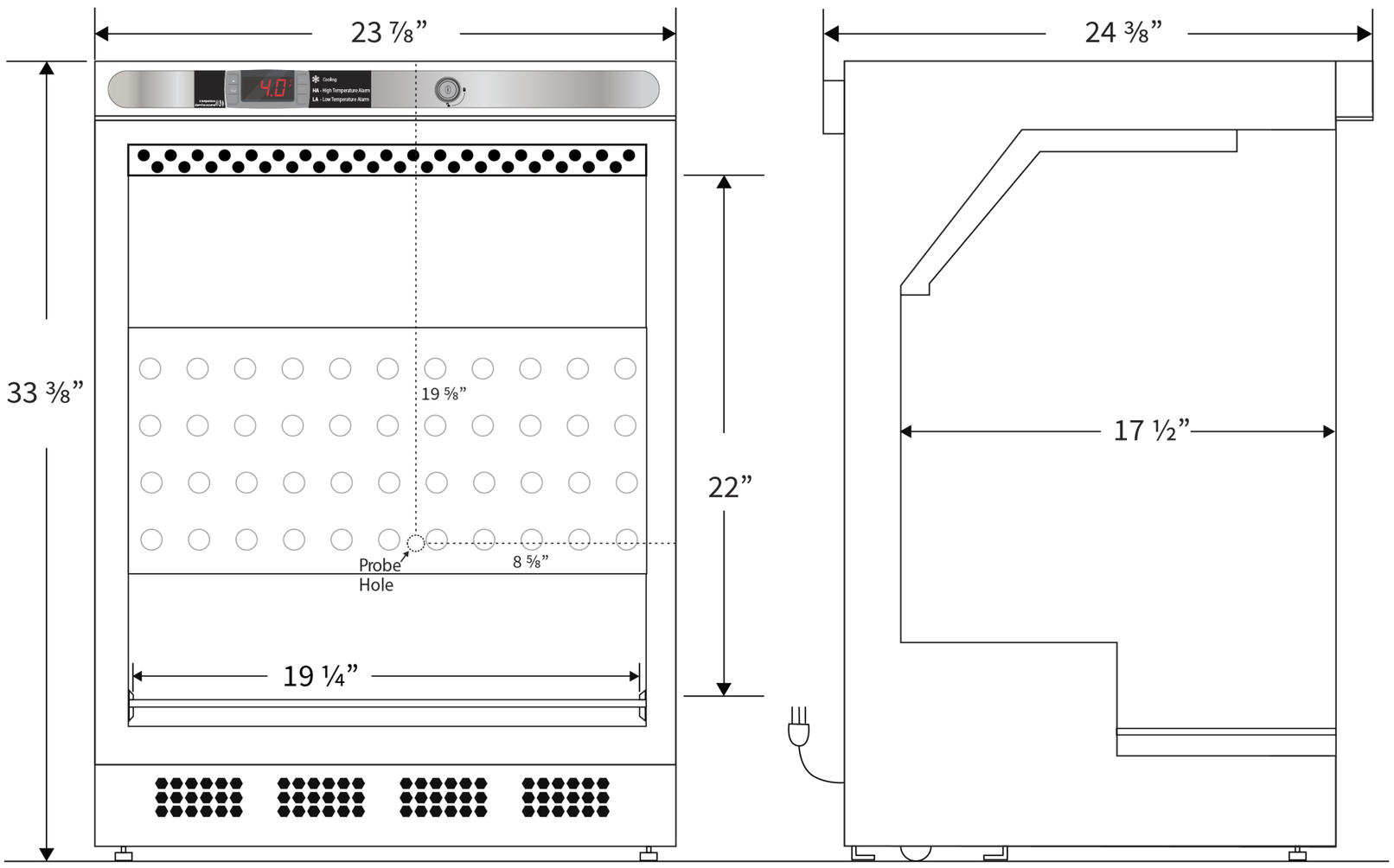
Images

Dimensions

	Width	Depth	Height	Door Swing	Total open Depth
Exterior	23 7/8"	24 3/8"	33 3/8"	23 1/2"	46"
Interior	19 1/4"	17 1/2"	22"		



Contact

Phone	866-674-7220
Email	info@bsilab.com
Rev_10052022	