

[illegible]

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		SYNERRY ROSUVASTATIN 5			Company Name/Country		Synerrry / Malaysia												
(Rosuvastatin Film-Coated Tablets USP 5 mg)																			
Component Type		Blister Foil			Pack Style		10T - Allu Allu Blister Foil												
Description		New Development Blister Foil Artwork as per Export-Order/Malaysia Country requirement, as per Change Control No. CC/PDD24/056.																	
Specifications : L x W x H		Layout/Drawing no.: (EPI 3015 273 003)			Site : Umedica		Artwork Language: English												
Full Size / Size		: NA			Type+ Blister +Strip Size+Adhesive+Tube Type		: 81 x 38 x 3 (TB) MM												
Size -Gatch-Cover+ Foil +BOPP		: 263 MM (262 MM to 264 MM)			Sachet Size/Pocket Size/Fold Size/Release Paper : NA														
Space between Labels		: NA			Transparent Paper Thickness					: NA									
GSM-of Paper+GSM of Aluminum Foil:		Between 63.5 & 71.6 g/m ²			Type of Paper / GSM		Type-of Foil +Fold +Coating GSM			: NA									
Thickness/Thickness of Foil		: 0.025 MM (24µ to 27µ)								: Between 6.0 & 8.0 g/m ²									
Cylinder		: 1 Nos			Lamination / Varnish / Material Grade					: NA									
Repeat/Tolerance : 38.975 MM ± 0.015 MM					Partition / Plate Size / Total GSM /					: NA									
(10 Repats = 389.6 ~ 389.9 MM) [Eyemark]					Sealing / Embossing / Core Diameter					: NA									
GSM/BF of Printed Shipper		: NA			Paper Grade / Flute Type / PLV of Shipper					: NA									
(Top/Middle/Bottom and Flute layer : NA																			
Printing Colour		: P-Orange 021 C [Orange Shade]								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 Demarcation (PDD/USP/In-house code)	Dosage Form (Rx/ strength)	Phar. U.C. No. / Neutral Code	Regulatory Agency Approval	Packing (Qty) / Presentation	Dosage details if required	Size/ (label claim) Design	Composition (H ⁺ , H ⁻ , G ⁺ or local product for oral products)	Item Code	OR Code/ (10/20) and colour as per requirement	Material and colour's specification	Storage Condition as per pharmaceutical registration	Registration No. / Other	Mfr. Name and Address (e-mail in the insert)	Marketed by/ Distributor's Name	Route of distribution (cases of Int./ Multidose Via matter)	Special Batch number/ Embossing	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.																			