SYNOCURE® 886 S 70

GENERAL INDUSTRY

SYNOCURE®

Product Application details	SYNOCURE [®] 886 S 70 is an acid functional acrylic resin designed to crosslink at room temperature with epoxy resins to give high solids content isocyanate-free two-pack coating systems. Coatings based on this resin are especially suitable for protection and maintenance in areas where rapid drying, hardness and abrasion resistance are required. SYNOCURE [®] 886 S 70 has been designed to react with economic bisphenol A type epoxies and still maintain good exterior durability.	
Performance Benefits	 Coatings formulation with VOC at or below 420g/l at application viscosity Fast drying Good exterior durability 	
Polymer Type	Solventborne Acrylic	
Sales Specifications	Solid Content at 125°C, % (ISO 3251)	68 - 72
	Viscosity at 25°C, mPa.s (ISO 3219)	3000 - 6000
	Colour, Gardner scale (ISO 4630)	5 max
	Acid value, mg KOH/g (ISO 2114)	44 - 52
	Volatile	2:1 xylene : n-butanol
	Flash point, °C (ISO 3679)	24
Other	Density / Specific Gravity at 20°C, g/ml (ISO 2811)	1.01
Characteristics ¹	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	RECOMMENDATIONS FOR USE SYNOCURE® 886 S 70 is designed for use with low viscosity epoxy resins of epoxy equivalent weight 180-190 (1) Active hydrogen equivalent weight of SYNOCURE® 886 S 70 is 1145 based on solid resin. A stoichiometric mixing ratio of 1/1 to 1.25 / 1 epoxy / active hydrogen equivalents is recommended although minor deviations from this will have little effect on performance. This isocyanate-free system is suitable for use with a wide range of both organic and inorganic pigments. As with other reactive two-component systems it is strongly recommended that all pigments are checked for stability with the system before commercialisation.	
	<u>SOLUBILITY</u> Aromatic hydrocarbons such as xylene together with minor proportions of esters and alcohols are the most suitable.	
	<u>OTHER ADDITIVES</u> Hindered amine light stabilisers (HALS) (2) are strongly recommended as additives for these acrylic/epoxy systems. SYNOCURE [®] 886 S 70 should only be used in applications consistent with the above recommendations. Proposals to use the resin in alternative systems should be discussed with Arkema before any action is taken.	
	Notes: (1) Araldite [®] GY250 (Hunstman) or Epikote™ Resin 828 (Momentive), (2) Tir total resin solids)	nuvin® 292 (Ciba) at 2% (based on

Product Safety	Please refer to the corresponding Safety Data Sheet.
Storage &	SYNOCURE [®] 886 S 70 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 12 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

