

Technical Data Sheet

PVdC coated PVC FILM

(Non Toxic Food/Pharma Grade Material)

Complex : 250 μ x 90 GSM

Color Range : Clear, Amber, Blue, Green, Orange, Peach, Red, White Opaque, Yellow & other colors upon request

Sr.No.	Properties	Test Method	Unit	Specification
1	Color & Appearance	In-house	---	As per standard sample
2	Thickness of Rigid PVC film	DIN 53370	microns	250 \pm 10%
3	Total Thickness	DIN 53370	microns	305 \pm 10%
4	GSM of Rigid PVC film	DIN 2286	gm/m ²	340 \pm 10%
5	GSM of PVDC film	DIN 2286	gm/m ²	90 \pm 3
6	Total GSM	DIN 2286	gm/m ²	430 \pm 10%
7	Dimensional Changes - Longitudinal	ASTM D1204	%	-5.0
	Dimensional Changes - Transversal			+2.0
8	Dart Impact Strength (from height of 1.52 mtrs)	ASTM D1709	gm	350 min.
9	Tensile Strength - Longitudinal	ASTM D882	kg/Sq.cm	450 min.
	Tensile Strength - Transverse			450 min.
10	Elongation - Longitudinal	ASTM D882	%	5 min.
	Elongation - Transverse			5 min.
11	WVTR @ 38°C & 90% RH	ASTM F1249	gm/m ² /24 hr.	0.4 max
12	OTR @ 23°C & 0% RH	ASTM D 3985	cc/m ² /24 hr.	1.5 max.
13	Forming Temperature	In-house	°C	110 -170
14	Toxicity Test	USP Current Edition	-	Non-toxic
15	Residue VCM	ASTM D 3749	ppm	1 max
16	Identification - PVC	FTIR	-	As per Standard IR spectrum
17	Identification - PVDC	In-house	-	Color of PVdC layer shall turn brown
18	Heat Seal Strength	In-house	Kg/cm	0.30 min.
19	Global migration	2005/79/EC	ppm	< 60
20	Inner Diameter of Core	In-house	mm	76 +/- 1
21	Width	In-house	mm	Std. +/- 1

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Regulatory Compliances:

- Commission Directive 2005/79/EC (including amendment 2004/19/EC): Plastic material and articles in contact with Food Regulations.
- Commission Directive 94/63/EC (including amendment 2004/12/EC): Packaging and packaging waste (providing limits in the heavy metal content)
- Council Directive 78/142/EEC – Specifying limit of vinyl chloride monomer content.
- European Pharmacopoeia – 3.1.11 (Current Edition) – Containers for Dry Dosage Forms for Oral Administration
- European Pharmacopoeia – 3.2.2 (Current Edition) – Plastic Containers and Closures for Pharmaceutical Use
- USP 661 (Current Edition) – Containers for Non-Sterile Solid & Liquid Dosage Forms.
- The additives used in these films comply with the quantitative and qualitative requirements of EC and 21 CFR Part 177.