

Linde Nitrous Oxide Plant Production Quality GRADE 2.5 (99.5%)

Sampling point	Purity / Impurities	Unit	Medical quality according to European Pharmacopoeia (Ph. Eur.) ¹⁾	Medical quality produced by the plant ³⁾
Purity in storage tank liquid phase	N ₂ O	% (V/V)	> 98,0	> 99,5
Impurities in storage tank liquid phase	N ₂	ppm (V/V)	not specified ²⁾	< 5000,0
	O ₂ + Ar	ppm (V/V)	not specified ²⁾	< 1000,0
	CO	ppm (V/V)	< 5,0	< 1,0
	CO ₂	ppm (V/V)	< 300,0	< 5,0
	H ₂ O	ppm (V/V)	< 67,0	< 10,0
	NO + NO ₂	ppm (V/V)	< 2,0	< 2,0

¹⁾ Analysis of gaseous phase, with containers (cylinders/ bundles) in the vertical position with the outlet valve uppermost, kept at room temperature for at least 6 hours before carrying out the analysis and only if the product temperature in the container is 15°C.

²⁾ There are no specification limits for analysis of N₂ and O₂ + Ar in Ph.Eur.

³⁾ Analysis of storage tank liquid phase.

Linde Nitrous Oxide UHP Production Quality GRADE 5.0 (99.999%)

Sampling point	Purity / Impurities	Unit	UHP raw material quality Before UHP purification	UHP (industrial quality) After UHP purification
Purity in storage tank liquid phase	N ₂ O	% (V/V)	> 99,8	> 99,999
Impurities in storage tank liquid phase	N ₂	ppm (V/V)	< 5000,0	< 2,0 ²⁾
	O ₂	ppm (V/V)	< 1000,0	< 1,0
	CO	ppm (V/V)	< 0,5	< 0,5 ²⁾
	CO ₂	ppm (V/V)	< 0,5	< 0,5
	H ₂ O	ppm (V/V)	< 1,0	< 1,0
	CH ₄	ppm (V/V)	< 1,0	< 1,0
	H ₂	ppm (V/V)	< 0,5	< 0,5
	NH ₃	ppm (V/V)	< 1,0	< 1,0
	NO-NO ₂	ppm (V/V)	< 1,0	< 1,0

¹⁾ UHP unit can remove N₂ and O₂ gases from the raw material produced by the production plant

²⁾ The impurities quantity in the final UHP product is equal with the impurities measured in the raw material.