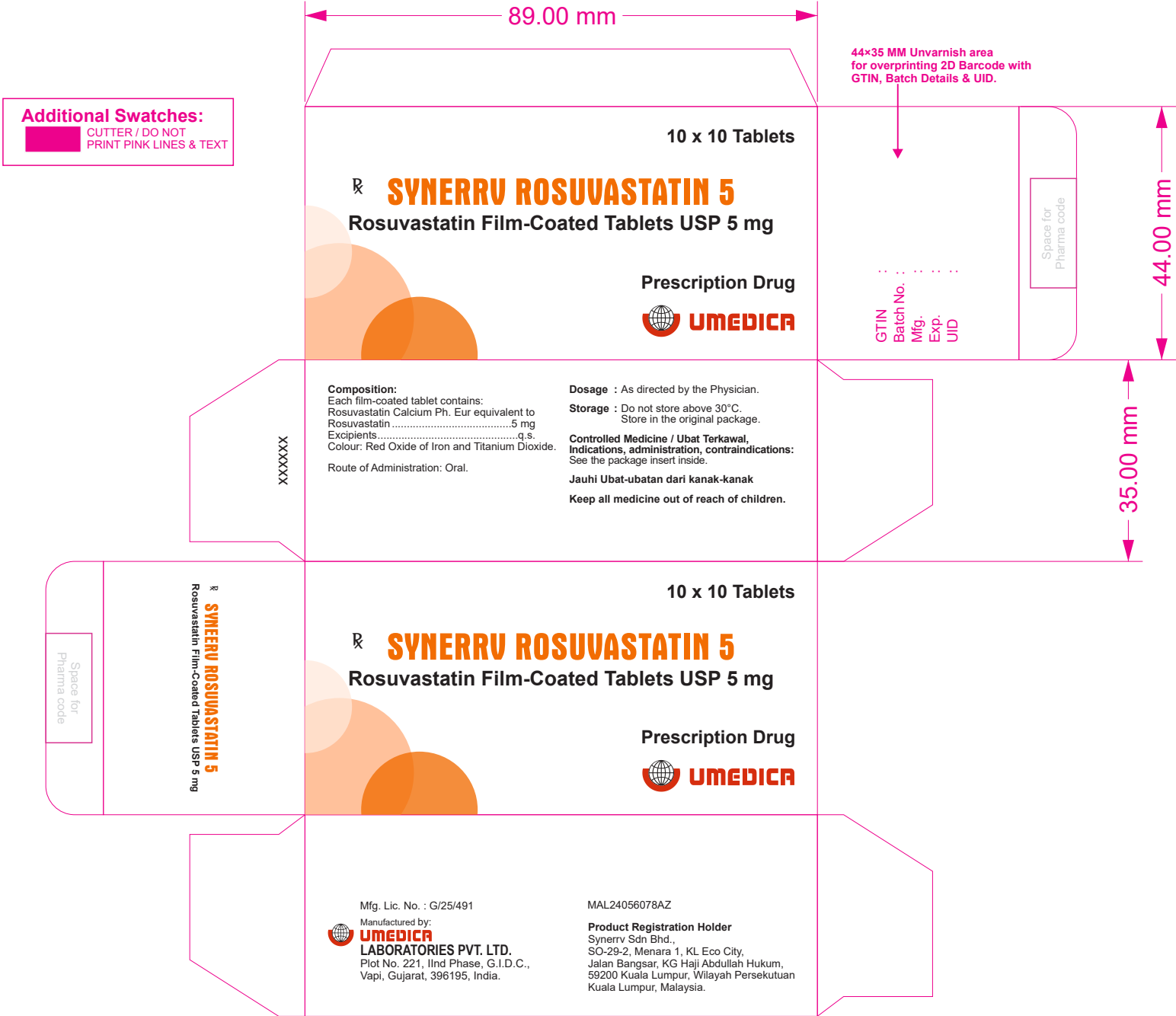




Format																					
Format Name		SPECIFICATION FOR ARTWORK APPROVAL																			
Format Number		FM/PDD/002-01-07		SOP Number		PDD/002															
New Artwork Code No. : XXXXXX				Old Artwork Code No.: NA		Artwork Effective Date:															
Product Name : SYNERRV ROSUVASTATIN 5 (Rosuvastatin Film-Coated Tablets USP 5 mg)				Company Name/Country : Synerrv / Malaysia																	
Component Type : Carton				Pack Style : 10×10T - Alu Alu Blister Pack																	
Description : New Development Carton Artwork as per Export-Order/Malaysia Country requirement.																					
Specifications : L×W×H				Layout/Drawing no.: (24-25-CTN-AUG-57)		Site : Umedica															
Artwork Language: English - Malay																					
Full-Size / Size : 89×44×35 MM				Type / Blister / Strip-Size / Adhesive / Tube-Type : Tuck In Flap																	
Size - Catch Cover / Foil / BOPP : NA				Sachet Size/Pocket Size/Fold Size/Release Paper : NA																	
Space between Labels : NA				Transparent Paper Thickness : NA																	
GSM of Paper /GSM of Aluminum-Foil: 300 GSM				Type of Paper /GSM : ITC Cyber XL Pac-GC1/JK Tuff Coated																	
Thickness/Thickness of Foil : NA				Type of Foil / Fold / Coating GSM : —																	
Cylinder : NA				Lamination/ Varnish /Material-Grade : UV Coated [Window Cut]																	
Repeat/Tolerance : NA				Partition / Plate Size / Total GSM / : NA																	
				Sealing / Embossing / Core Diameter : NA																	
GSM/BF of Printed Shipper : NA				Paper Grade / Flute Type / PLY of Shipper : NA																	
(Top/Middle/Bottom and Flute layer : NA																					
Printing Colour		P-Orange 021 C [Orange Shade]		P-485 C [Red Shade]		Black or as per Sample															
Approved by Export/Marketing :				[Approval attached with ADF]																	
CHECKLIST FOR CHECKING ARTWORK & SPECIFICATION																	Artwork Approved by Sign with Date				
Brand Name and Generic Name(As per IP/BP/USP/ In-house Pharmaco- peia)	Dosage Form and strength	Prefix Rx/ XRx	Mfg. Lic. No. / Neutral Code	Regulatory Agency Approval	Packing (Qty.) / Presentation	Dosage details if required	Size / Design	Composition /label claim/ Formulation	Schedule 'H', 'H1', 'G' or 'X' etc warning for local product / Red line for oral products	Item Code	QR Code/ Barcode (1D /2D)/ Pharma code as per requirement	Material and printing colours specifica- tion	Storage Condition as per pharmaco- peia/Buyer requirement/ Registration	Registration No. / Other requirement	Mfg. Name and Address, (Tel. No. and e-mail in the insert)	Marketed by/Distribut ors Name/Manu- facturer Name			Route of Admin. [In case of Inj./ Multidose Vial matter	Adequate Space (Batch details/2D barcode/ Embossing for Overprinting	
																				Prepared by PDD Department	Approved by PDD Department
																				PPIC Department	Production Packing Department
																				QC Department	
																				RA Department	
																				QA Department	
Note to Printer: • After receive the CDR File, Submit the print proof for approval. • The overprinting area should be unvarnished. • Machine proof must be verified against the approved hard copy in Light, Standard and Dark shade should be certified and signed by an authorized person. • Any deviation must be brought to the notice of the Packaging Development Department immediately in the writing. • For any clarification, please contact Packaging Development Department.																					