

CLAIMS

1. A method of inhibiting isomerization of a reducing saccharide in an aqueous solution containing said reducing saccharide upon a thermal treatment of said
5 aqueous saccharide solution, the method comprising the step of acidifying the aqueous saccharide solution prior to and/or in the course of the thermal treatment.
2. The method according to claim 1, wherein the aqueous saccharide solution is
10 acidified to a pH having a value of between about 1 to about 6, preferably to a pH having a value of between about 2 to about 5, and more preferably to a pH having a value of between about 3 to about 4.
3. The method according to claim 1 or 2, wherein the aqueous saccharide
15 solution is acidified by adding at least one acid to said aqueous saccharide solution.
4. The method according to claim 3, wherein the acid is an inorganic acid,
20 preferably selected from the group consisting of hydrochloric acid, sulfuric acid, sulfurous acid, phosphoric acid, boric acid, hydrofluoric acid, hydrobromic acid, perchloric acid, hydroiodic acid and carbonic acid.
5. The method according to claim 3, wherein the acid is an organic acid,
25 preferably selected from the group consisting of monocarboxylic acids, dicarboxylic acids, and tricarboxylic acids.
6. The method according to any one of claims 1 to 5, wherein the reducing sugar
30 is selected from the group consisting of aldoses, disaccharides and oligosaccharides, preferably a disaccharide selected from the group consisting of lactose and maltose.
7. The method according to any one of claims 1 to 6, wherein the thermal treatment comprises exposing the acidified aqueous solution to a temperature in the range of about 30 °C to about 150 °C.

8. A thermally treated aqueous solution containing at least one reducing saccharide, wherein the aqueous solution is obtained by a method according to any one of claims 1 to 7.
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9. The thermally treated aqueous solutions according to claim 8, wherein the aqueous solution is sterile.
10. The thermally treated aqueous solution according to claim 8 or 9, wherein the reducing saccharide is lactose, preferably present in a concentration of ≥ 10 mM, preferably in a concentration of ≥ 100 mM, more preferably in a concentration of ≥ 0.66 M, and most preferably in a concentration of ≥ 1 M.
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11. Use of a thermally treated aqueous solution containing at least one reducing saccharide according to any one of claims 8 to 10 in a biotechnological production process for producing a biological product.
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12. The use according to claim 11, wherein the biotechnological process is a selected from the group consisting of a biocatalytic production process and a fermentative production process.
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13. The use according to claim 12 for a fermentative production of a human milk oligosaccharide, wherein the thermally treated aqueous solution containing at least one reducing saccharide is a wet heat sterilized aqueous solution containing lactose.
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14. A method of biotechnological production of a biological product, the method comprises the steps of:
- providing at least one cell that is capable of producing the biological product;
 - 30 - cultivating the at least one cell in a fermentation broth containing and/or being supplemented with the thermally treated aqueous solution according to any one of claims 8 to 10 for the at least one cell to produce the biological product; and
 - 35 - optionally purifying the biological product from the fermentation broth.

15. The method according to claim 14, wherein the biological product is a human milk oligosaccharide, and wherein the thermally treated aqueous solution is a wet heat sterilized aqueous solution containing lactose.
- 5 16. The method according to claim 14 or 15, wherein the biological product is produced in an amount of ≥ 100 g/L in the fermentation broth, preferably in an amount of ≥ 150 g/L in the fermentation broth, more preferably in an amount of ≥ 200 g/L in the fermentation broth at the end of the fermentation process.
- 10 17. A biological product manufactured by a process according to claim 14 or 15, wherein the biological product is preferably selected from the group consisting of a human milk oligosaccharide, lactosucrose and lactobionic acid.
- 15 18. The use according to any one of claims 11 to 13, the method according to any one of claims 14 to 16, or the biological product according to claim 17, wherein the biological product is a human milk oligosaccharide being selected from the group consisting of 2'-fucosyllactose, 3-fucosyllactose, 2',3-difucosyl-lactose, lacto-*N*-triose II, lacto-*N*-tetraose, lacto-*N*-neotetraose, lacto-*N*-fucopentaose I, lacto-*N*-neofucopentaose I, lacto-*N*-fucopentaose II, lacto-*N*-fucopentaose III, lacto-*N*-fucopentaose V, lacto-*N*-neofucopentaose V, lacto-*N*-difucohexaose I, lacto-*N*-difucosylhexaose II, *para*-Lacto-*N*-fucosylhexaose, fucosyl-lacto-*N*-sialylpentaose b, fucosyl-lacto-*N*-sialylpentaose c, fucosyl-lacto-*N*-sialylpentaose c, disialyl-lacto-*N*-fucopentaose, 3-fucosyl-3'-sialyllactose, 3-fucosyl-6'-sialyllactose, lacto-*N*-neodifucohexaose I, 3'-sialyllactose, 6'-sialyllactose, sialyllacto-*N*-tetraoses LST-a, LST-b, LST-c, and disialyllacto-*N*-tetraose.
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