NITREX



GN01 NITREX Extra Sensitive



Latex-Free

Ultimate Tactile Sensitivity

Strong Chemical Resistance

Passes Viral Penetration Tests (F1671)

Tested for use with Chemotherapy Drugs

NITREX Extra Sensitive examination gloves are manufactured from a high quality nitrile formulation resulting in a strong performing glove with a high level of tactile sensitivity.

The advanced thin film technology utilised in the manufacture process results in a glove, which provides outstanding barrier protection against, viruses, bacteria, chemicals and has been tested for permeation performance against 13 chemotherapy drugs.

Specification

Glove details	Specification	
Material	Nitrile	
Length	Min. 240mm	
Protein Level	Nitrile contains no latex proteins	
Surface	Micro-Textured Fingers	
Shape	Ambidextrous	
Colour	Blue	
Sterilisation	Non Sterile	
Shelf Life	3 Years	

Physical Properties

Glove details	Properties
Force at Break (Newtons)	>6
Freedom from Holes	AQL 1.5
Length (mm)	Min. 240
Cuff Thickness (mm)	0.04
Palm Thickness (mm)	0.05
Finger Thickness (mm)	0.06

AQL (Acceptable Quality Level) refers to the maximum number of defective products that could be considered acceptable during the random sampling of an inspection, in this case freedom from holes in gloves.

The NITREX range of examination gloves has a freedom from holes of AQL 1.5. The lower the number, the fewer the holes and the higher the quality of gloves.

Sizing

Size	Palm width
Extra Small	75 ± 5mm
Small	85 ± 5mm
Medium	95 ± 5mm
Large	106 ± 5mm
Extra Large	116 ± 5mm

Quantities

Code	Box Qty	Case Qty
GN01E	200 Singles	10 Boxes
GN01S	200 Singles	10 Boxes
GN01M	200 Singles	10 Boxes
GN01L	200 Singles	10 Boxes
GN01X	200 Singles	10 Boxes

Re-order Codes

Size	MPC	NHS Code
Extra Small	GN01E	FTE1744
Small	GN01S	FTE1745
Medium	GN01M	FTE1746
Large	GN01L	FTE1747
Extra Large	GN01X	FTE1748

Quality Standards: Manufactured in a facility holding ISO 9001 and ISO 13485.

Endotoxin: Independent microbiology laboratories conduct routine monitoring of the product manufacturing facility to ensure the bacterial counts in all critical areas are controlled within our stringent specification.

Product standards: The product is tested and manufactured to the following standards: EN455-1, EN455-2, EN455-3, EN455-4.

Storage: Store in a cool, dry place away from sources of heat or direct sunlight and shielded from ozone and UV light.

Disposal: Dispose of gloves as clinical waste. Transit case and box can be recycled as paper or disposed of as clinical waste. Please follow local regulation and guidance.

Shelf life: Three (3) years from date of manufacture.

Country of origin: Malaysia

Notice: This product is free from natural rubber latex and is suitable for use in latex-free environments. Chemical accelerators are used in the manufacture process, which can cause skin reactions in some users. Stop use of the product if skin reactions occur.

Registering authority: In Europe the gloves are CE marked (notified body BSI, number 0086) indicating compliance with Council Directive 93/42/EEC.











ASTM D6978-05