15-11-3 polymirae

A Clyrell RC5056 Data Sheet



Food container

Clyrell RC5056作为高透明、高光泽无规共聚物(Random Copolymer),是一种适合用于食品容器、家庭用品等注 塑成型的产品。

Clyrell RC5056符合美国食品药品管理局(FDA)21 CFR 177.1520食品包装规定。

产品特点

高透明/刚性与韧性均衡/不包括荧光增白剂和BPA/可以作为食品容器使用

典型应用

婴儿奶瓶/家庭用品/注塑产品/食品容器/ISBM

物性项目 (a)	ASTM	測量
Melt Flow Rate (230 °C,2.16kg) (g/10min)	D 1238L	10
Flexural Modulus (kg/cm²)	D 790	14000
Tensile Strength at Yield (kg/cm²)	D 638	350
Elongation at Yield (%)	D 638	10
Izod Impact Strength(23°C) (kgcm/cm)	D 256	5
Rockwell hardness (R scale)	D 785	102
Vicat softening point (℃)	D 1525	134
HDT(0.46N/mm2) (℃)	D 648	100
Haze (%)	D 1003	9 (2mm)

- 1) The above values are only for guide and shall not be construed as specifications.
- 2) Before using a Polymirae product, users should make their own independent determination that the product is suitable for the intended use and can be used safely and legally.
- 3) SELLER MAKES NO WARRANTY; EXPRESS OR IMPLIED (INCLUDING ANY WARRANTY OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE OR ANY WARRANTY) OTHER THAN AS SEPARATELY AGREED TO BY THE PARTIES IN A CONTRACT.
- 4) The use of this product(s) is prohibited in:
- (i) any U.S. FDA Class I, Health Canada Class I, and/or European Union Class I medical devices, without prior notification to Seller for each specific product and application; or
- (ii) the manufacture of any of the following, without prior written approval by Seller for each specific product and application: U.S. FDA Class II Medical Devices; Health Canada Class II or Class III Medical Devices; European Union Class II Medical Devices; film, overwrap and/or product packaging that is considered a part or component of one of the aforementioned medical devices; packaging in direct contact with a pharmaceutical active ingredient and/or dosage form that is intended for inhalation, injection, intravenous, nasal, ophthalmic (eye), digestive, or topical (skin) administration; and tobacco related products and applications.
- 5) Additionally, the use of this product(s) is prohibited in:
- (i) U.S. FDA Class III Medical Devices; Health Canada Class IV Medical Devices; European Union Class III Medical Devices;
- (ii) applications involving permanent implantation into the body;
- (iii) life-sustaining medical applications; and
- (iv) lead, asbestos or MTBE related applications.
- 6) Users should review the applicable Material Safety Data Sheet before handling the product.
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