Application Data MPDiol® Glycol MPDiol Polyester Polyols for Urethane Coatings

Adipic Acid

Trimethylolpropane

General Information	characteristics of many of th these less likely to crystalliz made from MPDiol exhibit hy	aracteristics of diprimary diols but e branched diols. Polyol branching e upon sitting, increasing the shelf ybrid properties, somewhere betwee ring, flexibility, and toughness as co	g makes polymers resulting from life of the paint. Polyurethanes een a long chain and branched			
Key Features and Benefits	Faster Esterification Rates	Esterification rates for the same a	cid value are up to 30% faster with			
	Non-crystallizing Coatings: The unsymmetrical nature of the MPDiol molecule prevents close packing of the resin polymer chains. This is evidenced by longer shelf life of the product without hazing, cloudiness, or particulates developing in the solutions.					
	<i>Fast Curing</i> : The resulting polyester polyol is primary in nature and tends to react quickly with diisocyanates. MPDiol exhibits similar reactivity to that of 1,3-propandiol and the resulting films show performance equal to commonly used polyols.					
Polyester Polyol Synthesis and Propert Benefits	was performed at 213 °C. S up of each resin is summaria The primaries were used as as structural and spatial arr synthesis of diprimary versu propylene glycol (PG) based higher temperatures can so order to compensate for eva were of comparable acid nu yielded lower hydroxyl polyc chromatography (GPC) resu yielding broad bands sugges	e use of a catalyst. The esterification esized using this method. The make a study were primary and secondary. reactivity. Secondary diols were used ences in the amounts used in the ned from past experiences in making act slower than primary alcohol; ater amount of diol is necessary in ured that the polyols synthesized and trimethylpentane diol (TMPG) actions. Gel permeation y weight distributions. The increased the higher molecular weight products				
	Resin Components	Diprimary Diols moles	Primary/Secondary Diols			
	Diol	4.30	4.51			
	Isophthalic Acid	2.00	2.00			

2.00

0.70

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Table 1. Generic polyester polyol synthesis and resin composition used for comparative testing of commonly used primary and primary/secondary hydroxyl diols and MPDiol.

2.00

0.70



Acid Mn Mw **Diol Component** Hydroxyl Mw/Mn Number Number mg mg KOH/g KOH/g MPDiol 128 5.3 1260 2540 2.02 Ethylene glycol 162 6.2 920 2010 2.18 1.3-Propane diol 150 6.3 1100 2120 1.94 1.4-Butane diol 84 11 1840 4780 2.60 1.6 Hexane diol 1370 2620 1.91 131 6.8 Neopentyl glycol 130 8.8 1160 2200 1.89 Propylene glycol 132 6.6 920 1880 2.03 1.3-Butane diol 148 6.2 1050 2070 1.96 Trimethylpentane diol 94 11.0 1350 5240 3.87

 Table 2. Properties of resulting polyester polyols from several primary and primary/secondary hydroxyl glycols as compared to MPDiol

Substituting MPDiol one to one in this study provided a means for direct comparison and quantification of the structure reactivity correlation between MPDiol and other diols with similar characteristics. We measured the effects of diol selection on the synthesis rate and properties of the polyester polyol. The reactivity profiles of the diols were plotted to compare the reactivity of MPDiol to these. (See Figure 1) In this plot the increase in curvature is indicative of an increase in rate. The results suggest that ethylene glycol reacts the fastest under the esterification conditions followed by MPDiol and 1,3-propanediol. Even the branched nature of MPDiol does not interfere with its reactivity as evidenced by it reacting faster than linear 1,4-BDO and much faster than NPG

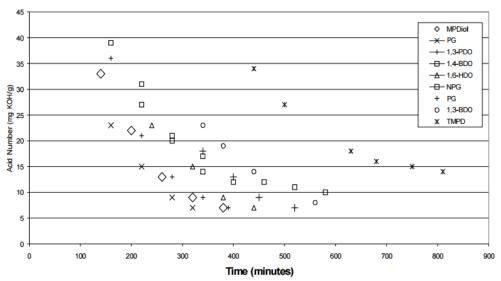


Figure 1. Polyesterification reactivity profile of commonly used primary and primary/secondary hydroxyl glycols plotted against MPDiol glycol for comparison.



A solvent compatibility or tendency for crystallize test was performed on the polyester polyols. Crystallization is evidenced by a cloudy, hazy or sometimes particulate in the resin. Over a period of three weeks only the MPDiol, 1,6-HDO, and all of the secondary/primary diol based polyesters remained clear. The results support the belief that branched polyols hinder close packing of the polyester chains and in this way inhibiting crystallization. A summary of the results is shown in Table 3. MPDiol is an attractive intermediate for use in polyester synthesis when combined with its reactivity and inherent compatibility with organic solvents it can reduce processing costs and increase shelf life of the final product.

Diol Component	Gardner-Holt Viscosity	Gardner Color	Clarity after 3 weeks
MPDiol	Z-4	1	Clear
Ethylene glycol	Z-5	1	Cloudy
1.3-Propane diol	Z-3	2	Hazy
1.4-Butane diol	Z-6	1	Cloudy
1.6-Hexane diol	Z-1	1	Clear
Neopentyl glycol	Z-6	1	Hazy
Propylene glycol	Z-5	1	Clear
1.3-Butane diol	Z-2	1	Clear
Trimethylpentane diol	Z-8	4	Clear

Polyurethane Polymers

Table 3. Three-week stability testing using 90% polyester polyol solids in 1:1 methyl ethyl ketone and propylene glycol methyl ether blend on a variety of polyester polyols including a MPDiol glycol based polyester.

Clear coat, two-pack aliphatic polyurethane coatings were prepared using the basic formulation shown in Table 4. Curing conditions were kept constant so that a direct comparison could be made between polyols. Methylethyl ketone (MEK) double rub performance of the films showed that these cured completely. The films showed good flexibility and resilience as well as toughness and hardness. The performance data suggests that the MPDiol based urethanes behave similar to 1,3-propandiol (1,3-PDO) since it cures as fast as 1,3-PDO and demonstrates comparable hardness. On closer inspection it is really a hybrid since it is a primary hydroxyl with a branched backbone. The film shows a high pencil hardness signifying it is as hard as a conventional branched system. As expected softer more resilient films than expected were obtained commensurate with conventional long chain polyols and evidenced by its sward hardness value.

Summary

MPDiol shows to be a viable alternative diol for the synthesis of polyester polyols. It affords colorless polyols while allowing for processing temperature latitudes that other diols cannot afford. The process time is shorter and urethane resin synthesis faster than similar diols. The resulting coating performance is comparable to commonly used diols and more stable in organic solutions than many of the diprimary diol commonly used for these applications. It is expected that the films should yield a weatherable and humidity stable coating. (See Application Data sheet titled "Weatherability of Melamine Cured Polyesters").



Formulation	Weight (gm)					
Part A						
Resin Solution	40.0					
Glycol Ether PM Acetate ¹	5.0					
Methyl Ethyl Ketone	5.0					
Xylenes	5.0					
FC-430 (Flow Control Additive) ²	0.28					
T-12 (Catalyst) ²	0.20					
Part B						
Desmodur® N-3390 (90% NVM) ⁴	NCO:OH 1.10: 1.00					
Polyol	Properties of Polyurethane Film ²					
	MEK 2X RUBS	Pencil Hardness	Sward Hardness			
Primary Diols						
MPDiol	>100	2H	8			
EG	>100	3H	20			
1,3-PDO	>100	2H	8			
1,4-BDO	>100	Н	2			
1,6-HDO	>100	F	2			
NPG	>100	Н	32			
Secondary Diols						
PG	>100	F	24			
PG 1,3-BD0	>100 >100	F H	24 16			

film properties for several primary/secondary hydroxyl based polyester polyols used in formulating polyurethane coatings.

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2593-V2-0511 Supersedes 2593-V2-0104