

GMP Solution

GMP Manufactured Process Chemical

Sodium Hydroxide Solution 25%, GMP Grade

Low Chloride, Low Iron, Made with WFI, BET Tested, GMP Manufactured

INTENDED FOR USE IN PHARMACEUTICAL GMP PROCESSES

High Purity Sodium Hydroxide 25% solution is intended for use in critical pharmaceutical processes, both upstream and downstream. This product is manufactured utilizing a proprietary, fully dedicated, validated GMP system that utilizes multiple manufacturing and purification steps to achieve high purity results without the use of pellets.

Lead Time: 1-2 Months

Minimum Order Quantity: 1000-liters

Na⁺OH⁻

Formula: NaOH F.W.: 40.00 g/mol

Density: 1.280 g/cm³ @ 20°C

Storage Temp: Ambient

CAS #: 1310-73-2 EC#: 215-185-5 UN: UN1824 ADR: 8,II

Merck Index: 14,08627

BIO PHARMA GRADE | Product Code: NAHY-4151 | Previously: NH4151

NaOH • F.W. 40.00 g/mol. • CAS# 1310-73-2



These are general specifications. BioSpectra will customize our products to meet your quality based requirements.

ANALYSIS	SPECIFICATIONS
Appearance and Color	Clear / Colorless Liquid
Assay (NaOH)	24.5 – 25.5%
Chloride (Cl)	≤ 5 ppm
Endotoxin	≤ 2.0 EU/mL
Heavy Metals (as Pb)	≤1ppm
Iron (Fe)	≤ 0.500 ppm

General Product Description:

The manufacturing of Bio Pharma Grade Sodium Hydroxide NAHY-4151 is performed at BioSpectra's Bangor, PA, US FDA registered, GMP facility and is conducted in a dedicated processing area using only dedicated equipment.

Molecular Formula: NaOH

Molecular Weight: 40.00 g/mol

CAS #: 1310-73-2

- Sodium Hydroxide 25% solution is a clear, colorless liquid.
- There are no known major food allergens (as defined by FDA and WHO) in the manufacture of this product.
- BioSpectra certifies that all Sodium Hydroxide 25% solution, NAHY-4151 manufactured at BioSpectra and its raw materials are not derived from or come in contact with animal parts, products and/or byproducts.
- Sodium Hydroxide 25% solution manufactured at BioSpectra and any raw materials used in the manufacture of Sodium Hydroxide 25% solution at BioSpectra are not subject to genetic modification.



Quality Assurance / Regulatory Support / Quality Control

Suitable for Research and Diagnostic Each Batch 100% Analyzed Management of Change Validated Analytical Methods Compendial Testing Trace Metals Analyzed Stability Testing Program BioSpectra Supply Chain Audit Trail Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Management of Change Validated Analytical Methods Compendial Testing Trace Metals Analyzed Stability Testing Program BioSpectra Supply Chain Audit Trail Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Validated Analytical Methods Compendial Testing Trace Metals Analyzed Stability Testing Program BioSpectra Supply Chain Audit Trail Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Compendial Testing Trace Metals Analyzed Stability Testing Program BioSpectra Supply Chain Audit Trail Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Trace Metals Analyzed Stability Testing Program BioSpectra Supply Chain Audit Trail Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Stability Testing Program BioSpectra Supply Chain Audit Trail Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
BioSpectra Supply Chain Audit Trail Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Product Origin Statement Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Customer Quality Audits Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Validated Manufacturing Process US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
US Manufactured at BioSpectra IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
IPEC cGMP Compliant Manufactured Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Customized Additional Specifications Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Multi-Compendial Testing Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Low Bioburden Low Endotoxin (LBLE) Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested
Enzyme Tested Suitable for use as Excipient Microbial / Endotoxin Tested ✓
Suitable for use as Excipient Microbial / Endotoxin Tested
Microbial / Endotoxin Tested ✓
M C 4 1 ED A D 4 1 E 1 4
Manufactured in FDA Registered Facility ✓
Customized Manufacturing Schedule
Custom Regulatory Packet
Accelerated Stability ✓ Video Conference access to BioSpectra Sites ✓
Complete access to Broduct Traceability
Access to Supply Chain Information

✓ indicates an attribute or level of compliance which is granted or available based on the purchase of the product grade.

Bio Pharma Grade: Intended for use as IPEC cGMP Compliant Chemical

LBLE: LBLE applies when product specifications include requirements for Bioburden Testing (TAMC/TYMC and/or Endotoxin).

LBLE stands for Low Bioburden, Low Endotoxin non-sterile products suitable for further use in parenteral manufacturing and other sterile applications.

GMP Compliance:

Bio Pharma Grade Sodium Hydroxide 25%, NAHY-4151 is suitable for use as a process chemical. It is manufactured in accordance with the IPEC-PQG Joint Good Manufacturing Practice Guide. This grade of Sodium Hydroxide 25% Solution is not suitable to be used as an Active Pharmaceutical Ingredient, Drug, Drug Product or Household Item.

Retest Date:

The recommended retest period for Sodium Hydroxide 25% solution is two years from the date of manufacture.

Storage and Shipping Conditions:

Ship and store in ambient temperature, there is no impact to the product within ambient conditions of 10-40°C. Store in a clean and dry area. Store in the original container. **Material will freeze at slightly lower temperatures**. Keep above 16°C to prevent freezing. Warming the product will allow for full dissolution of material.

GHS Classification:

Hazard Pictogram

Signal Word: Danger

Hazard Statements (GHS &CLP)

H290 - May be corrosive to metals.

H314 - Causes severe skin burns and eye damage.

H318 - Causes serious eye damage.

H402 - Harmful to aquatic life.

Stability and Reactivity:

Chemical Stability: Stable.

Possibility of Hazardous Reactions: Will not occur.

Incompatible Materials: Acids, organic materials, chlorinated

solvents, aluminum, phosphorus, zinc, tin.

Hazardous Decomposition Products: Sodium oxides.

Physical and Chemical Properties:

Appearance: Colorless liquid.

Odor: Odorless.

Odor threshold: Not Available.

pH: ~14

Boiling range: 112°C to 140°C Flash Point: Not flammable. Density: 1.280 g/cm³ at 20°C Solubility: Soluble in water.

Package Sizes:

940L totes, 200L drums, 19L pails, 10L pails, 4x4L case and 6x1L case.

