



(22) Date de dépôt/Filing Date: 1992/05/21

(41) Mise à la disp. pub./Open to Public Insp.: 1992/11/22

(45) Date de délivrance/Issue Date: 2002/07/16

(30) Priorité/Priority: 1991/05/21 (9110821.7) GB

(51) Cl.Int.⁵/Int.Cl.⁵ C07H 13/04, C07H 5/02

(72) Inventeur/Inventor:
Sankey, George H., GB

(73) Propriétaire/Owner:
Tate & Lyle Public Limited Company, GB

(74) Agent: MARKS & CLERK

(54) Titre : PROCEDE CONTINU POUR LA PREPARATION DE SUCROSE 6-ESTERS

(54) Title: CONTINUOUS PROCESS FOR THE PREPARATION OF SUCROSE 6-ESTERS

(57) **Abrégé/Abstract:**

A method is provided for the preparation of a sucrose 6-ester by: (i) reacting sucrose with a trialkyl orthoester or with a ketene acetal in an inert organic solvent in the presence of an acid catalyst to provide a sucrose alkyl 4, 6-orthoester, (ii) treating the sucrose alkyl 4, 6-orthoester under mild aqueous acidic conditions to provide a mixture of sucrose 4- and 6-monoesters, (iii) treating the mixture of esters with a base to convert the sucrose 4-ester into sucrose 6-ester, and (iv) neutralising the solution and isolating the sucrose 6-ester, an acidic ion exchange resin being used to catalyse preparation of the sucrose alkyl 4, 6-orthoester and to effect its hydrolysis. Sucralose is prepared by chlorinating the sucrose 6-esters prepared.

2069130

ABSTRACT

A method is provided for the preparation of a sucrose 6-ester by: (i) reacting sucrose with a trialkyl orthoester or with a ketene acetal in an inert organic solvent in the presence of an acid catalyst to provide a sucrose alkyl 4,6-orthoester, (ii) treating the sucrose alkyl 4,6-orthoester under mild aqueous acidic conditions to provide a mixture of sucrose 4- and 6-monoesters, (iii) treating the mixture of esters with a base to convert the sucrose 4-ester into sucrose 6-ester, and (iv) neutralising the solution and isolating the sucrose 6-ester, an acidic ion exchange resin being used to catalyse preparation of the sucrose alkyl 4,6-orthoester and to effect its hydrolysis. Sucralose is prepared by chlorinating the sucrose 6-esters prepared.

2069130

1

M&C FOLIO: P63523

WANGDOC: 0718p

Continuous Process for the Preparation of Sucrose 6-esters

This invention relates to an improved process for the preparation of sucrose 6-esters, which are key intermediates in one route to the production of sucralose (1,6-dichloro-1,6-dideoxy- β -D-fructofuranosyl 4-chloro-4-deoxy- α -D-galactopyranoside). Sucralose is a high intensity sweetener having a sweetness of several hundred times that of sucrose which is disclosed in our British Patent No. 1,543,167.

The preparation of sucralose involves the introduction of chlorine atoms into the 1'-and 6' - positions (i.e. displacement of two of the three primary hydroxyl groups) and at the 4-position (i.e. displacement of a secondary hydroxyl group). The third primary hydroxyl group, at the 6-position, must remain unaffected.

One important route to sucralose involves the preparation and subsequent chlorination of 2,3,6,3',4' -penta-O-acetyl sucrose in which the three hydroxyl groups to be displaced by chlorine atoms are unprotected while all the other hydroxyl groups are protected (see, for example, US Patent No. 4,362,869 and UK Patent GB 2 065 648 B).

An alternative, simpler route involves the preparation of a sucrose 6-ester which can be selectively chlorinated at the 4-, 1' - and 6' - positions and then de-esterified to provide sucralose. One such method of preparing sucrose 6-esters and converting them to sucralose is disclosed in our UK Patent GB 2 079 749 B, but this method produces a mixture of acylated sucrose derivatives with substituents at one or more of the primary positions, with a major proportion of the 6-ester.

A more selective method of preparing sucrose 6-esters, disclosed in our UK Patent GB 2 195 632 B, is based upon the preparation of a sucrose alkyl 4,6-orthoester by reacting sucrose with a trialkyl orthoacylate in the presence of an acid catalyst in an inert organic solvent, followed by hydrolysis of the orthoester to provide a mixture of sucrose 4- and 6-esters, which are then isomerised to provide a high yield of the required sucrose 6-ester. Ketene acetals, such as 1,1-dimethoxyethene, are useful alternatives to trialkyl orthoacylates in the preparation of sucrose alkyl 4,6-orthoesters, as disclosed in our UK Patent GB 2 255 975 B entitled Process for the preparation of Sucrose 6-esters, filed on 21st May 1991.

The method of GB 2 195 632 B is more efficient than that of GB 2 079 749 B, but involves several stages, each utilising a different acid or base which must be removed.

We have now found that the method of GB 2 195 632 B can be operated in a semi-continuous or continuous mode, using an acidic ion exchange resin to catalyse the preparation of the sucrose alkyl 4,6-orthoester and to hydrolyse it to a mixture of sucrose 4- and 6-esters, and then treating the hydrolysate with a base to convert the sucrose 4-ester to sucrose 6-ester. The process can be run continuously to produce a mixture of sucrose 4- and 6-esters, without the need to add or remove acid.

Thus, according to the present invention, there is provided a method for the preparation of a sucrose 6-ester by: (i) reacting sucrose with a trialkyl orthoester or with a ketene acetal in an inert organic solvent in the presence of an acid catalyst to provide a sucrose alkyl 4,6-orthoester, (ii) treating the sucrose alkyl 4,6-orthoester under mild aqueous acidic conditions to provide a mixture of sucrose 4- and 6-monoesters, (iii) treating the mixture of esters with a base to convert the sucrose 4-ester into sucrose 6-ester, and (iv) neutralising the solution and isolating the sucrose 6-ester, characterised in that an acidic ion exchange resin is used to catalyse preparation of the sucrose alkyl 4,6-orthoester and to effect its hydrolysis.

In general, the amount of trialkyl orthoester or ketene acetal used in step (i) can vary from about 1 to 2 ME per ME of sucrose, preferably from 1.1 to 1.5 ME.

Amongst the wide range of acidic ion exchange resins available, we have found that the most useful type to act as catalyst for preparation of the sucrose alkyl 4,6-orthoester is a macroreticular acid resin.

Choice of an acidic ion exchange resin for hydrolysis of the mixture of sucrose 4- and 6-esters is much less critical, and selection of the resin for this step is mainly a matter of cost and convenience.

Stages (i) and (ii) of the process can be run continuously by passing a solution of sucrose and the reagent in an inert organic solvent such as dimethylformamide through a column of macroreticular acid resin, continuously adding water to the eluate containing the sucrose alkyl 4,6-orthoester, and passing the aqueous solution through a second column of acidic ion exchange resin to effect the hydrolysis.

We have found that Amberlyst™ 15 (H⁺) macroreticular resin is suitable for the first stage and that it is convenient to use the same resin for stage (ii).

Conversion of the sucrose 4-ester to sucrose 6-ester can be carried out by adding a water soluble base such as potassium carbonate or sodium carbonate. Alternatively, an organic base such as tertiary butylamine can be used.

Use of a readily water soluble base such as potassium

2069130

5

carbonate lends itself to continuous operation more easily than a less soluble base such as calcium hydroxide. Thus, the eluate from stage (ii) can be collected in a holding tank, into which the base is metered to provide the appropriate concentration for migration of the ester group from the 4- to the 6- position.

The base must be neutralised before concentration and isolation of the sucrose 6-ester, in order to prevent degradation of the product. Neutralisation can be effected by addition of a suitable acid or, preferably, by passage through an acidic ion exchange resin to give a stream free from inorganic salts.

The sucrose 6-ester can be isolated by concentrating the neutralised solution to a syrup, dissolving the syrup in methanol, seeding, and collecting the crystalline product. After washing, e.g. with methanol, and drying, the sucrose 6-ester can be stored until required.

Sucrose 6-esters prepared according to the present invention can be used as intermediates for the production of sucralose. Thus, in a further embodiment of the present invention there is provided a process for the production of sucralose, by preparing a sucrose 6-ester, reacting the sucrose 6-ester with a chlorinating agent capable of selectively chlorinating the 4-, 1' - and 6' positions, optionally peresterifying the sucralose 6-ester so formed,

2069130

6

de-esterifying the sucralose ester, and recovering sucralose, in which the sucrose 6-ester is prepared by:

- (i) passing a solution of sucrose and a trialkyl orthoester or a ketene acetal in an inert organic solvent through a macroreticular acid resin to provide an eluate containing a sucrose alkyl 4,6-orthoester,
- (ii) continuously adding water to the eluate and passing the mixture through an acidic ion exchange resin to hydrolyse the sucrose alkyl 4,6-orthoester and to provide an eluate containing a mixture of sucrose 4- and 6- monoesters,
- (iii) treating the mixture of esters with a base to convert the sucrose 4-ester to sucrose 6-ester and (iv) neutralising the solution and isolating the sucrose 6-ester.

The invention is further illustrated by the following Example.

Example 1 Preparation of sucrose 6-acetate

A solution of sucrose (500 g) and trimethyl orthoacetate (225 ml; 1.2 ME) in dimethylformamide (2500 ml) was passed down a column of Amberlyst 15(H⁺) resin (25 g) at ambient temperature at a rate of 10 g/minute. Water was added continuously to the eluate issuing from the base of the column at a rate of 0.065 g water/g eluate and the diluted eluate was passed through a heat exchanger to cool the temperature from about 40° (heat of dilution) to about 25°.

2069130

7

The cooled solution from the heat exchanger was then passed down a second column of Amberlyst 15(H⁺) resin (25 g) at a rate of 10.65g/minute (i.e. at a rate which allowed the two columns to be run continuously in sequence).

A sample of the eluate from the second column (1851 g; equivalent to an input of 300 g of sucrose) was removed and treated with an aqueous 2% w/v solution of potassium carbonate (K₂CO₃ · 1.5H₂O; 37 ml). After one hour the solution was neutralised with dilute hydrochloric acid (0.5 molar; 16.2 ml) and then concentrated to a syrup (451.2 g) under reduced pressure at 50°. Analysis of the syrup showed a yield of crude sucrose 6-acetate of about 75% with sucrose (about 15%) as the main impurity.

The syrup was dissolved in hot methanol (600 ml), allowed to cool to 40° and then seeded with crystalline sucrose 6-acetate (1.5 g). Crystallisation was allowed to proceed overnight at ambient temperature and then the product was collected, washed with methanol (300 ml) and dried in vacuo at 40°. Yield 219.4 g; 50% molar; analysis: sucrose 6-acetate 82.8%, sucrose 4-acetate 0.5%, sucrose 4.7%, sucrose di-acetate 2.4%, methanol of crystallisation 7%.

Similar results can be obtained by using 1,1-dimethoxyethane (150 ml) in place of the trimethyl orthoacetate.

2069130

8

THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE DEFINED AS FOLLOWS:

1. In the method for the preparation of a sucrose 6-ester by: (i) reacting sucrose with a trialkyl orthoester or with a ketene acetal in an inert organic solvent in the presence of an acid catalyst to provide a sucrose alkyl 4,6-orthoester, (ii) treating the sucrose alkyl 4,6-orthoester under mild aqueous acidic conditions to provide a mixture of sucrose 4- and 6-monoesters, (iii) treating the mixture of esters with a base to convert the sucrose 4-ester into sucrose 6-ester, and (iv) neutralising the solution and isolating the sucrose 6-ester, the improvement which consists in that an acidic ion exchange resin is used to catalyse preparation of the sucrose alkyl 4,6-orthoester and to effect its hydrolysis in step (ii).
2. The method of Claim 1, in which the acidic ion exchange resin used is a macroreticular acid resin.
3. The method of Claim 1, in which the amount of trialkyl orthoester or ketene acetal used in step (i) is from about 1 to 2 mole equivalents (ME) per ME of sucrose.
4. The method of Claim 1, in which the amount of trialkyl orthoester or ketene acetal used in step (i) is from about 1.1 to 1.5 ME per ME of sucrose.

2069130

9

5. The method of Claim 1, in which the base used to convert the sucrose 4-ester into the sucrose 6-ester is selected from the group consisting of water soluble bases and organic bases.

6. The method of Claim 1, in which the base is neutralised after conversion of the sucrose 4-ester into the sucrose 6-ester by passing the basic solution through an acidic ion exchange resin.

7. The method of Claim 1, in which a solution of sucrose and a trialkyl orthoester or a ketene acetal in an inert organic solvent is passed through a column of macroreticular acid resin to provide an eluate containing a sucrose alkyl 4,6-orthoester, water being continuously added to the eluate before passing the aqueous solution through a second column of acidic ion exchange resin to provide a mixture of sucrose 4- and 6- monoesters.

8. The method of Claim 7, in which the amount of trialkyl orthoester or ketene acetal used is from about 1 to 2 ME per ME of sucrose.

9. The method of Claim 7, in which the amount of trialkyl orthoester or ketene acetal used is from about 1.1 to 1.5 ME per ME of sucrose.

2069130

10

10. The method of Claim 7, in which the base used to convert the sucrose 4-ester into the sucrose 6-ester is selected from the group consisting of water soluble bases and organic bases.

11. The method of Claim 7, in which the mixture of sucrose 4- and 6- monoesters thus produced is collected in a holding tank into which is metered a water soluble base to effect conversion of the sucrose 4-ester into the sucrose 6-ester.

12. The method of Claim 7, in which the base is neutralised after conversion of the sucrose 4-ester into the sucrose 6-ester by passing the basic solution through an acidic ion exchange resin.

13. A method for the production of sucralose, comprising preparing a sucrose 6-ester, reacting the sucrose 6-ester with a chlorinating agent capable of selectively chlorinating the 4-, 1' - and 6' positions, optionally peresterifying the sucralose 6-ester so formed, de-esterifying the sucralose ester, and recovering sucralose, in which the sucrose 6-ester is prepared by:

(i) passing a solution of sucrose and a trialkyl orthoester or a ketene acetal in an inert organic solvent through a macroreticular acid resin to provide an eluate containing a sucrose alkyl 4,6-orthoester,

(ii) continuously adding water to the eluate and passing the

2069130

11

mixture through an acidic ion exchange resin to hydrolyse the sucrose alkyl 4,6-orthoester and to provide an eluate containing a mixture of sucrose 4- and 6- monoesters, (iii) treating the mixture of esters with a base to convert the sucrose 4-ester to sucrose 6-ester, and (iv) neutralising the solution and isolating the sucrose 6-ester.

14. The method of Claim 13, in which the amount of trialkyl orthoester or ketene acetal used is from about 1 to 2 ME per ME of sucrose.

15. The method of Claim 13, in which the amount of trialkyl orthoester or ketene acetal used is from about 1.1 to 1.5 ME per ME of sucrose.

16. The method of Claim 13, in which the base used to convert the sucrose 4-ester into the sucrose 6-ester is selected from the group consisting of water soluble bases and organic bases.

17. The method of Claim 13, in which the mixture of sucrose 4- and 6- monoesters thus produced is collected in a holding tank into which is metered a water soluble base to effect conversion of the sucrose 4-ester into the sucrose 6-ester.

18. The method of Claim 13, in which the base is neutralised after conversion of the sucrose 4-ester into the sucrose 6-ester by passing the basic solution through an acidic ion exchange resin.