CONTIPHARMA PROTECT

a member of Bedelco group

2022 CATALOG

CONTIPHARMA
a member of Bedelco group
PROTECT

SARS-CoV-2 Antigen Test kit

Colloidal Gold Chromatographic Immunoassay

Component	Quantity	Contain	Weight	Dimension (mm)
Test card	25 packs	Test card x 25 pcs Desiccant x 25 pcs	300g	200x115x80
Extraction tube	1 pack	Extraction tube x 25 pcs		
Nozzle cap	2 packs	Nozzle cap x 25 pcs		
Swab	25 packs	Nasopharyngeal swab x 25 pcs		
Extraction	1 bottle	Extraction buffer x 8ml		
Instruction for use	1 copy	1 copy		

PRODUCT NAME

Severe Acute Respiratory Syndrome Coronavirus 2(SARS-CoV-2) Antigen Test Kit (Colloidal Gold Chromatographic Immunoassay)

SPECIFICATION

Ref. No.	Amount	Test cassette	Swab	Extraction buffer	Extraction tube
SC0232	1 T/Kit	1 pc	1 pc	1 x 0.4mL/vial	1 pc
SC0233	2 T/Kit	2 pcs	2 pcs	2 x 0.4mL/vial	2 pcs
SC0234	5 T/Kit	5 pcs	5 pcs	5 x 0.4mL/vial	5 pcs

INTENDED USE

The SARS-CoV-2 Antigen Test Kit is an immunochromatographic test system for the rapid, qualitative detection of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) antigen in human nasal swab specimens, can be used for diagnosis of coronavirus infection disease (COVID-19) in vitro, which is caused by SARS-CoV-2.

The SARS-CoV-2 Antigen Test kit provides preliminary test results, with negative results don't preclude SARS-CoV-2 infection. Cannot be used as the sole basis for treatment or other management decision. The test offers individuals the opportunity to take the nasal swab themselves under the supervision of a medical specialist.

PRINCIPLE

The SARS-CoV-2 Antigen Test Kit is based on colloidal gold immunochromatography method to detect SARS-CoV-2 N protein in respiratory secretions and other specimens. When the specimen is added into the test device, the specimen is absorbed into the device by capillary action, mixes with the gold-labeled antibody, and flows across the precoated membrane.

The SARS-CoV-2 antigen in specimen captured by the gold-labeled antibody S1a bound to antibody S1 immobilized in the Test Region (T) of the membrane, and this produces a colored test band that indicates a positive result.

When there is no SARS-CoV-2 antigen in the specimen or the concentration is lower than the detection limit of the test, there is not a visible colored band in the Test Region (T) of the device. This indicates a negative result.

To serve as a procedure control, a colored line will appear at the Control Region (C), if the test has been performed properly.

MATERIAL REQUIRED BUT NOT PROVIDED

Timer

Personal protective equipment, such a protective gloves, medical mask, goggles and lab coat.

Appropriate biohazard waste container and disinfectants.

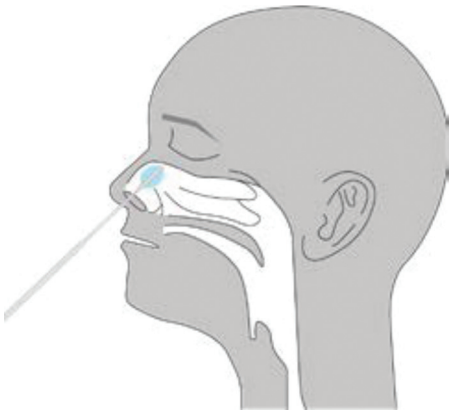
STORAGE CONDITION

The test card is stable for 12 months (while sealed in an aluminum foil bag) if stored at 2~30°C. When the test environment humidity is more than 60%, the test card needs to be used immediately after the opening of the aluminum foil bag. When the test environment humidity is less than 60%, the test card needs to be used within 1 hour after the opening of the aluminum foil bag.

SPECIMEN COLLECTION AND PREPARATION

Specimen collection:

1. It is applicable to the diagnosis of the Novel coronavirus from the samples of nasal swab. Use freshly collected samples for optimal test performance. Inadequate sample collection or improper sample handling may yield a false-negative result.
2. For the sampling of nasal swabs, insert the swab into one nostril of the patient. The swab tip should be inserted 2 to 4 cm deep, until a resistance can be felt. Then roll the swab 5 times over the mucous membrane inside the nostril to ensure that both mucus and cells are collected. Repeat this step with the same cotton swab in the other nostril so that sufficient sample is collected from both nostrils. Finally withdraw the swab from the nasal cavity.



REQUIREMENTS OF SPECIMENS

1. Remove the cap of an extraction buffer, add all the extraction buffer into the extraction tube.
2. Place the patient swab sample into the extraction tube. Roll the swab head against the inside of the extraction tube at least 3 times, and then wait for 1 minute.
3. Remove the swab while squeezing the swab head against the inside of the tube to expel as much liquid as possible from the swab. Dispose of the used swab in your biohazard waste.
4. Press the nozzle cap tightly onto the tube.

The extracted samples can be stored for no more than 1 hour at room temperature and for no more than 4 hours at a temperature in the range of 2 to 8 °C.

TEST PROCEDURE

1. Take out the test card from the aluminum foil bag and lay it flat on the test bench.
2. Add 3 drops (approximately 80µl) of extracted specimen to the specimen well of the test device.
3. Read the results within 15 minutes.

NOTE : The experiment should be done at 15~30°C.humidity 35%~85%.

INTERPRETATION OF RESULTS

1. The presence of two lines (Test and Control), regardless of the intensity of the test line, indicates a positive result.
2. A single Control Line indicates a negative result.
3. If the control line does not appear, the results are invalid and the test should be repeated.

Positive Négative Invalid

INSTRUCTIONS FOR ACTION ACCORDING TO TEST RESULT

1. In the event of a positive test result:
   - There is currently a suspicion of COVID-19 infection
   - Immediately contact a doctor/family doctor or the local health authority
   - Comply with local self-insulation guidelines
   - to have a PCR confirmation test carried out

2. In case of a negative test result:
   - Continue to comply with all applicable rules regarding contact with others and protective measures.
   - An infection may be present even if the test is negative.
   - In case of suspicion, repeat the test after 1 - 2 days, as the coronavirus cannot be detected accurately in all phases of an infection.

3. In case of an invalid test result:
   - Possibly caused by faulty test execution
   - Repeat the test
   - If test results remain invalid, contact a doctor or COVID 19 test center.

LIMITATION OF METHODOLOGY

1. The Novel SARS-CoV-2 Antigen Test Kit is an acute-phase screening test for qualitative detection. Sample collected may contain antigen concentration below the reagent's sensitivity threshold, so a negative test result does not exclude infection with novel coronavirus.
2. The Novel SARS-CoV-2 Antigen Test Kit detects viable and non-viable novel coronavirus antigen. Test performance depends on antigen load in the sample and may not correlate with cell culture performed on the same sample. A positive test does not rule out the possibility that other pathogens may be present, therefore, the results must be compared with all other available clinical and laboratory information to make an accurate diagnosis.
3. A negative test result may occur if the level of extracted antigen in a specimen is below the sensitivity of the test or if poor quality specimen is obtained.
4. Performance of the test has not been established for monitoring antiviral treatment of novel coronavirus.

5. Positive test results do not rule out co-infections with other pathogens.
6. Negative test results are not intended to rule in other coronavirus infection except the SARS-CoV-2.
7. Children tend to shed virus for longer periods of time than adults, which may result in differences in sensitivity between adults and children List.
8. A negative result may occur if the concentration of antigen in a specimen is below the detection limit of the test or if the specimen was collected or transported improperly, therefore a negative test result does not eliminate the possibility of SARS-CoV-2 infection, and should be confirmed by viral culture or PCR.

PERFORMANCE CHARACTERISTICS

1. SENSITIVITY & SPECIFICITY

Nasal swab specimens from 226 patients, which included 107 COVID-19 positive and 119 COVID-19 negative results confirmed by clinical diagnosis judgement. The result of clinical evaluation of SARS-CoV-2 Antigen Test Kit was as follows:

Method		PCR		SUM
SRAS-CoV-2 Antigen Test Kit	Result	Positive	Negative	
	Positive	102	1	103
	Negative	5	118	123
SUM		107	119	226
Sensitivity		95.33% (95%CI:91.31%~96.20%)		
Sensitivity		99.16% (95%CI:95.39%~99.85%)		
Accuracy		97.35% (95%CI:95.37%~97.74%)		

2. LIMIT OF DETECTION (LOD)

2019-nCoV Concentration	1 X 106 $TCID_{50}$/mL					
Dilution	1/100	1/200	1/400	1/800	1/1600	1/3200
Concentration in Dilution tested ($TCID_{50}$/ml)	$1X10^4$	$5X10^3$	$2.5X\ 10^3$	$1.25X10^3$	$6.25X10^2$	$3.125X10^2$
Rates of 20 replicates (%)	100(20/20)	100(20/20)	100(20/20)	100(20/20)	100(20/20)	10(2/20)
Limit of detection	$6.25X10^2TCID_{50}$/mL					

3. INTERFERENCE EXPERIMENT

The following substances were tested at the concentration shown, and no interference was found.

Mucin	100µg/mL	Acetylsalicylic acid	3.0 mM
Whole Blood	5% (v/v)	Ibuprofen	2.5 mM
Biotin	100µg/mL	Mupirocin	10 mg/mL
Neo-Synephrine (Phenylephrine)	5%(v/v)	Tobramycin	10µg/mL
Afrin Nasal Spray (Oxymetazoline)	5%(v/v)	Erythromycin	50uM
Saline Nasal Spray	5%(v/v)	Ciprofloxacin	50uM
Homeopathic	5%(v/v)	Ceftriaxone	110mg/mL
Sodium Cromoglycate	10 mg/mL	Meropenem	3.7µg/mL
Olopatadine Hydrochloride	10 mg/mL	Tobramycin	100µg/mL
Zanamivir	5 mg/mL	Histamine	100µg/mL
Oseltamivir	Hydrochloride	100µg/mL	1mmol/mL
Artemether-lumefantrine	10 mg/mL	Peramivir	100µg/mL
Doxycycline hyclate	50uM	Flunisolide	0.64nmol/ L
Quinine	50uM	Budesonide	0.3ng/mL
Lamivudine	150uM	Fluticasone	6µg/mL
Ribavirin	1 mg/mL	Lopinavir	8.2mg/mL
Daclatasvir	1 mg/mL	Ritonavir	417.8ng/mL
Acetaminophen	1 mg/mL	Abidor	N/A

4. CROSS-REACTIVITY

Virus/Bacteria/ Parasite	Strain	Source/Specimen type	Concentration	Result
SRAS-coronavirus	N/A	SINO/recombinant protein	25ug/mL	Negative
MERS-coronavirus	N/A		72 ug/mL	Negative
Adenovirus	Type 1	AMMS Inactivated culture virus	1,5E+06TCID$_{50}$/mL	Negative
	Type 3		7,5E+06TCID$_{50}$/mL	Negative
	Type 5		4,5E+06TCID$_{50}$/mL	Negative
	Type 7		1,0E+06TCID$_{50}$/mL	Negative
	Type 8		1,0E+06TCID$_{50}$/mL	Negative
	Type 11		2,5E+06TCID$_{50}$/mL	Negative
	Type 18		2,5E+06TCID$_{50}$/mL	Negative
	Type 23		6,0E+06TCID$_{50}$/mL	Negative
	Type 55		1,5E+06TCID$_{50}$/mL	Negative
Influenza A	H1N1 Denver	AMMS / Inactivated culture virus	3,0E+08TCID$_{50}$/mL	Negative
	H1N1 WS/33		2,0E+08TCID$_{50}$/mL	Negative
	H1N1 A/Mal/302/54		1,5E+08TCID$_{50}$/mL	Negative
	H1N1 New Caledonia		7,6E+08TCID$_{50}$/mL	Negative
	H3N2 A/ Hong Kong/8/68		4,6E+08TCID$_{50}$/mL	Negative

Virus/Bacteria/ Parasite	Strain	Source/Specimen type	Concentration	Result
Influenza B	Nevada/03/2011	AMMS / Inactivated culture virus	1,5E+08TCID$_{50}$/mL	Negative
	B/Lee/40		8,5E+08TCID$_{50}$/mL	Negative
	B/Taiwan/2/62		4,0E+08TCID$_{50}$/mL	Negative
Virus respiratoire syncytial	N/A	AMMS / Inactivated culture virus	2,5E+06TCID$_{50}$/mL	Negative
Legionella pneumophila	Bloomington-2	AMMS / Inactivated culture virus	1×10^5 PFU/mL	Negative
	Los Angeles-1		1×10^5 PFU/mL	Negative
	82A3105		1×10^5 PFU/mL	Negative
Mycobacterium tuberculosis	K	AMMS / Inactivated culture virus	1×10^5 PFU/mL	Negative
	Erdman		1×10^5 PFU/mL	Negative
	HN878		1×10^5 PFU/mL	Negative
	CDC1551		1×10^5 PFU/mL	Negative
	H37Rv		1×10^5 PFU/mL	Negative
Streptococcus pneumonia	4752-98 [Maryland (D1)6B-17]	AMMS / Inactivated culture virus	1×10^5 PFU/mL	Negative
	178 [Poland 23F-16]		1×10^5 PFU/mL	Negative
	262 [CIP 104340]		1×10^5 PFU/mL	Negative
	Slovakia 14-10 [29055]		1×10^5 PFU/mL	Negative
Streptococcus pyrogens	Typing strain T1 [NCIB 11841, SF 130]	AMMS / Inactivated culture virus	1×10^5 PFU/mL	Negative
Mycoplasma pneumoniae	Mutant 22	AMMS / Inactivated culture virus	1×10^5 PFU/mL	Negative
	FH strain of E aton Agent [NCTC10119]		1×10^5 PFU/mL	Negative
	36M129-B7		1×10^5 PFU/mL	Negative
Coronavirus	229E	AMMS / Inactivated culture virus	1,5E+06TCID$_{50}$/mL	Negative
	OC43		1,5E+06TCID$_{50}$/mL	Negative
	NL63		1,5E+06TCID$_{50}$/mL	Negative
	HKU1		1,5E+06TCID$_{50}$/mL	Negative
Human etapneumovirus 3 Type B1	Peru2-2002	AMMS / Inactivated culture virus	1,5E+06TCID$_{50}$/mL	Negative
Human Metapneumovirus	IA10-2003	AMMS / Inactivated culture virus	1,5E+06TCID$_{50}$/mL	Negative
(hMPV) 16 Type A1	Type 1	AMMS / Inactivated culture virus	1,5E+06TCID$_{50}$/mL	Negative
	Type 2		1,5E+06TCID$_{50}$/mL	Negative
	Type 3		1,5E+06TCID$_{50}$/mL	Negative
	Type 4A		1,5E+06TCID$_{50}$/mL	Negative
RhinoVIRUS A16	N/A	AMMS / Inactivated culture virus	1,5E+06TCID$_{50}$/mL	Negative

ATTENTION

1. For in vitro diagnostic use only.
2. Proper specimen collection storage and transit are critical to the performance of this test.
3. Use only once.
4. Do not touch the reaction area of test strip.
5. Do not use test kit beyond the expiration date.
6. Do not use the kit if the pouch is punctured or sealed not well.
7. Testing should be applied by professionally trained staff working in certified laboratories or clinics.
8. The test result should be interpreted by the physician along with clinical findings and other laboratory test results.
9. Dispose of test cards and items in contact with samples as medical waste after use.
10. Do not freeze.

INTERPRETATION OF ICONS

	Do not re-use		Temperature limit
IVD	In vitro diagnostics medical device	LOT	Batch code
	Contains sufficient for <n>test		Consult instructions for use
	Manufacturer		Date of manufacture
CE	CE mark		Use-by date
	Keep dry		This way up
	Fragile, handle with care		Stacking layer limit

EC REP Authorized representative in the European Community

GENERAL INFORMATION

Shenzhen Ultra-Diagnostics Biotec. Co., Ltd.
Room 701, No.71-3, Xintian Avenue, Xintian Community, Fuhai Street, Baoan District, Shenzhen, P.R.China 518103
Tel: +86-755-82599902
Fax: +86-755-82599221

EC REP

CMC Medical Devices & Drugs S.L.
C/ Horacio Lengo N°18, CP 29006, Málaga, Spain
Tel: +34951214054
Mail:info@cmcmedicaldevices.com

Issue Date: 24/03/2021

SARS-CoV-2 Antigen Test kit

PRODUCT NAME

NEWGENE - COVID-19 Antigen Detection Kit

SPECIFICATION

Multiple Sampling Methods : Nasal Swab / Nasopharyngeal Swab / Oropharyngeal Swab / Sputum (Saliva)
Multiple Packaging Specifications : 25 Tests/Box, 5 Tests/Box or 1 Test/Box
Global Recognition : Registered in 20+ Countries, Exported to 50+ Countries
Fast Detection : Results in 15 minutes
Superior Performance : High Sensitivity & Specificity

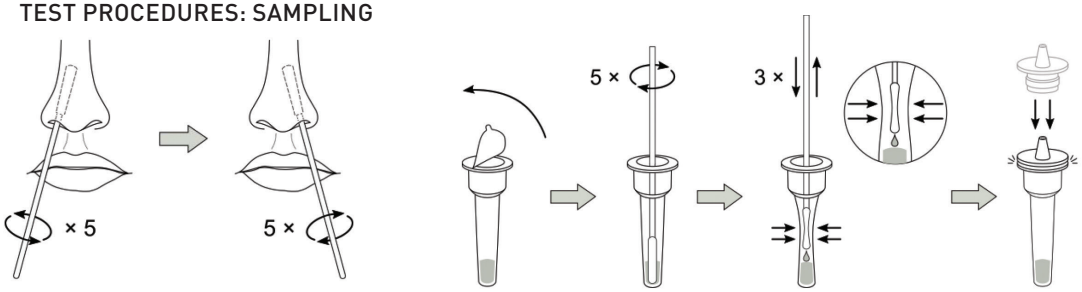
At present, NEWGENE COVID-19 Antigen Detection Kit has registered in many countries, including Germany, France, Italy, Switzerland, Belgium, Portugal, Czech, Denmark, Hungary, Greece, Poland, Moldova, Peru, Argentina, Ecuador, Kenya, Zimbabwe etc., and passed the clinical validation in national lab in Germany, Switzerland, Ecuador, Zimbabwe etc. The products show good performance in sensitivity and specificity compared with international brand products and have exported to more than 50 countries and regions.

PRODUCT INTRODUCTION

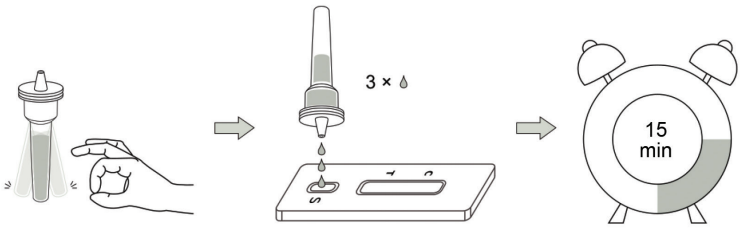
1. NASAL SWAB

Component	25 Tests/box	5 Tests/box	1 Test/box
Test Card	25	5	1
Sample Extraction Tube & Tube Cap	25	5	1
Sampling Swab: for Nasal Swab	25	5	1
Package Insert	1	1	1

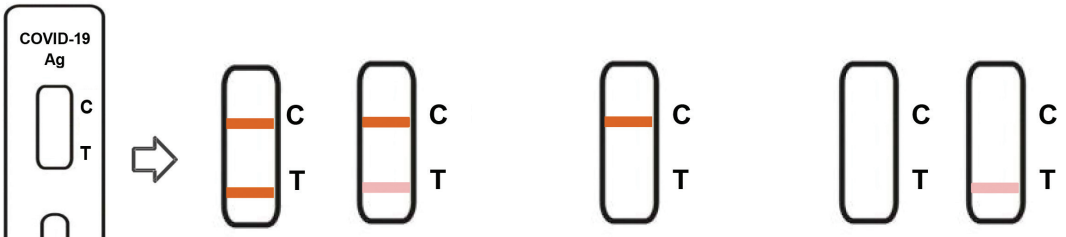
TEST PROCEDURES: SAMPLING



TEST PROCEDURES: DETECTION



INTERPRETATION OF RESULTS



Positive(+):
Red bands appear at both of T and C line in 15 to 30 minutes. A white band at the T line should be considered as a negative result.

Negative(-):
A red band appears at C line while no red band appears at T line in 15 to 30 minutes after sample loading.

Invalid:
As long as no red band appears at C line, it indicates that the test result is invalid, and should retest the sample with another test card.

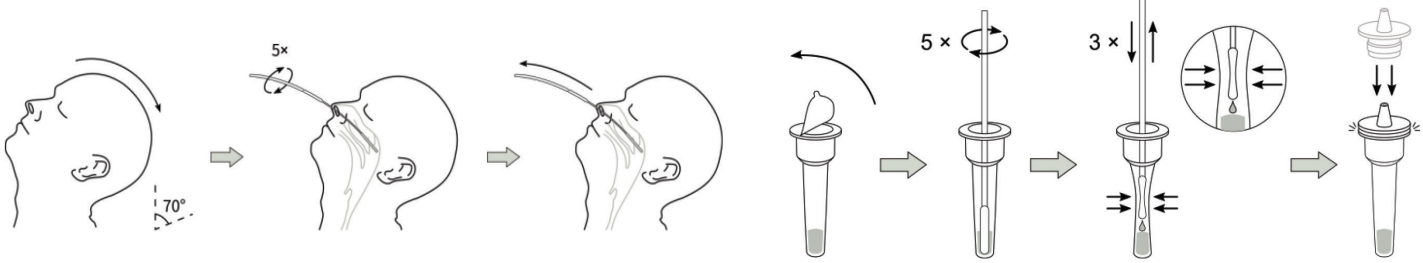
PERFORMANCE

Sensitivity	97.1%
Specificity	99.2%

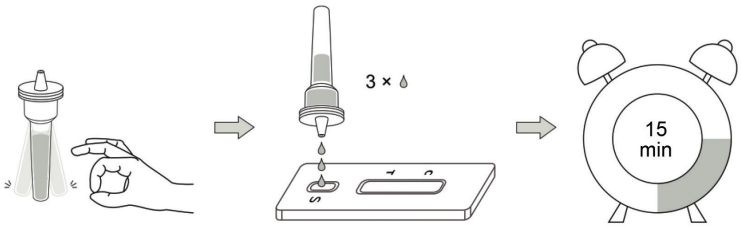
2. NASOPHARYNGEAL SWAB

Component	25 Tests/box	1 Test/box
Test Card	25	1
Sample Extraction Tube & Tube Cap	25	1
Sampling Swab: for Nasopharyngeal Swab	25	1
Package Insert	1	1

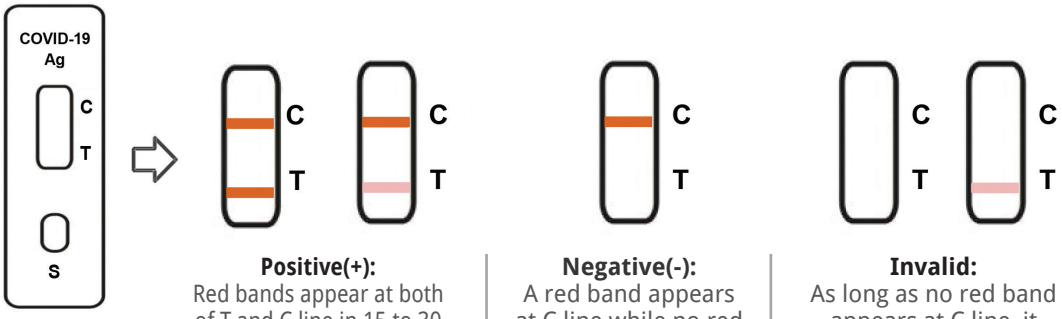
TEST PROCEDURES: SAMPLING



TEST PROCEDURES: DETECTION



INTERPRETATION OF RESULTS



Positive(+):
Red bands appear at both of T and C line in 15 to 30 minutes. A white band at the T line should be considered as a negative result.

Negative(-):
A red band appears at C line while no red band appears at T line in 15 to 30 minutes after sample loading.

Invalid:
As long as no red band appears at C line, it indicates that the test result is invalid, and should retest the sample with another test card.

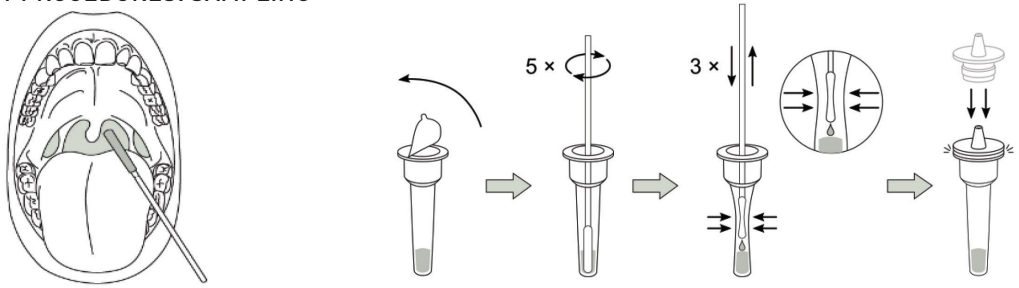
PERFORMANCE

Sensitivity	98,0%
Specificity	99.1%

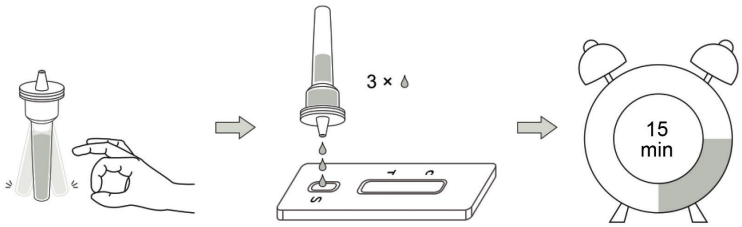
3. OROPHARYNGEAL SWAB

Component	25 Tests/box	1 Test/box
Test Card	25	1
Sample Extraction Tube & Tube Cap	25	1
Sampling Swab: for Oropharyngeal Swab	25	1
Package Insert	1	1

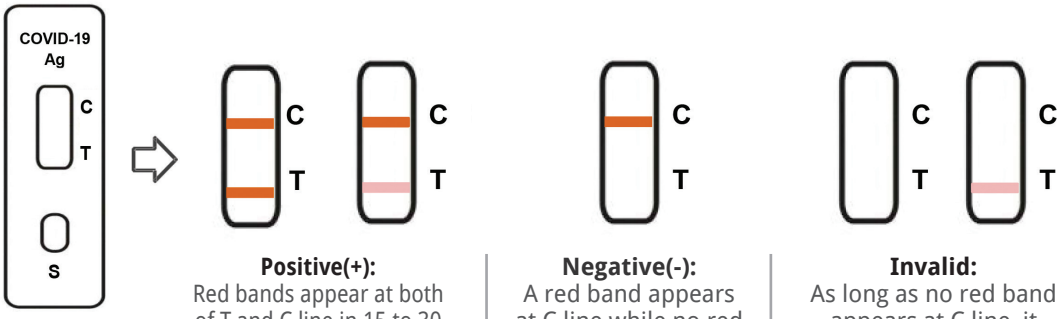
TEST PROCEDURES: SAMPLING



TEST PROCEDURES: DETECTION



INTERPRETATION OF RESULTS



Positive(+):
Red bands appear at both of T and C line in 15 to 30 minutes. A white band at the T line should be considered as a negative result.

Negative(-):
A red band appears at C line while no red band appears at T line in 15 to 30 minutes after sample loading.

Invalid:
As long as no red band appears at C line, it indicates that the test result is invalid, and should retest the sample with another test card.

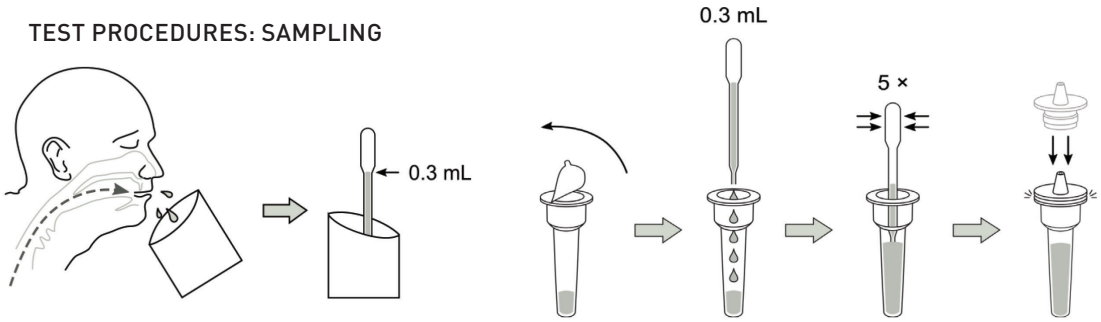
PERFORMANCE

Sensitivity	95,7%
Specificity	99.0%

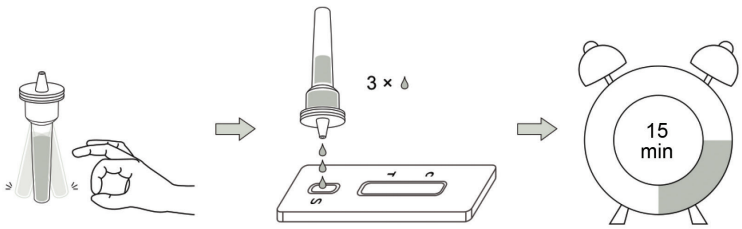
4. SPUTUM / SALIVA SAMPLE

Component	25 Tests/box	1 Test/box
Test Card	25	1
Sample Extraction Tube & Tube Cap	25	1
Paper Cup	25	1
Sputum Dropper	25	1
Package Insert	1	1

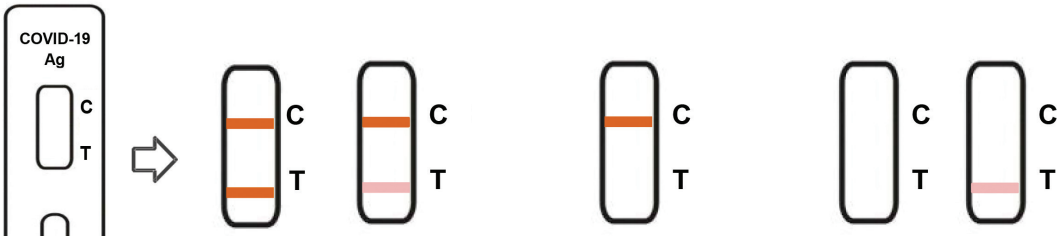
TEST PROCEDURES: SAMPLING



TEST PROCEDURES: DETECTION



INTERPRETATION OF RESULTS



Positive(+):
Red bands appear at both of T and C line in 15 to 30 minutes. A white band at the T line should be considered as a negative result.

Negative(-):
A red band appears at C line while no red band appears at T line in 15 to 30 minutes after sample loading.

Invalid:
As long as no red band appears at C line, it indicates that the test result is invalid, and should retest the sample with another test card.

PERFORMANCE

Sensitivity	97,3%
Specificity	99.2%

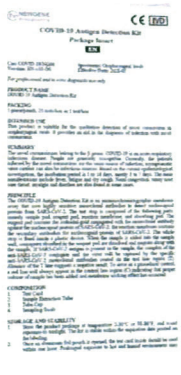
PACKAGING INFORMATION

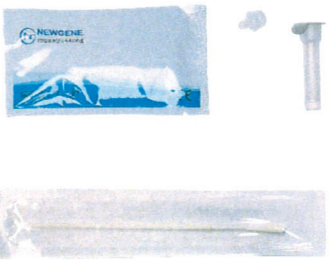
Nasopharyngeal Swab: NPS
Nasal Swab: NS
Oropharyngeal Swab: OS
Sputum / Saliva: S

BOXES

25 TESTS/BOX

Sample	NPS	NPS+S	NS	NS+S	OS	OS+S	S
Box (mm)	230*140*80						230*120*67
Box weight (kg)	0.34	0.38	0.32	0.36	0.35	0.4	0.39
Carton (mm)	585*485*425						510*490*360
Carton weight (kg)	1.5						1.3
PCS/Box	25						
Boxes/Carton	40						
PCS/Carton	1000						
Volume/Carton	0.12CBM						0.09CBM
NW/Carton (kg)	13.6	15.2	12.8	14.4	14	16	15.6
GW/Carton (kg)	15.1	16.7	14.3	15.9	15.5	17.5	16.9





5 TESTS/BOX

Sample	NS	
	Size (mm)	Weight (kg)
Inner box (mm)	193*85*42	0.081
Outer box (mm)	225*197*89	0.5
Carton	470*410*470	1.3
PCS/Inner Box	5	
Inner Boxes/Outer Box	5	
PCS/Carton	500	
Volume/Carton	0.09CBM	
NW/Carton (kg)	10	
GW/Carton (kg)	11.3	

1 TEST/BOX

Sample	S	NS	NS+S	NPS	OS	NPS+S	OS+S
Inner box (mm)	143*83*15			170*66*15			
Inner box weight (kg)	0.026	0.027	0.03	0.024	0.028	0.028	0.032
Outer box (mm)	305*197*88			277*182*112			
Outer box weight (kg)	0.78	0.80	0.88	0.73	0.83	0.83	0.93
Carton (mm)	630*420*470			590*570*395			
Carton weight (kg)	1.8			2.2			
PCS/Inner Boxes	1						
Inner Boxes/Outer Box	25						
PCS/Carton	500						
Volume/Carton	0.13CBM			0.133CBM			
NW/Carton (kg)	15.6	16	17.6	14.6	16.6	16.6	18.6
GW/Carton (kg)	17.4	17.8	19.4	16.8	18.8	18.8	20.4

INTERPRETATION OF ICONS

	Do not re-use		Store between 2-30°C
IVD	In vitro diagnostics medical device	LOT	Batch code
	Contains sufficient for <n>test		Consult instructions for use
	Manufacturer		Date of manufacture
CE	CE mark		Use-by date
	Keep dry		This way up
	Keep away from sunlight		Do not use if packaging is damaged

EC REP Authorized representative in the European Community

GENERAL INFORMATION

Shenzhen Ultra-Diagnostics Biotec. Co., Ltd.
Room 701, No.71-3, Xintian Avenue, Xintian Community, Fuhai Street, Baoan District, Shenzhen, P.R.China 518103
Tel: +86-755-82599902
Fax: +86-755-82599221

EC REP

CMC Medical Devices & Drugs S.L.
C/ Horacio Lengo N°18, CP 29006, Málaga, Spain
Tel: +34951214054
Mail:info@cmcmedicaldevices.com

CONTIPHARMA PROTECT
a member of Bedelco group

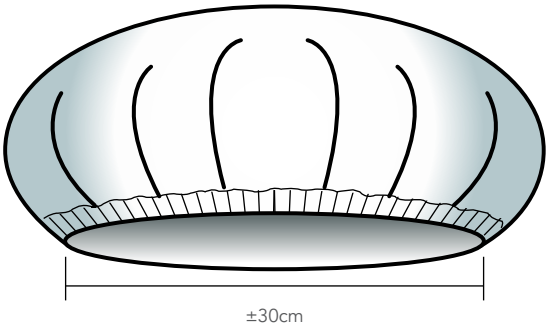
Clip cap blue/white

Ref 10001

Flexible elastic tightening, without pressure
Very light, pleasant to wear
Zero maintenance (single use)

STECHNICAL FEATURES

- Round model with **double elastic**
- Color: blue/white
- Single use
- Unwoven polypropylene
- Weight : 12 g/m²
- Packaging : bags of 100 pcs



APPLICATION

Use : These clip caps can be used in many jobs that require you to protect yourself from minor dirt and/or protect the environment from particles and micro-dust emitted by the user. Pharmaceutical industry, electronics, renovations, maintenance, hospitals, housekeeping, agri-food industry, cheese shops, dairies, factory visits, packaging ...

Cleanliness item : This product is not an individual protective equipment but a cleanliness item; do not use against hazards that could threaten the health or safety of the user. During activities in certain sensitive environments such as food, electronics, it is generally obligatory to remove any personal decorative objects: rings, necklaces, bracelets...

PP Pirat hat with mask

Ref 10002

SPÉCIFICATIONS

Polypropylene hood with built-in Type II mask.
The 25g polypropylene hood incorporates a 3-layer mask with good filtration capacity (type II). Its air permeability ensures great comfort.
It is particularly suitable for food companies and slaughterhouses where a medical mask is required.
A malleable nasal bar provides individual facial adaptation and latex-free elastics also allow allergy sufferers to use the product.

CONTIPHARMA PROTECT
a member of Bedelco group

CoverTex® Coverall C-1

SPECIFICATIONS

Descriptions : · elasticated 3 piece hood for optimum fit • elasticated cuffs, ankles and waist
· 2-way zip fastener.
· zip cover/stom flap.
· antistatic finish.

Material : Multilayer (SMMS) PP spunbond.

PSA category : Cat. III - Type 5 + 6.

EN ISO 13688:2013.

EN ISO 13982-1:2004 + A1:2010.

EN 1073-2:2002.

EN 1149-5:2008.

EN 14325:2004.

EN 13034:2005 + A1:2009 Type 6.

Certificate institute : Centro Tessile Cotoniero e Abbigliamento S.p.A (Centrocot), Piazza S. Anna 2, 21052 Busto Arsizio (VA) - Italy, Notified Body Number 0624.

Certificate number : CE 0672180430-00-00

Sizes : S-M-L-XL-2XL-3XL-4XL-5XL

Color : white
Packaging Unit : 50
Country of Origin : CN / VR China
Customs code : 62101098000

Overalls 35g

Ref 30001

SPÉCIFICATIONS

- Blue/White/Green disposable polypropylene coat for any type of use and effective protection
- 35g/m²
- Cord closure - tie behind
- Air-permeable
- Little absorbent
- Suitable for medical, agri-food, catering, industries
- One-size-fits-all
- Chest circumference: 690 mm x 2
- Length of the sleeve from the collar: 780 mm
- Width of the sleeve: 250 mm
- Total length: 1050 mm
- Packaging: 100 per carton

Overalls 35g white TU

Ref 30002

SPECIFICATIONS

- Disposable polypropylene coat for any type of use and effective protection
- 35 g/m²
- Sewn collar
- Velcro closure at the front
- With pockets
- Air-permeable
- Low-absorbing
- Suitable for medical, agri-food, catering, industry
- One-size-fits-all
- Chest circumference: 690 mm x 2
- Length of the sleeve from the collar: 780 mm
- Width of the sleeve: 250 mm
- Total length: 1050 mm
- Packaging: 100 per carton

Overalls PLP gr

Ref 30004

SPECIFICATIONS

- Color: white/blue
- Polypropylene 25 gr
- Tightened by elastic at the wrists
- Attached behind by cord
- One-size-fits-all

Velcro apron 25 g white

Ref 30006

SPECIFICATIONS

- Apron with Velcro front fastener
- Color: white
- Polypropylene 25g
- With collar
- Wrists tightened by elastic
- Size: M / L / XL

Coveralls

Ref 30007

White polypropylene/polyethylene coveralls with blue stripes, 65 grams, disposable, with hood and elastic at the waist at the wrists and ankles.

Size : L - XL - XXL

These coveralls offer an efficient and inexpensive clothing solution that protects against dry particles such as dirt and dust in the workplace. They are light, comfortable, skin breathes and they can be used in hospitals, food processing plants and general industrial applications.

Easy to endorse, anti-dust, antibacterial, antivirus
Certificates: on request

Overshoes

Ref 40001

SPECIFICATIONS

- Non-slip overshoe in unwoven propylene, non-sterile
- Elastic tightening of the ankles for perfect support
- Antiskid, reinforced and waterproof embossed sole
- Color: white with blue sole
- Dimensions: 16.5 x 41.5 cm
- Grammage: 30g/m²

Our overshoes provide an effective barrier against bacteria and viruses. Wearing these protections also reduces the risk of accidents at work.

ADVANTAGES :

- Ensure the cleanliness of the premises for optimal hygiene;
- Avoid external contamination;
- Protect shoes from splashes or messy splashes;
- Reduce work-related accident
- Protect employees and visitors passing through.

APPLICATION

- Agri-food industries
- Microelectronics sector
- Car line
- Space industry
- Hospitals and health facilities

GOLD TOUCH gloves

SPECIFICATIONS

· Marking of the glove: identification of the manufacturer or his representative, product reference, CE mark and size, EN-standard

CHARACTERISTICS

· Red nylon and black nitrile coated spandex glove.

CONFORMANCE

· CE cat.2 : GAnts with medium design : intermediate risks.
· All EC type examinations have been performed by CTC notified body nr (0075) Cert. nr 0075/032/162/01/19/0079.

PERFORMANCE NEIGHBOURS

Mechanical risks tested according to EN 388:2016

· Abrasion : 4
· Outage : 1
· Tear : 3
· Perforation : 1
· Cutting EN ISO 13997 : X

CLEANING AND MAINTENANCE

It is preferable to clean the gloves dry with a brush. The use of cleaning fluids or chemicals may alter the properties of the gloves, for which the manufacturer cannot be held responsible.

CONSERVATION AND STORAGE

The gloves should be stored in their original packaging in a dry place, away from sunlight and ultraviolet rays. When properly stored, the gloves will retain their properties for at least 2 years. It is impossible to predict their life span as it depends on the use made of them. The user should always check that the gloves are suitable for the intended use.

Mastery type gloves Guyard 1532 NAT LR

SPECIFICATIONS

- Mastery type glove in bovine leather
- Intermediate risks - Cat. II

FEATURES

- Full grain natural bovine leather.
- Elastic tightening on the back.
- American Cup.
- Chrome tanning.
- Length : to 22cm28 according to the size, (+0,5 /-0.5 mm).
- Available sizes : 8 to 11.

COMPLIANCE

- Declaration issued by Leitat (Spain) n 0162 EN 420 / 2003 + A1 / - 09EN / 3882016 - EN ISO 13997
- EU declaration 2016/425 available on demand

RECOMMENDED USES

- Gloves specially designed for small and medium handling.
- Work in a dry environment.

PERFORMANCE LEVELS:

- Abrasion: / 34
- Cut EN /388 2016 : 1 / 5
- Tear: / 24
- Perforation: / 24
- Cutting EN ISO : 13997A / F

PRECAUTIONS FOR USE

- Wear gloves on dry and clean hands. Pre-testing of gloves is recommended, as actual use conditions may differ from EU type testing.
- Before reuse, check the gloves for defects, holes, tears or abnormal wear.

PROTECTION LIMITS:

- Do not use for handling objects with a temperature > 50 C.
- Do not use for handling chemicals or liquids.

SAFETY, COMFORT & DEXTERITY

- Dexterity, level 5.
- The glove does not contain substances at levels known or suspected to have adverse effects on the hygiene or health of the user under the foreseeable conditions of use.

PACKAGING

- Bag: pairs10.
- Boxes: pairs100

MAINTENANCE AND STORAGE

- No particular maintenance is recommended. It is recommended to store the products in a cool, dark and ventilated place.
- The destruction at the end of life must be controlled and respect the European requirements of traceability and respect of the environment.

Guyard gloves 1545 150 pa cH

SPECIFICATIONS

- Mastery type glove in natural goat leather.
- Intermediate risks - Cat. II.

CHARACTERISTICS

- Full grain goat leather.
- Natural crust leather cuff 150 MM.
- Protection of the artery.
- Chrome tanning.
- Lengths: 35 to 39 cm according to the size.
- Available sizes: 8 to 11.

CONFORMANCE

- Declaration issued by Leitat (Spain) n°0162 - EN 420/2003 + A1/2009 - EN 388/2016 - EN 407 : 2004 - ISO 13997 - EN 12477 : 01/A1 : 05 TYPE B.
- EU declaration 2016/425 available on guyard-sa.fr.

USES OF RECEIVED USES

- Gloves designed for welding work.

PERFORMANCE NEIGHBOURS

· Abrasion :	2 / 4
· Cutting :	1 / 5 & A/F
· Tear :	2 / 5
· Perforation :	1 / 5
· Behaviour in fire :	4 / 4
· Contact heat :	1 / 4
· Convective heat :	1 / 4
· Radiant heat :	2 / 4
· Small splashes of liquid metal :	4 / 4
· • Large splashes of liquid metal :	X / 4

PRECAUTIONS OF THE LAW

· Wear gloves on dry and clean hands. Pre-testing of gloves is recommended as actual use conditions may differ from EU type testing.
· Before reuse, check the gloves for defects, holes, tears or abnormal wear.

PROTECTION LIMITES

· Do not use for handling objects with a temperature above 100°C.
· Do not use for handling chemicals.

SAFETY, COMFORT & DEXTERITY

· Dexterity, level 5.
· The glove does not contain substances at levels known or suspected to have adverse effects on the hygiene or health of the user under the foreseeable conditions of use.

CONDITIONNEMENT

· Bag : 10 pairs.
· Boxes: 100 pairs.

MAINTENANCE AND STORAGE

· No particular maintenance is recommended. It is recommended to store the products in a cool, dry and airy place, protected from light.
· The destruction at the end of life must be controlled and respect the European requirements of traceability and respect of the environment.

Guyard Welding Glove 1567 350 DM pH

SPECIFICATIONS

- Welder's glove in bovine crust.
- Intermediate risks - Cat. II.

CHARACTERISTICS

- Gloves in bovine crust.
- Red color.
- Fully lined with fleece.
- Sewn with 100% aromatic polyamide thread.
- Palm and thumb crust reinforcement, brown color.
- Available sizes: 9 to 10.

CONFORMANCE

- Declaration issued by Leitat (Spain) n°0162 - EN 420/03 +A1/09 - 407/04 - EN 388/16 - ISO 13997 - EN 12477/2001 TYPE A.
- EU declaration 2016/425 available on guyard-sa.fr.

RECEIVED USES

- MIG, MAG & ARC welding gloves.

PERFORMANCE NEIGHBOURS

· Abrasion :	4 / 4
· Cutting EN 388 / 2016 :	1 / 5
· Tear :	4 / 4
· Perforation :	4 / 4
· Cutting EN ISO 13997 :	B / F
· Behavior to fire :	4 / 4
· Contact heat :	1 / 4
· Convective heat :	3 / 4
· Radiant heat :	3 / 4
· Small splashes of liquid metal :	4 / 4
· Large liquid metal splashes :	X

PRECAUTIONS OF THE LAW

· Wear gloves on dry and clean hands. Pre-testing of gloves is recommended as actual use conditions may differ from EU type testing.
· Before reuse, check the gloves for defects, holes, tears or abnormal wear.

PROTECTION LIMITES

· Do not use for handling objects with a temperature above 100°C.
· Do not use for handling chemicals or liquids.

SAFETY, COMFORT & DEXTERITY

· Dexterity, level 1.
· These gloves do not contain substances at levels known or suspected to have adverse effects on the hygiene or health of the user under the foreseeable conditions of use.

CONDITIONNEMENT

· Bag : 12 pairs.
· Boxes: 60 pairs.

MAINTENANCE AND STORAGE

· No particular maintenance is recommended. It is recommended to store the products in a cool, dry and ventilated place, away from light.
· The destruction at the end of life must be controlled and respect the European requirements of traceability and respect of the environment.

Hppe Guyard MecaSafe 6550 30 knitted

SPECIFICATIONS

- HPPE knitted glove / glass fiber, polyurethane coating.
- Intermediate risks - Cat. II.

CHARACTERISTICS

- High density polyethylene knit 50.0%, fiberglass 25.0%, polyester 22.0%, spandex 3.0%.
- Grey polyurethane coating in the palm.
- Ventilated back.
- Elasticated cuff with colored overlay indicating size.
- Available sizes: 6 to 11.

CONFORMANCE

- Declaration issued by the CTC (France) n 00075 EN 420 1 03 + Al / 09 - EN 388 12016 - EN ISO 13997.
- EU declaration 2016/425 on request.

USES OF RECEIVED USES

- Handling of dry or slightly greasy cutting parts, glass.
- Industry, construction.

PERFORMANCE NEIGHBOURS

· Abrasion :	4 / 4
· Cutting EN 388 / 2016 :	4 / 5
· Tear :	4 / 4
· Perforation :	3 / 4
· Cutting EN ISO 13997 :	D / F

PRECAUTIONS OF THE LAW

- Before reuse, check the gloves for defects, holes, tears or abnormal wear.
- Wear gloves on dry and clean hands. Pre-testing of gloves is recommended as actual use conditions may differ from EU type testing.

PROTECTION LIMITES

- Do not use for handling objects with a temperature above 50°C.
- Do not use for handling chemicals or liquids.

SAFETY, COMFORT & DEXTERITY

- Dexterity, level 5.
- This PPE does not contain substances at levels known or suspected to have adverse effects on the hygiene or health of the user under the foreseeable conditions of use.

CONDITIONNEMENT

- Bag : 10 pairs.
- Boxes: 200 pairs.

MAINTENANCE AND STORAGE

- No particular maintenance is recommended.
- It is recommended to store the products in a dark, cool, dry and ventilated place.
- The destruction at the end of life must be controlled and respect the European requirements of traceability and respect of the environment.

Nitrile gloves « Purism » G001

Ref 50001

SPECIFICATIONS

G001 «Purism» examination gloves are suitable for work environments where there is possible contact with bodily fluids, microorganisms and chemicals. They do not contain natural rubber latex and are an excellent alternative for people with Type I allergies.

These are non-powder gloves that offer great flexibility, high sensitivity and resistance to chemicals. They are tear-resistant, ambidextrous and non-sterile.

SIZE

Size	Minimal length (mm)	Palm width (mm)	Weight (g)	Minimum thickness (mm)
XS	240	≤80	3.8 ± 0.2	0,05
S	240	80 ± 10	4.3 ± 0.2	0,05
M	240	95 ± 10	4.8 ± 0.2	0,05
L	240	110 ± 10	5.3 ± 0.2	0,05
XL	240	≥110	5.8 ± 0.2	0,05

STANDARD

EN455

MAIN COMMODITIES

- Butadiene nitrile rubber ≥90%
- Zinc oxide 4%
- Titanium dioxide 2.6%

APPLICATION

Medical, food, chemical industries,...

REMARKS

- Examination gloves are single-use products and should only be used for one procedure and/or one patient
- When wearing gloves, do not wear rings or other ornaments and cut nails gently
- Wear them indifferently on the left or right hand, please choose gloves adapted to the specifications of your hand
- Do not use if gloves are contaminated due to defective packaging
- Do not directly irradiate strong light such as sunlight or ultraviolet rays.

CONDITIONING

100-piece cardboard.

STORAGE

Gloves should be stored in a cool, dry and dark environment without corrosive gases.

Masks FFP2 / KN95

OUTSIDE

INSIDE

BRAND

PURISM

BACTERIAL FILTRATION EFFICIENCY

≥ 95%

SPECIFICATIONS

5-layer mask :
- First layer: 60g/m² of unwoven fabric
- Second and third layer: 30g/m² e high-efficiency unwoven fabric, blown by fusion
- Fourth layer: 50g/m² hot air cotton
- Fifth layer: 25g/m2 of unwoven fabric

Composition :
Non-woven wire-bound, non-woven smelted, hot air cotton, nose clip, earrings

Product performance :
perfect fit for each face and optimal protection. Suitable for the protection of certain non-oil particles, such as flu virus, dust, pollen, etc.

Recommended port time :
4-8 hours, to be changed between medical procedures. After contact with the used mask, be sure to disinfect your hands.

APPLICATION

Schools, high-speed trains, airports, public transport and other relatively densely populated places and family environment.

COMPLIANCE

EC marking according to European Directive 89/686/EEC
Standard 149:2001 - A1:2009 / GB2626-2006 KN95 in connection with REGULATION CE R2016/425 (Personal Protective Equipment)

CONDITIONING

5 pcs/bag, 4 bags/box, 48 boxes/cardboard, 960 pieces/cardboard,
Cardboard size: 57.5 × 33.5 × 38 cm - 8.35 kg

STORAGE

Store in a cool, dry and clean place, away from the fire, between -20 and 38 degrees C with a relative humidity of less than 30%.

STORAGE TIME

3 years

Particulate respirator FFP3 Purism

BRAND

PURISM

BACTERIAL FILTERING EFFICIENCY

≥ 95% (tested by gb2626-2019)

SIZE

13.8x11x5x5.2cm

SPECIFICATIONS

5-layer mask :
- non-woven PP fabric
- PP fusion blown fabric
- non-air woven fabric
- aluminum nose clamp
- nylon/polyurethane composite headband

APPLICATION

Purism FFP3 particle respirators provide effective respiratory protection for use in industries where workers will be exposed to solid particles (dust) and/or non-volatile liquid particles. Ideal for applications where a higher protective factor is required, providing protection against most gases, vapors and particles.

COMPLIANCE

CE Marking 2016/425
Standard EN 149/2001-a1/2009

CONDITIONING

20 pces / box, 20 boxes / cardboard, 400 pieces / cardboard,
Cardboard size: 67 × 28 × 36 cm - 11.7 kg

STORAGE

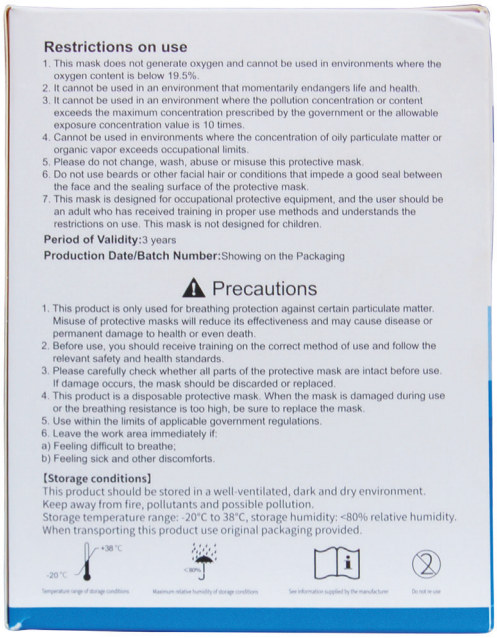
to be stored in a dry, clean, ventilated and non-corrosive gaseous area, away from ignition sources or combustible materials

STORAGE TIME

3 years


Purism
Particulate Respirator
Model: C001
CE 0370
FFP3 NR
EN 149:2001+A1:2009
Regulation(EU) 2016/425
20PCS
Restrictions on use
Precautions

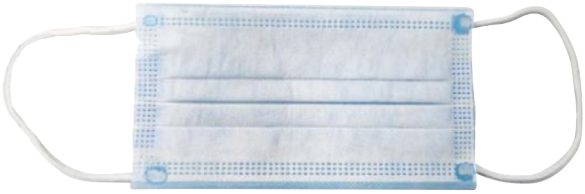

Wearing method
1. Hold the mask with your fingertips at the position of the nose clip to allow the headband to hang freely.
2. Put the mask around your chin with the nose clip facing upwards and wear the lower headband on the neck below the ears.
3. Put the upper headgear on the top of the head.
4. Use the index fingers of both hands to move from the middle of the nose clip to both sides and press down the nose clip until it is close to the bridge of the nose. One-handed operation is not recommended.
5. Before entering the work area, you must check the tightness of the mask and face, the inspection method is as follows:
a) Cover the mask with both hands, taking care not to affect the position of the mask on the face;
b) Take a deep breath. If you feel a leak near the bridge of the nose, adjust the nose clip again according to step 4;
c) If the leak is on the edge of the mask, the headgear should be readjusted.
Warning! If the mask and face cannot be in close contact, do not enter the work area. Please contact your supervisor.
Removal method
Referring to step 2 of the wearing method, hold the protective mask with one hand to maintain the wearing position, and use the other hand to pull down the headband and pull it over the top of the head. Continue to hold the mask with your hand, then pull the headband over the top of your head and remove the mask.
LOT
3years
生产日期 2020/08/22
00098521
Shandong Daddy's Choice Health Science and Technology Co.,Ltd
Address: Tiandingfeng road Garden street Economic Development Zone
Linyi County Dezhou City Shandong Province 251500 CN
Tel: 0534-8271567, 0534-8271568;
Website: www.daddychoice.com.cn
Email: sales@daddychoice.com
MADE IN CHINA
6 973601 650148


Restrictions on use
1. This mask does not generate oxygen and cannot be used in environments where the oxygen content is below 19.5%.
2. It cannot be used in an environment that momentarily endangers life and health.
3. It cannot be used in an environment where the pollution concentration or content exceeds the maximum concentration prescribed by the government or the allowable exposure concentration value is 10 times.
4. Cannot be used in environments where the concentration of oily particulate matter or organic vapor exceeds occupational limits.
5. Please do not change, wash, abuse or misuse this protective mask.
6. Do not use beards or other facial hair or conditions that impede a good seal between the face and the sealing surface of the protective mask.
7. This mask is designed for occupational protective equipment, and the user should be an adult who has received training in proper use methods and understands the restrictions on use. This mask is not designed for children.
Period of Validity:3 years
Production Date/Batch Number:Showing on the Packaging
Precautions
1. This product is only used for breathing protection against certain particulate matter. Misuse of protective masks will reduce its effectiveness and may cause disease or permanent damage to health or even death.
2. Before use, you should receive training on the correct method of use and follow the relevant safety and health standards.
3. Please carefully check whether all parts of the protective mask are intact before use. If damage occurs, the mask should be discarded or replaced.
4. This product is a disposable protective mask. When the mask is damaged during use or the breathing resistance is too high, be sure to replace the mask.
5. Use within the limits of applicable government regulations.
6. Leave the work area immediately if:
a) Feeling difficult to breathe;
b) Feeling sick and other discomforts.
【Storage conditions】
This product should be stored in a well-ventilated, dark and dry environment.
Keep away from fire, pollutants and possible pollution.
Storage temperature range: -20°C to 38°C, storage humidity: <80% relative humidity.
When transporting this product use original packaging provided.
-20°C
+38°C
<80%

CONTIPHARMA PROTECT
a member of Bedelco group

Disposable surgical mask Purism Type IIR

OUTSIDE



INSIDE

BRAND

PURISM

BACTERIAL FILTRATION EFFICIENCY

≥ 99%

SPECIFICATIONS

3-layer mask :

- First layer: waterproof non-woven fabric (new non-woven fabric): effectively inhibits and insulates harmful particles and gases
- Second layer: high-density filtering layer (blown non-woven fabric)
- Third layer: layer in direct contact with the skin, in new non-woven fabric, resistant to moisture, allowing easy, non-toxic, non-stimulating, soft and skin-friendly breathing

Composition :
unwoven-bound fabric, fusion-blown non-woven fabric, earrings and nasal bridge bar.

Recommended port time :
4-8 hours, to be changed between medical procedures. After contact with the used mask, be sure to disinfect your hands.

APPLICATION

for medical personnel during operations. It covers the user's mouth, nose and jaw, providing a physical barrier to prevent the direct passage of pathogens, microorganisms, bodily fluids and particles. Don't use in a traumatic environment.

COMPLIANCE

EC marking according to European Directive 89/686/EEC
Standard EN 14683 Type IIR

CONDITIONING

10 pcs/ bag, 5 bags / box, 48 boxes / cardboard, 2400 pieces / cardboard,
Cardboard size: 47.5 × 41.5 × 56 cm - 11.7 kg

STORAGE

to be stored in a dry, clean, ventilated and non-corrosive gaseous area, away from ignition sources or combustible materials

STORAGE TIME

3 years

Tech Total - Flex S3

Ref 70109

SPECIFICATIONS

Usage : Indoor/Outdoor
Type : High
Activity : Construction / Clean industry / Electronic industry, ESD
Upper : Waterproof ON MICRO TD® microfiber, Richelieu assembly without side seams
Lining : AIR SYSTEM® black, integral to the toes, «slipper» effect
Tongue : Waterproof gusset in ON STEAM® microfiber, AIR SYSTEM® lined and 10 mm foam padded
Malleolus guard : ON STEAM® microfiber with AIR SYSTEM® lining and 10 mm foam padding
Closure : Round polyester eyelets and laces in black
End cap : Ultralight and exra wide 200 joules Composite LIGHT SYSTEM®
Anti-perforation plate : FLEX SYSTEM® composite, non-magnetic and athermal
Insole : Thermoformed bi-density foam, activated carbon, antistatic and ESD, FRECH TECH® antibacterial
Semelle : PU 2D, TECH SOLE® ankle flexor, FO hydrocarbon resistant, antistatic, HI-CI
Cramping : Mixed industry indoor/outdoor, unhooked heel, cambered crampons, SRC standard
Weight (Kg/pair) : 1.200
Colors : Black upper and dark grey/black sole
Sizes : 35-49
Standards : EN ISO 20345:2011 SRC / EN 61340-4-3 / DGUV 112-191
EC Certificate of Conformity n°: 0075/020/161/08/17/1266
Specificities : Acid and welding splash resistant microfiber (UNE EN ISO 20349/2011)
Specificity 2 : Retro-reflective piping on the heel
Specificity 3 : Large choice of sizes : from 35 to 49 !

SPECIFIC FEATURES

- Lightweight composite tip 200 joules
- Resistance to water absorption and penetration of the stem (WRU)
- Electrical discharge Standards EN 61340-5-1

CORDURA

on steam

Doublure ARTICO

GENERAL FEATURES

- Antistatic properties (A)
- Energy absorption at heel (E)
- Abrasion resistance
- Cleated sole
- Sole insulation against heat (HI)
- Sole insulation against cold (CI)
- Extra large toe cap 200 joules
- Hyper breathable lining
- Puncture resistance 1100 N (P)

Tech Power - Flex S3

Ref 70804

SPECIFICATIONS

Usage : Indoor/Outdoor
Type : Low
Activité : Construction / Warehousing, logistics, transport / Electronic industry, ESD
Tige : Waterproof greased nubuck leather and CORDURA® canvas
Doublure : ARTICO® black color, integral to the toes, «slipper» effect
Languette : Waterproof gusset in ON STEAM® microfiber, ARTICO® lined and 10 mm foam padded
Protège malléoles :
Fermeture : Perforations and round laces in polyester with «Quick lock» system
Embout : Ultralight and extra wide 200 joules Composite LIGHT SYSTEM®.
Plaque antiperforation : FLEX SYSTEM® composite, non-magnetic and athermal
Insole : Thermoformed dual density foam, activated carbon, antistatic and ESD, FRESH TECH® antibacterial
Outsole : PU 2D, TECH SOLE® ankle flexor, FO hydrocarbon resistant, antistatic, HI-CI
Cramping : Mixed industry indoor/outdoor, unhooked heel, cambered crampons, SRC standard
Weight (Kg/pair) : 1.200
Colors : Black upper and dark grey/black sole
Sizes : 35-48
Standards : EN ISO 20345:2011 SRC / EN 61340-5-1 / DGUV 112-191
CE certificate of conformity n° : 0075/020/161/08/17/1266 - EXT 09/09/17
Specifics : Quick lock system
Specifics 2 : Reflective piping on the heel

SPECIFIC FEATURES

- Lightweight composite tip 200 joules
- No metal parts, non-magnetic
- Resistance to water absorption and penetration of the stem (WRU)

CORDURA

on steam

Doublure ARTICO

GENERAL FEATURES

- Antistatic properties (A)
- Energy absorption at heel (E)
- Abrasion resistance
- Cleated sole
- Sole insulation against heat (HI)
- Sole insulation against cold (CI)
- Extra large toe cap 200 joules
- Hyper breathable lining
- Puncture resistance 1100 N (P)

Tech Runner - Flex S3

Ref 70805

SPECIFICATIONS

Suitable for :	Inside/Outside
Type :	Ankle shoe
Activity :	Construction / Storage, Logistic, Transport / Electronic Industry, ESD
Upper :	Water resistant full supple leather and CORDURA® material
Lining :	ARTICO® black color, up to the toes, slipper feeling
Tongue :	Bellows tongue in ON STEAM® microfibre, lining ARTICO® and padded with 10mm thick foam
Ankle protection :	CORDURA® material, ARTICO® lining and padded with 10mm thick foam
Fastening :	Round polyester black laces with «Quick lock» system
Toe cap :	Ultralight and extra wide 200 joules, Composite LIGHT SYSTEM®
Midsole :	Composite FLEX SYSTEM®, anti-magnetic and thermal insulated
Insole :	Thermoformed in active carbon dual density foam, antistatic, ESD and antibacterial FRESH TECH®
Sole :	PU 2D, TECH SOLE® ankle anti-twist system, oil resistant FO, antistatic, HI-CI
Sole pattern :	Multi-purpose cleated indoor/outdoor outsole, well-defined heel, cleated arch, SRC standard
Weight (Kg/pair) :	1.200
Colors :	Black upper and dark grey/light grey outsole
Sizes :	35-48 (2-13)
Standards :	EN ISO 20345:2011 SRC / EN 61340-5-1 / DGUV 112-191
EC Certificate of Conformity N° :	0075/020/161/08/17/1266 - EXT 08/09/17
Special features :	Quick fastening system «Quick lock»
Special features 2 :	Reflective piping on the heel part

SPECIFIC FEATURES

- Integral 100% composite (Air System + Light System + Flex System)
- Metal free, non magnetic
- Water absorption and penetration resistance of upper (WRU)
- Electro Static Discharge Standard EN 61340-5-1

CORDURA
on steam
Doublure ARTICO

GENERAL FEATURES

- Antistatic properties (A)
- Heel energy absorption (E)
- Abrasion resistance
- Crampon sole
- Heat-insulated sole (HI)
- Cold-insulated sole (CI)
- 200 joule extra-wide toecap
- Hyper-ventilated lining
- Perforation-resistance 1100 N (P)

NORA MULTIRALF WHITE

SPECIFICATIONS

Sole : midsole in expanded material: cold insulation and shock absorber
End cap : steel
Sizes : 36-49/50

① Steel tips
② Midsole in expanded material
③ Anti-puncture sole Optimized
④ sole
⑤ Mondpoint® Quality
⑥ Gommaforte® measuring system

CONTIPHARMA PROTECT
a member of Bedelco group

MTS NORA MULTIRAF WHITE

SPECIFICATIONS

· Foamed material midsole : cold insulation and shock absorber
· Size : 36-49/50
· Steel end cap
· Spacer made of expanded material
· Anti-puncture sole
· Optimized sole
· Mondopoint ® measuring system
· Gommaforte® Quality

NORMS

· EN345 S4

NORAMAX FOOD - WHITE

SPECIFICATIONS

Upper : polyurethane - easy cleaning
Lining : anti-bacterial lining
Insole : Ortholite® cushioned comfort foam insole
Sole : easy to clean outsole, anti-dirt - slip resistance: SRC
Anti-perforation plate : steel
Tip : metal
Sizes : 36-49
Height : 33 cm (reference: size 42)

NORMS

· EN ISO 20345:2011 S4 CI SRC

SAFRON - WHITE

SPECIFICATIONS

· EN345 Gommaforte®
· Mélange de caoutchouc et de matières thermoplastiques
· Haute résistance à l'usure
· Embout acier 200 joules
· Pointures : 36-47

NORMS

· EN ISO 20345 :2011 – S4

CONTIPHARMA PROTECT
a member of Bedelco group

Tech Constructor - S3

Ref 70114

SPECIFICATIONS

Usage :	Outdoor
Type :	High
Activity :	Construction / Electronic industry, ESD
Upper :	Water-repellent greased nubuck grain leather and CORDURA® canvas
Lining :	ARTICO® black, integral to the toes, «slipper» effect
Tongue :	Waterproof gusset in ON STEAM® microfiber, ARTICO® lined and 10 mm foam padded
Malleolar protection :	CORDURA® fabric, ARTICO® lining and 10 mm foam padding
Closure :	Steel laces and round polyester laces in black
End cap :	Ultra-light and extra wide 200 joules Composite LIGHT SYSTEM®.
Anti-puncture plate :	Stainless steel
Insole :	Thermoformed dual density foam, activated carbon, antistatic and ESD, FRESH TECH® antibacterial
Outsole :	PU 2D, TECH SOLE® ankle flexor, FO hydrocarbon resistant, antistatic, HI-CI
Cramping :	Mixed industry indoor/outdoor, unhooked heel, cambered crampons, SRC standard
Weight (Kg/pair) :	1.200
Colors :	Black upper and dark grey/black sole
Sizes :	35-48
Standards :	EN ISO 20345:2011 SRC / EN 61340-5-1 / DGUV 112-191
EC Certificate of Conformity No :	0075/020/161/08/17/1266 - EXT 14/09/17
Special features :	End cap reinforcement made of specific abrasion-resistant material (secure, waterproof stitching)
Specificity 2 :	Model specially designed for Construction and Public Works

SPECIFIC FEATURES

- Lightweight composite tip 200 joules
- Resistance to water absorption and penetration of the rod (WRU)
- Static discharge Standards EN 61340-5-1

CORDURA

on steam

Doublure ARTICO

GENERAL FEATURES

- Antistatic properties (A)
- Heel energy absorption (E)
- Abrasion resistance
- Crampon sole
- Heat-insulated sole (HI)
- Cold-insulated sole (CI)
- 200 joule extra-wide toecap
- Hyper-ventilated lining
- Perforation-resistance 1100 N (P)

Luna+ - S2

Ref 15209

SPECIFICATIONS

Suitable for :	Indoor
Type :	Low shoe
Activity :	Food industry, Catering, Medical / Chemical Industry
Upper :	Dry-Tek® material, thickness 2.2mm, acid resistant, washable and breathable
Lining :	Cambrelle Amicor+® lining with "Refreshing" microcells perfume system
Tongue :	Dry-Tek® material
Ankle protection :	
Fastening :	Pull on style with elastic strap
Toe cap :	Composite 200 joules Light System®
Midsole :	
Insole :	Antibacterial and antimicrobial foam
Sole :	PU 2D, oil resistant FO, antistatic and HI-CI insulate
Sole pattern :	Special non-slip crampons for smooth and greasy floors, SRC standard
Weight (Kg/pair) :	0.800
Colors :	White upper and white/light grey outsole
Sizes :	35-49 (2-14)
Standard :	EN ISO 20345:2011 SRC
EC Certificate of Conformity N° :	0075/020/161/04/13/0269
Special features :	Upper without side stitching: no water infiltration risks
Specificity 2 :	Dry-Tek® material resistant to 12 acids and corrosive fluids (standard EN 13832-2)
Specificity 3 :	Washable up to 60°C

SPECIFIC FEATURES

- 200 joules light composite toecap
- Metal free, non magnetic
- Water absorption and penetration resistance of upper (WRU)

Cambrelle+ with Amicor Plus

GENERAL FEATURES

- Antistatic properties (A)
- Heel energy absorption (E)
- Abrasion resistance
- Crampon sole
- Heat-insulated sole (HI)
- Cold-insulated sole (CI)
- 200 joule extra-wide toecap
- Hyper-ventilated lining
- Perforation-resistance 1100 N (P)

CONTIPHARMA PROTECT
a member of Bedelco group

MOLIÈRE WHITE STRINGS

SPECIFICATIONS

Upper : coated leather
Lining : Cambrelle®
Tip : Latex foam
Insole : PU 2D
Sizes : 35-48

NORMS

· EN ISO 20345 :2011 – S2

PYTHON S3

Ref 11125

SPECIFICATIONS

· Classical ankle boot
· Upper in printed black leather
· Ankle padded collar in synthetic material on 10 mm foam
· Waterproof padded and lined bellows tongue in synthetic PU material on 10mm foam
· Lined in "Air System"® on foam, grey colour
· Reflective piping on the outside counter
· Fastening by laces through triangles
· Polyamide/polyester black/beige laces
· Full insole in latex foam, antistatic and antibacterial
· Double density polyurethane antistatic sole
· Multi purpose cleated outsole
· Anti-slip factor: 0.27
· Shock absorption in the heel
· 200 joules resistance steel toe cap
· Steel anti-puncture midsole
· Approx. weight per pair: 1,2 kg (pair)
· Colours: upper black with black sole
· Sizes: from 35 to 48

APPLICATIONS

· Multi purpose

NORMS

· Conform to the EN ISO 20345:2011- SRC - Safety footwear
· Tested by the C.T.C. (Centre Technique du Cuir)
· P.P.E category 2 (intermediate risks)
· CE certificate of conformity N° : 0075/020/161/05/13/0428 – EXT 05/05/13

RAVEN S2 SRC MOCCASIN

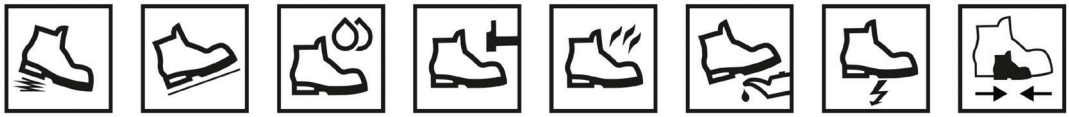
SPECIFICATIONS

Safety moccasin with insert. Antistatic PU sole with anti-slip density resistant to oils and mechanical wear, energy absorption under the heel. Resistant microfiber upper, permeable lining and antibacterial features.

Anti-perference :

Upper : Micro fiber
End cap : Metal
Sole : Single density polyurethane
Lining : Polyester mesh
Size : 35-36-37-38-39-40-41-42-43-44-45-46-47-48
Shade : 80 - white

FEATURES



STANDARDS

EN ISO 20345:2011 / S2 SRC

ED COMFORT PLUG EAR PLUG

Ref 60001

SPECIFICATIONS

Disposable earplugs made of very soft PU foam, ergonomic design for an easier insertion in the ear canal and a perfect adhesion to its walls for a great comfort of use. EN 352-2.

FEATURES

- Earplugs Attachment System.
- Passive Disposable Protection.
- System.

STANDARDS

EN 352-2:2002.

Protect yourself
Protect lives

CONTIPHARMA PROTECT
a member of Bedelco group srl

Rue de l'Environnement 22
4100 Seraing
Belgium

Mobile : +32 4 330 18 60
Email : info@bedelco.com
Web : www.contipharma-protect.com