

3M™ CoTran™ 9702 Membrane

Controlled Caliper Ethylene Vinyl Acetate (EVA) Membrane

Description:

This intact (continuous) membrane has very uniform caliper. Its surface texture improves both web handling and bonding between the membrane and the adhesive. The product is supplied in roll form for use in the fabrication of transdermal, topical and transmucosal drug delivery systems.

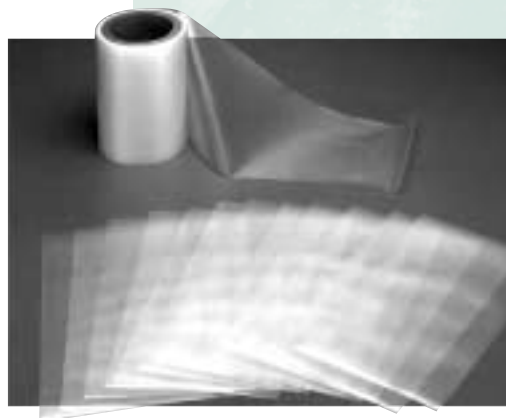
- 9% vinyl acetate
- Translucent
- Uniform 2 mil caliper, crossweb and downweb
- Heat-sealable
- Can be laminated directly to adhesives
- Manufactured for use in pharmaceutical products

Film Properties:

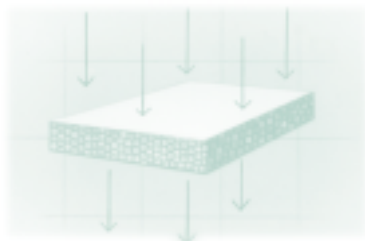
Test data presented are average values and are not intended to be used for specification purposes.

Properties:	Nominal Values U.S.	Nominal Values Metric
Caliper	2 mils	50.8 μm
Vinyl Acetate Content	9 %	9 %
Tensile		
MD	4.4 lbs./in.	19.6 N/25.4 mm
CD	2.0 lbs./in.	8.9 N/25.4 mm
Elongation		
MD	200 %	200 %
CD	300 %	300 %
MVTR		
(g/m ² /24 hr)	52.8	52.8

Note: 3M test methods and product specifications available upon request.



Membranes



3M Drug Delivery Systems

3M™ CoTran™ 9702 EVA Membrane

General Information:

	Nominal Values U.S.	Nominal Values Metric
Length:	2500 yds. on 3 in. I.D. cores	2286 m on 76 mm I.D. cores
Approximate roll diameters:	2500 yds. – 16 in. 1800 yds. – 13 in. 1000 yds. – 10 in.	2286 m – 41 cm 1650 m – 33 cm 914 m – 25 cm
Width:		
Maximum	48 in.	122 cm
Sample Roll:	25 yds. × 6 in.	23 m × 15 cm
Minimum Order Quantity:	134 yd ²	111 m ²
Maximum Per Roll:	2100 yd ²	1743 m ²

Traceability:

Each roll and shipment will contain a lot number which is traceable through production records to input materials and process conditions. Material traceability will be in compliance with United States current Good Manufacturing Practices.

Product Safety:

Product safety is characterized by the following evaluations:

USP Plastics Class V Extraction

- Acute Systemic Toxicity
- Intracutaneous Irritation

Storage Conditions:

The product should be stored at temperatures between 32–86°F (0–30°C) and relative humidity less than 75 percent. The product will maintain the stated performance, for a period of two years from the manufactured date, when stored under recommended conditions.

Product Use:

Customer is solely responsible for determining the suitability of Transdermal Components for the intended use and performing any necessary safety testing.

For More Information, Please Call:

U.S.A. (800) 643-8086

Europe 44 (1509) 613227

Japan (03) 3709-9671

www.3M.com/DDS



See 3M General Conditions of Sale for Transdermal Components for warranty information and other terms of sale.

3M

Drug Delivery Systems

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Minimum 10% Post-Consumer Fiber

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