ENCOR® 3250

ENCOR[®] 3250 is an acrylic copolymer emulsion, free from APE surfactants. Product ENCOR[®] 3250 is well suited to be used in formulations for upholstery, where it provides a high **Application details** softness, an easy plate release and a very good embossing retention. ENCOR[®] 3250 finds also use in printing applications. Clear, soft and tack-free film • Performance High flexibility **Benefits Polymer** Acrylic Copolymer • Type Solid Content at 105°C, % (ISO 3251) 36 - 38 Sales pH (ISO 976) 6.5 - 8.0 **Specifications** Viscosity at 23°C, mPa.s (Brookfield RVT, 100rpm, sp2) (ISO 2555) 20 - 100 Stabilizing system А Minimum Film Formation Temperature, °C (ISO 2115) < 5 Tq (DSC), °C -10 Other Density / Specific Gravity at 23°C, g/ml (ISO 2811) 1.05 Characteristics¹ Average Particle size, nm (ISO 13321) 140 1 The data provided for these properties are typical values, intended only as guides, and should not be construed as sales specifications ENCOR[®] 3250 can be used alone or in combination with other acrylic and/or polyurethane **Formulation** binders. Guidelines



ARKEMA COATING RESINS

Product Safety	Please refer to the corresponding Safety Data Sheet.
Storage & Handling	ENCOR [®] 3250 should be stored indoors in the original, unopened and undamaged container, in a dry place at storage temperatures between 5°C and 30°C. Exposure to direct sunlight should be avoided. The product is protected to prevent any microbial deterioration during normal conditions of storage but care should be taken to avoid accidental contamination during subsequent handling and processing. In the above mentioned storage conditions the shelf life of the resin will be 6 months from the

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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 persent 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

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