



All Lakeland® CleanMax® Apparel is:

- IEST-RP-CC003 Category I Particle Cleanliness
- Latex and silicone-free
- Compatible with ISO Class 4-8 cleanrooms and all controlled environments
- Individually-packaged and protective outer bag for ante areas
- Resistant to viral, blood, and body fluid penetration
- Resistant to blood borne pathogens
- Chemical penetration resistance to oils and bleach



Bound Seams

CleanMax® features bound seams, which are precisely sewn with an additional outer binding. This increases seam strength and provides a better barrier from particulates than simple serged seams.

Tunneled Elastic Wrists

CleanMax® features tunneled elastic for better sealing properties.

Lakeland Industries has spent over 40 years as a leader in protecting people, products, and environments. Our CleanMax® products are composed of a high-quality microporous laminate material that is lightweight, breathable, and impervious to liquids, harsh chemicals, and microorganisms.

Both sterile and non-sterile configurations of CleanMax® meet IEST-RP-C003 Category 1 particulate cleanliness standards and are ready for immediate use in ISO Class4-8 cleanrooms.

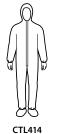
All sterile configurations are gamma radiation sterilized to a 10-6 Sterility Assurance Level. These garments provide excellent comfort as well as superior protection, so you can easily don and doff your garments to reduce excursions and risk of contamination.

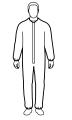


CleanMax® Configurations



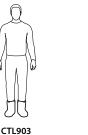
CTL191













CTL850

Clean Manufactured Non-Sterile Garments

Frock - CTL191CM

- · Mandarin collar Zipper closure
- No pockets
- Tunneled elastic wrists with thumb loops Sizes: M – 5X Case Pack: 30

Clean Sterile

included

Coverall – CTL414CM Zipper closure

- Attached hood and boots
- Tunneled elastic on wrists (with thumb loops), and back half of waist

Sizes: M – 5X Case Pack: 25 Clean Manufactured Sterile Garments STERILE R Irradiation Sterilization

Coverall – CTL414CS

Coverall – CTL417CM

CTL417

- Zipper closure Tunneled elastic on wrists (with thumb loops), ankles, and back half of waist
- Sizes: M 5X Case Pack: 25

Coverall – CTL428CM Hood - CTL713CM · Zipper closure Covers shoulders

 Attached hood Tunneled elastic on wrists (with thumb loops), ankles, and back half of waist

Sizes: M - 5X Case Pack: 25

Boot Cover -CTL903CMP

· Ties to customize fit • 19" high Case Pack: 100 Non-skid Vinyl sole

Sleeve CTL850CMP-18

- Tunneled elastic top Bound seams
 - Tunneled elastic Thumb loops
- Sizes: S/M, L/XL, 2X Size: 18" length Case Pack: 50 pair Case Pack: 50 pair

- Zipper closure Tunneled elastic on
- boots Tunneled elastic on wrists (with thumb loops), and back half

of waist

Coverall – CTL417CS

- Zipper closure Certificate of Radiation Attached hood and
- Gamma radiation indicator dots on each İPA resistant ink

Sizes: M - 5X Case Pack: 25

Coverall – CTL428CS

- Zipper closure wrists (with thumb loops), ankles, and back half of waist
- Sizes: M 5X Case Pack: 25

Hood - CTL713CS Covers shoulders One size

One size

- Attached hood Tunneled elastic on wrists (with thumb
- loops), ankles, and back half of waist Sizes: M - 5X Case Pack: 25

Boot Cover -CTL903CSP Tunneled elastic ton

- · Ties to customize fit • 19" high Case Pack: 100 • Non-skid Vinyl sole
 - Sizes: S/M, L/XL, 2X Case Pack: 50 pair

Sleeve CTL850CSP-18

- Bound seams
- Tunneled elastic Thumb loops
- Size: 18" length Case Pack: 50 pair

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Disposable Cleanroom Suits: Tips for Cleanroom Apparel Selection

Confidence in your cleanroom starts with understanding how to select the right disposable apparel for your unique needs. In just a few minutes, our team of cleanroom industry experts will work with you to determine the type of garment required for your application and environment, and discuss how we can help you protect your team effectively with clean-manufactured garments.

Applications for CleanMax® Cleanroom Apparel CleanMax® Clean Manufactured Sterile

- Aseptic or Terminally Sterile Cleanroom Environments
- ISO Class 4-8 Cleanroom
- Sterility assurance level of 10⁻⁶ SAL

CleanMax® Clean Manufactured Non-Sterile

• ISO Class 4-8 or below Non-Aseptic Cleanrooms or **Controlled Environments**

Garment Configurations

Apparel	ISO 8	ISO 7	ISO 6	ISO 5 Non-Sterile	ISO 5 Sterile (Aseptic)	ISO 4	ISO 3	ISO 1 & 2
Hair cover	R	R	R	R	R	R	R	AS
Barrier gloves	AS	AS	AS	AS	R	R	R	R
Facial cover	AS	AS	AS	R	R	R	R	AS
Hood	AS	AS	AS	R	R	R	R	AS
Frock	R	R	AS	AS	NR	NR	NR	NR
Coverall	AS	AS	R	R	R	R	R	R
Shoe cover	R	R	AS	AS	NR	NR	NR	NR
Boot	AS	AS	R	R	R	R	R	R
Typical Frequency of Change*	2X/week	2X/week	3X/week	1X/day	Per Entry	Per Entry	Per Entry	Per Entry

Chart shows Lakeland® garments relevant to ISO 5. Recommendations from IEST-RP-CC003. **R** = Recommended, **NR** = Not Recommended, **AS** = Application Specific

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CleanMax[®] Features and Benefits

Clean Manufacturing Difference

Clean manufacturing helps ensure that operators and PPE are not adding particulates to the environment.

Smooth Storm Flaps

Covering the zipper further helps protect the critical chest and front area of the coverall from potential particle breakthrough.

Secure Thumb Loops

Elastic wrists with thumb loops help secure coveralls and frocks in place to prevent potential skin exposure during wear.

Chemical Penetration Resistance

CleanMax® offers chemical penetration resistance to oils, bleach, and other chemicals.

Premium Packaging for Peace of Mind

CleanMax® configurations are individually packaged and expertly folded to prevent excessive wrinkling and the potential for excessive excursions.



Bound Seams Throughout

CleanMax® features bound seams, which are precisely sewn with an additional outer binding, to provide a better barrier from breakthrough and protection from strike-through.

Smooth Surface Area

CleanMax® is smoother than other brands, meaning particulates are less likely to harbor on the garment's surface.

Cuffed Ankles for Efficient Donning

The cuffed ankle on CleanMax® provides six inches of freedom when you step into your coverall.



Download the **Disposable Cleanroom Garment Guide**



CleanMax® Physical Properties

Physical Property	Test Method	Units	CleanMax [®] Sterile Test Results
Basis Weight	ASTM D3776	oz/y²	1.55 oz/y²
Grab Tensile MD	ASTM D5034	lbs.	14.07 lbs.
Grab Tensile XD	ASTM D5034	lbs.	8.72 lbs.
Trapezoidal Tear MD	ASTM D1117	lbs.	8.8 lbs.
Trapezoidal Tear CD	ASTM D1117	lbs.	5.0 lbs.
Ball Burst	ASTM D3787	lbs.	19.0 lbs.
Air Permeability	ASTM D737	cfm	<0.562 cfm/ft ²
Water Vapor Transmission	ASTM 96-80	g/m2-24hrs	663.38
Bacterial Filtration Efficiency	ASTM F2101	%	99.999%
Particle Filtration Efficiency	ASTM F2299	%	99.999%

CleanMax® Penetration and Resistance Properties

Physical Property	Test Method	Units	Test Results
Synthetic Blood Penetration	ASTM F 1670	Time to Penetration (> 60 minutes)	Pass
Viral Penetration Resistance	ASTM F 1671	Time to Penetration (> 60 minutes)	Pass
Resistance to Penetration by Blood and Bodily Fluids using Synthetic Blood	ISO 16603	Pressure in kPa	Pass –no strike-through at 20kPa
Resistance to Penetration by Blood Borne Pathogens	ISO 16604	Pressure in kPa	Pass –no strike-through at 20kPa
Resistance to Permeation of Chemotherapy Drugs	ASTM D6978	Minimum Breakthrough Time >240 minutes	Pass*

^{*}Tested drugs include Cisplatin, Cyclophosphamide, Cyclosporin A, Doxorubicin Hydrochloride, Etosposide (Toposar), Flourouracil, Methotrexate, Mitomycin C, Paclitaxel

CE Testing

CleanMax® Physical Properties – CE Test Data

Physical Property	Test Method	CE Class
Abrasion Resistance	EN 530 method 2	Class 2
Flex Cracking	ISO 7854 method B	Class 4
Trapezoidal Tear (MD/CD)	ISO 9073-4	3/2
Tensile Strength (MD/CD)	ISO 13934-1	2/1
Puncture Resistance	EN 863	Class 1
Seam Strength	ISO 13935-2	3

CleanMax® Resistance to Penetration by Infectious Agents – CE Test Data

Physical Property	Test Method	CE Class
Resistance to Penetration by Blood Borne Pathogens	ISO 16604	Class 6 of 6
Resistance to Biologically-Contaminated Aerosols	ISO 22611	Class 3 of 3
Resistance to Dry Microbial Contact	ISO 22612	Class 3 of 3
Resistance to Wet Bacterial Penetration	EN 14126 Annex A / ISO 22610	Class 6 of 6











CleanMax® Bulk Packaging!

Bulk packaging in a single overlay bag is available for CleanMax® non-sterile frocks, coveralls, boots, and hoods.

Does Your Cleanroom Apparel Meet Current IEST Standards?

Download our free white paper to find out:



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Lakeland's general purpose isolation gowns are constructed from spun bonded polypropylene fabric. Designed with the wearer's comfort in mind, this gown is both lightweight and breathable while offering protection from dirt, dry particulates, and light splashes in non-hazardous environments.

Key features and benefits:

- Spun bonded polypropylene
- · Lightweight, breathable, and comfortable fit
- · Wrap-around design with neck and waist ties for full coverage
- Self-certified as Category I PPE
- Latex and silicone free

Style Number: C2192IG

Sizes: Universal Fit (S-L) and XL

Case Pack: 100/Case (5 bags with 20 per case)



General Purpose Isolation Gown -C21921G

- Wrap around design Neck and Waist ties
- · Elastic wrists
- Sizes: Universal fit (S-L) Case Pack: 100



AAMI LEVEL 2 NON-SURGICAL CE CERTIFIED ISOLATION GOWN

For use when exposure to fluids is expected to be low to moderate

Lakeland's AAMI Level 2 Non-surgical CE Certified Isolation Gown is constructed from SMMS fabric. Designed with the end-user's comfort in mind, this non-surgical gown is both lightweight and breathable. Lakeland's Isolation Gown offers basic protection when exposure to fluids is expected to be low to moderate.



EN1149-5:2018 EN1149-1:2006

Key features and benefits:

- SMMS
- Lightweight, breathable, and comfortable fit
- Wrap around design with neck and waist ties for full coverage
- Elastic cuff at wrist
- Taped seams
- Latex and silicone free
- Non-sterile

Style Numbers: C8192TIG (Universal fit), C8192TIG-XL

Sizes: Universal Fit (S-L) and XL

Color: Blue

Case Pack: 100/Case (5 bags with 20 per case)



AAMI Level 2 **CE Certified**

- **Isolation Gown** C8192TIG Wrap around
- design Neck and Waist ties
- Elastic wrists
- Sizes: Universal fit (S-L) and XL Case Pack: 100

Liquid Barrier Performance Classification of Protective Apparel and Drapes Intended for Use in **Health Care Facilities - ANSI/AAMI PB70:2012**

	Test Method	Requirement	Lakeland AAMI Level 2 Isolation Gown Test Results	Anticipated Risk of Exposure
LEVEL 1				
Requirements at 4% AQL	AATCC 42:2017 "Impact Penetration"	≤ 4.5 g	Average Fabric - 0.0 g	Low
LEVEL 2				
Requirements at 4% AQL	AATCC 42:2017 "Impact Penetration"	≤ 1.0 g	Average Fabric - 0.0 g	Moderate
Requirements at 4% AQL	AATCC 127:2018 "Hydrostatic Pressure"	≥ 20 cm	Average Fabric - 59.09 cm	Moderate

Physical Properties

Physical Property	Test Method	CE Minimum Requirements	Lakeland Isolation Gown Test Results	CE Class According to EN 14325
Abrasion Resistance	EN530	>10	>100 <500	Class 2
Tear Resistance	ISO9073-4	MD/XD >10	MD 65N / XD41N	Class 3
Tensile Strength	ISO13934-1	MD/XD >30	MD 92N / XD 59N	Class 1
Puncture Resistance	EN863	>5N	8N	Class 1
Liquid Test - Penetration (%) Sulphuric Acid 30% and Sodium Hydroxide10%	ISO6530	<10%	0%	Class 3
Liquid Test - Repellency (%) Sulphuric Acid 30% and Sodium Hydroxide10%	ISO6530	>80%	>95%	Class 3

This gown should not be used in a surgical setting. For use when exposure to fluids is expected to be low to moderate. This gown should not be used in a clinical setting where Level 3 or 4 protection is warranted, or in any setting involving invasive procedures or where there is a high risk of contamination. This product has not been FDA cleared or approved. This product has been authorized by the FDA under the EUA for use as PPE in healthcare settings by healthcare personnel. This AAMI Level 2 Non-surgical isolation gown may help protect healthcare professionals and/or patients from the transfer of SARS-COV-2 virus in low or minimal risk level situation to prevent the spread of COVID-19. This product is only authorized for the duration of the declaration. This product should not be used in the presence of a high intensity heat source or flammable gas.

CleanMax® Sterility Documentation with Lakeland



Accessible 24/7 with a Simple Scan!

Why a QR Code?

As a cleanroom operator or quality assurance professional, accessing sterility documentation for your protective clothing should be simple.

Lakeland recognized the need for our customers to quickly find and store sterility documentation and developed a simple QR code system for our CleanMax® product.



How Does It Work?

The three-step approach includes:

- Locating the QR code on any CleanMax® garment label, packaging, or shipping box.
- Scanning the QR code with a smart phone or tablet.
- Viewing and printing or saving the sterility document associated with your lot number.

Where Can I Get a Sample?

You can request a sample of Lakeland CleanMax® by calling Customer Service at 800-645-9291 or emailing sales@lakeland.com. Our product specialists are standing by to help you find the right product for your application and would be happy to answer any questions you may have.

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Why Packaging Matters for Your Controlled Environment

Cleanroom PPE packaging is critical to the safety of your staff and the integrity of your controlled environment. It's not enough to have clean and sterile manufacturing processes, the packaging process must be designed to:

- Reduce excursions
- · Decrease gowning time and error
- Safeguard against contamination

Lakeland® CleanMax is packaged to ensure maximum garment protection

- Each package features an easy-open strip, allowing for less handling when removing the PPE from packaging.
- Garments are folded and left unzipped to allow for easier donning.
- Garments are individually packaged to maintain cleanliness and avoid excessive wrinkling.
- The packaging can be wiped down with IPA without the fear of the print smearing or creating cross contamination.
- Individually-packaged garments are placed in a sealed, heavy-duty outer bag to protect the integrity of the PPE and to facilitate staged entry into the gown room.
- Coveralls and frocks are positioned so that wearers can only grab the inside of the garment, right below the collar area, eliminating any contamination of the exterior surface.
- As the PPE is pulled out of the packaging, the folded piece will fall easily so that it's ready for the next step of the donning process.
- For customers with less critical environments, bulk packaging is available in a single overlay bag on CleanMax® Clean Manufactured Non-Sterile frocks, coveralls, boots, and hoods.

Availability is Essential

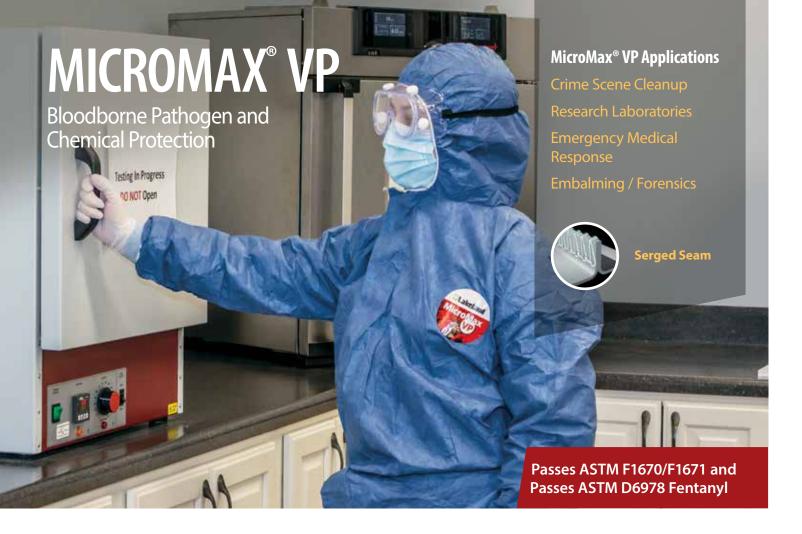
Product availability is critical for your controlled environment. It's unacceptable to lose valuable production hours because your supplier is unable to deliver PPE when you need it. Our team at Lakeland, along with our network of exceptional distributors, will help you find the right PPE for your application and ensure you get your order when you need it.





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Ideal for use in crime labs, for crime scene clean-up, and by emergency response personnel, MicroMax® VP is specifically designed to protect when the risk of blood, body fluids, blood borne pathogens, and viral contamination is greatest.

- Protective hood
- Seamless front reduces risk of contaminant exposure
- Taped storm flap protects zipper
- Elastic back for more comfortable fit
- Passes ASTM F1670/F1671 for Blood and Viral Protection

MicroMax® VP Fabric Physical Properties

Physical Property	Test Method	Units	Test Results
Material Thickness	ASTM D1777		15 mil
Material Weight	ASTM D3776		80 gsm
Tensile Strength MD	ASTM D5034	lbs.	36.30 lbs.
Tensile Strength CD	ASTM D5034	lbs.	24.15 lbs.
Elongation MD	ASTM D5034	%	59 Avg.
Elongation CD	ASTM D5034	%	71 Avg.
Water Vapor Transmis- sion Rate	ASTM E96		16 g/sq. meter/ 24 hrs. avg.
Bursting Strength Hydraulic Method	ISO 13938-1		29.4 psi avg.
Burn Test 45°	CPSC16 CFR 1610		Pass
Surface Resistance Requirement for BS EN1149-5:2008 is \leq 2.5 \times 10 ⁹ Ω .	EN1149-1	Ω	The test sample meets the requirement 2.4 X 10 ⁸

MicroMax® VP Fabric Liquid Penetration Data

Physical Property	Test Method	Test Results
Liquid Penetration Using Synthetic Blood	ASTM F1670	Pass
Viral Penetration using фX174 bacteriophage suspension	ASTM F1671	Pass

MicroMax® VP Fabric ASTM F903 **Liquid Penetration Data**

Physical Property	Test Method	Test Results
Methanol	ASTM F903	Pass
Ethyl Acetate	ASTM F903	Pass
Sulfuric Acid (97%)	ASTM F903	Pass
Tetrahydrofuran	ASTM F903	Pass
Sodium Hydroxide	ASTM F903	Pass
Acetone	ASTM F903	Pass
Hydrofluoric Acid	ASTM F903	Pass
Acetonitrile	ASTM F903	Pass

MicroMax® **VP** - Premium Protection from High-Risk Contaminants



Protective hood

Seamless front reduces the risk of possible contamination from liquid penetration

Taped storm flap keeps contaminants away from zipper

Elastic back gives a more comfortable fit and helps prevent rip-outs

Available with attached boots to help prevent cross-contamination during

MicroMax® VP Configurations



- Zipper closure Attached hood
- · Attached hood
- Elastic wrists Sizes: S – 5X



 Elastic wrists · Elastic ankles Sizes: S – 5X

Critical Protection - Tested Performance

Hazard Type	Lakeland® Brand	Test Method	Test Results
Blood Borne Pathogens	MicroMax® VP	ASTM F1671 Viral Penetration φX174 Bacteriophage Suspension	Pass

Fentanyl - Testing per ASTM D6978

Lakeland® Brand	Test Drug and Concentration	Minimum Breakthrough Detection Time (Specimen 1/2/3) (Minutes)	Steady State Permeation Rate (Specimen 1/2/3) (µg/cm2/minute)	Other Observations
MicroMax® VP*	Fentanyl Citrate Injection, 100 mcg/2mL	>240	NA	Slight swelling; no degradation

* MicroMax[®] VP fabric holds out liquid Fentanyl, but is only recommended for Fentanyl in powder form due to serged seam construction

Users should also ensure the gloves they are using for chemotherapy have been tested against the most recent standards. The current standard for exam gloves used in chemotherapy is ASTM D6978-05 "Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.

Prior to ASTM D6978-05 many exam gloves were tested against ASTM F739 "Resistance of Protective Clothing Materials to Permeation by Liquids or Gases Under Conditions of Continuous Contact". ASTM D6978-05 uses ASTM F739 as a test method, but has a chemical permeation requirement that is 10 times more stringent than what is required by ASTM F739. Users of gloves tested under ASTM D6978-05 have a higher level of confidence that the gloves they are using are tested to the current, more stringent ASTM standard.

Protecting your products, people, and controlled environments.













In the USA

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