

COUNTY OF ROCKLAND
Department of General Services
Purchasing Division

Contract Award Notification

Title: **Personal and Protective Equipment (PPE)**

Contract Period: November 16, 2020 through May 15, 2021 with 4 six-month options
Ext thru 11/15/21 w/ 3 six-month, Ext thru 5/15/22 w/ 2 six-month
options; **Ext thru 11/15/22 w/ six-month option**

Original Date of Issue: November 16, 2020

Date of Revision: **02/24/2022**

BID No: **RFB-RC-2020-089**

Catalog: **Health and Hospitals**

Authorized Users: County Agencies, All Political Subdivisions

Address Inquiries To:

Name: Michele Phillips
Title: Purchaser I
Phone: 845-364-2984
Fax: 845-364-3809
E-mail: phillipm@co.rockland.ny.us

Description

This contract is to provide personal and protective equipment supplies.

Contract #	Vendor Number	Contractor & Address	Telephone No.
Bid 20-089-A	0000026650	S2C Focus Inc. 527 Mallard Point Drive North Aurora, IL 60542 Contact: Rishi Raj rraj@s2cfocus.com	630-701-3625 FAX: 630-206-0391
Bid 20-089-B	0000026654	Green Dream International 32 W. 8 th Street Suite 607 Erie, PA 16501 Contact: Varand Vartanian info@gdicompany.com	814-616-7800 FAX: 202-204-8444

Contract #	Vendor Number	Contractor & Address	Telephone No.
Bid 20-089-C NOT RENEWED	0000026657	Nationwide Instruction for Cardiovascular Education 1 Gate Court Dix Hills, NY 11746 Contact: Dr. Christopher S. Byron chris.byron@niceheart.com	516-369-4958 FAX: 631-643-2780
Bid 20-089-D NOT RENEWED	0000026655	PPE Trade and Support 12855 Runway Road # 1301 Los Angeles, CA 90094 Contact: Janet Lee contact@ppetradesupport.com	404-759-8409
Bid 20-089-E NOT RENEWED	0000026656	Golden Circle Group 1125 Crenshaw Boulevard 2 nd Floor Los Angeles, CA 90094 Contact: Jim Leatherman jim@goldencircle.group	562-810-3367
Bid 20-089-F	0000026660	US Health Express 18701 Arenth Avenue City of Industry, CA 91748 Contact: Jianping Liu Ph.D. jianping.liu@jointown-intl.com	773-960-1822 FAX: 626-810-2005
Bid 20-089-G Rescinded	0000026662	Initium Health 1401 Wewatta Street Suite 103 Denver, CO 80202 Contact: James Corbett james@initiumhealth.org	303-928-8511
Bid 20-089-H NOT RENEWED	0000007926	Kenton Healthcare Box 120 Springfield, TN 37172 Contact: Nari Sadarangani kentron@kentronmedical.com	615-668-1147

PERSONAL PROTECTIVE EQUIPMENT (PPE)
Revised November 23, 2021

COUNTY OF ROCKLAND
DGS – PURCHASING DEPARTMENT
BLDG. A, 6TH FLOOR, 50 SANATORIUM ROAD
POMONA, NY 10970
TELEPHONE NO.: 845-364-3820
FAX NO.: 845-364-3809

BONFIRE ITEM #	LINE NO.	DESCRIPTION	ITEM NUMBER	EST. QTY.	UNIT	UNIT PRICE	MFG.& PRODUCT CODE	PACKING INFORMATION	SUPPLIER
BONFIRE ITEM #		SECTION I - N95 AND KN95 MASKS - As needed quantities - Submit unit pricing for the minimum order quantities listed to furnish and deliver the specified masks in accordance with the specifications					MANDATORY FIELDS	MANDATORY FIELDS	
#1-1	1	Mask, N95 -Aura Industrial Particulate Respirator, MFG. 3M MODEL 9205+ or approved equal, Must be NIOSH approved PACKAGING: 440/Case MINIMUM ORDER QTY. 1,000-4,999	47562	1000	Price Per Mask	Pending	Pending	Pending	Pending
#1-2	2	Mask, N95 -Aura Industrial Particulate Respirator, MFG. 3M MODEL 9205+ or approved equal, Must be NIOSH approved PACKAGING: 440/Case MINIMUM ORDER QTY. 5,000-9,999	47562	5000	Price Per Mask	Pending	Pending	Pending	Pending
#1-3	3	Mask, N95 -Aura Industrial Particulate Respirator, MFG. 3M MODEL 9205+ or approved equal, Must be NIOSH approved PACKAGING: 440/Case MINIMUM ORDER QTY. 10,000 - 24,999	47562	10000	Price Per Mask	Pending	Pending	Pending	Pending
#1-4	4	Mask, N95 -Aura Industrial Particulate Respirator, MFG. 3M MODEL 9205+ or approved equal, Must be NIOSH approved PACKAGING: 440/Case MINIMUM ORDER QTY. 25,000 - 49,999	47562	25000	Price Per Mask	Pending	Pending	Pending	Pending
#1-5	5	Mask, N95 -Aura Industrial Particulate Respirator, MFG. 3M MODEL 9205+ or approved equal, Must be NIOSH approved PACKAGING: 440/Case MINIMUM ORDER QTY. 50,000 - 99,999	47562	50000	Price Per Mask	Pending	Pending	Pending	Pending

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#1-6	6	Mask, N95 -Aura Industrial Particulate Respirator, MFG. 3M MODEL 9205+ or approved equal, Must be NIOSH approved PACKAGING: 440/Case MINIMUM ORDER QTY. 100,000 Plus	47562	100000	Price Per Mask	Pending	Pending	Pending	Pending
#1-7	7	Mask, N95 -Particulate Respirator, MFG. 3M MODEL 8210 or approved equal, Must be NIOSH approved for Healthcare Professionals PACKAGING:: 20/Bx, 8 BX/CA MINIMUM ORDER QTY: 1,000 - 4,999	47562	1000	Price Per Mask	Pending	Pending	Pending	Pending
#1-8	8	Mask, N95 -Particulate Respirator, MFG. 3M MODEL 8210 or approved equal, Must be NIOSH approved for Healthcare Professionals PACKAGING:: 20/Bx, 8 BX/CA MINIMUM ORDER QTY: 5,000 - 9,999	47562	5000	Price Per Mask	Pending	Pending	Pending	Pending
#1-9	9	Mask, N95 -Particulate Respirator, MFG. 3M MODEL 8210 or approved equal, Must be NIOSH approved for Healthcare Professionals PACKAGING:: 20/Bx, 8 BX/CA MINIMUM ORDER QTY: 10,000 - 24,999	47562	10000	Price Per Mask	Pending	Pending	Pending	Pending
#1-10	10	Mask, N95 -Particulate Respirator, MFG. 3M MODEL 8210 or approved equal, Must be NIOSH approved for Healthcare Professionals PACKAGING:: 20/Bx, 8 BX/CA MINIMUM ORDER QTY: 25,000 - 49,999	47562	25000	Price Per Mask	Pending	Pending	Pending	Pending
#1-11	11	Mask, N95 -Particulate Respirator, MFG. 3M MODEL 8210 or approved equal, Must be NIOSH approved for Healthcare Professionals PACKAGING:: 20/Bx, 8 BX/CA MINIMUM ORDER QTY: 50,000 - 99,999	47562	50000	Price Per Mask	Pending	Pending	Pending	Pending

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#1-12	12	Mask, N95 -Particulate Respirator, MFG. 3M MODEL 8210 or approved equal, Must be NIOSH approved for Healthcare Professionals PACKAGING:: 20/Bx, 8 BX/CA MINIMUM ORDER QTY: 100,000 Plus	47562	100000	Price Per Mask	Pending	Pending	Pending	Pending
#1-13	13	Mask, N95 -Aura Health Care Particulate Respirator and Surgical Mask, MFG. 3M MODEL 1870+ or approved equal, Must be NIOSH approved for health care use PACKAGING: 20/Bx, 6 BX/CA MINIMUM ORDER QTY: 1,000-4,999	47562	1000	Price Per Mask	Pending	Pending	Pending	Pending
#1-14	14	Mask, N95 -Aura Health Care Particulate Respirator and Surgical Mask, MFG. 3M MODEL 1870+ or approved equal, Must be NIOSH approved for health care use PACKAGING: 20/Bx, 6 BX/CA MINIMUM ORDER QTY: 5,000-9,999	47562	5000	Price Per Mask	Pending	Pending	Pending	Pending
#1-15	15	Mask, N95 -Aura Health Care Particulate Respirator and Surgical Mask, MFG. 3M MODEL 1870+ or approved equal, Must be NIOSH approved for health care use PACKAGING: 20/Bx, 6 BX/CA MINIMUM ORDER QTY: 10,000 - 24,999	47562	10000	Price Per Mask	Pending	Pending	Pending	Pending
#1-16	16	Mask, N95 -Aura Health Care Particulate Respirator and Surgical Mask, MFG. 3M MODEL 1870+ or approved equal, Must be NIOSH approved for health care use PACKAGING: 20/Bx, 6 BX/CA MINIMUM ORDER QTY: 25,000-49,999	47562	25000	Price Per Mask	Pending	Pending	Pending	Pending
#1-17	17	Mask, N95 -Aura Health Care Particulate Respirator and Surgical Mask, MFG. 3M MODEL 1870+ or approved equal, Must be NIOSH approved for health care use PACKAGING: 20/Bx, 6 BX/CA MINIMUM ORDER QTY: 50,000 - 99,999	47562	50000	Price Per Mask	Pending	Pending	Pending	Pending

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#1-18	18	Mask, N95 -Aura Health Care Particulate Respirator and Surgical Mask, MFG. 3M MODEL 1870+ or approved equal, Must be NIOSH approved for health care use PACKAGING: 20/Bx, 6 BX/CA MINIMUM ORDER QTY: 100,000 Plus	47562	100000	Price Per Mask	Pending	Pending	Pending	Pending
#1-19	19	Mask, KN95-Particulate Respirator, Non-Woven polypropylene, cone shape, latex free. Mfg Jinhu Jiadaifu Medical Supplies Co, Ltd (KN95) unit of measure per box of 20 @ \$0.39=7.80 EACH BOX Min order: 1,000 - 4,999	47562000034	1000	Price Per Box				NOT RENEWED
#1-20	20	Mask, KN95-Particulate Respirator, Non-Woven polypropylene, cone shape, latex free. Mfg Jinhu Jiadaifu Medical Supplies Co, Ltd (KN95) unit of measure per box of 20 @ \$0.39=7.80 EACH BOX Min order: 1,000 - 4,999	47562000035	5000	Price Per Box				NOT RENEWED
#1-21	21	Mask, KN95-Particulate Respirator, Non-Woven polypropylene, cone shape, latex free. Mfg Jinhu Jiadaifu Medical Supplies Co, Ltd (KN95) unit of measure per box of 20 @ \$0.39=7.80 EACH BOX Min order: 1,000 - 4,999	47562000036	10000	Price Per Box				NOT RENEWED
#1-22	22	Mask, KN95-Particulate Respirator, Non-Woven polypropylene, cone shape, latex free. Mfg Jinhu Jiadaifu Medical Supplies Co, Ltd (KN95) unit of measure per box of 20 @ \$0.39=7.80 EACH BOX Min order: 1,000 - 4,999.	47562000037	25000	Price Per Box				NOT RENEWED
#1-23	23	Mask, KN95-Particulate Respirator, Non-Woven polypropylene, cone shape, latex free. Mfg Jinhu Jiadaifu Medical Supplies Co, Ltd (KN95) unit of measure per box of 20 @ \$0.39=7.80 EACH BOX Min order: 1,000 - 4,999	47562000038	50000	Price Per Box				NOT RENEWED

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#1-24	24	Mask, KN95-Particulate Respirator, non-woven polypropylene, cone shape, latex free, MFG. Guangzhou Powecom Labor GB2626-2006 or or approved equal MUST BE ON FDA EUA LIST AS SPECIFIED PACKAGING: 10/Bag 100 BAGS //CA MINIMUM ORDER QTY: 100,000 Plus	47562000039	100000	Price Per Box	NOT RENEWED			
		SECTION II- VARIOUS PPE SUPPLIES. Quantities are estimated, submit unit pricing for each item to furnish and deliver the specified supplies in accordance with the specifications.							
#2-1	25	Gown- Disposable, 100% polypropylene, fluid resistant, tie back, elastic cuffs, full length, color: yellow, AAMI PB70 Level 1 protection or approved equal Packaging: 5/Pk, 20PK/CA	47562000040	1000	Price Per Bag	\$ 9.00	Wuhan Orient Honest International Trade Co., Ltd. - GWN1-103 RAGLAN SLEEVE - SIZES XS to 3X - NEED TO INDICATE SIZE TO BE ORDERED	10 pcs/bag - 100 pcs/carton	Green Dream International LLC
#2-2	26	Gown- Disposable, 100% polypropylene, fluid resistant, tie back, elastic cuffs, full length, color: yellow, AAMI PB70 Level 2 protection or approved equal PACKAGING: 5/Pk, 20PK/CA	47562000041	1000	Price Per Bag	\$ 10.00	Wuhan Orient Honest International Trade Co., Ltd. - GWN2-214 RAGLAN SLEEVE SIZES XS to 3X - NEED TO INDICATE SIZE TO BE ORDERED	10 pcs/bag - 100 pcs/carton	Green Dream International LLC

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#2-3	27	Gown- Disposable, 50% polypropylene/50% polyethylene, fluid resistant, tie back, elastic cuffs, full length, color: blue, AAMI PB70 Level 3 protection or approved equal PACKAGING: 5/Pk, 20PK/CA	47562000042	1000	Price Per Bag	\$ 11.50	Wuhan Orient Honest International Trade Co., Ltd. - GWN3-305 RAGLAN SLEEVE SIZES XS to 3X - NEED TO INDICATE SIZE TO BE ORDERED	10 pcs/bag - 100 pcs/carton	Green Dream International LLC
#2-4	28	Gown- Disposable, blue, polyethylene film, lightweight, single layer, low-density, impervious, for clinical and laboratory settings, Full-length wraparound, tear-away collar, thumb loop, ASTM F1670, ASTM1671, PolyCo 10400, individually wrapped with dispenser, or approved equal PACKAGING: 20/Bx, 5BX/CA Size: Regular	4756200003	100	Price Per Box	Pending	Pending	Pending	Pending
#2-5	29	Gown- Disposable, blue, polyethylene film, lightweight, single layer, low-density, impervious, for clinical and laboratory settings, Full-length wraparound, tear-away collar, thumb loop, ASTM F1670, ASTM1671, PolyCo 10500, individually wrapped with dispenser or approved equal PACKAGING: 15/Bx, 5BX/CA Size: Large	47562	100	Price Per Box	Pending	Pending	Pending	Pending

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BONFIRE ITEM #	LINE NO.	DESCRIPTION	ITEM NUMBER	EST. QTY.	UNIT	UNIT PRICE	MFG.& PRODUCT CODE	PACKING INFORMATION	SUPPLIER
#2-6	30	Gown- Disposable, blue, polyethylene film, lightweight, single layer, low-density, impervious, for clinical and laboratory settings, Full-length wraparound, tear-away collar, thumb loop, ASTM F1670, ASTM1671, PolyCo 10600, individually wrapped with dispenser or approved equal PACKAGING: 15/Bx, 5BX/CA Size: X-Large	47562	100	Price Per Box	Pending	Pending	Pending	Pending
#2-7	31	Gown- Disposable, blue, Latex-Free polyethylene film, heavyweight gown, fluid resistant, Over head style, open back, apron tie, thumb loop, ASTM F1670, Medline CRI5020 and NONTH180 or approved equal PACKAGING: 15/Bx, 5BX/CA Size: Universal	47562000043	100	Price Per Box				NOT RENEWED
#2-8	32	Gown- Disposable isolation gown, yellow, sms, fluid resistant, non sterile, elastic cuff, tie back, long sleeve, knee length, Procure PC150 or approved equal PACKAGING: 10 Pk, 5 PK/CA SIZE: Regular	47562		Price Per Box	Pending	Pending	Pending	Pending
#2-9	33	Gown- Disposable isolation gown, yellow, sms, fluid resistant, non sterile, elastic cuff, tie back, long sleeve, knee length, Procure PC150 or approved equal PACKAGING: 10 Pk, 5 PK/CA Size: X-Large	47562		Price Per Box	Pending	Pending	Pending	Pending
#2-10	34	Gloves- Nitrile examination gloves, non-sterile, 4 mil. Thickness, 9.5" length, Latex-FREE, Powder-free, ambidextrous, Better Touch 807.2 or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: Small	47541000032	50	Price per Case	N/A	N/A	N/A	Rescinded Award
#2-11	35	Gloves- Nitrile examination gloves, non-sterile, 4 mil. Thickness, 9.5" length, Latex-FREE, Powder-free, ambidextrous, Better Touch 807.3 or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: Medium	47541000033	50	Price per Case	N/A	N/A	N/A	Rescinded Award

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#2-12	36	Gloves- Nitrile examination gloves, non-sterile, 4 mil. Thickness, 9.5" length, Latex-FREE, Powder-free, ambidextrous, Better Touch 807.4 or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: Large	47541000034	1000	Price per Case	N/A	N/A	N/A	Rescinded Award
#2-13	37	Gloves- Nitrile examination gloves, non-sterile, 4 mil. Thickness, 9.5" length, Latex-FREE, Powder-free, ambidextrous, Better Touch 807.5 or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: X-Large	47541000035	1000	Price per Case	N/A	N/A	N/A	Rescinded Award
#2-14	38	Gloves- Nitrile examination gloves, non-sterile, 4 mil. Thickness, 9.5" length, Latex-FREE, Powder-free, ambidextrous, Better Touch 807.6, or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: XX-Large	47562000033	10	Price Per Box	Pending	Pending	Pending	Pending
#2-15	39	Gloves- Black Nitrile, medical grade examination gloves, Latex-FREE, Powder-free, ambidextrous, 4mil palm, 5mil finger, Better Touch 792.2 or approved equal PACKAGING: 100/box 10bx/ca Size: Small	47541	50	Price Per Box	Pending	Pending	Pending	Pending
#2-16	40	Gloves- Black Nitrile, medical grade examination gloves, Latex-FREE, Powder-free, ambidextrous, 4mil palm, 5mil finger, Better Touch 792.3 or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: Medium	47541	50	Price Per Box	Pending	Pending	Pending	Pending
#2-17	41	Gloves- Black Nitrile, medical grade examination glove, Latex-FREE, Powder-free, ambidextrous, 4mil palm, 5mil finger, Better Touch 792.4 or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: Large	47541	1000	Price Per Box	Pending	Pending	Pending	Pending

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BONFIRE ITEM #	LINE NO.	DESCRIPTION	ITEM NUMBER	EST. QTY.	UNIT	UNIT PRICE	MFG.& PRODUCT CODE	PACKING INFORMATION	SUPPLIER
#2-18	42	Gloves- Black Nitrile, medical grade examination gloves, Latex-FREE, Powder-free, ambidextrous, 4mil palm, 5mil finger, Better Touch 792.5 or approved equal PACKAGING: 100/Bx, 10 BX/CA Size: X-Large	47541	1000	Price Per Box	Pending	Pending	Pending	Pending
#2-19	43	Gloves- EMS Latex examination gloves-15 Mil extra thick, powder-free, non-sterile, ambidextrous, textured fingers, single use only, Better Touch 811.2, or approved equal PACKAGING: 50/Bx, 10 BX/CA Size Small	47541	10	Price Per Box	Pending	Pending	Pending	Pending
#2-20	44	Gloves- EMS Latex examination gloves-15 Mil extra thick, powder-free, non-sterile, ambidextrous, textured fingers, single use only, Better Touch 811.3 or approved equal PACKAGING: 50/Bx, 10 BX/CA Size: Medium	47541	20	Price Per Box	Pending	Pending	Pending	Pending
#2-21	45	Gloves- EMS Latex examination gloves-15 Mil extra thick, powder-free, non-sterile, ambidextrous, textured fingers, single use only, Better Touch 811.4 or approved equal Packaging: 50/Bx, 10 BX/CA Size: Large	47541	20	Price Per Box	Pending	Pending	Pending	Pending
#2-22	46	Gloves- EMS Latex examination gloves-15 Mil extra thick, powder-free, non-sterile, ambidextrous, textured fingers, single use only, Better Touch 811.5 or approved equal PACKAGING: 50/Bx, 10 BX/CA Size: X-Large	47541	20	Price Per Box	Pending	Pending	Pending	Pending

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#2-23	47	Face Shield-disposable, full face coverage, anti-fog, elastic band, 1.3"-1.5" thick foam, latex-free, polymer film, non-sterile, pre-assembled, Nikomed, USA Inc N1288-FSDISP (12.6"X8.6") or approved equal PACKAGING: 10/Pk, 10 PK/CA	47555000012	500	Price Per Cs	\$ 115.20	6/BLIDER-PACK-240 / CASE	CATCH MY STYLE 10165	S2C Focus Inc.
#2-24	48	Surface Disinfecting wipes- registered with EPA as effective against Covid-19, Bleach free, PDI Professional Disposables- Germicidal Super Sani-cloth large (6X6.75 in) - Q55172-160/Cn, 12CN/CA, Lysol Disinfecting wipes code 1920090650-75/Cn or approved equal	34594	1000	Price Per Container	Pending	Pending	Pending	Pending
#2-25	49	Sanitizing hand wipes (Individual Packets)- Purell 62% Ethyl alcohol, Gojo Industries- 9021-1M or PDI Professional Disposables- Antimicrobial Sani-Hands, 65.9% Ethyl alcohol, D43600 or approved equal PACKAGING: 100/Box, 10 BX/CA	34594	500	Price Per Box	Pending	Pending	Pending	Pending
#2-26	50	Sanitizing hand wipes- P.A.W.S Antimicrobial hand wipes , 65.9% Ethyl alcohol-160/CN or PDI Professional Disposables- Sani-Hands 70% Ethyl alcohol P13472-135/CN or approved equal PACKAGING: 12 CN/CA	34594	500	Price Per Container	Pending	Pending	Pending	Pending
#2-27	51	Hand Sanitizer Gel, 2 oz bottle, 70% Ethyl alcohol, Purell Advantage 9605-24 or approved equal PACKAGING: 2 oz/Bt, 24 BT/CA	43573	1000	Price Per Bottle	Pending	Pending	Pending	Pending
#2-28	52	Hand Sanitizer Gel, 8 oz bottle with pump, 70% Ethyl alcohol, Purell Advantage 9652-12 or approved equal PACKAGING: 8 oz/Bt, 12 BT/CA	43573000001	1000	Price Per Cs	NOT RENEWED			NOT RENEWED

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#2-29	53	Non-Contact Infrared Thermometer- Clinical use, automatic power-off, measure in Fahrenheit, provides body temperature reading in one second from 1.2"-2" from forehead, LCD digital display screen, high temperature alarm, FDA approved, Berrcom JXB-178 or approved equal	25726000001	30	EACH	\$ 8.50	Guangdong Health & Health Medical Equipment	1 Piece/Pack	US Health Express Corp

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 BLDG. A., 6TH FLOOR, 50 SANATORIUM RD, POMONA, NY 10970
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**TITLE: PERSONAL PROTECTIVE EQUIPMENT
 (PPE)**

RFB-RC-2020-089

PURCHASES BY OTHER

LOCAL GOVERNMENTS, SCHOOL DISTRICTS, AND NON PROFIT AGENCIES

As per the New York State General Municipal Law, all political subdivisions of New York State are allowed to make purchases through the resulting contract(s). As per Rockland County Procurement Policy, Non Profit Agencies approved to participate in New York State's Contract Extension Program are authorized to make purchases through the resulting contract(s).

1. The County of Rockland shall make all contract award information available to other political subdivisions and non profit agencies through our website: www.rcpurchasing.com
2. Any other political subdivision or Rockland County non profit agency will issue purchase orders directly to vendors within the specified contract period referencing the County's contract and shall be liable for any payments due on such purchase orders; and shall accept sole responsibility for any payment due.
3. All purchases shall be subject to audit and inspection by the other political subdivisions and Rockland County non profit agencies for which the purchase was made.
4. No officer, board or agency of a county, town, village, or school district shall make any purchase through the County when bids have been received for such purchase by such officer, board or agency, unless such purchase may be made upon the same terms, conditions and specifications at a lower price through the County.
5. All Bidders shall be on notice that as a condition of the award of a County contract, the successful bidder shall accept the award of a similar contract with any other political subdivision in New York State and Rockland County non profit agencies authorized to use New York State's contracts, if called upon to do so. A listing of approved Rockland County non profit agencies is available on the Purchasing Division's website at www.rcpurchasing.com. The County, however, will not be responsible for any debts incurred by the participants pursuant to this or any other agreement.
6. Necessary deviations from the County's specifications in the award of a participant contract, whether such deviations relate to quantities, or delivery points shall be resolved between the successful bidder and the other political subdivisions and Rockland County non profit agencies.

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 TELEPHONE: 845-364-3820 / TELEFAX: 845-364-3809

**TITLE: PERSONAL PROTECTIVE EQUIPMENT
(PPE)**

RFB-RC-2020-089

SPECIFICATIONS

1. SCOPE

- 1.1. This solicitation is seeking competitive pricing from qualified supplier(s) to furnish and deliver Personal Protective Equipment (PPE) supplies as specified. It is anticipated that orders will be placed by the County of Rockland and any authorized agency on an as needed basis. (see page 5 - PURCHASE BY OTHERS).
- 1.2. Bidders must provide all of the requested certifications, documentation and information with their bid submission as required. Failure to comply may result in your bid being deemed non-responsive and removed from further consideration for award.

2. INSURANCE REQUIREMENTS

- 2.1. It is recommended that each bidder review the insurance requirements. If a potential bidder cannot meet the coverages limits, and requirements as stated, they should not submit a bid.
- 2.2. The County of Rockland requires awarded suppliers to provide a current insurance certificate (see sample certificate for coverage and limits required) There will be no exception to the coverages, limits and requirements as shown on the Sample Certificate. If the bidder cannot meet the requirements they will be deemed non-responsible and removed from further consideration for award.
- 2.3. The County of Rockland must be listed as additional. In addition to the liability certificate required, a valid NYS Wkr's Compensation and NYS Disability Certificate or Attestation of Exemption is required (see Insurance specifications).
 - 2.3.1. In an effort to expedite the award of this contract, bidders may submit the required Insurance Certificates with their bid. However, the apparent low bidder will be given five (5) business days from notice of award to supply these forms or the award will be rescinded.

<p style="text-align: right;">PAGE: 8</p> <p style="text-align: center;">COUNTY OF ROCKLAND - DGS-PURCHASING BLDG. A., 6TH FLOOR, 50 SANATORIUM RD, POMONA, NY 10970 TELEPHONE: 845-364-3820 / TELEFAX: 845-364-3809</p>	
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2.4. SECTION I & II - As needed Quantities

2.4.1. Orders shall be placed by any authorized agency on an as needed basis (“PURCHASES BY OTHERS”).

2.4.2. Unit pricing provided shall include FOB Destination pre-paid and allowed to the ship to location of the ordering agency authorized under this solicitation.

3. TERM

3.1. SECTION I & II – As needed quantities

3.1.1. Awarded supplier shall hold the pricing awarded under this section for a six (6) month term with the option to renew for one (1) additional six (6) month term. Option terms are exercised by mutual agreement between the County of Rockland and the Supplier.

4. QUANTITIES – SECTION I & II - As needed quantities

4.1. SECTION I - Minimum order quantities are listed in this section. Orders to be placed on an as needed basis.

4.1.1. The successful bidder shall be required to furnish quantities.

4.1.2. There is no guarantee of quantities as orders will be placed by agencies issuing individual Purchase Orders for quantities needed.

4.2. SECTION II – Quantities listed are estimated with no minimum order quantity. The successful bidder shall be required to furnish quantities ordered even if they are greater or less than quantities listed.

4.2.1. There is no guarantee of quantities as orders will be placed by agencies issuing individual Purchase Orders for quantities needed.

5. PURCHASE ORDERS

5.1. SECTION I & II – Supplier shall accept Purchase Orders from any authorized ordering agency.

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(PPE)**

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6. PAYMENT TERMS

6.1. SECTION I & II

6.1.1. Upon delivery, inspection and acceptance ordering agency will process payment in accordance with the terms provided on the Purchase Order.

7. PPE and Supplies

7.1. MASKS

7.2. Masks offered under this section shall not appear on the following:

7.2.1. Offered Masks must not appear on the current Emergency Use Authorization(EUA) Listing of removed respirator models (See Attachment A)

7.2.2. Offered masks must not have been deemed counterfeit or appear to be counterfeit in accordance with the guidelines set forth by the CDC. (URL Link below)

7.2.2.1. <https://www.cdc.gov/niosh/npptl/usernotices/counterfeitResp.html>

7.3. The County is requesting unit pricing on two types of masks. N95 and KN95

<p style="text-align: right;">PAGE: 10</p> <p style="text-align: center;">COUNTY OF ROCKLAND - DGS-PURCHASING BLDG. A., 6TH FLOOR, 50 SANATORIUM RD, POMONA, NY 10970 TELEPHONE: 845-364-3820 / TELEFAX: 845-364-3809</p>	
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7.4. N95 Particulate Filtering Facepiece Respirators

7.4.1. Specified Masks as manufactured by 3M, Honeywell, Makrite or approved equal. If submitting an alternate, bidder must provide a side by side comparison showing their product meets or exceeds the specified product. (see Attachment B – Technical Specification Sheet)

7.4.1.1.N95 Masks offered must be on the CDC NIOSH approved list/Certified Equipment List as noted below:

7.4.1.2.Bidder must supply Mfg. FDA Certification with their bid.

Mfg	Model	CDC APPROVAL LISTING URL LINK
3M or approved equal	9205+	https://wwwn.cdc.gov/niosh-cel/ NIOSH Approval Number TC-84A-8590
3M or approved equal	1870+ Surgical	https://www.cdc.gov/niosh/npptl/topics/respirators/disp_part/respsource3surgicaln95.html
3M or approved equal	8210	https://www.cdc.gov/niosh/npptl/topics/respirators/disp_part/default.html

7.4.2. N95 masks offered must be approved for healthcare use or have be on the FDA Emergency Use Authorization list

<p style="text-align: right;">PAGE: 11</p> <p style="text-align: center;">COUNTY OF ROCKLAND - DGS-PURCHASING BLDG. A., 6TH FLOOR, 50 SANATORIUM RD, POMONA, NY 10970 TELEPHONE: 845-364-3820 / TELEFAX: 845-364-3809</p>	
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7.5. KN95 – Disposable Filtering Facepiece Respirator – Non-NIOSH Approved

7.5.1. Specified masks as Mfg. by Guangzhou Powecom Labor GB2626-2006- or approved equal.

If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. (see Attachment B – Technical Specification Sheet). Failure to meet this requirement may deem your bid non-responsive.

7.5.2. Offered mask must be on the FDA Emergency Use Authorization (EUA).

7.5.3. <https://www.fda.gov/medical-devices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/personal-protective-equipment-euas>

7.6. Gowns – Offered Gowns must not be on the FDA recall list. All gowns offered must meet the Standards of Gowns as recognized by the FDA:

7.6.1. <https://www.fda.gov/medical-devices/personal-protective-equipment-infection-control/medical-gowns>

7.6.2. Disposable 100% Polypropylene

7.6.2.1. Fluid Resistant

7.6.2.2. American National Standard - AAMI PB70 STANDARD as standard recognized by the FDA

7.6.2.3. Seeking pricing on Level 1, Level 2 and Level 3

7.6.2.4. Blue

7.6.2.5. Full Length

7.6.2.6. Tie Back

7.6.2.7. Elastic Cuffs

7.6.2.8. Pricing offered in accordance with the proposal pages. County is seeking pricing for:

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- Level 1: *Minimal risk*, to be used, for example, during basic care, standard isolation, cover gown for visitors, or in a standard medical unit
- Level 2: *Low risk*, to be used, for example, during blood draw, suturing, in the Intensive Care Unit (ICU), or a pathology lab
- Level 3: *Moderate risk*, to be used, for example, during arterial blood draw, inserting an Intravenous (IV) line, in the Emergency Room, or for trauma cases

7.6.3. Disposable Polyethylene Film

7.6.3.1. Specified PolyCo 10400 or approved equal– Sized Regular, Large, Extra Large or equivalent. If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. (see Attachment B – Technical Specification Sheet). Failure to meet this requirement may deem your bid non-responsive.

7.6.3.2. ASTM F1670

7.6.3.3. American National Standard - AAMI PB70 STANDARD – Levels 1 & 2 (**see above**)

7.6.4. Disposable Isolation Gown. Specified Mfg. Procure PC150 or equal

7.6.4.1. Disposable isolation gown, yellow, sms, fluid resistant, non sterile, elastic cuff, tie back, long sleeve, knee length

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7.7. Gloves – Specified Mfg. Central Association for the Blind and Visually Impaired or equal

7.7.1. If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. Failure to meet this requirement may deem your bid non-responsive.

7.7.2. Nitrile Gloves – Specified Better Touch (4 mil) or approved equal

7.7.2.1. Latex-Free

7.7.2.2. Powder-Free

7.7.2.3. Ambidextrous

7.7.2.4. 4 mil Thickness

7.7.2.5. 9.5” length

7.7.2.6. Non-Sterile

7.7.3. Black Nitrile Gloves – Specified Better Touch (4-mil – palm) or approved equal

7.7.3.1. Medical Grade Examination Gloves

7.7.3.2. Latex-Free

7.7.3.3. Powder-Free

7.7.3.4. Ambidextrous

7.7.3.5. 4 mil palm, -5 mil finger Thickness

7.7.4. EMS Latex Examination Gloves Specified Better Touch (15 mil) or approved equal

7.7.4.1. 15 mil Extra Thick

7.7.4.2. Powder-Free

7.7.4.3. Non-Sterile

7.7.4.4. Textured Fingers

7.7.4.5. Single Use Only

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7.8. Face Shields -Specified mfg. Nikomed, USA Inc. N1288 – FSDISP or approved equal

- 7.8.1. Full Face Coverage (below the chin)
- 7.8.2. Pre-Assembled
- 7.8.3. Elastic Band
- 7.8.4. 1.3”-1/5” Thick Foam,
- 7.8.5. Latex-Free
- 7.8.6. Polymer Film
- 7.8.7. Non-Sterile
- 7.8.8. If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. (see Attachment B – Technical Specifications Sheets). Failure to meet this requirement may deem your bid non-responsive and removed from further consideration for award.

7.9. Surface Disinfecting Wipes – Specified Mfg. PDI or approved equal

- 7.9.1. PDI – Q55172 or approved equal. EPA Registration 9480-4
 - 7.9.1.1. Active Ingredient -Quaternary ammonium; Isopropanol (Isopropyl alcohol)
 - 7.9.1.2. Super Sani-Cloth Germicidal Disposable Wipe
 - 7.9.1.3. Surface Types - Hard Nonporous (HN); Food Contact Post-Rinse Required (FCR)
 - 7.9.1.4. Use Sites: Healthcare; Institutional
 - 7.9.1.5. Why product is on the N List: Kills a harder-to-kill pathogen than SARS-CoV-2 (COVID-19); Emerging viral pathogen claim
 - 7.9.1.6. Disinfecting directions for following Pathogens - Rhinovirus 39; Adenovirus
- 7.9.2. Offered product must be on the EPA List N Disinfectants for use against SARS-CoV-2 (COVID-19). Bidder shall provide EPA Registration Number with their bid. Failure to comply may result in your bid being deemed non-responsive and removed from consideration for award.

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- Link to site: <https://www.epa.gov/pesticide-registration/list-n-disinfectants-use-against-sars-cov-2-covid-19>
- If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. Failure to meet this requirement may deem your bid non-responsive and removed from further consideration for award.

7.10. Sanitizing Hand Wipes individual packets and cartons – PDI or approved equal

- 7.10.1. Individual Packets – PDI D43600 or approved equal
- 7.10.2. 70% Ethyl Alcohol or no less than 62%
- 7.10.3. Safe to use before eating or drinking
- 7.10.4. Kills 99.99% germs
- 7.10.5. Meets OSHA BLOODBORNE PATHOGEN STANDARD 29 CFR Part 1910.1030 (d)(2)(iv)
- 7.10.6. Latex- Free
- 7.10.7. If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. Failure to meet this requirement may deem your bid non-responsive and removed from further consideration for award.

7.11. Sanitizing Hand Wipes – Mfg. PDI or approved equal. Medium Size Canister

- 7.11.1. Medium Size Container 135 Count – PDI P13472 or approved equal
- 7.11.2. 6" x 7.5 P
- 7.11.3. 70% Ethyl Alcohol or no less than 65.9%
- 7.11.4. Safe to use before eating or drinking
- 7.11.5. Kills 99.99% germs
- 7.11.6. Meets OSHA BLOODBORNE PATHOGEN STANDARD 29 CFR Part 1910.1030 (d)(2)(iv)
- 7.11.7. Late Free
- 7.11.8. If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. Failure to meet this requirement may deem your bid non-responsive and removed from further consideration for award.

7.12. Hand Sanitizer Gel – Specified Purell Advantage 9605-24 and 9652-12 or approved equal

- 7.12.1. 70% Ethyl Alcohol no less than 60%

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7.13. Non-Contact Infrared Thermometer for Clinical Use

- 7.13.1. Specified Berrcom JXB-178 or approved equal
- 7.13.2. Unit size: 187*120*57 mm(LxWxH)
- 7.13.3. Unit weight(without battery): 80g
- 7.13.4. Temperature display resolution: 0.1C(0.1F)
- 7.13.5. Consumption: ≤300mW
- 7.13.6. Memory: 32 sets
- 7.13.7. Normal using condition:
- 7.13.8. Ambient temperature: 10°C ~ 40°C (50°F ~ 104°F)
- 7.13.9. Relative humidity: ≤85%
- 7.13.10. Audible alarm if temperature is more than 38°C (100.4°F)
- 7.13.11. It can be displayed in either Celsius or Fahrenheit
- 7.13.12. Longevity use 100,000 readings
- 7.13.13. Eliminates cross contamination Hygienic and easy to use
- 7.13.14. Accuracy: ± 0.3°C (0.6°F)
- 7.13.15. Response Time: 1 sec
- 7.13.16. Suggested Age: Newborn and Up
- 7.13.17. Automatic power-off: <30 secs
- 7.13.18. If submitting an alternate, bidder must provide, with their bid, a side by side comparison showing their product meets or exceeds the specified product. Failure to meet this requirement may deem your bid non-responsive and removed from further consideration for award.

8. DELIVERY

- 8.1. Product pricing to include FOB destination pre-paid and allowed, delivery to be within 15 days after receipt of purchase order.

9. NON-RESTRICTIVE USE OF BRAND NAME OR EQUAL SPECIFICATIONS

- 9.1. The use of a brand name is for the purpose of describing the standard of quality, performance, and characteristics desired and is not intended to limit or restrict competition.

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10. MANUFACTURER BRAND, PART NUMBER AND PACKAGING

- 10.1. Actual manufacturer brand name and part numbers being bid are to be indicated in the part number column. Bidders are to indicate actual manufacturer's part number in this column even if bidding as specified. If the actual manufacturers brand name and product code on any item line being bid is left blank, bid may be rejected.
- 10.2. Bidders must submit SDS sheets with their bid, when applicable.
 - 10.2.1. SDS sheets submitted must meet the current Hazard Communication Standard issued by the Occupational Safety & Health Administration (OSHA).

11. SAMPLES

- 11.1. The apparent low bidder must provide a sample of the offered product within two (2) business days from date of request. Sample must be sent ATTN: Ann Marie Curley, CPPB, Assistant Director of Purchasing, Purchasing Division, 50 Sanatorium Road, Bldg. A, 6th Floor, Pomona, NY 10970.
- 11.2. Bidders may wish to send samples prior to the solicitation scheduled close date and time. Samples must be sent to the address above, the solicitation number and supplier name must be clearly visible on the package.

12. REFERENCES

- 12.1. Bidders must provide a minimum of three (3) references where the offered product has been delivered and accepted. See Certificate of Experience. Failure to submit this form may result in your bid submission being deemed non-responsive and removed from further consideration for award.

13. PROPOSAL PAGES

- 13.1. **Bid will be rejected if the following is not adhered to:**
- 13.2. The Manufacture Name, Brand Name, Manufacturer Product Code and Packaging Columns **must** be filled in. All required documentation as stated in the specifications must be submitted at time of bid.

14. AWARD

- 14.1. Award will be made on a line by line basis to the lowest responsive responsible bidder (s) that meet the stated requirements.
 - 14.1.1. The County of Rockland reserves the right to make a no award on a line by line basis.



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 BID ITEM #'S 1-19 THROUGH 1-23
 KN95 MASKS CERTIFICATIONS
 PPE TRADE COMPANY LLC

ppetradesupport.com
 contact@ppetradesupport.com
 404-759-8409

Jinhua Jiadaifu Medical Supplies Co., Ltd. (KN95)

<https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/ciRL/rl.cfm?lid=654956&lpcd=MSH>

The screenshot shows the FDA's Establishment Registration & Device Listing page. The header includes the FDA logo and navigation links for Home, Food, Drugs, Medical Devices, Radiation-Emitting Products, Vaccines, Blood & Biologics, Animal & Veterinary, Cosmetics, and Tobacco Products. The main content area displays the following information for the establishment:

- Proprietary Name:** Disposable medical masks; disposable surgical masks; KN95 masks
- Classification Name:** RESPIRATOR, SURGICAL
- Product Code:** MSH
- Device Class:** 2
- Regulation Number:** 876.4040
- Medical Specialty:** Ear
- Registered Establishment Name:** srzrzi
- Establishment Operations:** Manufacturer, Specification Developer
- Other Information:** 10063163, Jinhua Jiadaifu Medical Supplies Co., Ltd.

The page also includes a footer with accessibility information, contact details, and various links.

<https://www.fda.gov/medicaldevices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/personal-protective-equipment-euas>

Jiangsu Yimao Filter Media Co., Ltd	9570K, 9560K	* IFUG?
Jiaxing Yinuo Busway Co., Ltd.	YJP2	
Jingzhou Strong Sciences & Technology Development Co., Ltd.	STAS502, A9507	* STAS502: IFUC • A9507: IFU VS
Jinhua Ai Kou Protective Equipment Co.,	AKQ03	* IFUGT
Jinhua Jiadaifu Medical Supplies Co., Ltd.	Disposable Non-Medical Face Mask (KN95)	* IFUC?
Jinwells (Tianjin) Science and Technology Co., Ltd.	JWS-1, JWS-2	* JWS-1: IFU S * JWS-2: IFU GS
Jinyi (Tianjin) Medical Technology Co., Ltd.	JYOS01	
Unifan Shandong Technology Co., Ltd.	S9-KN95	* IFUG?

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RID ITEM #'S 1-13 THROUGH 1-23
KN95 MASKS CERTIFICATIONS
PPE TRADE COMPANY

CERTIFICATION OF REGISTRATION

2020

This certifies that:

JINHUA JIADAIFU MEDICAL SUPPLIES CO. LTD
Dongxi industrial zona, bailonggiao town, wuchang district,
jinhua, Zhejiang, 321000, CHINA

Was registered with US Food and Drug Administration, Center for Devices and Radiological Health, pursuant to the Code of Federal Regulations 21 CFR 807, by F&W (Shanghai) Certification Co., Ltd.

Owner/Operator Number: 10063163
Device Listing#: See annex
Expiration Date: December. 31, 2020

2028F&W will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration terminates after issuance of this certificate. F&W makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate holder's device or establishment by the US Food and Drug Administration. F&W assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.9, "Registration of a device establishment or assignee. A registration number does not in any way denote approval of the establishment or its products. Any representation that makes an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding." The US Food and Drug Administration does not issue a certificate of registration nor does the FDA Food and Drug Administration recognize a certificate of registration. F&W is not affiliated with the FDA Food and Drug Administration.



Registration Co., Ltd.
on behalf of
NM (Shanghai) Certification Co., Ltd.

Executive Director

Dated: March 20, 2020

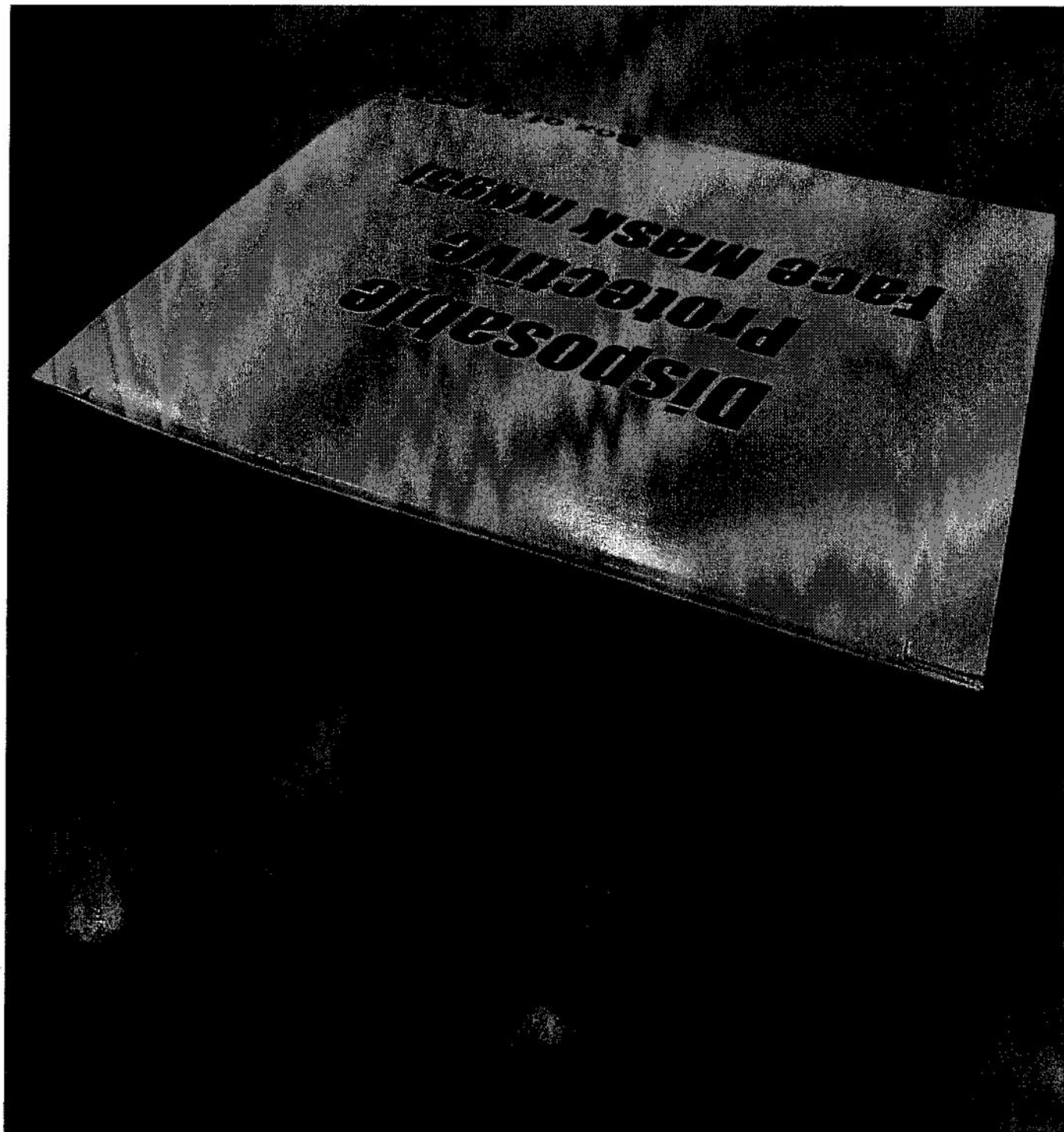
MW (Shanghai) Certification Co., Ltd. TEL: 821-62960771

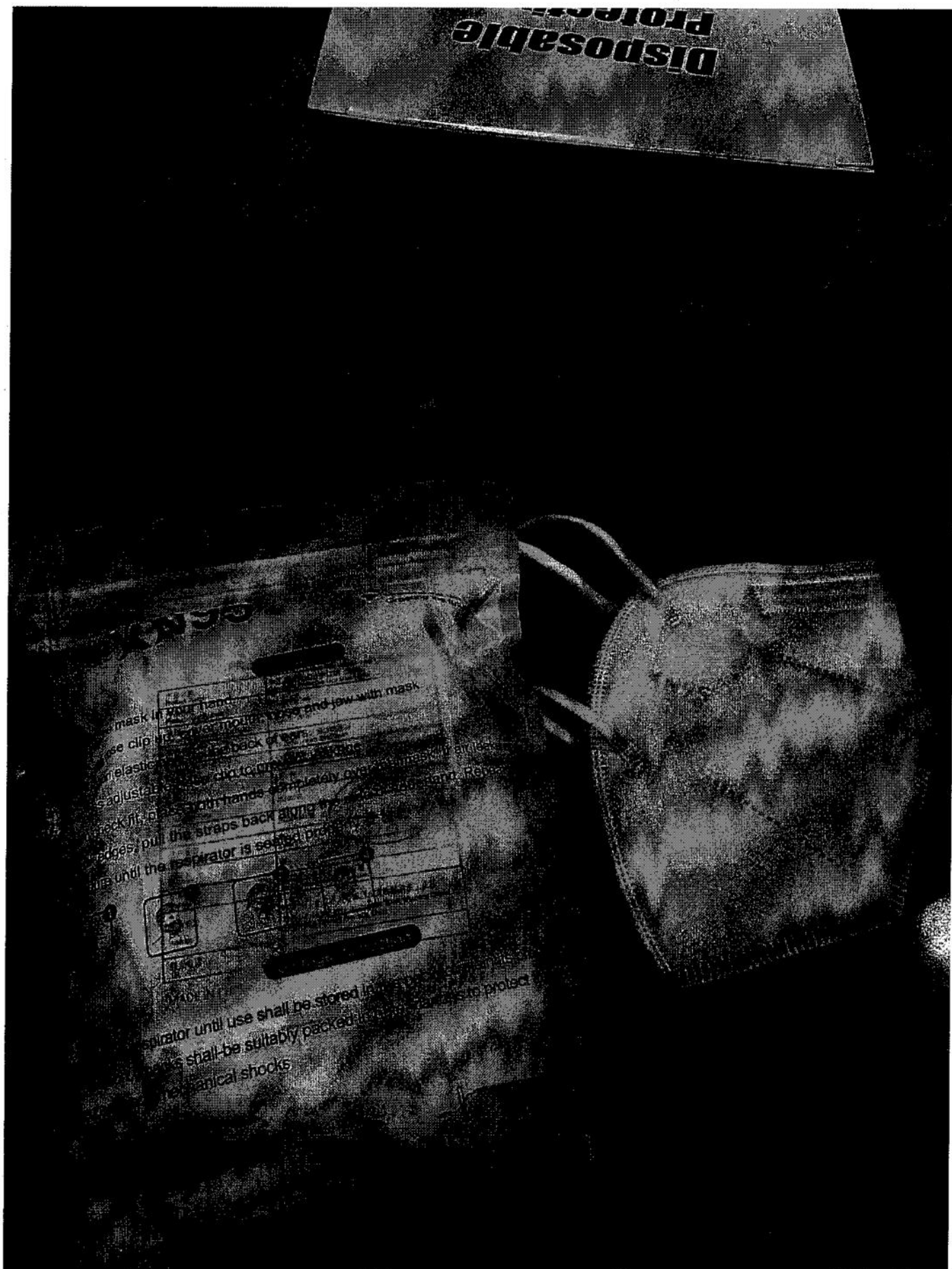
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KN95 MASKS CERTIFICATIONS
PPE TRADE COMPANY LLC

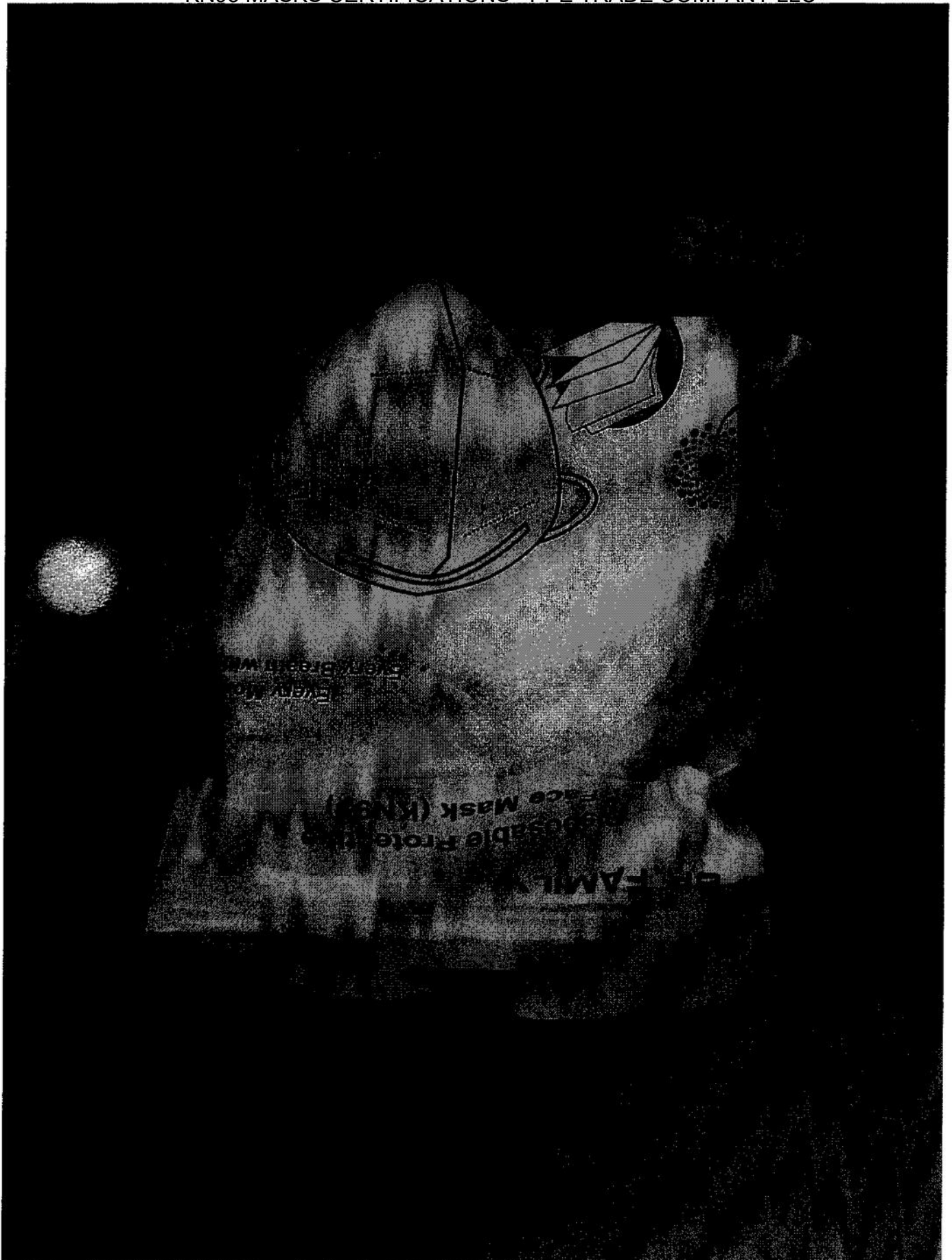




RFB-RC-2020-089 PPE

BID ITEM #'S 1-19 THROUGH 1-23

KN95 MASKS CERTIFICATIONS - PPE TRADE COMPANY LLC



Appendix A: Authorized Imported, Non-NIOSH Approved Respirators Manufactured in China (Updated: July 7, 2020)

The table below includes a list of non-NIOSH respirators authorized by this Umbrella EUA for emergency use during the COVID-19 public health emergency.



As stated in the EUA, authorized respirators should be used in accordance with CDC's recommendations. For the most current CDC recommendations on optimizing respirator use, please visit CDC's webpage: [Strategies for Optimizing the Supply of N95 Respirators](#).

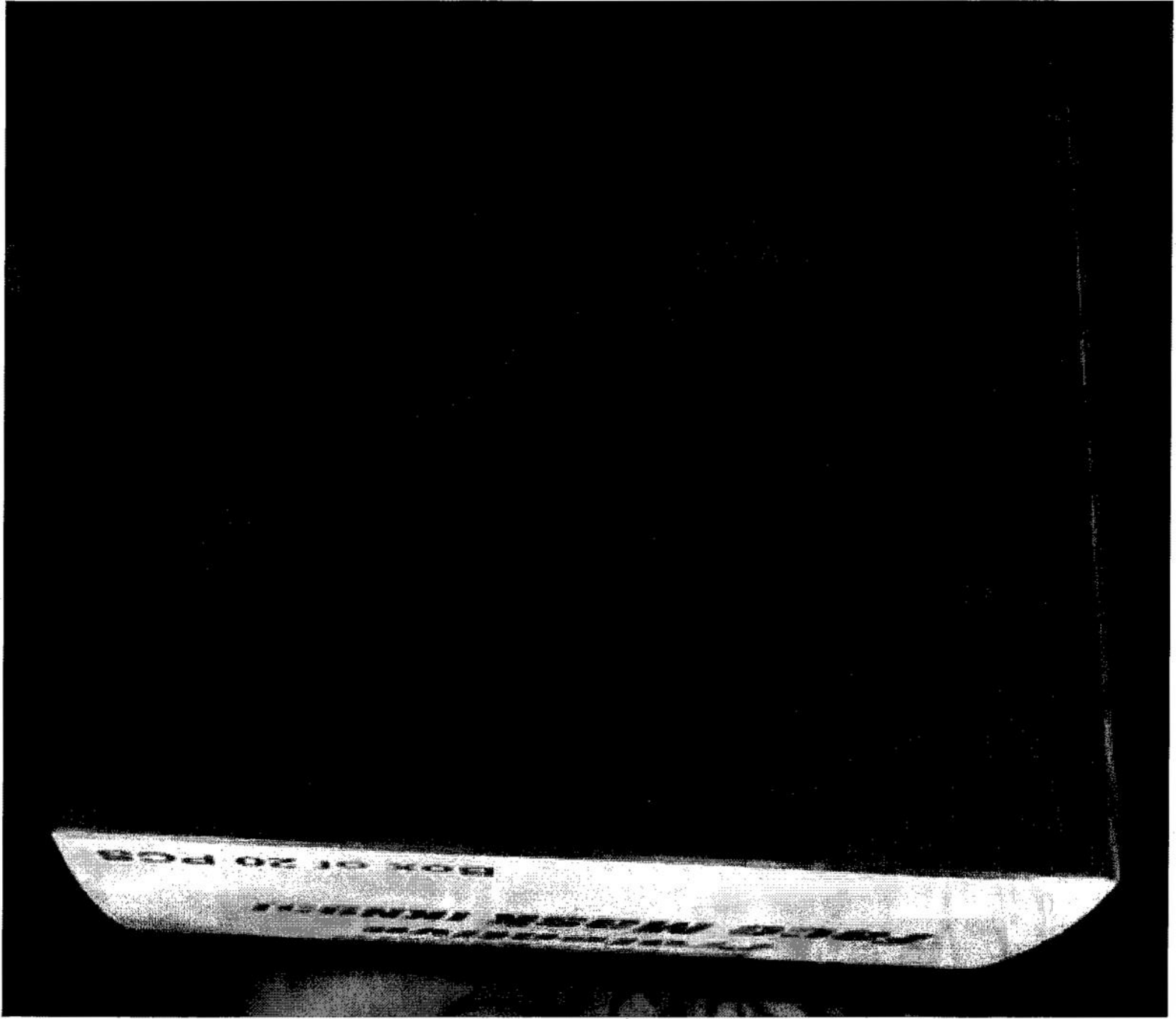
Search:

Show entries

Manufacturer	Respirator Model(s)	\$ instructions for Use
3M	9001, 9002, 9501, 9501+, 9501V+, 9502, 9502+, 9502V+, 9505+, 9541, 9541V, 9542, 9542V, 9552, 9552V	<ul style="list-style-type: none"> 9001: IFU Gf 9002: IFU Gf 9501: IFU Of 9501+: IFUGf 9501V+: IFU G? 9502: IFU Of 9502+: IFU Of 9502V+: IFU Gf 9505+: IFU Gf 9541: IFU Gf 9541V: IFU GT 9542: IFU Gf 9542V: IFU Gf 9552: IFU C? 9552V: IFU Gf
Allmed Medical Products Co., Ltd	LP220002	<ul style="list-style-type: none"> IFU Gf
Anhui Zhongke Duling Commercial Appliance Co. Ltd	M-9501	
AOK Tooling Ltd. (aka Shenzhonghai Medical)	20130040, 20130045A, 20180021, 20130038, 20190019, 910	<ul style="list-style-type: none"> 20130040: IFU Gf 20130045A: IFU Gf 20180021: IFU Gf 20130038: IFU Gf 20190019: IFU Gf

RFB-RC-2020-089 PPE
 BID ITEM #S 1-19 THROUGH 1-23
 KN95 MASKS CERTIFICATIONS
 PPE TRADE COMPANY LLC

Hunan Kangweining Medical Devices Co., Ltd.	YH-1 (non-sterile) and YH-11 (sterile)	
Jiande Chaomei Daily Chemicals Co.	F-Y3-A	• IFU 
Jiangsu Jiaao Medical Technology Co., Ltd.	JA95-1 Filtering half mask	in IFUGf
Jiangsu Yimao Filter Media Co., Ltd	9570K	• IFUGf
 Jinhua Jiadaifu Medical Supplies Co., Ltd.	Disposable Non-Medical Face Mask (KN95)	7 • IFUGf
Lanshan Shendun Technology Co., Ltd.	SD-KN95	
Liaoning Dalian Jieying Energy Saving Environmental Protection Technology Development Co., Ltd.	JY-FH-F JY-FH	



Box of 20 PCS

FOLD OVER

RFB-RC-2020-089 PPE
BID ITEM #'S 1-19 THROUGH 1-23
KN95 MASKS CERTIFICATIONS - PPE TRADE COMPANY LLC

NPPTL COVID-19 Response: International Respirator Assessment

Manufacturer: Jinhua Jiadaifu Medical Supplies Co., Ltd.
Model Tested: Disposable Non-medical Face Mask (KN95)
Date Tested: June 19, 2020

These findings pertain to the Jinhua Jiadaifu Medical Supplies Co., Ltd., Disposable Non-medical Face Mask (KN95). The packaging for this product indicates that it meets GB2626-2006 (the Chinese standard for Respiratory Protective Equipment – Non-Powered Air-Purifying Particle Respirator).

Thirty respirators were submitted for evaluation. The respirators were sampled into groups often for evaluation. The samples were tested using a modified version of NIOSH Standard Test Procedure (STP) TEB-APR-STP-0059. This modified assessment plan can be found [here](#).

No certificate of approval was provided with the samples received; therefore, the authenticity of the claims cannot be validated.

The maximum and minimum filter efficiency was 99.88% and 99.54%, respectively. All thirty respirators measured more than 95%.

While the above-listed product classification has similar performance requirements to NIOSH-approved devices, NIOSH does not have knowledge about the sustained manufacturer quality system and product quality control for these products. NIOSH also does not have knowledge about the product's handling and exposures after leaving its manufacturer's control.

In addition, this product is an ear loop design. Currently, there are no NIOSH-approved products with ear loops; NIOSH-approved N95s have head bands. Furthermore, limited assessment of ear loop designs, indicate difficulty achieving a proper fit. While filter efficiency shows how well the filter media performs, users must ensure a proper fit is achieved.

This assessment is not a part of the NIOSH respirator approval process and will in no way lead to or preclude NIOSH approval through the official approval process. This assessment was developed as an assessment of the filter efficiency for those respirator's represented as certified by an international certification authority, other than NIOSH, to support the availability of respiratory protection to US healthcare workers due to the respirator shortage associated with COVID-19. Only particulate filter efficiency was assessed.

The results provided in this letter are specific to the subset of samples that were provided to NPPTL for evaluation.

These results will be used to update the CDC guidance for [Crisis Capacity Strategies \(during known shortages\)](#).

Evaluation of International Respirators

Test: Modified TEB-APR-STP-0059

Date Tested: June 19, 2020

Report Prepared: June 20, 2020

Manufacturer: Jinhua Jiadaifu Medical Supplies Co., Ltd.

Item Tested: Disposable Non-medical Face Mask (KN95) (Sample Group 1 of 3)

Country of Certification: China (GB2626-2006)

Pictures have been added to the
end of this report.

Filter	Flow Rate (Lpm)	Initial Filter Resistance (mmHzO)	Initial Percent Leakage (%)	Maximum Percent Leakage (%)	Filter Efficiency
1	85	12.1	0.20	0.20	99.80
2	85	12.0	0.24	0.24	99.76
3	85	11.6	0.26	0.26	99.74
4	85	11.4	0.25	0.25	99.75
5	85	11.2	0.24	0.24	99.76
6	85	12.2	0.22	0.22	99.78
7	85	11.3	0.27	0.27	99.73
8	85	12.2	0.15	0.15	99.85
9	85	13.1	0.20	0.20	99.80
10	85	11.4	0.25	0.25	99.75
Minimum Filter Efficiency: 99.73			Maximum Filter Efficiency: 99.85		

- The test method utilized in this assessment is not the NIOSH standard test procedure that is used for certification of respirators. Respirators assessed to this modified test plan do not meet the requirements of STP-0059, and therefore cannot be considered equivalent to N95 respirators that were tested to STP-0059.
- Respirators tested may not be representative of all respirators with the same certification mark. NIOSH has no control over suppliers and distributors of respirators certified by other national or international parties.
- This assessment is not a confirmation that it conforms with any or all of its specifications in accordance with its certification mark.
- This assessment was not a part of the NIOSH approval program. These results do not imply nor preclude a future approval through the NIOSH respirator approval program.

NPPTL COVID-19 Response: International Respirator Assessment

RFB-RC-2020-089 PPE
BID ITEM #'S 1-19 THROUGH
1-23
KN95 MASKS
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PPE TRADE COMPANY LLC

Test: Modified TEB-APR-STP-0059

Date Tested: June 19, 2020

Report Prepared: June 20, 2020

Manufacturer: Jinhua Jiadaifu Medical Supplies Co., Ltd.

Item Tested: Disposable Non-medical Face Mask (KN95) (Sample Group 2 of 3)

Country of Certification: China (GB2626-2006)

Filter	Flow Rate (Lpm)	initial Filter Resistance (mmHzO)	Initial Percent Leakage (%)	Maximum Percent Leakage [%]	Filter Efficiency
11	85	12.2	0.12	0.12	99.88
12	85	12.5	0.28	0.28	99.72
13	85	13.3	0.16	0.16	99.84
14	85	12.0	0.16	0.16	99.84
15	85	13.4	0.19	0.19	99.81
16	85	12.9	0.16	0.16	99.84
17	85	12.3	0.15	0.15	99.85
18	85	12.8	0.46	0.46	99.54
19	85	13.9	0.19	0.19	99.81
20	85	12.6	0.14	0.14	99.86
Minimum Filter Efficiency: 99.54			Maximum Filter Efficiency: 99.88		

- The test method utilized in this assessment is not the NIOSH standard test procedure that is used for certification of respirators. Respirators assessed to this modified test plan do not meet the requirements of STP-0059, and therefore cannot be considered equivalent to N95 respirators that were tested to STP-0059.
- Respirators tested may not be representative of all respirators with the same certification mark. NIOSH has no control over suppliers and distributors of respirators certified by other national or international parties.
- This assessment is not a confirmation that it conforms with any or all of its specifications in accordance with its certification mark.
- This assessment was not a part of the NIOSH approval program. These results do not imply nor preclude a future approval through the NIOSH respirator approval program.

NPPTL COVID-19 Response: International Respirator Assessment

RFB-RC-2020-089 PPE
BID ITEM #'S 1-19 THROUGH
1-23
KN95 MASKS CERTIFICATIONS
PPE TRADE COMPANY LLC

Test: Modified TEB-APR-STP-0059

Date Tested: June 19, 2020

Report Prepared: June 20, 2020

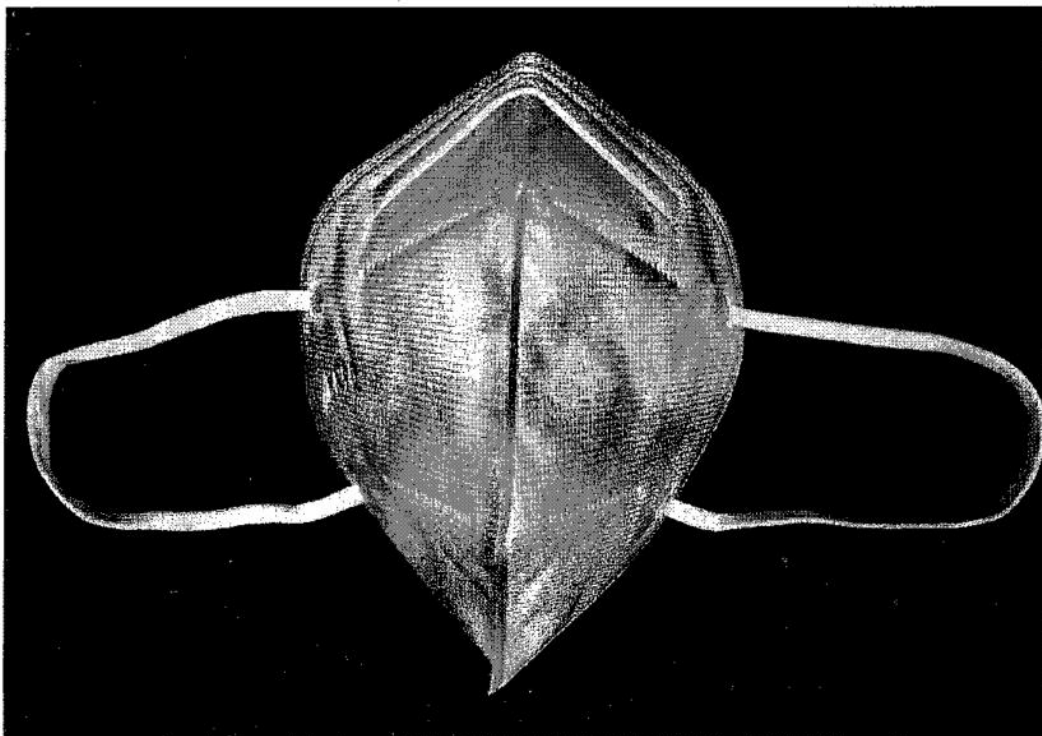
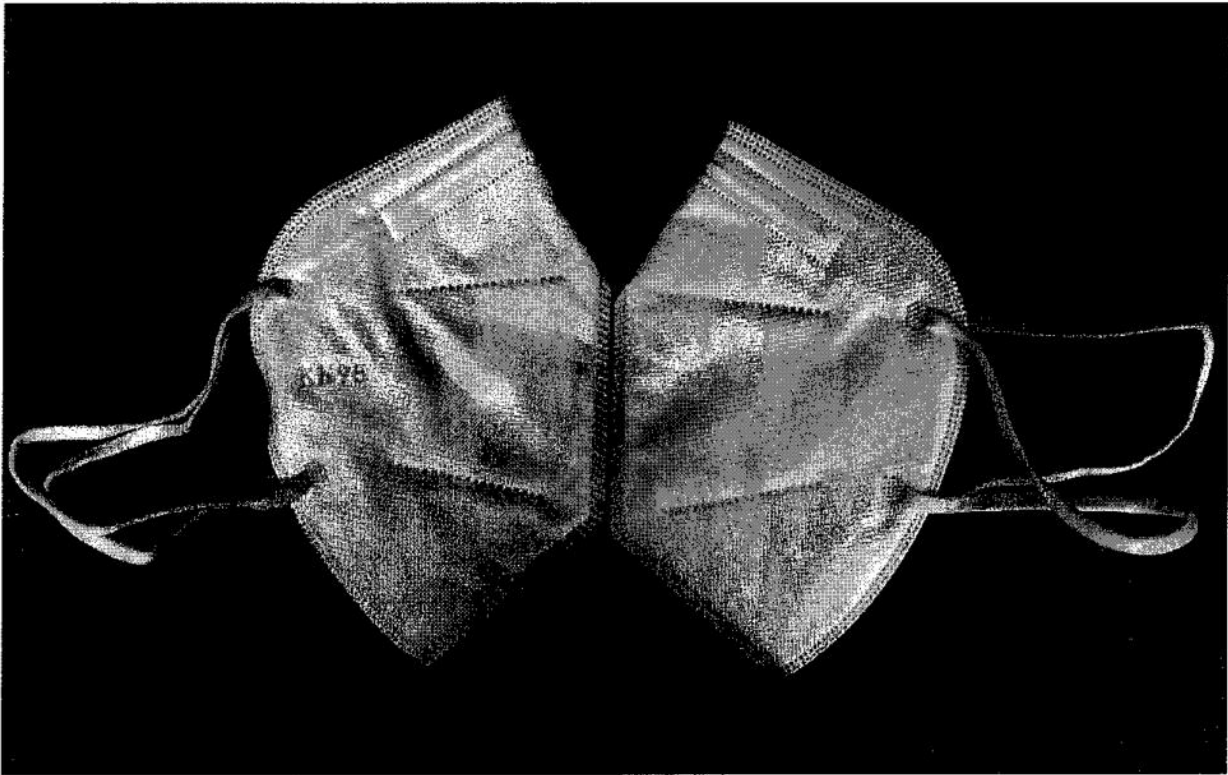
Manufacturer: Jinhua Jiadaifu Medical Supplies Co., Ltd.

Item Tested: Disposable Non-medical Face Mask (KN95) (Sample Group 3 of 3)

Country of Certification: China (GB2626-2006)

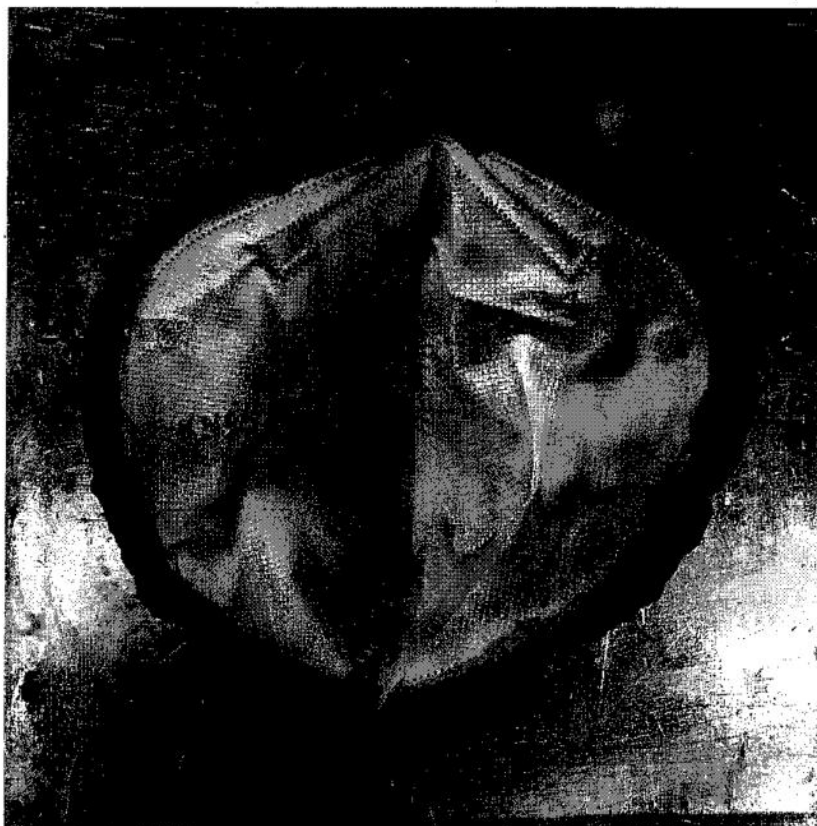
Filter	Flow Rate (Lpm)	Initial Filter Resistance (mmH ₂ O)	Initial Percent Leakage (%)	Maximum Percent Leakage (%)	Filter Efficiency
21	85	12.4	0.34	0.34	99.66
22	85	11.4	0.20	0.20	99.80
23	85	12.7	0.22	0.22	99.78
24	85	14.5	0.19	0.19	99.81
25	85	11.9	0.20	0.20	99.80
26	85	10.9	0.28	0.28	99.72
27	85	11.9	0.19	0.19	99.81
28	85	11.3	0.21	0.21	99.79
29	85	13.4	0.20	0.20	99.80
30	85	11.8	0.17	0.17	99.83
Minimum Filter Efficiency: 99.72			Maximum Filter Efficiency: 99.83		

- The test method utilized in this assessment is not the NIOSH standard test procedure that is used for certification of respirators. Respirators assessed to this modified test plan do not meet the requirements of STP-0059, and therefore cannot be considered equivalent to N95 respirators that were tested to STP-0059.
- Respirators tested may not be representative of all respirators with the same certification mark. NIOSH has no control over suppliers and distributors of respirators certified by other national or international parties.
- This assessment is not a confirmation that it conforms with any or all of its specifications in accordance with its certification mark.
- This assessment was not a part of the NIOSH approval program. These results do not imply nor preclude a future approval through the NIOSH respirator approval program.



NPPTL COVID-19 Response: International Respirator Assessment

RFB-RC-2020-089 PPE
BID ITEM #'S 1-19 THROUGH
1-23
KN95 MASKS
CERTIFICATIONS
PPE TRADE COMPANY LLC



UNIVERSAL

CERTIFICATION

NB 2163

EU TYPE EXAMINATION CERTIFICATE

Certificate No: 2163-PPE-724

Respiratory protective devices, filtering half masks to protect against particles manufactured by

Jinhua Jiadaifu Medical Supplies Co., Ltd.

Dongxi Industrial Zone, Bailongqiao Town, Wucheng District, Jinhua City, Zhejiang Province,
China

are tested and evaluated according to

**EN 149:2001 + A1:2009 Respiratory Protective Devices -
Filtering Half Masks to Protect Against Particles -
Requirements, Testing, Marking**

Based on the type examination conducted with the evaluation of test report Sj technical file
according to Personal Protective Equipment Regulation (EU) 2016/425 Annex 5, it is approved
that the product meets the requirements of the regulation,

Product Definition

Brand Name: DR FAMILY Model: JDF-01
Filtering half mask
Classification: FFP2 NR

Here by the manufacturer is allowed to use notified body number (2163) and can fix CE mark, as
shown below, on the Category III product models given above, with;

- Issuing an appropriate EU Declaration of Conformity according to Personal Protective Equipment Regulation (EU) 2016/425 Annex 9.
- Ongoing successful performance in fulfilment of the requirements set out in Personal Protective Equipment Regulation (EU) 2016/425 and harmonised standards, ensured by assessments based on Annex 7 (Module C2) or Annex 8 (Module D) of the regulation no later than 1 year from the beginning of Serial production.

This certificate is initially issued on 09/06/2020 and will be valid for 5 years, if there is no
change in the relevant harmonised standard affecting the essential health and safety
requirements.



Suat KACMAZ

Suat KACMAZ
UNIVERSAL CERTIFICATION
Director



Verify the validity with the QR code

mmf



NB 2163

CERTIFICATE OF CONFORMANCE

Certificate No: 2163-PPE-724/01

Respiratory protective devices, filtering half masks to protect against particles manufactured by

Jinhua Jiadaifu Medical Supplies Co., Ltd.

Dongxi Industrial Zone, Bailongqiao Town, Wucheng District, Jinhua City, Zhejiang Province, China

Continues to fulfil the requirements of

EN 149:2001 + A1:2009 Respiratory Protective Devices - Filtering Half Masks to protect Against Particles - Requirements, Testing, Marking

Based on the evaluation of test reports and internal quality control audit reports according to EN 149+A1:2009 and Personal Protective Equipment Regulation (EU) 2016/425 Annex VII (Module C2). This certificate implies that the manufactured products shown below are in conformance with the approved EU Type Examination model and meets the requirements of the regulation.

Product Definition

Model	Glass	EU Type Examination Certificate		
		Serial Nr.	Date	Issuing NB Nr.
DR-FAMILY / XDF-01	FFP2 NR	2163-PPE-724	06.2020	2163

Hereby the manufacturer is allowed to use notified body number (2163) and can fix CE mark, as

shown below, on the Category II product models given above, with:

- Issuing an appropriate EU Declaration of Conformity according to **Personal Protective Equipment Regulation (EU) 2016/425 Annex 9**.
- Taking all measures necessary so that the manufacturing process and its monitoring ensure the Homogeneity of production and conformity of the manufactured PPE with the type described in the type examination certificate.

This certificate is issued on 09/06/2020 and will be valid for one year, until 08/06/2021 if the manufacturer makes no major change in the product designs and manufacturing processes affecting the product performance on the essential health and safety requirement.



2163

As

Suat KACMAZ
UNIVERSAL CERTIFICATION
Director

Verify the validity with the QR code



FvHSP


a

mm

mm

17

RFB-RC-2020-089 PPE
 BID ITEM 1-24 KN95 MASKS
 FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC





Certificate Of Registration

GUANGZHOU NAN QI XING NONWOVEN CO., LTD
 North Side Of Beihuan Road, Wanbian Waga, Shiqi Town, Panyu District, Guangzhou, Guangdong, China
 Has Completed VM1 The U.S. Food And Drug Administration Pursuant To 21 CFR Part 807: Establishment Registration And Device Listing

Owner/Operator No: 1M63420
 Broker Listing: SM Next Major


Huawin will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate. Huawin makes no other representations or warranties, nor does this certificate make sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. Huawin assumes no liability to any person or entity in connection with the foregoing.

The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Huawin is not affiliated with the U.S. Food and Drug Administration.


Mam
 Issue date: Apr 17, 2020
 Expire Date: Dec 31, 2020

Shenzhen Huawin Testing Certification Co., Ltd.
 Add: 7F, U Center No.743, Zhongli Road, Bao'an, Shenzhen, China
 Http://www.huawinlab.com E-mail: info@huawinlab.com



Certificate Of Registration

Device Util/Mfg:	Cert No.	Proprietary Name	Model No.
Ijuting No. 03831 22	Q&R	Disposable medical mask	Y-1, Y-2
0333 123	KWA	Disposable medical mask	Y-1, Y-2
0380065	LYU	Disposable medical mask	Y-1, Y-2
D389285	OEA	Disposable Protective Clothing for medical use	Planar ear loop type 160cm, 185cm, 170cm, 175cm, 180cm, 185cm
D389283	MSH	KN95 Respirator Mask	KN95
		N95 Respirator Mask	N95
		Child Respirator Mask	NC-1



Manager/V
 Issue date: Apr 17, 2020
 Expire Date: Dec 31, 2020

Shenzhen Huawin Testing Certification Co., Ltd.
 Add: 7F, U Center No.743, Zhongli Road, Bao'an, Shenzhen, China
 Http://www.huawinlab.com E-mail: info@huawinlab.com

RFB-RC-2020-089 PPE
BID ITEM 1-24 KN95 MASKS
FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC

No. FZ2008490

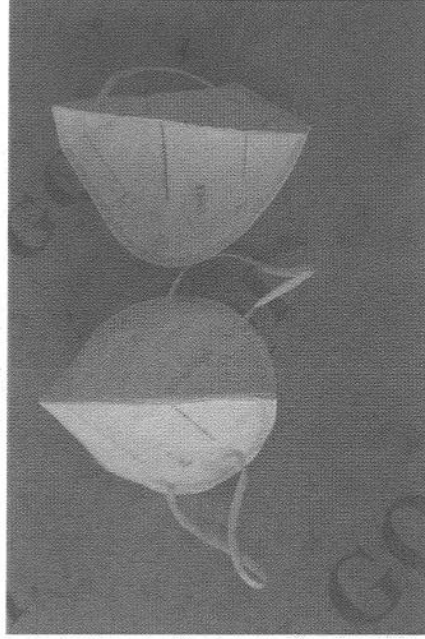
& 测试报告
TEST REPORT

M 4 W it 5 W



照/其他检测标准

(01)



样品描述 Sample description

型号规格或其它说明 Type, Specification or other description

FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC

GUANGDONG TESTING INSTITUTE OF PRODUCT QUALITY SUPERVISION



TEST REPORT

m i m pi s n

mm Name of Sample	mm n Sealing	kn95	mm Sample number
mm Trade mark	mm Commissioned by	mm 委托方地址 Address of client	mm Types, Specification
mm Inspected unit	mm Sampling unit	mm 生产日期 Date of manufacture	mm Testing Purpose
mm 抽样地点 Location of sampling	mm 抽样日期 Sampling date	mm 抽样基数 Basic quantity of sampling	mm 产品编号/型号 Product No./batch No
mm Date of sampling	mm 收货日期 Date of receiving	mm 检验地点 Location of testing	mm Commission
mm 检测依据 Testing reference	mm 判定依据 Judging reference	mm 有效期 Validity	mm 生产日期 Date of manufacturing
mm 检测结论 Remarks	mm 检测结果 Test results	mm Test results are attached as below.	mm 有效期 Validity
mm 检测结论 Remarks	mm 检测结果 Test results	mm Test results are attached as below.	mm 有效期 Validity

14

IMS!mm

Test Report

(Electronic feisicii)

Verification Website: www.gtgc.net.cn

Verification Code: OEST-0641-04

No:20R002928

Issue Date: 2020-06-09

Applicant: GUANGZHOU NAN QI XING NONWOVEN CO., LTD

Address: NORTH SIDE OF BEIERHUAN ROAD, WENBIAN VILLAGE, SHIJI TOWN, PANYU DISTRICT, GUANGZHOU

Information confirmed by applicant:

Filtering half mask

Quantity: 100 pieces

Manufacture's name: GUANGZHOU NAN QI XING NONWOVEN CO., LTD

Standard Adopted:

Client Requirement

Date Received/Date Test Started: 2020-06-03

Conclusion:

Penetration test against solid particulates

M

Note: "M"-Meet the standard's requirement "F"-Fail to meet the standard's requirement "—" -No comment

Remark:

Penetration test against solid particulates was judged by 42 CFR Part 84 <Approval of respiratory protective devices> as per client's requirement.

All the tested items are tested under the standard condition (except for indication).

Copies of the report are valid only re-stamped.

The experiment was carried out at No.1, Zhujiang Road, Panyu District, Guangzhou, Guangdong, P.R.China.

Approved By:

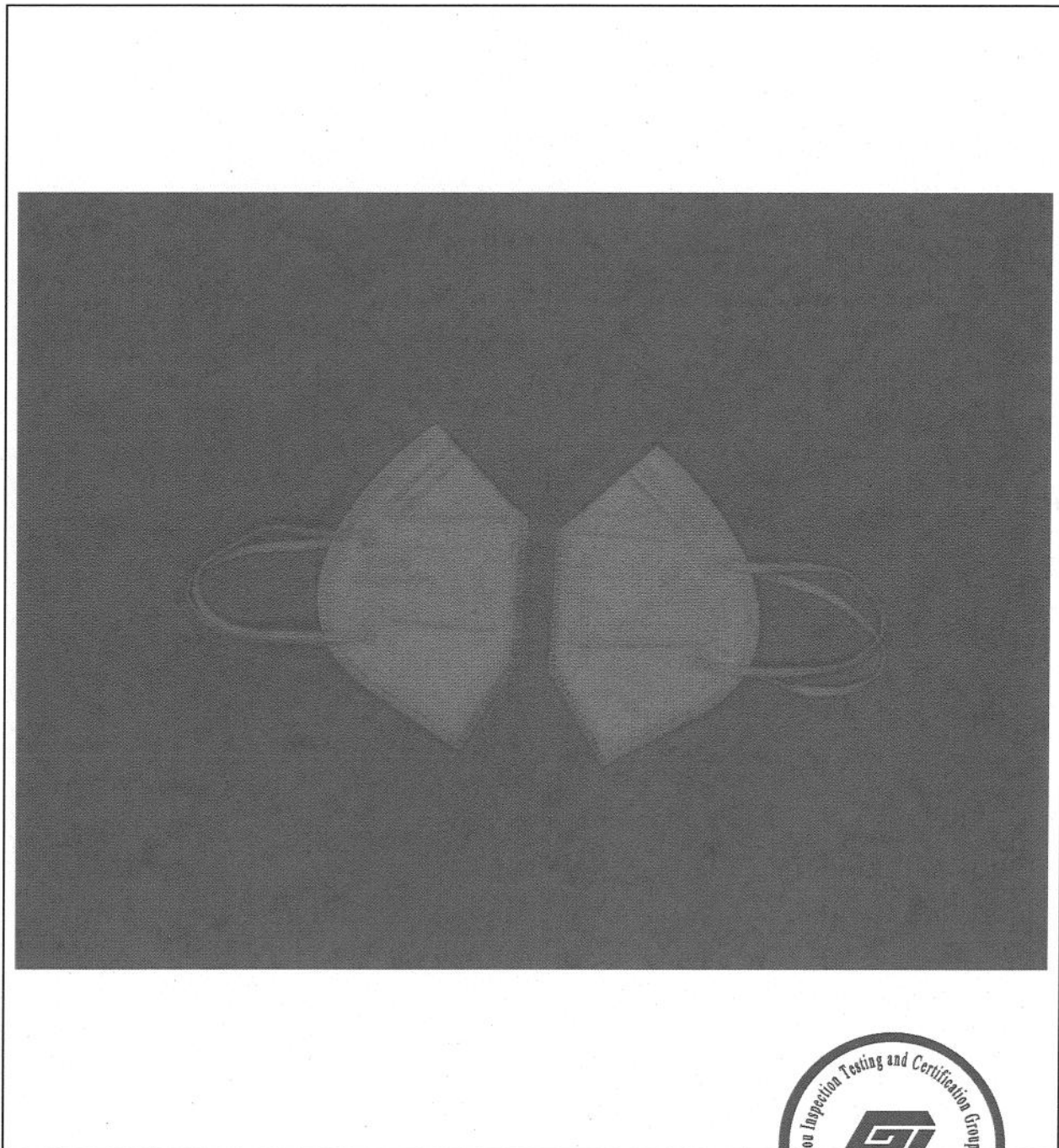
ZiShan Guo Senior Engineer*2.1skm Gtte*

Page 1 of 4

Test Report

(Electronic version)

No: 20R002928



RFB-RC-2020-089 PPE
BID ITEM 1-24 KN95 MASKS
FDA CERTIFICATIONS - GLOBAL CIRCLE GROUP LLC

Test Report

(Electric version)

No: 20R002928

Penetration test against solid particulates

Test Method: NIOSH Procedure No.RCT-APR-STP-0057,0058,0059

General:

This STP describes the test method to be used for the determination of particulate filter efficiency level for N95 series filters against solid particulates for non-powered, Air-purifying respirators test in sufficient detail to allow a person knowledgeable in the appropriate technical field to conduct the test and determine whether or not the product meets the established requirements.

Test equipment:

TSI Model 8130 automated filter tester

The environmental conditions of the laboratory and test condition:

Pretreatment: Temperature and humidity chamber capable of maintaining $(38 \pm 2.5)^\circ\text{C}$ and $(85 \pm 5)\%$ relative humidity for $(25 \pm 1)\text{h}$

Test environment temperature: 23.0°C , relative humidity: 36.1%

Flow rate: 85L/min

Aerosol particle: NaCl

Count median diameter: 0.075 μm

RFB-RC-2020-089 PPE
BID ITEM 1-24 KN95 MASKS
FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC



Test Report

(Electronic version)

No: 20R002928

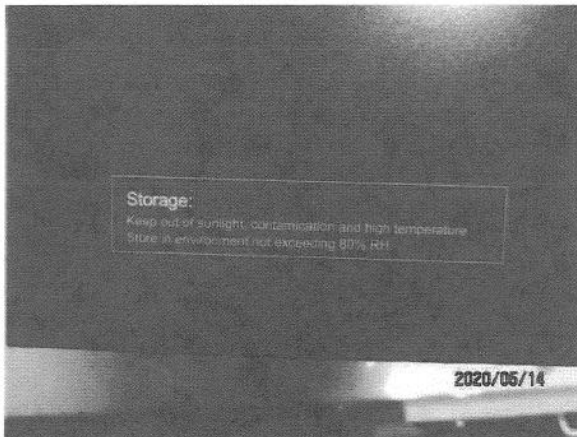
Results:

Sample	Filter efficiency (%)	Requirement (%)	Conclusion
1	99.839	<p>≥95</p> <p>Client Requirement</p>	Pass
2	99.784		
3	99.817		
4	99.852		
5	99.704		
6	99.775		
7	99.621		
8	99.798		
9	99.549		
10	99.605		
11	99.698		
12	99.712		
13	99.663		
14	99.574		
15	99.490		
16	99.875		
17	99.812		
18	99.598		
19	99.647		
20	99.682		

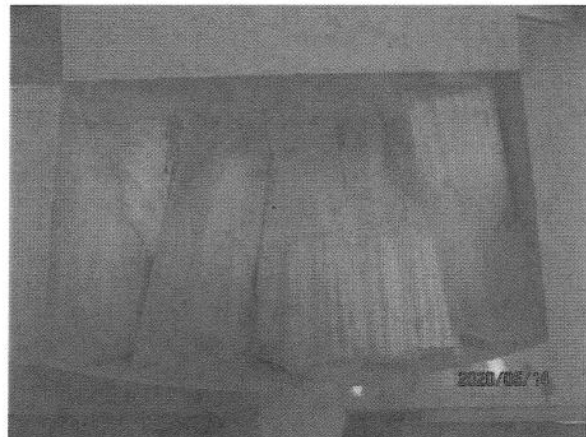
RFB-RC-2020-089 PPE
BID ITEM 1-24 KN95 MASKS
FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC



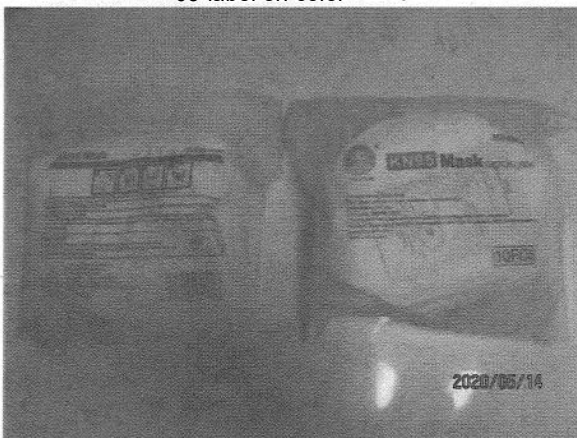
—End of Report—



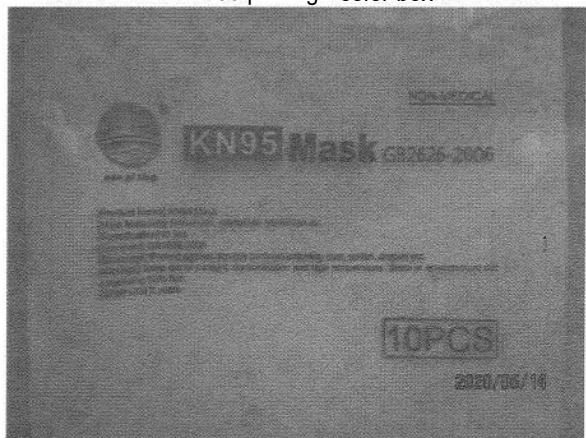
05 label on color box-6



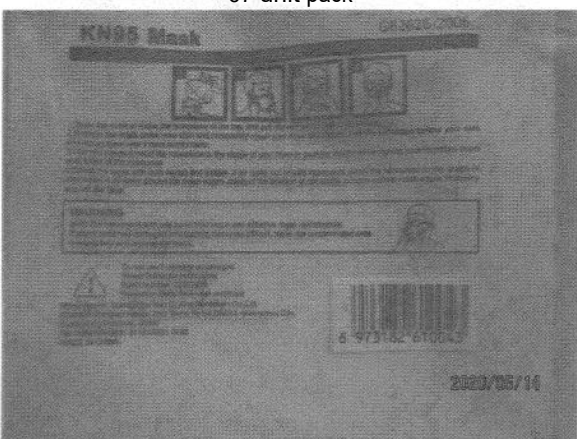
06 package-color box



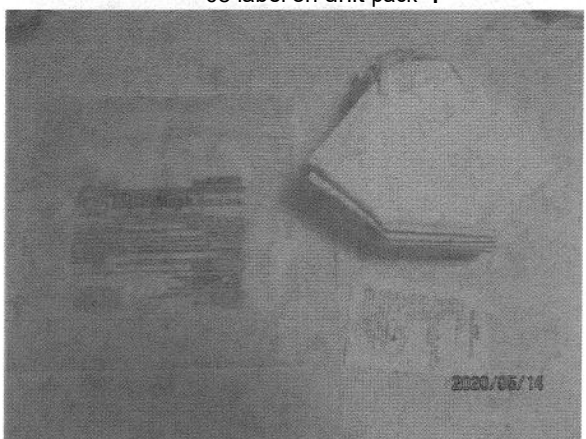
07 unit pack



08 label on unit pack-1



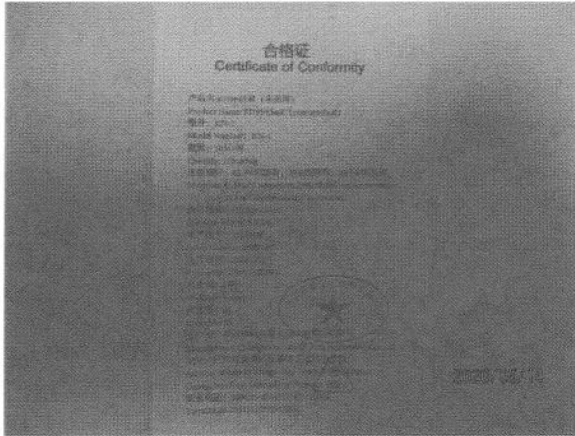
08 label on unit pack-2



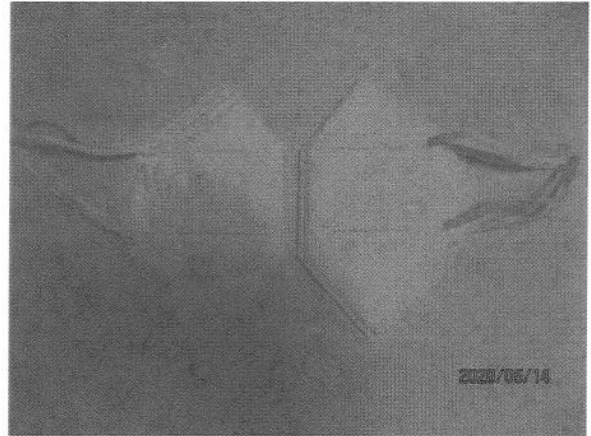
09 package-unit pack

RFB-RC-2020-089 PPE
BID ITEM 1-24 KN95 MASKS
FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC

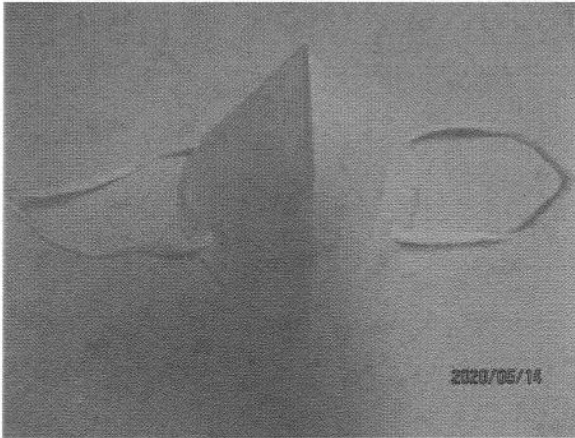




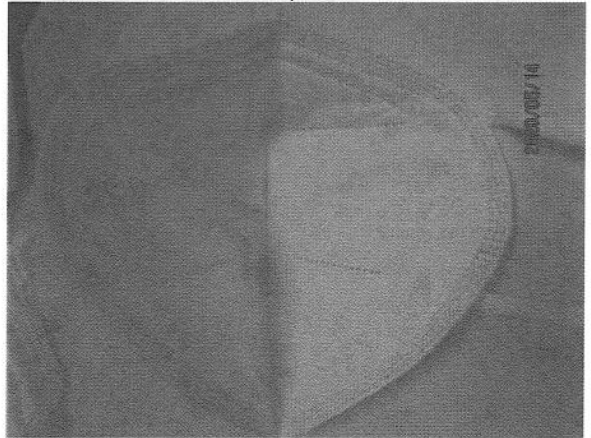
10 inner sheet



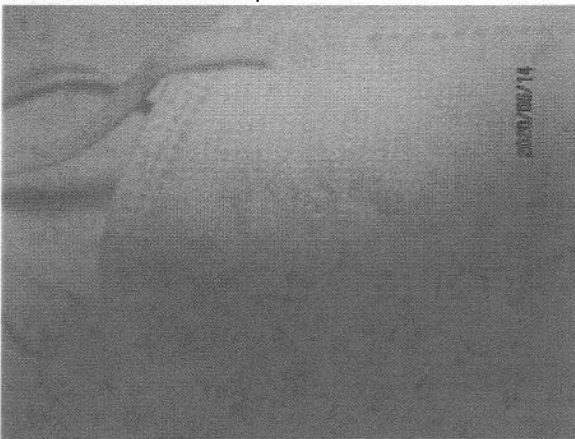
11 product-1



11 product-2



11 product-3



12 marking on product

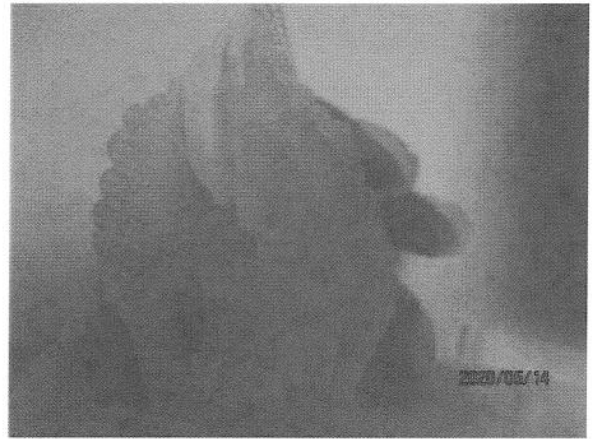


13 barcode check-1

RFB-RC-2020-089 PPE
BID ITEM 1-24 KN95 MASKS
FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC



13 barcode check-2



14 construction check
N/A



15 tension check for mask band

N/A

RFB-RC-2020-089 PPE
BID ITEM 1-24 KN95 MASKS
FDA CERTIFICATIONS - GOLDEN CIRCLE GROUP LLC

As part of the government's continuous quality assessment of these respirators, the FDA, working with CDC NIOSH, conducted additional assessments and found that some of the respirators authorized under the April 3, 2020 EUA did not meet the expected performance standards. In response, the FDA revised and reissued the EUA on May 7, 2020 to among other revisions, revise the third criterion for eligibility. Additionally, the FDA, in collaboration with CDC NIOSH, is increasing surveillance and sampling of all respirators imported from China - all respirator shipments from China that come into the U.S. will be subject to random sampling and testing by CDC NIOSH to determine whether the respirator meets the expected particulate filtration standards.

On June 6, 2020 the FDA further revised the Scope of Authorization of this EUA, to among other changes, further revise the third eligibility criterion, revise the second eligibility criterion, and remove decontaminated respirators from the scope of authorized products such that authorized respirators listed in Appendix A will no longer be authorized if they are decontaminated.

On October 15, 2020, the FDA reissued the Emergency Use Authorization (EUA) for Non-NIOSH-Approved Disposable Filtering Facepiece Respirators Manufactured in China to authorize for emergency use only those respirators listed in the EUA's Appendix A as of the date of this reissuance. Effective immediately, this EUA no longer includes the three eligibility criteria that were included in the previous June 6, 2020 authorization letter, meaning the FDA will no longer be reviewing requests and adding new respirator models to Appendix A based on those criteria. Further explanation of the revisions can be found in the Frequently Asked Questions (FAQs) about Non-NIOSH-Approved Filtering Facepiece Respirators (/medical-devices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/faqs-euas-non-niosh-approved-respirators-during-covid-19-pandemic).

- EUA Letter of Authorization - Umbrella EUA: Non-NIOSH-Approved Disposable Filtering Facepiece Respirators Manufactured in China (/media/136664/download) (Reissued October 15, 2020)
- Non-NIOSH Approved Disposable Filtering Facepiece Respirators Manufactured in China EUA FAQs (/medical-devices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/faqs-euas-non-niosh-approved-respirators-during-covid-19-pandemic)
- Appendix A: Authorized Respirators, Non-NIOSH Respirators Manufactured in China (Updated October 15, 2020)
- Respirator Models No Longer Authorized (Updated October 15, 2020)

Appendix A: Authorized Imported, Non-NIOSH Approved Respirators Manufactured in China (Updated: October 15, 2020)

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The table below includes a list of non-NIOSH respirators authorized by this Umbrella EUA (/media/i36664/download) for emergency use during the COVID-19 public health emergency.

As stated in the EUA, authorized respirators should be used in accordance with CDC's recommendations. For the most current CDC recommendations on optimizing respirator use, please visit CDC's webpage: Strategies for Optimizing the Supply of N95 Respirators (<https://www.cdc.gov/coronavirus/2019-ncov/hcp/respirators-strategy/index.html>).

Search:

Show entries

Manufacturer	Respirator		Instructions for Use
	Model(s)		
GUANGZHOU BIOFIL AIR PURIFICATION MATERIALS CO.,LTD	MY3D2		<ul style="list-style-type: none">IFU (http://www.baifeier.com.cn/en/article-28481-41890.html) D? (http://www.fda.gov/about-fda/website-policies/website-disclaimer)
Guangzhou Bofeite Safety Protective Supplies Co., Ltd.	HT9510V, HT9510		
Guangzhou Carrot Mall Network Technologies Co., Ltd.	IRYS-01		<ul style="list-style-type: none">IFU (http://www.ivrou.com/images/Intended_Use_and%200ther_Instructions_of_IVROU.pdf) (2? (http://www.fda.gov/about-fda/website-policies/website-disclaimer)
Guangzhou Harley Commodity Company Limited	L-103V KN95		
Guangzhou Nan Qi Xing Non-Woven Co., Ltd.	KN-1 Respirator		<ul style="list-style-type: none">IFU (https://www.gznqx.com/china-kn95-china-masks-customized-nanqixing) GT (http://www.fda.gov/about-fda/website-policies/website-disclaimer)
Guangzhou Powecom Labor Insurance Supplies Co., LTD	KN95		<ul style="list-style-type: none">IFU (http://www.powecom.com/eng_product.html) C (http://www.fda.gov/about-fda/website-policies/website-disclaimer)



5979 VALLEY VIEW BLVD, LAS VEGAS, NV 89118, (702) 790-7264 | GLOBALSANITIZERS.COM

Issued 5/20/2020
Revision: 1.0

SAFETY DATA SHEET

1. IDENTIFICATION

Product Identifier: Hand Sanitizer Gel
Supplier: Global Sanitizer Technologies
5979 Valley View Blvd
Las Vegas, NV 89118
(702) 790-7264
Contact Person: Safety Manager
Emergency Contact: 1-800-424-9300 (CHEMTREC USA)



2. HAZARD(S) IDENTIFICATION

Physical Hazard:	Flammable Liquids	Category 2
	Carcinogenicity	Category 2
	Aspiration Hazard	Category 1
	Specific Target Organ Toxicity (repeated exposure)	Category 2
	Specific Target Organ Toxicity (single exposure)	Category 3
	Skin Irritation	Category 3
	Eye Irritation	Category 2A
	Chronic Aquatic Toxicity	Category 2

Pictograms



Signal Word: DANGER

Hazard Statement: Highly flammable liquid and vapor.
May be fatal if swallowed and enters airways.
Harmful if inhaled.
May cause cancer.
Suspected of damaging fertility or the unborn child.
May cause damage to organs.

Precautionary Statements

General: Read and understand label before use.
Keep out of reach of children.
Have SDS in hand if medical advice is required.

Prevention:

Wear eye and/or face protection.
Wear protective gloves.
Keep away from heat/sparks/open flames/hot surfaces.
No smoking.
Use explosion proof electrical, material handling equipment and tools.
Avoid and guard against static discharge.
Ground/Bond container and receiving equipment.

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Response: IF SWALLOWED - Do not induce vomiting / seek medical attention
 IF ON SKIN/HAIR - Rinse with water or shower
 IF ON CLOTHING - Remove contaminated clothing / Rinse effected area or shower
 IN CASE OF FIRE - Use alcohol resistant foam, carbon dioxide, dry powder or water fog

Storage: Store in well ventilated space, keep cool

Disposal: Dispose of contents and container in accordance with all local/regional/national/international regulations

Hazard(s) not otherwise classified: None known

3. COMPOSITION / INFORMATION ON INGREDIENTS

Chemical Name	Common Name and Synonyms	CAS Number	% (v/v)
Ethanol	Alcohol, Ethyl Alcohol	64-17-5	70%
Water	Water, Aqua	7732-18-5	29.37%
Glycerin	Vegetable Glycerin	56-81-5	1%
Hydroxyethyl Cellulose	Hydroxyethylcellulose	102-71-6	<1%
Triethanolamine	Triethanolamine, Trolamine	9004-62-0	<1%
Aloe Vera Extract	Aloe Vera	85507-69-3	<1%
Citric Acid	Citric Acid, 2-Hydroxy-1	77-92-9	<1%

4. FIRST AID MEASURES

Eyes: Immediately flush eyes for 15 minutes
 Vapor exposures 1,000 ppm to 10,000 ppm may cause temporary irritation.
 Continuous tear in occurs at levels greater than 15,000 ppm.
 Direct eye contact causes moderate to sever irritation.

Ingestion: Do NOT induce vomiting, rinse mouth, get medical attention / advice immediately.
 First acts as a stimulant, but increased volume can produce stupor
 May cause irritation to the gastrointestinal tract with nausea, vomiting and abdominal pain.
 May cause headaches, tremors, fatigue, central nervous system depression, narcosis, or coma

Inhalation: Move to fresh air, IF NOT BREATHING - admister artifical respiration
 Difficulty breathing - administer oxygen, seek medical attention
 Excessive inhalation is irritating to the eyes and upper respiratory tract and may cause symptoms of intoxication
 Aspiration into lungs may cause pulmonary edema and chemical pneumonitis. May also cause unconsciousness, coma, respiratory failure or death.
 Recovery from inhalation of concentrations less than 10,000 ppm for brief periods occurs in a few minutes.

Skin Contact: Wash area with soap and water for 15 minutes
 Remove affected clothing and wash before wearing again
 Seek medical attention if irritation develops
 May cause redness and/or a mild burning sensation of the skin with acute exposure to the liquid
 Remove natural oils and fats from skin resulting in dermatitis

Most important symptoms or effects and any symptoms that are acute or delayed: Irritation of eyes, nose and throat
 Unconsciousness
 Decreased motor function
 Skin irritation
 Respiratory failure

5. FIRE FIGHTING MEASURES

Suitable Extinguishing Equipment:	Carbon Dioxide Polar Solvent Foam Alcohol Resistant Foam Dry Chemical Extinguishers Large Quantities of Dilluge Water
Non-Suitable Extinguishing Equipment:	Ordinary Foam Water spray may be suitable for small fires, but not large fires Avoid solid water stream, as this may scatter the media and spread the fire
Specific Hazards that may develop from the chemical during the fire:	Highly Flammable Explosive vapor / air mixtures may form Sensitive to static shock Vapor may carry and ignite by distant source Vapor may cause flash fire
Recommendations on special protective equipment or precautions for firefighters:	Self-contained breathing apparatus Protective clothing Face mask Evacuate if loud sound emitting from vents Water spray may be used on unopened containers to keep cool

6. ACCIDENTAL RELEASE MEASURES

Personal precautions and protective equipment to prevent contamination of skin, eyes and clothing:	Ventilate area Remove sources of ignition Do not handle spill without proper Personal Protective Equipment (PPE) Stay out of low areas or confined space Use explosion proof equipment and non-sparking tools
Emergency procedures for evacuation:	Isolate Hazard Area Remove all non essential personnel Ensure ignition sources are removed or isolated Refer to Section 8 for Personal Protective Equipment (PPE)
Methods and materials used for containment:	If possible, stop leak SMALL SPILLS - Absorb spill with sand or other absorbant material and place container for proper disposal LARGE SPILLS - Dike around spills to avoid entry into sewers, water system, ground Absorb with appropriate material

7. HANDLING AND STORAGE

Precautions for safe handling, recommendations for incompatible chemicals, minimizing release of the chemical into the environment, advice on general hygiene practices:	Wear Personal Protective Clothing Avoid contact with skin Avoid breathing in fumes / vapors Avoid contact with eyes Ground / bond container and equipment Use explosion proof equipment Use non-sparking tools
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Recommendations on conditions for safe storage, including incompatibilities. Advice on specific storage requirements:

Store in cool areas
Ensure container are closed tightly
Provide adequate ventilation
Store away from open flame
Do not store near oxidizing agents

8. EXPOSURE CONTROLS / PERSONAL PROTECTIVE EQUIPMENT

OSHA Permissible Exposure Limits (PELs):

Chemical Name	CAS#	OSHA PEL	ACGIH TLV
Ethanol	64-17-5	1000 ppm, 1800 mg/m3	1000 ppm, 1800 mg/m3

Engineering Controls:

Adequate general local exhaust ventilation shall be provided. Refer to ACGIH Industrial Ventilation

Personal Protection Equipment Recommendations:

EYES - Safety glasses. Opportunity for splash-wear goggles and / or face shield
SKIN - Wear chemical resistant gloves. Impervious boots, apron, or coveralls as needed in areas of unusual exposure.

Special PPE Requirements:

Gloves - PVC or Neoprene
Clothing - Fire Rated if exposure deems necessary
NIOSH approved respirators are required if exposure limits are exceeded.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance:	Colorless liquid
Upper/Lower Flammability or Explosive Limits:	Upper - 3.3% / Lower 19.0%
Odor:	Ethanol
Vapor Pressure @ 20°C:	59.3
Vapor Density @ 78°C:	1.59
Odor Threshold:	<1 ppm
pH:	Neutral
Relative Density @ 20°C:	790.0 kg/m3
Melting Point:	< -173°F
Freezing Point:	< -113.89°C
Solubility:	Water (slight)
Initial Boiling Point:	173°F
Flash Point:	50-55°F (Closed cup)
Evaporation Rate:	Varies with conditions - Rapid
Flammability (Solid, Gas):	Flammable Liquid and Vapor
Partition Coefficient n-Octanol/Water:	Not Available
Auto Ignition Temperature:	363°C
Decomposition Temperature:	Not Available
Viscosity:	Not Available

10. STABILITY AND REACTIVITY

Reactivity:	Not Available
Chemical Stability:	Stable under normal / intended use
Possibility of Hazardous Reactions:	Hydrogen Peroxide (Fire), Strong Oxidizers (Explosion or violent reaction)
Conditions to Avoid:	Heat, Spark, Open Flame, Static Electricity
Incompatible Materials:	Oxidizers, Peroxides
Hazardous Decomposition:	Carbon Monoxide, Oxides of Nitrogen

11. TOXICOLOGY INFORMATION

Routes of Exposure:	INHALATION: May cause irritation to nose, throat, lungs. May cause headaches. INGESTION: Irritation of mouth, throat, and stomach SKIN CONTACT: May cause dermatitis EYE CONTACT: Irritation
Chronic Effects:	DELAYED: Defating of skin, anemia, leukemia, low white blood cell count, sever degeneration of the peripheral nervous system, lung disfunction, pneumatocele formation IMMEDIATE: Nervousness, fatigue, nausea, labored breathing, blurred vision, irritation of nose, throat and mucous membrane
Numerical Measures of Toxicity:	ACUTE TOXICITY - May cause lung damage if swallowed. Oral Rat - LD50: 7060mg/kg = adjusted LD 25807 ppm, derived value 2581 ppm Inhalation Rat - LC50: 20363 ppm = adjusted .5 hr 40727 ppm, derived value 4073 ppm
Symptoms (Lowest to Most Severe):	1) Irritation of eyes, nose, throat, lungs, skin. May cause headaches. 2) Vomiting, Nausea, blurred vision, fatigue 3) Unconsciousness, corneal damage, narcosis, cyanosis 4) Could be fatal if ingested into airways
Chemical Listings:	Not Available

12. ECOLOGICAL INFORMATION

Ecotoxicity:	Fathead Minnow	LC50 = 14,765714	Not Toxic
	Clawed Toad	EC50 = 10,513,200	Not Toxic

Persistence and Degradability:	Degradation is expected to occur rapidly. Unlikely to be persistent in nature.
Bioaccumulative Potential:	Readily Biodegradable
Mobility in Soil:	Very High
Other Adverse Effects:	Not Available

13. DISPOSAL CONSIDERATIONS

Appropriate Disposal Containers:	Sealable containers in accordance with applicable local, regional, national and international regulations.
Disposal Methods:	EPA RCRA (40 CFR 261.21) Flash point below 140°F - Ignitable Hazardous Waste - Waste Code #D001
Waterways:	Do not allow chemical or container to enter waterways or ditches
Landfills and/or Incineration:	Incinerate material under controlled conditions in approved incinerator

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14. TRANSPORT INFORMATION

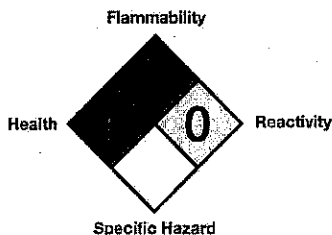
UN Number: UN 1170
UN Proper Shipping Name: Ethanol
Transport Hazard Class: 3
Packing Group: II
Environmental Hazards: Maritime Transport (Not recognized as marine pollutant by Department of Transportation 49 CFR 172.101)
Transport in Bulk: Follow Federal, State and local regulations
Special Precautions: DANGER. Follow safety instructions, SDS and emergency procedures.

15. REGULATORY INFORMATION

OSHA: OSHA Hazard Communication Standard 29 CFR 1910.1200
All components are on the United States of America EPA TSCA Inventory List
SARA 302: Not Listed
SARA 311/312: Not Listed
SARA 313: Not Regulated

16. OTHER INFORMATION

NFPA:
4 Extreme
3 Serious
2 Moderate
1 Slight
0 Minimal



ISSUE DATE: 4/14/2020
REVISION: 1.0

The information above is believed to be accurate and represents the best information currently available to us. However, we make no warranty of merchantability or any other warranty, expressed or implied, with respect to such information, and we assume no liability resulting from its use. Users are responsible to make their own investigation to determine the suitability of the information for their particular purposes or use.

In no event shall Global Sanitizer Technologies be liable for any claims, losses, or damages of any third party or for lost profits or any special, indirect, incidental, consequential, or exemplary damages, howsoever arising, even if Global Sanitizer Technologies, Inc. has been advised of the possibility of such damages.

Viral Penetration ASTM Method F 1671 Final Report

Test Article: BLUE CO-POLY GOWN PRODUCT #8888K2
 Study Number: 1098183-S01
 Study Received Date: 17 Sep 2018
 Testing Facility: Nelson Laboratories, LLC
 6280 S. Redwood Rd.
 Salt Lake City, UT 84123 U.S.A.
 Test Procedure(s): Standard Test Protocol (STP) Number: STP0062 Rev 16
 Deviation(s): None

Summary: This test method was performed to evaluate the barrier performance of protective materials which are intended to protect against blood borne pathogen hazards. Test articles were conditioned for a minimum of 24 hours at $21 \pm 5^\circ\text{C}$ and 30-80% relative humidity (RH), and then tested for viral penetration using a ΦX174 bacteriophage suspension. At the conclusion of the test, the observed side of the test article was rinsed with a sterile medium and assayed for the presence of ΦX174 bacteriophage. The viral penetration method complies with ASTM F1671; sampling was at the discretion of the sponsor. All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Number of Test Articles Tested: 1
 Number of Test Articles Passed: 1
 Test Article Side Tested: Outside
 Test Article Preparation: Cut from the Middle Portion of Gown at Random
 Exposure Procedure: B (Retaining Screen: Woven Polyester Mesh, with >50% Open Area)
 Compatibility Ratio: 1.0
 Environmental Plate Results: Acceptable

Results:

Test Article Number	Pre-Challenge Concentration (PFU/mL)	Post-Challenge Concentration (PFU/mL)	Assay Titer (PFU/mL)	Visual Penetration	Test Result
1	2.3×10^8	2.2×10^8	$<1^a$	None Seen	Pass
Negative Control	2.3×10^8	2.2×10^8	$<1^a$	None Seen	Acceptable
Positive Control	2.3×10^8	2.2×10^8	TNTC ^b	Yes	Acceptable

^a A value of <1 plaque forming unit (PFU)/mL is reported for assay plates showing no plaques.

^b TNTC = PFUs were too numerous to count.



Study Director

Jennifer Jorgenson, B.S.

Study Completion Date



1098183-S01

801-290-7500

nelsonlabs.com

sales@nelsonlabs.com

nrg

FRT0062-0001 Rev 9

Page 1 of 1

ASTM Method F 1670 Synthetic Blood Penetration Final Report

Test Article: BLUE CO-POLY GOWN PRODUCT #8888K2
Purchase Order: 913188
Study Number: 1098184-S01
Study Received Date: 17 Sep 2018
Testing Facility: Nelson Laboratories, LLC
6280 S. Redwood Rd.
Salt Lake City, UT 84123 U.S.A.
Test Procedure(s): Standard Test Protocol (STP) Number: STP0061 Rev 09
Deviation(s): None

Summary: This test method was performed to evaluate the resistance of protective materials to penetration by synthetic blood under conditions of continuous liquid contact. Protective materials' pass/fail determinations are based on visual detection of synthetic blood penetration. Test articles were conditioned for a minimum of 24 hours at $21 \pm 5^{\circ}\text{C}$ and 30-80% relative humidity (RH) and then tested for liquid penetration using synthetic blood. The synthetic blood penetration method complies with ASTM F1670; sampling was at the discretion of the sponsor. All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Number of Test Articles Tested: 1
Number of Test Articles Passed: 1
Test Article Side Tested: Outside
Test Article Preparation: Cut from Middle Portion of the Gown
Exposure Procedure: B (Retaining Screen: Woven Polyester Mesh, with >50% Open Area)

Results:

Test Article Number	Synthetic Blood Penetration	Result
1	None Seen	Pass
Negative Control	None Seen	Acceptable
Positive Control	Yes	Acceptable

Study Director

Jennifer Jorgenson, B.S.

26 Sep 2018
Study Completion Date

1098184-S01