

K993347

11/19/99

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1.0. 510K SUMMARY as required by: 807.92(c)

2.0 APPLICANT :

NAME M/s. SRLANUSHAM RUBBER INDUSTRIES
PVT.LTD.

ADDRESS PIONEER MANIKANDAN BUILDINGS
VADASERY, NAGERCOIL,
TAMILNADU, INDIA-629001.

PH.NO. : 91-4652-33091,

FAX NO : 91-4652-32871.

CONTACT PERSON : MR. N. PARAMASIVAN
MANAGING DIRECTOR.

3. DEVICE TRADE NAME : NIL
COMMON NAME : Patient Examination Glove (Nitrile)

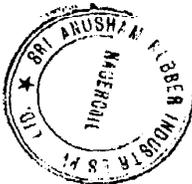
4. Legally marketed device to which the company claiming equivalence:
Class I Patient Examination Gloves (Nitrile) 80LZA that meets all the
requirements of ASTM D3578 - 95.

5. DESCRIPTION OF THE DEVICE :

Class I Patient Examination Gloves (Nitrile) 80LZA that meets all
the requirements of ASTM D3578- 95.

6. Intended use of the Device:

Examination glove (Nitrile) is a disposable device made of Nitrile Latex intended for medical
purpose that is worn on the examiners hand or finger to prevent contamination between patient
and examiner.



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**7.0 TECHNOLOGICAL CHARACTERISTICS OF THE DEVICE
COMAPARED TO PREDICATE DEVICE.**

Measured Parameters of Examination gloves (Nitrile) manufactured by Anusham Latex,			ASTM D3578 Requirement for Examination glove (Nitrile)
Characteristics	SIZE	Value	
1. Length	EX-S	235-240 mm	220 mm minimum
	S	235-240 mm	220 mm minimum
	M	235-240 mm	230 mm minimum
	L	235-240 mm	230mm minimum
2. Width	EX S	70MM	70 +/- 6 mm
	S	82 mm	80 +/- 6 mm
	M	93 mm	95 +/- 6 mm
	L	107 mm	111 +/- 6mm
3. Thickness	EX S	0.10mm	0.08 mm minimum
	S	0.10mm	0.08 mm minimum
	M	0.10mm	0.08 mm minimum
	L	0.10mm	0.08 mm minimum

PHYSICAL PROPERTIES

CHARACTERISTICS	BEFORE AGEING		AFTER AGEING	
	SARI VALUE *	ASTD 3578 REQUIREMENT	SARI VALUE	ASTD 3578 Requirement
Tensile Strength	18 – 20 mpa	14 mpa min	16 – 18 mpa	14 mpa min
Elongation at break %	750 – 800%	700% min	650-700%	500% min

SARI – SRLANUSHAM RUBBER INDUSTRIES .



PERFORMANCE REQUIREMENT:

Characteristics	Related defects	Level followed By		AQL followed by SARI	AQL Value as per ASTM D3578.
		SARI	As per ASTM D3578		
Freedom from Holes	Holes	S4	S4	1.5	4
Dimension	Width , Length Thickness.	S2	S2	4	4
Physical Property	Tensile Strength, Elongation at Break.	S2	S2	4	4

POWDER CONTENT

SARI VALUE	ASTM REQUIREMENT
120 +/- 20 mg / glove	-

PROTEIN CONTENT:

SARI VALUE	FDA REQUIREMENT
30 +/- 10 ppm	Value not fixed.

MOISTURE CONTENT:

SARI VALUE	FDA REQUIREMENT
0.8% max	No value fixed

BIOCOMPATABILITY:

SARI GLOVE	FDA REQUIREMENT
Biologically Compatible	Biologically Compatible



8.0 Performance Data:

The performance test data of the Nitrile examination glove manufactured by Sri Anusham Rubber Industries Pvt Ltd is given below.

Measured Parameters of Examination gloves (Nitrile) manufactured by Anusham Rubber industries Pvt. Ltd.,		
Characteristics	SIZE	Value
1. Length	EX-S	235-240 mm
	S	235-240 mm
	M	235-240 mm
	L	235-240 mm
2. Width	EX S	70MM
	S	82 mm
	M	93 mm
	L	107 mm
3. Thickness	EX S	0.10mm
	S	0.10mm
	M	0.10mm
	L	0.10mm

PHYSICAL PROPERTIES

CHARACTERISTICS	Before Ageing	After Ageing
Tensile Strength	18 - 20 mpa	16 - 18 mpa
Elongation at break %	750 - 800%	650-700%

INSPECTION LEVEL OF AQL:

Characteristics	Related defects	Level	AQL
Freedom from Holes	Holes	S4	1.5
Dimension	Width , Length Thickness.	S2	4
Physical Property	Tensile Strength, Elongation at Break.	S2	4



POWDER CONTENT : 120 +/- 20 mg per glove

PROTEIN CONTENT: 30 +/- 10 ppm

MOISTURE CONTENT: .0.8% max

BIOCOMPATABILITY: Biologically Compatible.

9. Clinical Data : NA

10. CONCLUSION OF PERFORMANCE TEST DATA:

The Examination gloves (Nitrile) manufactured by Sri. Anusham Rubber Industries Pvt.Ltd,

- Meet or exceed the ASTM D3578
- Meet FDA Pin hole Requirement.
- Meet labelling claim as shown by the data in 6

11. ANY OTHER INFORMATION:

Any other information required by FDA regarding product safety and effectiveness will be provided on request.





DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV 19 1999

Mr. N. Paramasivan
Managing Director
SRI Anusham Rubber Industries Private Limited
Pioneer Manikandan Buildings
Vandasery, Nagercoil-629001
Tamil Nadu, S. India

Re: K993347
Trade Name: Nitrile Examination Glove-Powdered
Regulatory Class: I
Product Code: LZA
Dated: September 28, 1999
Received: October 5, 1999

Dear Mr. Paramasivan:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Pre-market Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Good Manufacturing Practice for Medical Devices: General (GMP) regulation (21 CFR Part 820) and that, through periodic GMP inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in

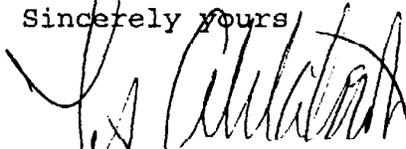
Page 2 - Mr. Paramasivan

the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours



Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

3.0 INDICATIONS FOR USE:

PAGE No

APPLICANT : SRLANUSHAM RUBBER INDUSTRIES PVT. LIMITED

510(K) No. : K993347

DEVICE NAME : NITRILE EXAMINATION GLOVES, *POWDERED*

INDICATIONS FOR USE:

Nitrile Examination Glove is a disposable device made of Nitrile Latex intended for medical purpose, that is worn on the examiners hand or finger to prevent contamination between patient and Examiner.



Chin S. Lin

(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices

Device Number K 993347

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