

SUPER GELKYD® 6008 WD 45

ARCHITECTURAL COATINGS

ARKEMA COATING RESINS

Product Application details

SUPER GELKYD® 6008 WD 45 is a pumpable thixotropic alkyd especially designed for use in high durability exterior coatings.

It combines the proven durability of SYNOLAC® 6005 WDA 70 with the rheological control of the SUPER GELKYD® technology. The result is a flexible thixotropic resin supplied in bulk that can be used as sole binder or alternatively blended with SYNOLAC® 6005 WDA 70.

The resin is supplied in low aromatic white spirit to enable the formulation of coatings that do not require adverse environmental labelling.

SUPER GELKYD® 6008 WD 45 can be formulated with polar solvents, alkyds (whatever their polarity) and a wide range of difficult pigments, which are not compatible with polyamide-based products.

The resin is primarily recommended for use in thixotropic exterior decorative coatings woodstains, where it may be incorporated as the major proportion of the binder according to the structure required.

Polymer Type

- Thixotropic Alkyd

Sales Specifications

Solid Content at 125°C, % (ISO 3251)	44 - 46
Viscosity at 25°C, mPa.s (at 2500 s-1) (ISO 3219)	300 - 500
Colour, Gardner scale (ISO 4630)	8 max

Other Characteristics¹

Appearance	Slightly hazy, pumpable thixotropic resin
Volatile	Low aromatic white spirit
Flash point, °C (ISO 3679)	40
Density / Specific Gravity at 25°C, g/ml (ISO 2811)	0.88
Type of fatty acid	Linoleic rich
Fatty Acid content, %	52
Gel Strength at 25°C, internal method LQ 37, g.cm	80 - 200

¹ The data provided for these properties are typical values, intended only as guides, and should not be construed as sales specifications

Formulation Guidelines

ACTIVATION

The full structure is attained only with the addition of polar solvent, at the end of paint manufacture. Propylene glycol is the preferred activating solvent. Normally between 2-3% of propylene glycol, calculated on total solid resin content, is sufficient to fully develop the structure. In thixotropic paints based on polyamide technology, the presence of highly polar solvents such as alcohols would destroy structure, the presence of alcohols in SUPER GELKYD® 6008 WD 45 based coatings is crucial to structure and development.

DRIERS

SUPER GELKYD® 6008 WD 45 requires metal driers to accelerate the autoxidation process. A suitable combination of driers for use in systems containing SUPER GELKYD® 6008 WD 45 is: 0.06% cobalt, 0.09% zirconium and 0.1% calcium calculated as metals on solid resin. Depending on the formulation (clear, pigmented, etc...) and on the application, the loading of each drier may be increased or reduced in order to achieve the appropriate drying/hardness profile. The use of an anti-skinning agent is essential to prevent in-can skinning of the finished product.

STORAGE REMARK

The aspect end of this grade can differ from drum to drum, depending on the production and storage/transportation history. This is inherent to SUPER GELKYD® 6008 WD 45 and does not alter the application properties after activation.

SUPERGELKYD®
BY ARKEMA

Product Safety

Please refer to the corresponding Safety Data Sheet.

Storage & Handling

SUPER GELKYD® 6008 WD 45 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.

In the above mentioned storage conditions the shelf life of the resin will be 6 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

ARKEMA
INNOVATIVE CHEMISTRY