

【DESCRIPTION】

【Disclosure Title】

**POLYURETHANE UREA ELASTIC YARN HAVING IMPROVED DYEABILITY
AND MANUFACTURING METHOD THEREFOR**

5 **【Technical Field】**

The present disclosure relates to a polyurethane urea elastic yarn having improved dyeability and a method for manufacturing the same, and in particular, to a polyurethane urea elastic yarn having improved dyeability for an acid dye
10 and a method for manufacturing the same.

【Background Art】

A polyurethane fiber having elastic properties has been widely used in the textile industry. A 'spandex' fiber is used as a general term meaning such an elastic fiber, and generally
15 refers to a synthetic fiber in which a polyurethane bond is 85 % or more in the structure forming the fiber.

Such a polyurethane fiber is generally prepared by a polymerization reaction first of reacting a polyol, such as a high molecular weight diol compound with an excess diisocyanate
20 compound to obtain a prepolymer having an isocyanate group on both ends of the polyol, and a polymerization reaction second of dissolving the prepolymer in a proper solvent and then adding a diamine-based or diol-based chain extender to the solution for reacting.

A polyurethane fiber may be used in combination with various other fibers such as acryl, wool, cotton and silk, and have been generally used for purposes requiring elasticity such as underwear, swimwear and stockings.

5 Due to such unique properties having elasticity, a polyurethane fiber has been actively used in various applications, and as the application expands, new additional properties have been continuously required for existing polyurethane fibers. A number of polyurethane fibers having
10 more enhanced heat resistance and resilience have been developed so far, and demands for polyurethane fibers having high thermosetting properties and polyurethane fibers having enhanced dyeability are also gradually increasing.

A polyurethane fiber generally has a problem of poor
15 dyeability, and general technologies for enhancing dyeability for an acid dye include 1) a method of introducing a tertiary nitrogen atom into a polyurethane polymer chain (Japanese Patent Publication No. S62-23097 B), 2) a method of introducing a salt of an organic acid or inorganic acid and a tertiary
20 amine to a polyurethane polymer chain (Japanese Patent Publication No. S50-17520 B), 3) introducing a tertiary or quaternary nitrogen atom to an end of a polyurethane polymer chain (Japanese Patent Publication No. S44-16386 B), 4) a method of using a low molecular diamine as a chain extender

(Japanese Patent Application Laid-Open Publication No. S59-108021 A), and the like.

However, gelation readily occurs in the reaction in the above methods 1) and 2), and the method 3) has a problem in
5 that the amount of the introduced nitrogen atom is difficult to control. In addition, the method 4) of using a low molecular diamine as a chain extender may enhance dyeability, but has a problem of decreasing softness and elasticity.

In addition thereto, Korean Patent Application Laid-Open
10 Publication Nos. 10-2009-0118997 A and 10-2005-0070652 A have tried to improve dyeability by dyeing with a specific blue acid dye containing a polymer having a maleimide structure formed with a maleimide unit or adding a talc compound, respectively, however, there is a limit to sufficiently enhance dyeability of
15 polyurethane.

【Disclosure】

【Technical Problem】

The present disclosure is directed to providing a polyurethane urea elastic yarn having improved dyeability for
20 an acid dye, and a method for manufacturing the same.

【Technical Solution】

In view of the above, a polyurethane urea elastic yarn having improved dyeability for a dye according to one embodiment of the present disclosure is obtained by reacting,

(a) a polyol including a first polyol and a second polyol; (b) at least one of diisocyanates; (c) at least one of diamine chain extenders; and (d) at least one of amine chain terminators, wherein the second polyol is polyethylene glycol, and the first polyol and the second polyol are materials different from each other.

In addition, the second polyol is mixed in 5.0 mol% to 20.0 mol% with respect to the total amount of polyol.

In addition, the polyurethane urea elastic yarn has an amine-terminated number of 20.0 meq/kg to 45.0 meq/kg.

A method for manufacturing a polyurethane urea elastic yarn having improved dyeability according to the present disclosure includes,

(a) preparing a polyurethane prepolymer by bringing a polyol including a first polyol and a second polyol into contact with at least one of diisocyanates; (b) adding a solvent to the prepolymer in the step (a); (c) bringing the product in the step (b) into contact with at least one of diamine chain extenders and at least one of amine chain terminators; and (d) preparing a polyurethane urea elastic yarn by spinning the product in the step (c), wherein the second polyol is polyethylene glycol, and the first polyol and the second polyol are materials different from each other.

In addition, the second polyol is mixed in 5.0 mol% to 20.0 mol% with respect to the total amount of polyol.

In addition, the polyurethane urea elastic yarn has an amine-terminated number of 20.0 meq/kg to 45.0 meq/kg.

A fabric according to the present disclosure is a fabric having enhanced dyeability for an acid dye by knitting the
5 polyurethane urea elastic yarn according to one embodiment of the present disclosure and nylon.

【Advantageous Effects】

By mixing polyethylene glycol in a step of preparing a polyurethane prepolymer, accessibility of an acid dye to a
10 polyurethane urea elastic yarn is enhanced by enhancing hydrophilicity of the polyurethane urea elastic yarn, and as a result, dyeability of the polyurethane urea elastic yarn for the acid dye can be enhanced, and an advantage of dye color deepening of a fabric obtained by knitting the polyurethane
15 urea elastic yarn of the present disclosure with nylon can be expected as well.

【Mode for Disclosure】

Hereinafter, the present disclosure will be described in detail.

20 The present disclosure relates to a polyurethane urea elastic yarn having improved dyeability, the polyurethane urea elastic yarn including a product obtained by reacting (a) a polyol including a first polyol and a second polyol; (b) at least one of diisocyanates; (c) at least one of diamine chain
25 extenders; and (d) at least one of amine chain terminators,

wherein the second polyol is polyethylene glycol, and the first polyol and the second polyol are materials different from each other.

In addition, the present disclosure relates to a method
5 for manufacturing a polyurethane urea elastic yarn having improved dyeability, the method including (a) preparing a polyurethane prepolymer by bringing a polyol including a first polyol and a second polyol into contact with at least one of diisocyanates; (b) adding a solvent to the prepolymer in the
10 step (a); (c) bringing the product in the step (b) into contact with at least one of diamine chain extenders and at least one of amine chain terminators; and (d) preparing a polyurethane urea elastic yarn by spinning the product in the step (c), wherein the second polyol is polyethylene glycol, and the first
15 polyol and the second polyol are materials different from each other.

In the present disclosure, the polyol includes a first polyol and a second polyol. In other words, a first polyol and a second polyol are mixed to be used as the polyol, and the
20 first polyol and the second polyol are materials different from each other.

In the present disclosure, polytetramethylene ether glycol, polypropylene glycol, polycarbonatediol and the like may be used as the first polyol.

In the present disclosure, the second polyol is polyethylene glycol. Polyethylene glycol has excellent hydrophilicity, and therefore, hydrophilicity of the polyurethane urea elastic yarn is enhanced by mixing the polyethylene glycol in the step of preparing a polyurethane prepolymer, which resultantly enhances accessibility of an acid dye to the polyurethane urea elastic yarn, and dyeability of the polyurethane urea elastic yarn for the acid dye may be enhanced.

10 In the present disclosure, the polyethylene glycol is preferably mixed in 5.0 mol% to 20.0 mol% with respect to the total amount of polyol. When using the polyethylene glycol in less than 5.0 mol%, enhancement of dyeability for an acid dye may not be expected, and using the polyethylene glycol in
15 greater than 20.0 mol% may cause serious decline in the properties of the elastic yarn and a decrease in the process applicability.

In the present disclosure, 4,4'-diphenylmethane diisocyanate, hexamethylene diisocyanate, toluene diisocyanate, butylene diisocyanate, hydrogenated p,p-methylene diisocyanate
20 and the like may be used as the diisocyanate, and one or more compounds thereof may be used.

The polyol and the diisocyanate are brought into contact with each other to prepare a polyurethane prepolymer, and a
25 solvent is added thereto. Herein, the solvent is preferably

dimethylacetamide. After that, the result is brought into contact with at least one of chain extenders and at least one of chain terminators to prepare a polyurethane urea solution. The prepared polyurethane urea solution, a product, goes
5 through a fiber spinning process such as dry spinning or melt spinning to prepare the polyurethane urea elastic yarn.

In the present disclosure, the chain extender is at least one of diamines. Examples of the usable diamine include hydrazine, ethylenediamine, 1,2-propanediamine, 1,3-
10 propanediamine, 1,2-butanediamine (1,2-diaminobutane), 1,3-butanediamine (1,3-diaminobutane), 1,4-butanediamine (1,4-diaminobutane), 1,3-diamino-2,2-dimethylbutane, 4,4'-methylenebis-cyclohexylamine, 1-amino-3,3,5-trimethyl-5-aminomethylcyclohexane, 1,6-hexanediamine, 2,2-dimethyl-1,3-
15 diaminopropane, 2,4-diamino-1-methylcyclohexane, N-methylaminobis(3-propylamine), 2-methyl-1,5-pentanediamine, 1,5-diaminopentane, 1,4-cyclohexanediamine, 1,3-diamino-4-methylcyclohexane, 1,3-cyclohexane-diamine, 1,1-methylene-bis(4,4'-diaminohexane), 3-aminomethyl-3,5,5-
20 trimethylcyclohexane, 1,3-pentanediamine (1,3-diaminopentane), m-xylylenediamine and mixtures thereof, but are not limited thereto. Ethylenediamine is preferred as the chain extender.

In the present disclosure, the chain terminator is at least one of amines. Examples of the usable amine may include
25 diethylamine, cyclohexylamine, n-hexylamine and mixtures

thereof, but are not limited thereto. Diethylamine is preferred as the chain terminator.

In the polyurethane urea elastic yarn of the present disclosure, the yarn preferably has an amine-terminated number
5 of 20.0 meq/kg to 45.0 meq/kg. The amine end of the yarn serves as a dyeing site bondable to a dye, and when the yarn has an amine-terminated number of less than 20.0 meq/kg, a sufficient dyeability enhancing effect may not be expected due to the lack of dyeing sites. In addition, when the yarn has an
10 amine-terminated number of greater than 45.0 meq/kg, dyeability is sufficient, however, the amine-terminated number excessively increases, which may cause a serious problem in storage stability of a polymer.

In addition, in order to prevent discoloration and
15 property decline of the polyurethane urea caused by ultraviolet rays, atmospheric smog, a heat treatment process associated with spandex processing, and the like, a sterically-hindered phenol-based compound, a benzofuranone-based compound, a semicarbazide-based compound, a benzotriazole-based compound, a
20 polymeric tertiary amine stabilizer and the like may be properly combined and added to the spinning solution in the present disclosure. In addition to the above-mentioned components, pigments or dyes with a blue color, a complementary color of a yellow color, may be added to the spandex fiber of
25 the present disclosure to reduce yellowishness of the yarn.

Furthermore, the polyurethane urea elastic yarn of the present disclosure may include additives such as titanium dioxide or magnesium stearate in addition to the above-described components.

5 The polyurethane urea elastic yarn of the present disclosure may be knitted with nylon to prepare a fabric, and such a knitted fabric has an excellent dye color deepening effect by having improved dyeability for an acid dye.

Hereinafter, the present disclosure will be specifically
10 described with reference to examples, however, the following examples and experimental examples are just illustrating one embodiment of the present disclosure, and the scope of the present disclosure is not limited to the following examples and experimental examples.

15 Example 1

To 33.8 kg of polytetramethylene ether glycol (PTMG, molecular weight 1800), 5.0 mol% of polyethylene glycol (molecular weight 2000) was mixed, and after adding 8.8 kg of 4,4'-diphenylmethane diisocyanate thereto, the result was
20 reacted while stirring for 120 minutes at 90 °C in a nitrogen gas stream to prepare a polyurethane prepolymer having isocyanate on both terminals. The prepolymer was cooled to room temperature, and 66.1 kg of dimethylacetamide was added thereto as a solvent to obtain a polyurethane prepolymer
25 solution.

Subsequently, 1.1 kg of ethylenediamine as a chain extender and 0.1 kg of diethylamine as a chain terminator were dissolved in 15.6 kg of dimethylacetamide, and the result was added to the prepolymer solution at 10 °C or lower to obtain a polyurethane urea solution in which the polyurethane urea solution solid has an amine-terminated number of 63 meq/kg.

In addition, 1.5 weight% of triethylene glycol-bis-3-(3-tertiary-butyl-4-hydroxyphenyl)propionate as an antioxidant, 4 weight% of hydrotalcite ($\text{Mg}_4\text{Al}_2(\text{OH})_{12}\text{CO}_3 \cdot 3\text{H}_2\text{O}$) coated with 1 weight% of melamine polyphosphate as an inorganic chlorine resistance agent, and 0.5 weight% of titanium dioxide as a light resistance agent were mixed to the polyurethane urea solution to prepare a polyurethane urea spinning solution.

The spinning solution obtained as above was spun at a rate of 900 m/min by dry spinning to manufacture a polyurethane urea elastic yarn with 40 denier, 3 filaments, and a yarn amine-terminated number of 33 meq/kg.

Example 2

In the process of preparing the prepolymer of Example 1, 10.0 mol% of polyethylene glycol (molecular weight 2000) was mixed to 32.0 kg of polytetramethylene ether glycol (PTMG, molecular weight 1800), and after adding 8.7 kg of 4,4'-diphenylmethane diisocyanate thereto, the result was reacted while stirring for 120 minutes at 90 °C in a nitrogen gas stream to prepare a polyurethane prepolymer having isocyanate

on both terminals. The preparation process after that was the same as in Example 1.

Example 3

In the process of preparing the prepolymer of Example 1,
5 20.0 mol% of polyethylene glycol (molecular weight 2000) was
mixed to 28.2 kg of polytetramethylene ether glycol (PTMG,
molecular weight 1800), and after adding 8.7 kg of 4,4'-
diphenylmethane diisocyanate thereto, the result was reacted
while stirring for 120 minutes at 90 °C in a nitrogen gas
10 stream to prepare a polyurethane prepolymer having isocyanate
on both terminals. The preparation process after that was the
same as in Example 1.

Comparative Example 1

A polyurethane urea elastic yarn was prepared in the same
15 manner as in the preparation process of Example 1 except that
polyethylene glycol was not mixed in the step of preparing a
prepolymer.

Comparative Example 2

In the process of preparing the prepolymer of Example 1,
20 4.0 mol% of polyethylene glycol (molecular weight 2000) was
mixed to 34.2 kg of polytetramethylene ether glycol (PTMG,
molecular weight 1800), and after adding 8.8 kg of 4,4'-
diphenylmethane diisocyanate thereto, the result was reacted
while stirring for 120 minutes at 90 °C in a nitrogen gas
25 stream to prepare a polyurethane prepolymer having isocyanate

on both terminals. The preparation process after that was the same as in Example 1.

Comparative Example 3

In the process of preparing the prepolymer of Example 1,
5 21.0 mol% of polyethylene glycol (molecular weight 2000) was
mixed to 27.8 kg of polytetramethylene ether glycol (PTMG,
molecular weight 1800), and after adding 8.7 kg of 4,4'-
diphenylmethane diisocyanate thereto, the result was reacted
while stirring for 120 minutes at 90 °C in a nitrogen gas
10 stream to prepare a polyurethane prepolymer having isocyanate
on both terminals. The preparation process after that was the
same as in Example 1.

Experimental Example

Each of the spandex manufactured in the examples and the
15 comparative examples was dyed with a black color acid dye, and
the degree of dyeability (darkness) was measured as a
brightness (L*) value after dyeing, and comparatively
evaluated.

The evaluation results are compared and shown in the
20 following Table 1.

Herein, the L* value, which is the degree of dyeability,
was measured by measuring reflectivity of the yarn using a
spectrophotometer, and the measured value was calculated using
a calculation formula of the CIE Lab color difference formula.

25 [Table 1]

	Polyethylene Glycol Content	L* Value	Modulus [g]	Spinning Workability
Example 1	5.0 mol%	34	7.6	Favorable
Example 2	10.0 mol%	10	7.4	Favorable
Example 3	20.0 mol%	6	7.0	Favorable
Comparative Example 1	Not Added	76	8.1	Favorable
Comparative Example 2	4.0 mol%	52	7.9	Favorable
Comparative Example 3	21.0 mol%	5	6.6	Poor

* L* value: in the L*a*b* color difference values, the L* value is a value representing brightness, and when dyeing with the same dye, a lower L* value is obtained as the color is darker.

5 From the above-described experimental results, it was identified that significantly superior dyeability was obtained in the examples according to the present disclosure compared to in the comparative examples. In addition, it was identified that there was a critical significance for the amount of the
10 polyethylene glycol used according to the present disclosure.

Comparative Example 3 had problems in that modulus of the yarn decreased due to the excessive addition of polyethylene glycol, and the use in the process was difficult due to poor workability during the spinning.

15 Comparative Example 2 had a problem in that the L* value of the yarn significantly increased due to the insufficient addition of polyethylene glycol, which led to poor dyeability.

【CLAIMS】

【Claim 1】

A polyurethane urea elastic yarn having improved dyeability, the yarn comprising (a) a polyol including a first
5 polyol and a second polyol; (b) at least one of diisocyanates;
(c) at least one of diamine chain extenders; and (d) at least one of amine chain terminators,

wherein the second polyol is polyethylene glycol; and

the first polyol and the second polyol are materials
10 different from each other.

【Claim 2】

The polyurethane urea elastic yarn having improved dyeability of Claim 1, wherein the second polyol is mixed in
5.0 mol% to 20.0 mol% with respect to the total amount of
15 polyol.

【Claim 3】

The polyurethane urea elastic yarn having improved dyeability of Claim 1, which has an amine-terminated number of
20.0 meq/kg to 45.0 meq/kg.

20 **【Claim 4】**

A method for manufacturing a polyurethane urea elastic yarn having improved dyeability, the method comprising:

(a) preparing a polyurethane prepolymer by bringing a polyol including a first polyol and a second polyol into
25 contact with at least one of diisocyanates;

(b) adding a solvent to the prepolymer in the step (a);
(c) bringing the product in the step (b) into contact with at least one of diamine chain extenders and at least one of amine chain terminators; and
5 (d) preparing a polyurethane urea elastic yarn by spinning the product in the step (c),
wherein the second polyol is polyethylene glycol; and
the first polyol and the second polyol are materials different from each other.

10 **【Claim 5】**

The method for manufacturing a polyurethane urea elastic yarn having improved dyeability of Claim 4, wherein the second polyol is mixed in 5.0 mol% to 20.0 mol% with respect to the total amount of polyol.

15 **【Claim 6】**

The method for manufacturing a polyurethane urea elastic yarn having improved dyeability of Claim 4, wherein the polyurethane urea elastic yarn has an amine-terminated number of 20.0 meq/kg to 45.0 meq/kg.

20 **【Claim 7】**

A fabric with enhanced dyeability for an acidic dye by knitting the polyurethane urea elastic yarn of any one of Claims 1 to 3 and nylon.

【Claim 8】

A fabric with enhanced dyeability for an acidic dye by knitting the polyurethane urea elastic yarn manufactured using the method of any one of Claims 4 to 6 and nylon.