

> APPLICATION BULLETIN

# Mevopur<sup>™</sup> Healthcare Colorants and Formulations and Mevopur<sup>™</sup> Healthcare Functional Additives for Bormed<sup>™</sup> Resins

Mevopur<sup>™</sup> Healthcare Colorants and Formulations and Mevopur<sup>™</sup> Healthcare Functional Additives help medical device, diagnostics and pharmaceutical packaging sectors meet the growing challenges of product consistency, compliance and reliability. They are adapted to your manufacturing process and available as either a 'ready-to-use' formulation or a concentrate that is diluted in the Bormed<sup>™</sup> resin of choice.

## **MEVOPUR COLORANTS**

- Standard colors in PP and PE to shorten time-to-market
- Custom colors matched to Pantone, RAL or other color reference
- White colorants for pharmaceutical packaging for LDPE, HDPE and PP with opacity to reach USP chapter <671>

## **MEVOPUR FUNCTIONAL ADDITIVES**

- Antistatic formulation with good flow/impact for injection molding
- Amide- and amine-free antistatic for film in contact with API (Active Pharmaceutical Ingredients)
- UV blocking for transparent PP and PE packaging to meet USP chapter <671>

- Non-migrating friction reduction/processing aid for PE and PP
- Laser marking/welding additive for fast, reliable marking/assembly
- MVTR (Moisture Vapor Transmission Rate) reduction for HDPE to improve shelf-life
- Thermal protection of the polymer during processing
- Gamma/e-beam sterilization protection for PP (up to 50kGy) with color compensation for reduced yellowing

## **REGULATORY SUPPORT**

- Raw materials tested to:
  - ISO 10993-1 and USP <87> <88> biological evaluation
  - European Pharmacopeia 3.1.3/3.1.5 (polyolefin)
  - USP <661.1> (polyethylene)
  - ICH Q3D elemental impurities
- Registered Drug Master File (Type III) and/or Device Master File
- Food contact established with FDA/EU\*



\* FDA/EU compliance information available upon request Bormed<sup>™</sup> is a registered trademark of Borealis



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It is the responsibility of the medical device manufacturer and the person placing the medical device on the market to ensure compliance of the medical device with all applicable laws and regulations, including the suitability of all raw materials and components used for its manufacture.

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