SILIPORITE® NK 30 AP POWDER

Product Data Sheet

PRODUCT BENEFITS

SILIPORITE® NK 30 AP is a synthetic zeolite type A of crystalline structure in Potassium-Sodium Aluminosilicate form having an effective pore opening approximately of 3 angströms (3A).

SPECIFICATIONS

	Unit	Values	Test control
Particles retained on 20 µm mesh size	% weight	2.0 max	009
Delta T	°C	33 min	004
Loss on ignition (1 hour at 950°C)	% weight	2.0 max	001
Water adsorption capacity at 23°C +/- 2°C - 10 % Relative Humidity	% weight	21.5 min	002B

TYPICAL PHYSICAL PROPERTIES

	Unit	Values	Test control
Pot-life at 20°C	Min	30 min	422
Viscosity Brookfield (50 % wt castor-oil paste)	mPa-s	10 000	422
Desorption N ₂	cm ³ / 100 g	15	015
Settled density		0.50 – 0.85	005



SILIPORITE® NK 30 AP POWDER

APPLICATIONS

SILIPORITE® NK 30 AP can be used for all purposes where a drying agent, easy to use and of high efficiency is needed.

- Buildina:
 - Insulating Glass Units to avoid water condensation while minimizing Nitrogen desorption.
 - Polyurethane systems (paints, coating, sealant) for the application where foaming should be avoided.
- Fine Chemicals / Pharmaceuticals:
 - Water adsorption in chemical equilibrium reactions.

STANDARD PACKAGING

SILIPORITE® NK 30 AP is delivered in waterproof drums, with inner polyethylene bag, or in bag as

follows:

Bags Net weight: 20 kg / 1 200 kg
Metallic drum Net weight: 80 kg / 640 kg
Big bag Net weight: 900 kg

FOR MORE INFORMATION

Contact your ARKEMA representative or ARKEMA's Molecular Sieves Department:

Telephone: +33 (0) 1 49 00 38 00 Fax: +33 (0) 1 49 00 38 03

E-mail: molecular.sieves@arkema.com

Version: 7 - March, 2017

Registered trademark of ARKEMA.2015 ARKEMA. All rights reserved.

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/responsibleproduct-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.

