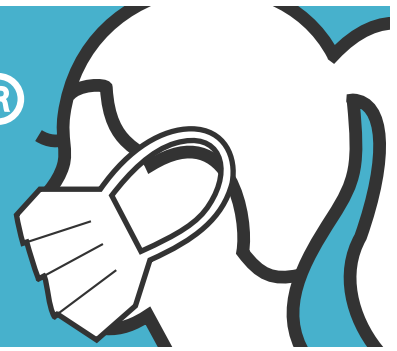


PROTECT[®] AND CARE



PRODUCT FILE

Nitrile medical examination gloves

Non sterile 3.5 g

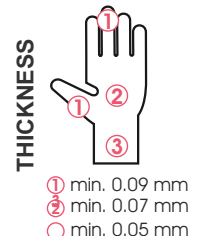
Ref. A00136

Specifications

- Single use
- Suitable for HACCP and medical purposes
- Prevent contamination & protect against viruses, bacteria & fungi
- 100% Nitrile: stronger than other glove materials such as latex, natural rubber or vinyl.
- Are flexible and offer much greater tactile sensitivity than standard diagnostic gloves.
- Good puncture and tear resistance
- Hypo allergenic: - Latex free
- Powder free (less chance of irritation.)
- Impermeable when used in wet or greasy environments
- Comfortable fit:
 - fitted with roll cuffs for easy donning and doffing
 - ambidexterous
- The roughened fingertips ensure a secure grip, even in wet environments

Sizes:

	Length mm	Palm width mm
S 6-7	240	80 +/-10
M 7-8	240	95 +/-10
L 8-9	240	110 +/-10
XL 9-10	240	+110 +/-10



- Clear size indication with color
- Packaging with barcode and QR code
- Info in 27 languages: NL, FR, EN, DE, ES, IT, PT, BG, CS, DA, EL, FI, HU, HR, KK, LT, LV, NO, PL, RO, RU, SK, SL, SR BS, SV, TR & UK

Packaging

Dispenser box of 100 pcs/box - 10 boxes/master box = 1000 gloves (500 pairs)

EN ISO 374-1:2016
+A1:2018
TYPE B

GESCHIKT VOOR HACCP EN MEDISCHE DOELEINDEN
APPROPRIE POUR HACCP ET SOINS MEDICALES
SUITABLE FOR HACCP AND MEDICAL PURPOSES
FÜR HACCP UND MEDIZINISCHE ZWECKE GEEIGNET

Res AP (2004/4
Regulation (EU) 10/2011
Ref. EN 1186: Part 2, Part 3
& Part 14 incl. Olive oil

Permeation Performance Level

Letter	Chemical	CAS	Permeation Level EN ISO 374-1:2016+A1:2018 Type B	Degradation Level EN ISO 374-4:2019
K	40% Sodium Hydroxide	1310-73-2	6	-8.3%
P	30% Hydrogen Peroxide	7722-84-1	2	34.1%
T	37% Formaldehyde	50-00-0	4	34.3%

Measured breakthrough
time (minutes)

1	2	3	4	5	6
>10	>30	>60	>120	>240	>480

In compliance with: MDR (EU) 2017/745 Class I
PPE Regulation (EU) 2016/425: Category III
EN ISO 21420:2020
EN 455-1-2020, EN 455-2-2015, EN 455-3-2015
ASTM D 6978, ASTM F 1671

EU Type Examination and Module C2 Notified Body 2777
SATRA TECHNOLOGY Europe Limited, Bracetown Business Park,
Clonee Dublin 15, D15 YN2P, Republic of Ireland

Test acc. to ASTM D 6978 - Test Chemotherapy

TEST CHEMOTHERAPY DRUG AND CONCENTRATION	MINIMUM BREAKTHROUGH DETECTION TIME (Specimen 1/2/3) (Minutes)
Carbimide (BCNU) 3.3 mg/ml (3,300 ppm)	14.7 (15.1, 14.7, 16.8)
Cisplatin 1.0 mg/ml (1,000 ppm)	240 min.
Cyclophosphamide (Cytoxin) 20 mg/ml (20,000 ppm)	240 min.
Cytarabine 100 mg/ml (100,000 ppm)	240 min.
Doxorubicin (IDC) 10.0 mg/ml (10,000 ppm)	240 min.
Doxorubicin hydrochloride 2.0 mg/ml (2,000 ppm)	240 min.
Etoposide (Etopos) 20.0 mg/ml (20,000 ppm)	240 min.
Fluorouracil 50.0 mg/ml (50,000 ppm)	240 min.
Irinotecan 50.0 mg/ml (50,000 ppm)	240 min.
Methotrexate 25 mg/ml (25,000 ppm)	240 min.
Mitomycin C 0.5 mg/ml (500 ppm)	240 min.
Mitomycin 2.0 mg/ml (2,000 ppm)	240 min.
Paclitaxel (Taxol) 6.0 mg/ml (6,000 ppm)	240 min.
Thioplep 10.0 mg/ml (10,000 ppm)	58.8 (110.0, 58.8, 67.0)
Vincristine Sulfate 1.0 mg/ml (1,000 ppm)	240 min.

ISO 374-5:2016

TESTED IN ACCORDANCE TO:
EN ISO 374-5:2016 and EN ISO 374-4:2019
Degradation results indicate the change in puncture resistance
of the gloves after exposure to the challenge chemical
PROTECTION AGAINST BACTERIA, VIRUSES & FUNGI- PASS

EN ISO 374-5

EN ISO 21420

EN 455
1+2+3

CE 2777

AQL 1.0