SYNOLAC® 8000 BA 75 (FORMERLY SYNOLAC® 4000 BA 75) INDUSTRIAL WOOD

		IG RESINS
Product Application details	ommended for use in-coating, roller or and drying, quickly lcohols as diluents, on.	
Polymer Type	Solventborne Alkyd	
	Solid Content at 125°C, % (ISO 3251)	74 - 76
Sales	Reduced Viscosity at 20°C, s (4mm, 60% in Butyl acetate) (DIN 53 211)	45 - 50
Specifications	Iodine Colour index, (60% % in Butyl acetate) (DIN EN 1557)	3 max
	Acid value, mg KOH/g (ISO 2114)	12 max
	Volatile	Butyl acetate
	Flash point, °C (ISO 3679)	24
	Density / Specific Gravity at 20°C, g/ml (ISO 2811)	<u>1.04</u>
Characteristics ¹	Type of fatty acid Fatty Acid content, %	Peanut oil 40
		10
	Note: Acid value and/or Hydroxyl value quoted relative to solid resin 1 The data provided for these properties are typical values, intended only as guides, and should not be construed as sales specifications	
Formulation Guidelines	SOLUBILITY SYNOLAC [®] 8000 BA 75 is completely soluble in aromatic hydrocarbons, esters, ketones, glycol ethers and their esters. It is partially soluble in alcohols and aliphatic hydrocarbons. <u>COMPATIBILITY</u> SYNOLAC [®] 8000 BA 75 is compatible with many non-drying, short oil alkyd resins, some polyester resins, like SYNOLAC [®] 680 BA 70, no-plasticized urea and melamine formaldehyde resins, maleic and ketone resins, phthalic acid esters as plasticizers and nitrocellulose. It is incompatible with drying medium to long oil alkyds, modified alkyds, hydrocarbon resins, high melting maleic resins.	
	OTHER ADDITIVES SYNOLAC [®] 8000 BA 75 may be pigmented with all commonly used inorganic and organic pigments. Matt finishes are easily obtained using silica matting agents, like Acematt [®] OK 500 (1) or Syloid [®] Silica ED 30 (2), preferably in combination with a polyethylene or polypropylene wax, like Crayvallac [®] WN-1495 (3), Crayvallac [®] WN-1135 (3) or wax dispersions like Crayvallac [®] WS-8050 (3).	
	Notes: (1) Evonik Industries AG, (2) Grace, (3) Arkema	



ARKEMA COATING RESINS

Product Safety	Please refer to the corresponding Safety Data Sheet.
Storage &	SYNOLAC [®] 8000 BA 75 should be stored indoors in the original, unopened and undamaged container, in a dry place at a temperature not exceeding 30°C. Exposure to direct sunlight should be avoided.
Handling	In the above mentioned storage conditions the shelf life of the resin will be 9 months from the shipping date

The statements, technical information and recommendations contained herein are believed to be accurate as of the date hereof. Since the conditions and methods of use of the product and of the information referred to herein are beyond our control, Arkema expressly disclaims any and all liability as to any results obtained or arising from any use of the product or reliance on such information; NO WARRANTY OF FITNESS FOR ANY PARTICULAR PURPOSE, WARRANTY OF MERCHANTABILITY OR ANY OTHER WARRANTY, EXPRESSED OR IMPLIED, IS MADE CONCERNING THE GOODS DESCRIBED OR THE INFORMATION PROVIDED HEREIN. The information provided herein relates only to the specific product designated and may not be applicable when such product is used in combination with other materials or in any process. The user should thoroughly test any application before commercialization. Nothing contained herein constitutes a license to practice under any patent and it should not be construed as an inducement to infringe any patent and the user is advised to take appropriate steps to be sure that any proposed use of the product will not result in patent infringement. See SDS for Health & Safety Considerations.

Arkema has implemented a Medical Policy regarding the use of Arkema products in medical devices applications that are in contact with the body or circulating bodily fluids (http://www.arkema.com/en/social-responsibility/responsible-product-management/medical-device-policy/index.html) Arkema has designated medical grades to be used for such medical device applications. Products that have not been designated as medical grades are not authorized by Arkema for use in medical device applications that are in contact with the body or circulating bodily fluids. In addition, Arkema strictly prohibits the use of any Arkema products in medical device applications that are implanted in the body or in contact with bodily fluids or tissues for greater than 30 days. The Arkema trademarks and the Arkema name shall not be used in conjunction with customers' medical devices, including without limitation, permanent or temporary implantable devices, and customers shall not represent to anyone else, that Arkema allows, endorses or permits the use of Arkema products in such medical devices.

It is the sole responsibility of the manufacturer of the medical device to determine the suitability (including biocompatibility) of all raw materials, products and components, including any medical grade Arkema products, in order to ensure that the final end-use product is safe for its end use; performs or functions as intended; and complies with all applicable legal and regulatory requirements (FDA or other national drug agencies). It is the sole responsibility of the manufacturer of the medical device to conduct all necessary tests and inspections and to evaluate the medical device under actual end-use requirements and to adequately advise and warn purchasers, users, and/or learned intermediaries (such as physicians) of pertinent risks and fulfill any postmarket surveillance obligations. Any decision regarding the appropriateness of a particular Arkema material in a particular medical device should be based on the judgment of the manufacturer, seller, the competent authority, and the treating physician.

Arkema Coating Resins

420, rue d'Estienne d'Orves 92705 Colombes Cedex - France arkema.com - **arkemacoatingresins.com**

