

Advanced Antiviral & Antibacterial Protection for Paper & Boards

SWISS TECHNOLOGY INSIDE

CONSUMER CONCERNS TODAY

MAAK

- Consumers have become much more aware of how infectious viruses and dangerous microbes spread due to COVID-19 pandemic.
- Consumers are taking preventive measures to protect themselves & their families.
- Many studies confirm that consumers are feeling anxious, sad, scared, and overwhelmed.
- MAAK takes care of all the above concerns through its anti viral fighting abilities.

CONSUMER DEMAND FOR ANTIVIRAL

- The Covid-19 Pandemic has thrown up challenges to mankind in almost all spheres of life.
- The major challenge being the prevention of transmission of infections from various surface like Interior hard surfaces, textiles, paper products, Wooden and plywood surfaces etc (Herein after referred to as SURFACES).
- Day to day interactions with above surfaces have surprisingly gone up, due to unavoidable touch from persons who may be carrying the Covid-19 virus.
- The awareness for hygiene has increased considerably.
- The above surfaces provide a large hosting surface area for viruses
- After the coronavirus lockdown, personal protection is a topic of discussion as life carefully returns to normal.



HOW MAAK ANTIVIRAL PROTECTION WORKS

- MAAK ANTIVIRAL PROTECTION is a custom blend of Sanitized T20-19, a patented Swiss quaternary silane technology.
- Creates a highly-cationic charge density on the treated surface, deactivating the spread of the virus on contact.
- Acts quickly to prevent the transmission of enveloped and non-enveloped viruses
- Provides antiviral protection

Q.A.

Provides antibacterial protection.



HOW MAAK ANTIVIRAL PROTECTS THE WORLD FROM COVID-19





LIFE OF THIS ANTIVIRAL COATING IS AS LONG AS THE COATED SURFACE EXISTS . THE CORONAVIRUS STARTS GETTING KILLED FROM THE 10TH MINUTE AND 99.2% OF THE VIRUS IS KILLED BY THE 30TH MINUTE. THIS "KILL CYCLE" ENSURES THAT THE VIRUS DOES NOT PROPOGATE ITSELF DURING THE LIFE CYCLE OF THE COATED SURFACES, THEREBY MINIMIZING THE SPREAD OF VIRUS FROM PERSON TO PERSON.

> A SMALL CONTRIBUTION FROM MAAK IMPEX FOR A SAFER WORLD

MAAKIM

Consumer Friendly & Environmentally Safe



Fast Facts





Global Registration & Compliance

- In compliance with the requirements of the
 - BPR: Regulation (EU) No 528/2012 (the Biocidal Products Regulation, BPR).
 - REACH: Regulation (EC) No 1907/2006 (REACH)
- Registered with the U.S. Environmental Protection Agency as Sanitized[®] Brand T 20-19
- Compliance for Bluesign, Oeko-Tex and ZDHC MRSL.
- The active substance is AOX-free and is readily degradable in biological waste water plants according to OECD 301A
- The treated articles were tested for skin sensitisation according to the international norm OECD 406 and passed successfully the Repeated Insult Patch Test (RIPT)





Ø ZDHC



bluesian



Claims & Compliance

- MAAK ANTIVIRAL PROTECTION is a performance-based trademark.
- MAAK ANTIVIRAL PROTECTION is fast-acting antiviral and antibacterial agent and the treated articles performance verified to qualify to use the MAAK ANTIVIRAL PROTECTION trademark.
- Testing for MAAK ANTIVIRAL PROTECTION the trademark utilizes AATCC-100 modified for viruses.



MAAK ANTIVIRAL PROTECTION Benefits

- Swiss Antimicrobial Technology with all the safety and compliance certificates.
- Doesn't change mechanical or visual properties of the coated surface.
- Antiviral
- Antibacterial
- Safe to touch
- Safe for the Environment
 - Swiss Technology
 - Peace of Mind

SWISS TECHNOLOGY INSIDE • Trust of the customer





ApplicationAreas

- All Paper Products like boxes, bags, News print, Text books etc.,
- Plywood boxes and products
- Pinewood boxes and Products
- Interior surfaces such as Table tops, desk, chairs, Wooden counters, walls etc.,

FORM NO: 10-A001 Rev No : 02 Date: JULY 2020

SAFETY DATA SHEET

COMPANY

1. IDENTIFICATION OF THE PRODUCT AND

1.1. Product identifier	MAAK ANTIVIRAL PROTECTION		
1.2. Relevant identified uses of the substance or	Antibacterial and Antiviral finish for textiles.		
mixture and uses advised against			
	<u> </u>		
2. HAZARD IDENTIFICATION			11.
2.1. Classification of the substance or mixture*		Product is a Mixture. Non- flammal	-
		dangerous substance according to C	GHS. It is a corrosive liquid
		and can cause irritation.	
*Regulation (EC) No. 1907/2006 REACH and as per D	PD: Dir	ective 1999/45/FC	
2.2 Hazard statement			
Signal word	Danger	N	
	skin rea	hazard - H315 – Causes skin irritation. H3 action. H318 – Causes serious eye damage category – 1A	,
2.3. Label elements**	he pro	oduct need not be labeled in accordance	e with EC directives
**Classification and marking according to Regulation (EC) No. 1272/2008 REACH			
2.4. Other hazards		HMISCLASSIFICATION	NFPA RATING
		Health Hazard: 1	Health Hazard: 1
		Flammability: 0	Fire: 0
		Physical Hazards: 0	Reactivity Hazard: 0

MAAK IN * 2.4.

FORM NO: 10-A001

Rev No : 02 Date : JULY 2020

3.COMPOSITION/INFORMATION ON INGREDIENTS

3.1 Chemical Identity	An aqueous suspension of Quaternary ammonium- silane
3.2 C.A.S No of active ingredient	41591-87-1
4.FIRST AID MEASURES:	
4.1. Description of first aid measures	In all cases of doubt, or when symptoms persist, seek medical advice. If unconscious place in recovery position and seek medical

	4.1. Description of first aid measures	In all cases of doubt, or when symptoms persist, seek medical advice. If unconscious place in recovery position and seek medical advice. In case of unconsciousness give nothing by mouth, place in recovery position and seek medical advice. No mouth-to-mouth or mouth-to-nose resuscitation. Use Ambu bag or ventilator.
	4.2. Skin	Take off immediately all contaminated clothing. After contact with skin, wash immediately with plenty of water and soap. Do not use solvents or thinners.
	4.3. Eyes	Keep eyelids open, wash out with plenty of clean, fresh water and seek medical advice. Remove contact lenses, if present and easy to do. Continue rinsing.
	4.4. Inhalation	Remove casualty to fresh air and keep warm and at rest. In case of irregular breathing or respiratory arrest provide artificial respiration. No mouth-to-mouth or mouth-to- nose resuscitation. Use Ambu bag or ventilator.
	4.5. Ingestion 5.FIRE FIGHTING MEASURES	If swallowed, rinse mouth with water (only if the person is conscious). Seek medical advice immediately. Keep victim calm. Do NOT induce vomiting.
1	5.1. Extinguishing media	Water spray jet, Foam, Carbon dioxide, Extinguishing powder.
	5.2. Special exposure hazards / unusual hazards	Dense black smoke occurs during fire. Inhaling hazardous decomposing products can cause serious health damage.
F	5.3. Advice for firefighters	Wear a self-contained breathing apparatus and chemical protective clothing. Cool closed containers that are near the source of the fire. Do not allow water used to extinguish fire to enter drains, ground or waterways.
	5.4 Unsuitable extinguishing media	Strong water jet.

6.ACCIDENTAL RELEASE MEASURES	FORM NO: 10-A001 Rev No : 02 Date = JULY 2020
6.1. Personal precautions	Wear personal protective suits. Chemical goggles or full faced shield. Chemical resistant rubber or neoprene gloves, NIOSH approved positive pressure air supplied respirator Avoid contact with eyes and skin. Avoid breathing dust/fume/gas/mist/vapours/spray.
6.2. Environmental precautions 6.3. Clean up procedure	Observe local byelaws. Do not allow to enter into surface water or drains. If the product contaminates lakes, rivers or sewages, inform competent authorities in accordance with local regulations. For small spills use Mop, wipe or soak with
NAAK I.	cloth or absorbents. E.g. sand, kieselguhr saw dust, etc and dispose according to regulations. Large spills should be contained to prevent spreading. Do not allow product to enter lakes, sewers, streams, ponds, estuaries, oceans, or other waters unless permitted by law.
6.4 Reference to other sections 7.HANDLING AND STORAGE	Prevent spreading over wide area (by containment)
7.1. Advice on safe Handling	General ventilation is required. Avoid contact with eyes and skin. Avoid breathing dust/fume/gas/mist/vapours/spray. Provide for sufficient ventilation; if possible, use resp. install internal exhaust systems. When using do not eat, drink or smoke. Take off immediately all contaminated clothing. Thorough skin- cleansing after handling the product
7.2. Conditions for safe storage	Store in a well-ventilated and dry room at temperatures between 10 °C and 30 °C. Protect from heat and direct sunlight. Make sure spills can be contained, e.g. in sump pallets or kerbed areas. Keep/Store only in original container.
7.3. Incompatible packaging materials.	Advised to use original containers only. Avoid materials prone for rusting. Do not store together with: Alkali (lye), Oxidizing agent, Reducing agent.

FORM NO: 10-A001 Rev No : 02 Date : JULY 2020

8.EXPOSURE CONTROLS AND PERSONAL PROTECTION

8.1. Exposure controls	Engineering controls: Use mechanical local exhaust at
8.1. Exposure controls	point of vapor or mist release. Ensure that existing
	ventilation is sufficient to prevent exceeding the
	recommended PEL/TLV levels.
Personal protective equipment	
Respiratory protection	Generally, not required. In the case of aerosol-mist
	formation protection is essential. Where adequate
	ventilation is not available, use NIOSH- approved
9. PHYSICAL AND CHEMICAL PROPERTIES	respirator with organic filter.
<u></u>	
Hand protection	For prolonged or repeated handling, the following glove
	material must be used: NBR (Nitrile rubber). Thickness of
	the glove material > 0,4 mm; Breakthrough time
	(maximum wearing time) > 480 min
Eye protection	Goggles giving full protection required.
Lyeprotection	Goggles giving full protection required.
Hygiene measures	After contact clean skin thoroughly with water and soap
	or use appropriate cleanser. Immediately remove any
	contaminated clothing, shoes or stockings.

9.1. Information on basic	physical and chemical propert	ties
Form		Liquid
Color		Off white to white
		Characteristic
Odor		
	pH (1% diluted) at 25deg C	NA
	pH at 25 deg 🔨 💙	7.0-9.0
	Boiling point	100 degree Celsius.
6	Melting point	Not applicable , N/E
Flash point		Not applicable
~ ()/I	Flammability	Nonflammable
	Auto flammability	N/E
	of active ingredients	
	Explosive properties	Not applicable
	Oxidizing properties	Not applicable
	Vapor pressure	N/E
	Specific gravity	Approx. 1.0.
	Solubility in water	miscible
	Solids in %	Approx. 4 wt %
	Organic solvent	Approx 5 wt %
Water		91 wt %

FORM NO: 10-A001 Rev No : 02 Date : JULY 2020

9.2. Other Information	
Vapor density	Not determined
Evaporat ion rate, (Butyl	Not determined
acetate =1)	
Conductivity	Not determined
10. STABILITY AND REACTIVITY If stored and handled in accordance with standard indu	Not determined strial practice no hazardous reactions are known.
10.1. Stability	Stable under NTP, sensitivity to light prior to curing on textiles or other substrates.
10.2. Reactivity	No specific hazard to be mentioned
10.3. Possibility of hazardous reactions	Keep away from strong acids, strong bases and strong oxidizing agents to avoid exothermic reactions
10.4. Conditions to avoid	Protect from heat and direct sunlight
10.5. Materials to avoid contact	No incompatible chemicals known.
11700199129169198866669005	Hazardous decomposition by products may form with exposure to high temperatures.
11.1. Information on toxicological effects	
On Skin	Causes skin burns. May cause an allergic skin reaction.
Eyes	Causes serious eye damage.
Ingestion	Oral, LD50, Rat: > 5000 mg/kg.
Other effects	No data available
Chronic health effects	No data available
Teretogenedity	Reproductive toxicity No indications of human reproductive toxicity exist. Germ cell mutagenicity; evaluation: No indications of human germ cell mutagenicity exist.
Carcinogenic	Carcinogenicity: No indications of human carcinogenicity exist.

9.1. Information on basic physical and chemical prope	erties	
Form	Liquid	
Color	Off white to white	
Odor	Characteristic	
pH (1% diluted) at 25deg C	NA	
pH at 25 deg C	7.0 – 9.0	
Boiling point	100 degree Celsius.	
Melting point	Not applicable , N/E	
Flash point	Not applicable	
Flammability	Nonflammable	
Auto flammability of active ingredients	N/E	
Explosive properties	Not applicable	
Oxidizing properties	Not applicable	
Vapor pressure	N/E	
Specific gravity	Approx. 1.0.	
Solubility in water	miscible	
Solids in %	Approx. 4 wt %	
Organic solvent	Approx 5 wt %	
Water	91 wt %	

FORM NO: 10-A001 Rev No : 02 Date : JULY 2020

12.ECOLOGICAL INFORMATION

12.1. Toxicity	Fish toxicity, LC50, Oncorhynchus mykiss (Rainbow trout): 1.73 mg/l
12.2. Persistence & degradability	Toxicological data are not available.
12.3. Bioaccumulative potential	Toxicological data are not available.
12.4. Mobility	Toxicological data are not available.
12.5. Behaviour in sewage	Toxicological data are not available.
12.6. Other adverse effects	The substances in the mixture do not meet the PBT/vPvB criteria according to REACH, annex XIII

13.DISPOSAL CONSIDERATIONS

13.1. Waste treatment method	
Product disposal	Do not allow to enter into surface water or drains. This material and its container must be disposed of in a safe way. Waste disposal according to directive 2008/98/EC, covering waste and dangerous waste. Observe in addition any national regulations.
Packaging disposal	Handle contaminated packages in the same way as the substance itself.
14.TRANSPORT INFORMATION	
14.1 Classification for ROAD and Rail	Corrosive, Irritant symbol

Not applicable Not applicable

Not applicable

Corrosive, Irritant

transport: Proper shipping name

UN number

Class

Packing group

a change broup

14.2 Classification for SEA transport (IMO- Corrosive , Irritant

IMDG): Proper shipping name

14.3 Classification for AIR transport

(IATA/ICAO): Proper shipping name

15. REGULATORY INFORMATION	FORM NO10-A001 Rev No : 02 Date : JULY 2020	C
15.1. Health and safety Information	Precautionary Statements P261 - Avoid breathing vapors P280 - Wear protective gloves/protective clothing P302 + P352 - IF ON SKIN: Wash with plenty of soap and water P305 +P351+P338: IF IN EYES: : Rinse cautiously with water for several minutes. Remove contact lenses, if present and Continue rinsing. P310 - Immediately call a POISON CENTER or doctor/ physician. P501 - Dispose of contents/container to industrial incineration plant	
15.2 Product Risk classification Product Risk Phrase	Risk phrase applicable are as follows 1.R36 - irritating to eyes 2.R43 - May cause sensitization by skin contact. 3.R41 - Risk of serious damage to eyes. 4. R52 - Harmful to aquatic organisms	
15.3 Product safety phrase	Safety phrases are as follows 1 S2 -Keep out of the reach of children 2.S3/7-Keep container tightly closed in a cool place 3.S26 - In case of contact with eyes, rinse immediately with plenty of water and seek medical advice 4.S29/35- Do not empty into drains; dispose of this material and its container in a safe way 5 S61-Avoid release to the environment. Refer to special instructions/safety data sheet.	

REV NO: 00 DATE: MAY 2020 TECHNICAL DATA SHEET

Quaternary Ammonium-Silane Antimicrobial Formulation

General Information

MAAK ANTIVIRAL PROTECTION is a ready-to-use formulation of an EPA registered organo-functional silane based antimicrobial agent that uses the well tested, polymeric binding properties and antimicrobial attributes of quaternary ammonium-silane (quat-silane) chemistry to create antimicrobial treated articles. This unique binding system provides a durable, broad spectrum, non-leaching coating that reacts on and with the coated surface. This treatment creates a surface that is inhospitable for microbes and is effective against mold, mildew and algae as a static agent. When incorporated into industrial and household products, and consumer textiles during the manufacturing process or in use, MAAK ANTIVIRAL PROTECTION inhibits the growth of microbes to offer protection from offensive odors and product deterioration, increased durability and product freshness.

It is supplied in dilute, ready-to-use, liquid form, making it easy to mix with compatible alternative finishing agents (like anti-wrinkle resins, fluorocarbons, wicking agents and softeners, water proof coatings, Paints, polishes, paper coatings etc.,).

Special features

MAAKIMI

- Silane binding technology provides excellent durability
- Broad spectrum activity: controls odor-causing microorganisms
- Thermal stability: high tolerance in manufacturing
- Easy to apply on natural and absorbing surfaces
- Maintains aesthetics and freshness to the treated surface.

REV NO: 00

DATE: MAY 2020

• Can be applied by pad, spray, or exhaustion process

- Dermatologically tested
- Active substance is free from AOX
- Easily bio-degradable
- EPA approved base ingredient (No. 91742-3)
- Base ingredient complies with EU BPR and BLUESIGN certification
- Base ingredient accepted for OEKO-TEX Standard 100

Physical Properties

Composition	Silane-functional tetraalkylammonium compound in water	
Appearance (Visual)	Off white to white	
Ionic Nature	Cationic	
Solubility in Water	Miscible	
pH Value	7.0-9.0	
Quat-Silane	3.5-4.5	
concentration, wt. %		
Ecology/Toxicology	The usual hygiene and safety rules for handling chemicals	
	should be observed in storage, handling & use. Follow SDS.	

Shelf Life

Application

MAAK ANTIVIRAL PROTECTION may be applied to both organic and inorganic surfaces.

TECHNICAL DATA SHEET

REV NO: 00 DATE: MAY 2020

Directions for Use: Use standard coating methods such as padding, saturation, spray, foam, or exhaust applied as a dilute aqueous solution to give **1.0% to 5.0%** percent by weight of active ingredients.

As MAAK ANTIVIRAL PROTECTION is a dispersion, the active may settle on storage. It will be redispersible on proper shaking. It is always advised to shake the container well before use.

For pad and exhaust applications, aqueous bath solutions can be prepared by simply adding the **MAAK ANTIVIRAL PROTECTION** to water with stirring. Aqueous solutions should remain with agitation for 1 hour prior to use, however, lower time frames may be acceptable depending on substrate. After applying treatment, the surface should be allowed to dry at temperatures to a maximum of 160°C (320°F) to effectively complete curing of the siloxane bonds and to remove excess water, solvents and/or traces of volatile solvents from hydrolysis.

Additional binder (1-2 wt%) can be added for achieving extended durability for cotton and polyester fabrics. Additional non-ionic wetting agent (max 0.5 wt%) can be used for towels and other articles where absorbency is important.

Application Procedures

Padding-Drying

Padding

Padding bath temperature: Approx. 20-40 °C Bath pH: 6.0-8.0 (can be optionally adjusted with acetic acid, check pH after adding the product)

TECHNICAL DATA SHEET

REV NO: 00 DATE: MAY 2020

MAAK ANTIVIRAL PROTECTION

TECHNICAL DATA SHEET

Exhaust - Semi-Hydro - Tumble Dry

REV NO: 00 DATE: MAY 2020

Exhaust: 15 - 20 Mins

Exhaust at bath temperature: approx. 20 °C – 40 °C

Bath pH: 6.0-8.0 (can be optionally adjusted with acetic acid, check pH after adding the product)

Bath MLR: 1:6 - 1:8

Tumble drying temperature: 60-100 °C (drying time depends on quality of fabric/garment)

Dip spin – Tumble drying

Dip spin: 15 - 20 Mins

Dip spin at bath temperature: approx. 20 °C – 40 °C

Bath pH: 6.0-8.0 (can be optionally adjusted with acetic acid, check pH after adding the product)

MLR: 1:1

Tumble drying temperature: 60-100 °C (drying time depends on quality of fabric/garment)

Approved applications for MAAK ANTIVIRAL PROTECTION are the preservation of non-food contact

<u>coatings</u> and films, and industrial and household woven and nonwoven fibers and

textiles. From mattresses and linens, to sports apparel and footwear, to fabrics used in hygienic environments, MAAK ANTIVIRAL PROTECTION is an ideal solution.

For use in fibers: non-food contact uses in industrial and household woven and non-woven fibers such as bedding, apparel, footwear, wall and floor coverings, carpets, draperies, wiping cloths, brushes, filters, insulation, tents, awnings, and traps. Use **MAAK ANTIVIRAL PROTECTION** in the treatment bath at levels that provide 1.0 to 5.0 active ingredient on the fiber (depending on end-use claims and expectations).

Use MAAK ANTIVIRAL PROTECTION in a well-ventilated area, free of sparks and open flames. Standard city water may be used, provided it is free of high concentrations of metal ions.

Antimicrobial Activity

Substrates treated with MAAK ANTIVIRAL PROTECTION are noted for their proven, outstanding skin tolerance and are safe for human and the environment. The MAAK ANTIVIRAL PROTECTION active ingredient and the proprietary silane technology binding protocol incorporated within the XTS-18 formulation provide a reliable and durable bacteriostatic effect against both Gram positive and Gram negative bacteria, yeasts and fungi. MAAK ANTIVIRAL PROTECTION both covalently and ionically binds to itself and fiber surfaces creating a surface modification in which odor causing organisms cannot adhere and colonize without changing other physical properties of the final fabric. MAAK ANTIVIRAL PROTECTION has been tested for efficacy against a variety of bacterial and fungal organisms. Please note that approval of this product by United States Environmental Protection Agency limits efficacy claims made for antimicrobial treated articles to non-pathogenic organisms.

Antimicrobial and anti-odor performance has been demonstrated using industry standard test techniques including ASTM E3160-18, ASTM E2149, JIS L1902, ISO 20743, ASTM E3162-18, AATCC TM100 and IACM0710. Standard ASTM E3162-18 method can be followed as wash protocol. All tests performed and verified by the International Antimicrobial Council and test results are available upon request.

Storage, Handling and Disposal

Please refer to the Safety Data Sheet for this product for precise instructions. The processing and use of industrial chemicals require adequate technical and professional knowledge. In general, avoid eye and skin contact, and wear correct personal protective equipment. Avoid prolonged inhalation of MAAK ANTIVIRAL **PROTECTION** vapors.

Store and use the MAAK ANTIVIRAL PROTECTION in a well-ventilated area, away from sparks or open flames.

It should be stored at ambient conditions in the original container, tightly sealed. Protect from frost and heat. **Do not freeze**.

ANTIVIRAL REPORT FOR PAPER & BOARD SURFACES



TEST REPORT NUMBER : MUM 55063 / 2020



X PVT LTL

Testtex India Laboratories Pvt. Ltd.

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www.testtex.com

Tested samples will be retained for 60 days after issue of test report unless otherwise agreed with the customer. Testing Textiles, Leather Products, Footwear, Water, Food, Packaging Material, Construction Materials & Consumer Goods TTI/REC30/TR/TEXTILES

Testtex

TEST	REPORT NUMBE	CR : MUM 55063 / 2020)			MUMB	AI
NAM	OF CLIENT . M	S MOHAMAD AMIN	ABDUL RAHM.	AN SAIT.			
				ate Of Sample	Submission : 30.09.2	020	
Addre	ss: #3275, 12 TH N	fain, HAL 2 nd Stage, In	diranagar, D	ate Of Test Sta	rt: 12.10.2	020	
	Bangalore - 50	50038		ate Of Test Co	mpletion : 14.10.2	020	
			D	ate Of Reportin	ng: 15.10.20	020	
		om rugated Paper Board (Method for Antimicrol	-				
Purpose of Test Organ		viral Finishes on Fabric erichia coli bacteriopha					
Test Organ Test Condi	isms : Esch tions :	erichia coli bacteriopha					
Test Organ Test Condi I. Conta	isms : Esch tions : ct Time	erichia coli bacterioph: : 10 & 30 Minutes					
Test Organ Test Condi 1. Conta 2. Incuba	isms : Esch tions : ct Time ntion Temp	erichia coli bacterioph : 10 & 30 Minutes : 37°C +/- 2°C	age MS2 ATCC				
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra	isms : Esch tions : ct Time ttion Temp dizer Used	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing l	age MS2 ATCC				
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra 4. Media	isms : Esch tions : ct Time titon Temp lizer Used and Reagent	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing t : TSA Agar	age MS2 ATCC				
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra 4. Media	isms : Esch tions : tt Time tion Temp dilzer Used and Reagent tion Period	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing t : TSA Agar : 48 Hours	age MS2 ATCC		Recovered MS2	Formula	Percent
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra 4. Media	isms : Esch tions : tt Time tion Temp lizer Used and Reagent tion Period Recovered	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing l : TSA Agar : 48 Hours Recovered MS2	age MS2 ATCC broth	15597			
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra 4. Media 5. Incuba	isms : Esch tions : tt Time tion Temp dilzer Used and Reagent tion Period	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing l : TSA Agar : 48 Hours Recovered MS2	age MS2 ATCC broth Formula	15597 Percent	Recovered MS2 Phage after contact time =	Formula	
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra 4. Media 5. Incuba	isms : Esch tions : ct Time tion Temp lizer Used and Reagent tion Period Recovered MS2 Phage	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing l : TSA Agar : 48 Hours Recovered MS2 Phage after contact time = 10 min	age MS2 ATCC broth Formula	15597 Percent	Recovered MS2 Phage after contact time = 30 min	Formula	
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra 4. Media 5. Incuba	isms : Esch tions : tt Time tion Temp lizer Used and Reagent tion Period Recovered MS2 Phage after contact	erichia coli bacteriophi : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing t : TSA Agar : 48 Hours Recovered MS2 Phage after contact time =	age MS2 ATCC broth Formula	15597 Percent reduction	Recovered MS2 Phage after contact time =	Formula	reduction
Test Organ Fest Condi Conta Co	isms : Esch tions : ttime tion Temp lizer Used and Reagent tion Period Recovered MS2 Phage after contact time = 0 hr	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing l : TSA Agar : 48 Hours Recovered MS2 Phage after contact time = 10 min	age MS2 ATCC broth Formula	Percent reduction 90.5 %	Recovered MS2 Phage after contact time = 30 min	Formula	reduction
Test Organ Test Condi 1. Conta 2. Incuba 3. Neutra 4. Media 5. Incuba	isms : Esch tions : tt Time tition Temp litzer Used and Reagent tition Period Recovered MS2 Phage after contact time = 0 hr (Pfu/sample)	erichia coli bacterioph: : 10 & 30 Minutes : 37°C +/- 2°C : D/E Neutralizing l : TSA Agar : 48 Hours Recovered MS2 Phage after contact time = 10 min (Pfu/sample)	age MS2 ATCC broth Formula 100(B-A)/B=R]	15597 Percent reduction	Recovered MS2 Phage after contact time = 30 min (Pfu/sample)	Formula [100(B-A)/B=R]	reduction

Note :

MAAKIM

0% - Not Acceptable <50% - Insignificant >50% - Significant

>95% - Acceptable & Significant Where,

R = Percentage of Bacteria / Reduction

A = the no. of bacteria recovered from the inoculated treated test specimen swatches in the jar over the desired contact time. B = no. of bacteria recovered from inoculated treated test specimen swatches in the jar immediately after 0 contact time. Remark : Sample shows antiviral activity.

AUTHORISED SIGNATORY

Testtex India Laboratories Pvt. Ltd.

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Tirupur

Noida

Tested samples will be retained for 60 days after issue of test report unless otherwise agreed with the customer, Tested samples will be retained to be out at the root of test report unless outerwise agreed with the customer. Testing Textiles, Leather Products, Footwar, Water, Food, Packaging Material, Construction Materials & Consumer Goods TTI/REC30/TR/TEXTILES

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ANTIVIRAL REPORT FOR PLYWOOD XTS FINISH - 80 GPL

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TEST REPORT NUMBER : MUM57316/2020



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			TESTR	EPORT		MUMBAI	
REPORT N	NUMBER : MUM	57316/2020					
ULR No. :	TC5835202000361	71P		NCAT			
NAME OF	CLIENT : M/S M	OHAMAD AMIN A	BDUL RAIM	Date Of Sample Sub	mission : 27-10)-2020	
Address :	#3275, 12" Mair	, HAL 2 Stage, In	diranagar,	Date Of Test Start :			
	Bangalore - 5600	38		Date Of Test Comp		2020	
				Date Of Reporting :		.010	
				Date Of Reporting .	04-11-2020		
Cell No :	erson : Mr. Amcen +91 9845050098 ar@ficuspax.com	Rahaman					
Sample Des	scription : Ply Woo	d, XTS 80 GPL					
Sample Des	Test Mathod for	ntimicrobial Activ	ity of Hard Non	-Porous Surfaces	IS Z 2801: 20	12	
Name of Test :	I est Michion for i						
Purpose of Tes	t : Antivira	l Finishes on Hard	Non-Porous Su	rfaces			
Test Organism	e : Escherie	chia coli bacterioph	age MS2 ATCC	15597			
Test Organism	5 Thorney						
Test Condition	ns :						
1 Contact T		: 10 & 30 Minutes :	at 35°C +/- 1°C				
1. Contact T	lime	: 10 & 30 Minutes : : 35°C +/- 1°C	at 35°C +/- 1°C	5			
2. Incubatio	Time on Temp	: 10 & 30 Minutes : : 35°C +/- 1°C : SCDLP	at 35°C +/- 1°C				
2. Incubatio 3. Neutraliz	lime In Temp er Used	: 35°C +/- 1°C	at 35°C +/- 1°C	5			
2. Incubatio 3. Neutraliz 4. Media an	lime on Temp er Used ad Reagent	: 35°C +/- 1°C : SCDLP	at 35°C +/- 1°C				
2. Incubatio 3. Neutraliz 4. Media an 5. Incubatio Sample	lime on Temp er Used ad Reagent	: 35°C +/- 1°C : SCDLP : TSA Agar	at 35°C +/- 1°C Log of 0 hr	Count after 10 min (Cfu / sample)	Log of 10 min	R = [Log B – Log C)]	% Reducti
2. Incubatio 3. Neutraliz 4. Media an 5. Incubatio Sample	Yime n Temp er Used d Reagent <u>on Period</u> Parameter	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr		Count after 10 min (Cfu/			
2. Incubatio 3. Neutraliz 4. Media an 5. Incubatic Sample Sentification Treated	Time on Temp er Used od Reagent on Period	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after Ohr (Cfu / Sample)	Log of 0 hr	Count after 10 min (Cfu / sample)	min	R = [Log B - Log C)] 0.864842098	% Reducti 86.34 %
2. Incubatio 3. Neutraliz 4. Media an 5. Incubatic Sample Jentification Treated	Yime n Temp er Used di Reagent on Period Parameter MS2	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr (Cfu / Sample) 133000	Log of 0 hr 5.123851641	Count after 10 min (Cfu / sample) 86000 630000	min 4.934498 5.799341	0.864842098	86.34 %
2. Incubatio 3. Neutraliz 4. Media an 5. Incubatio Sample dentification	Yime n Temp er Used di Reagent on Period Parameter MS2	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr (Cfu / Sample) 133000	Log of 0 hr 5.123851641	Count after 10 min (Cfu / sample) 86000	min 4.934498		
2. Incubatio 3. Neutraliz 4. Media an 5. Incubatic Sample lentification Treated Un-Treated Sample	Time in Temp er Used d Reagent <u>on Period</u> Parameter MS2 Bacteriophage	: 35°C 4/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr (Cfu / Sample) 133000 141000	Log of 0 hr 5.123851641 5.149219113	Count after 10 min (Cfu / sample) 86000 630000 Count after 30 min (Cfu /	min 4.934498 5.799341 Log of 30	0.864842098 R = [Log B – Log	86.34 %

Un-Treated Note : A Value of 2.0 or above is considered "antimicrobial" by JIS

Where, R = Value of antimicrobial activity A = Average of the number of viable cells of bacteria immediately after inoculation on the untreated test piece. B = Average of the number of viable cells of bacteria on the untreated piece after 24 hrs. C = Average of the number of viable cells of bacteria on the treated piece after 24 hrs. Parayte's complete hours antiviral activity.Remark : Sample shows antiviral activity.

Authorised Signatory

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ANTIVIRAL REPORT FOR PINEWOOD XTS FINISH - 80 GPL



TEST REPORT NUMBER : MUM57315 / 2020



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TEST	REPORT
REPORT NUMBER : MUM57315/2020	MUMB/
ULR No. : TC583520200036170P	
NAME OF CLIENT : M/S MOHAMAD AMIN ABDUL RAIR	MAN SAIT.
Address : #3275, 12 ¹¹¹ Main, HAL 2 nd Stage, Indiranagar,	Date Of Sample Submission : 27-10-2020
Address: #3275, 12 ¹¹¹ Main, HAL 2 ⁿ¹ Stage, Indiranagar, Bangalore - 560038	Date Of Test Start : 02-11-2020
•	Date Of Test Completion : 04-11-2020
	Date Of Reporting : 04-11-2020
Contact Person : Mr. Ameen Rahaman Cell No : +91 9845050098 E - Mail: ar@ficuspax.com	

Sample Description : Pine Wood, XTS 80 GPL Name of Test : Test Method for Antimicrobial Activity of Hard Non-Porous Surfaces – JIS Z 2801: 2012

: Antiviral Finishes on Hard Non-Porous Surfaces Purpose of Test

; Escherichia coli bacteriophage MS2 ATCC 15597 Test Organisms

- Test Conditions
- : 10 & 30 Minutes at 35°C +/- 1°C 1. Contact Time
- : 35°C +/- 1°C 2. Incubation Temp : SCDLP
- 3. Neutralizer Used
- 4. Media and Reagent : TSA Agar
- 5. Incubation Period : 48 Hours

Sample Identification	Parameter	Count after 0hr (Cfu / Sample)	Log of 0 hr	Count after 10 min (Cfu/ sample)	Log of 10 min	$\mathbf{R} = [\mathbf{Log} \ \mathbf{B} - \mathbf{Log} \ \mathbf{C})]$	% Reduction
Treated		13100	5.117271296	28000	4.447158		
Un-Treated	MS2 Bacteriophage	13500	5.130333768	510000	5.70757	1.260412145	94.50 %

Sample Identification	Parameter	Count after 0hr (Cfu / Sample)	Log of 0 hr	Count after 30 min (Cfu / sample)	Log of 30 min	R = [Log B – Log C)]	% Reduction
Treated	MS2	13100	5.117271296	7800	3.8920946	4.02171925	99,99 %
Un-Treated	Bacteriophage	13500	5.130333768	82000000	7.9138139		

Note : A Value of 2.0 or above is considered "antimicrobial" by JIS

Where, R = Value of antimicrobial activity

A = Average of the number of viable cells of bacteria immediately after inoculation on the untreated test piece. B = Average of the number of viable cells of bacteria on the untreated piece after 24 hrs.C = Average of the number of viable cells of bacteria on the treated piece after 24 hrs.Remark : Sample shows antiviral activity.

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ANTIVIRAL REPORT FOR WOOD WITH VENEER FINISHED WITH POLISH SURFACES

TEST REPORT NUMBER : MUM 61127 / 2020

TEST REPORT NUMBER : MUM 61127 / 2020





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esttex TEST REPORT REPORT NUMBER : MUM61127 / 2020 MUMBAI ULR No. : TC583520200039996P NAME OF CLIENT : M/S MOHAMAD AMIN ABDUL RAHMAN SAIT Address : #3275, 12TH Main, HAL 2nd Stage, Indiranagar, Date Of Sample Submission : 14-12-2020 Bangalore - 560038 Date Of Test Start : 21-12-2020 Date Of Test Completion: 23-12-2020 Date Of Reporting: 24-12-2020 Contact Person : Mr. Ameen Rahaman Cell No : +91 9845050098 E - Mail: ar@ficuspax.com Sample Description : Wood With Veneer Finished With Polish Name of Test : Test Method for Antimicrobial Activity of Hard Non-Porous Surfaces - JIS Z 2801: 2012 Purpose of Test : Antiviral Finishes on Hard Non-Porous Surfaces **Test Organisms** : Escherichia coli bacteriophage MS2 ATCC 15597 Test Conditions 1. Contact Time : 10 & 30 Minutes at 35°C +/- 1°C 2. Incubation Temp : 35°C +/- 1°C 3. Neutralizer Used : SCDLP Media and Reagent 4. : TSA Agar Incubation Period : 48 Hours 5. Count after 10 Sample Identification Count after 0hr Log of 10 Log of 0 hr min (Cfu/ $\mathbf{R} = [\mathbf{Log} \ \mathbf{B} - \mathbf{Log} \ \mathbf{C})]$ % Reduction Parameter (Cfu / Sample) min sample) 142000 Treated 5.152288344 148000 5.170262 MS2 0.156074146 30.18 % Bacteriophage 158000 5.198657087 212000 5.326336 **Un-Treated** Count after 30 Log of 30 R = [Log B - LogCount after 0hr Sample Identification Log of 0 hr min (Cfu/ % Reduction Parameter (Cfu / Sample) min C)] sample) 142000 5.152288344 5200 3.7160033 Treated MS2 2.096910013 99.2 % Bacteriophage 158000 5.198657087 650000 5.8129134 Un-Treated Note : A Value of 2.0 or above is considered "antimicrobial" by JIS Where, R = Value of antimicrobial activity A = Average of the number of viable cells of bacteria immediately after inoculation on the untreated test piece. **B** = Average of the number of viable cells of bacteria on the untreated piece after 24 hrs. **C** = Average of the number of viable cells of bacteria on the treated piece after 24 hrs. Remark : Sample shows antiviral activity. AUTHORISED SIGNATORY Testtex India Laboratories Pvt. Ltd. H.O. & CENTRAL LABORATORY: 4218588

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ANTIVIRAL REPORT FOR WOOD FINISHED WITH PAINT

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TEST REPORT NUMBER : MUM 61126 / 2020

TEST REPORT NUMBER : MUM 61126/2020



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	T NUMBER : MU					MUMBAI	
	D. : TC58352020003						
NAME	OF CLIENT : M/S	MOHAMAD AMI	NABDUL RAH				
Addre	Bangalore - 56	lain, HAL 2 nd Stage	, Indiranagar,	Date Of Sample :		4-12-2020	
	Dangalore - 50	0058		Date Of Test Star			
				Date Of Test Cor			
				Date Of Reportin	g: 24-12-2020)	
Cell N	t Person : Mr. Amo o : +91 9845050098 il: <u>ar@ficuspax.com</u>						
Sample	Description : W	OOD Finished W	ith Paint				
				on-Porous Surfaces -	IIS 7 2801- 2	2012	
					310 20 2001. 2	.012	
Purpose of	Test : Antivi	iral Finishes on Har	d Non-Porous S	urfaces			
Test Organi	- Fasha						
Test Organ	isms : Esche	richia coli bacteriop	hage MS2 ATC	C 15597			
Test Condit	ions :						
Test Condit 1. Contac		: 10 & 30 Minutes	at 35°C +/- 1°C				
1. Contac		: 10 & 30 Minutes : 35°C +/- 1°C	s at 35°C +/- 1°C				
1. Contac 2. Incuba 3. Neutral	t Time tion Temp lizer Used	: 35°C +/- 1°C : SCDLP	at 35°C +/- 1°C				
 Contac Incuba Neutral Media : 	t Time tion Temp lizer Used and Reagent	: 35°C +/- 1°C : SCDLP : TSA Agar	s at 35°C +/- 1°C				
 Contac Incuba Neutral Media : 	t Time tion Temp lizer Used	: 35°C +/- 1°C : SCDLP	s at 35°C +/- 1°C				
 Contac Incuba Neutral Media: Incubal 	t Time tion Temp lizer Used and Reagent	: 35°C +/- 1°C : SCDLP : TSA Agar	s at 35°C +/- 1°C Log of 0 hr	Count after 10 min (Cfu /	Log of 10 min	R = [Log B - Log C)]	% Reductio
 Contac Incuba Neutral Media Incuba 	t Time tion Temp lizer Used and Reagent tion Period Parameter	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours		Count after 10		R = [Log B – Log C)]	% Reductio
1. Contac 2. Incuba 3. Neutra 4. Media 5. Incubal Sample dentification Treated	t Time tion Temp lizer Used and Reagent tion Period	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr (Cfu / Sample)	Log of 0 hr	Count after 10 min (Cfu / sample)	min	R = [Log B – Log C)] 0.166331422	% Reductio 31.81 %
1. Contac 2. Incuba 3. Neutral 4. Media : 5. Incubat Sample dentification Treated Un-Treated	t Time tion Temp lizer Used and Reagent tion Period Parameter MS2	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr (Cfu / Sample) 135000 146000	Log of 0 hr 5.130333768 5.164352856	Count after 10 min (Cfu / sample) 225000 330000 Count after 30	min 5.352183 5.518514	0.166331422	31.81 %
Contac Incuba Neutra Neutra Neutra Incuba Sample dentification Treated Un-Treated Sample	t Time tion Temp lizer Used and Reagent tion Period Parameter MS2	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr (Cfu / Sample) 135000	Log of 0 hr 5.130333768	Count after 10 min (Cfu / sample) 225000 330000	min 5.352183		
Contac Lorotac Lorotac Lorotac Lorotac Somple Sample dentification Treated Un-Treated	t Time tion Temp lizer Used and Reagent tion Period Parameter MS2 Bacteriophage	: 35°C +/- 1°C : SCDLP : TSA Agar : 48 Hours Count after 0hr (Cfu / Sample) 135000 146000 Count after 0hr	Log of 0 hr 5.130333768 5.164352856	Count after 10 min (Cfu / sample) 225000 330000 Count after 30 min (Cfu /	min 5.352183 5.518514 Log of 30	0.166331422 R = [Log B - Log	31.81 %

Note R = Value of antimicrobial activity

A = Average of the number of viable cells of bacteria immediately after inoculation on the untreated test piece.

B = Average of the number of viable cells of bacteria on the untreated piece after 24 hrs.

C = Average of the number of viable cells of bacteria on the treated piece after 24 hrs.

Remark : Sample shows antiviral activity.

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