

SNAP® 2709

ARCHITECTURAL COATINGS

(FORMERLY E 20509)

ARKEMA COATING RESINS

Product Application details

SNAP® 2709 is a low MFFT acrylic emulsion with a morphology specifically designed to allow an excellent film formation. It is particularly adapted to the formulation of trim enamels for exterior and interior applications.

SNAP® 2709 is formulated without alkylphenolethoxylates or formaldehyde.

SNAP® 2709 has been developed to be used as binder for both interior and exterior trim enamels. Its specific & controlled structure ensures a stable minimum film formation temperature as well as a very good film formation with no coalescing agent even at low temperatures.

In terms of film properties, SNAP® 2709 allows to get a coating exhibiting a good compromise of flexibility and hardness and a very good block resistance. The final properties of the coating are achieved extremely rapidly (within 24 hours of drying) and are stable with time as no coalescing agent (VOC or non VOC) is used. SNAP® 2709 is a binder of choice for durable exterior coatings.

Performance Benefits

- Excellent blocking resistance (even at high temperatures)
- Excellent hardness development
- Excellent wet and dry adhesion (especially onto aged alkyds)
- Excellent flexibility
- Easy controlled rheology
- Good pigment compatibility

Polymer Type

- Acrylic Emulsion

Sales Specifications

Solid Content % (ISO 3251)	44 - 46
pH (ISO 976)	7.5 - 8.5
Viscosity at 23°C, mPa.s (Brookfield RVT , 20rpm) (ISO 2555)	500 max

Other Characteristics¹

Stabilizing system	A / NI
Minimum Film Formation Temperature, °C (ISO 2115)	5
Density / Specific Gravity, g/ml (ISO 2811)	1.06
Average Particle size, nm (ISO 13321)	90 - 100

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

Formulation Guidelines

RHEOLOGY

SNAP® 2709 is easy to formulate as its reactivity to thickeners is balanced and controlled. It is possible to achieve the desired rheological profile (more or less Newtonian) depending on the targeted application.

This controlled thickeners reactivity allows to optimise the levelling of SNAP® 2709-based paints and varnishes.

Rheology can be adjusted using associative thickeners such as HEUR thickeners : Polyurethane thickeners such as Coapur™ XS 22 (1), Coapur™ 830W (1), Coapur™ 2025 (1) which are efficient; Hydrophobically Modified PolyEther thickeners, e.g. Aquaflo® NHS 300 (2), Aquaflo® NMS 450 (2), are also suitable. HASE thickeners, such as Rheotech™ 2000 (1) may also be suitable for satin paints.

OTHER ADDITIVES

Defoamers such as Byk®-022 (3), Byk®-028 (3), Byk®-093 (3), FoamStar® ST 2438 (4) as well as FoamStar® ED 2522 (4) are suitable.

SNAP® 2709 formulations pH should be adjusted with ammonia : it is not recommended to formulate it with non-volatile amines or soda.

Notes: (1) Coatex, (2) Ashland Specialty Ingredients, (3) Byk, (4) BASF

Product Safety

Please refer to the corresponding Safety Data Sheet.

Storage & Handling

SNAP® 2709 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 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

The logo for ARKEMA, with the word in a bold, sans-serif font. The 'A', 'R', 'K', 'E', and 'M' are in dark blue, while the 'A' at the end is in green.