

FDA Food Contact

February 2026, Version 3.0
Previous version obsolete



Applicable to: Sultron™ LSG PPSU R5100 black BK937 stock shapes

FDA

We hereby provide the following information based on the compliance status of the materials used at present by Mitsubishi Chemical Advanced Materials for the manufacture of the product mentioned above, as set out in the FDA regulations that apply in the United States of America, for plastic materials and articles intended to come in contact with foodstuffs:

The product complies with the requirements of the Food Contact Notification (FCN) Numbers 83 and 1215 on "Poly (oxy(1,1'-biphenyl) -4,4'- diyoxy -1,4-phenylenesulfonyl -1,4-phenylene)" of the American Food and Drug Administration's (FDA) Inventory of Effective Notifications for F for the use as articles or components of articles intended for repeated use in contact with food, with all food types I to IX, and under conditions of use B to H as defined in Tables 1 and 2 of 21 CFR 176.170 (c).

This product has not been evaluated for the use in contact with infant formula and/or human (breast) milk.

USDA

USDA does not approve materials. USDA requires that articles are made of FDA food contact compliant materials.

EU 10/2011

The product is not being manufactured or tested according to EU 10/2011 requirements. For EU 10/2011 compliance, specific "Food Grade" items are available to purchase as manufactured and tested according to the regulation. These specific items will have "...FG..." in the product names, are labelled with the "fork and glass" icon and will arrive with Declarations of Compliance (DoC).

NSF

The product is not NSF listed or NSF approved. NSF 51 compliance is based on the FDA compliance of the raw materials. If NSF 51 compliance is needed for the product, the farthest end user must contact NSF to begin the certification process. NSF will then send Mitsubishi Chemical Advanced Materials a "Request for Formulation" form, which will be filled out and returned to NSF.

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The products should neither be used in invasive or implantable medical devices that are intended to be introduced in and remain inside the human body or replace an epithelial surface or the surface of the eye for more than 24 hours (or 30 days for Ketron™ LSG PEEK-CLASSIX™ white respectively), nor should they be used as critical components of medical devices that are essential to the continuation of human life.

The products do not fall within the applicable regulations of a medical device.

This statement is subject to change without notice and becomes obsolete in case of any changes to the regulation referenced, or in case of any changes to the product's composition, or in case a new version is issued. New versions of this statement will either be published on our website or are provided to you upon request. Please always consult either our website or contact your MCAM sales representative to request the current version of this statement. MCAM undertakes no express or implied obligation to communicate the expire of previous versions nor the issue of new versions of this statement.

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PEEK-CLASSIX™ is a trademark of Invivio Ltd., Technology Centre, Hillhouse International, Thornton Cleveleys, Lancashire FY5 4QD, United Kingdom.

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