

Styrene Allyl Alcohol

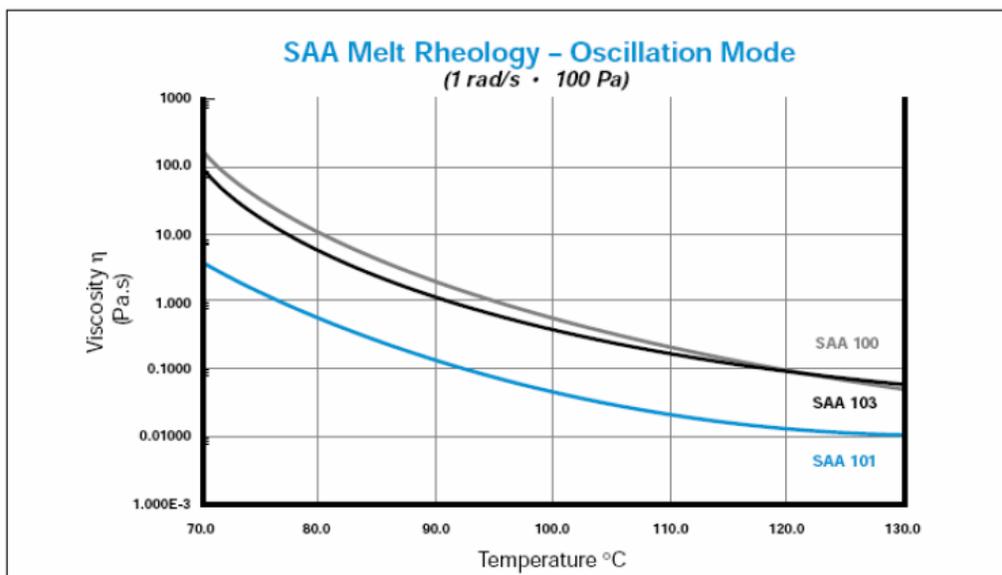
SAA-100™ and SAA-101™ Copolymers
Hydroxy Functional Resins for Toners

- Features**
- Low Viscosity & Melt Temperatures
 - High Functionality
 - High Thermal Stability

- Benefits**
- Improved Adhesion
 - Brilliant Color
 - Heat Resistance
 - Water Resistance

Typical Properties

COMPOSITION	SAA-100	SAA-101
Bound Styrene, mole %	70	60
Bound Allyl Alcohol, mole %	30	40
PHYSICAL PROPERTIES		
Physical Form	Pastilles	Pastilles
Appearance	White	White
Color, APHA (30% in MEK)	40	40
Molecular Weight, GPC (Number Average, Mn)	1,500	1,200
Molecular Weight, GPC (Weight Average, Mw)	3,000	2,500
Pd (Polydispersity)	2.0	2.1
Hydroxyl Number (mg KOH/gm)	210	255
Equivalent wt.	267	220
Acid Number (mg KOH/gm)	<0.1	<0.1
Tg, °C	62	57
TS (Softening Point), °C	79	73
T1, °C	90	78
T2, °C	103	95
T1/2, °C	105	98
Viscosity (cps @ 100°C)	317	338
Specific Gravity	1.05	1.09



Before using a product sold by a company of the LyondellBasell family of companies, users should make their own independent determination that the product is suitable for the intended use and can be used safely and legally.

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.

This product(s) may not be used 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; tobacco related products and applications, electronic cigarettes and similar devices, and pressure pipe or fittings that are considered a part or component of a nuclear reactor. Additionally, the product(s) may not be used in: (i) U.S. FDA Class III Medical Devices; Health Canada Class IV Medical Devices; European 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. All references to U.S. FDA, Health Canada, and European Union regulations include another country's equivalent regulatory classification.

Users should review the applicable Safety Data Sheet before handling the product.

Alkylate, Duopac, Duoprime, Filmex, MPDIOL, Polymeg, SAA-100, SAA-101, TBAC, Tebol, T-Hydro, and Tufflo are trademarks owned or used by the LyondellBasell family of companies.

Duopac, Duoprime, Filmex, MPDIOL, Polymeg, Tebol, T-Hydro and Tufflo are registered in the U.S. Patent and Trademark Office.

Houston, Texas, USA | Tel: +1 713 309 7200 or toll-free within USA +1 888 777 0232

Rotterdam, The Netherlands | Tel: +31 10 275 5500

Hong Kong, China | Tel: +852 2882 2668

© LyondellBasell Industries Holdings, B.V. 2011

www.lyondellbasell.com

2593-V2-0511
Supersedes 2593-V2-0104