

METHOD OF MANUFACTURE OF POLYACROLEIN

This invention relates to a method of manufacture of polyacrolein and, in particular, to a method of manufacture of polyacrolein for use in antimicrobial compositions.

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Background

The use of polyacrolein in antimicrobial applications has been described in U.S. Patent 5,290,894, Australian Application 11686/95 and its European counterpart EP 667358, in our International patent publication WO 96/38186 (PCT/AU96/00328) and more recently we have described a method of improving the activity of acrolein polymers in our International patent publication WO 01/60874 (PCT/AU00/00107).

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Many of the most stable compositions of polyacrolein for use as antimicrobials are formed by heating the polymer in air. Indeed Australian Application No 11686/85 (Werle et al assigned to Degussa) teaches that only those preparations dried using the process of the invention described therein with strong air current and final air temperatures of $>60^{\circ}\text{C}$, preferably at 75°C are alkali soluble.

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Our International Publication WO 01/60874 describes a method of improving antimicrobial activity of acrolein polymers in which the solid polymers are oxidized in air and the oxidized polymers are heated in an alcohol solution in the presence of added alkali. The presence of carboxylic acid groups formed by oxidation in air is believed to improve the solubility of polyacrolein compositions. We have now found that polyacrolein polymer of high solubility; antimicrobial activity and stability may be formed without the requirement for this aerial oxidation step.

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Summary

Accordingly, the present invention provides a method of manufacture of soluble, microbiologically active and stable polyacrolein comprising: (a) polymerising acrolein in the presence of base to form a polymer of acrolein; (b) dissolving the polymer of acrolein in an alcohol selected from monoalcohols and polyols, optionally with addition of water to form an alcohol solution of the polymer of acrolein; (c) heating the alcohol solution of acrolein; and (d) mixing aqueous base with the polymer of acrolein.

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It is preferred that the aqueous base is mixed with the alcohol solution of the polymer of acrolein after heating of the alcohol solution of the polymer of acrolein.

5 The addition of base after heating, as opposed to before heating, is preferred as the addition of base after formulation affords solutions which maintain their biocidal activity for long periods upon storage.

10 The alcohol solution of the polymer of acrolein is preferably heated for sufficient time so that when rendered alkaline by the addition of aqueous base it does not precipitate when further diluted by a factor of one in ten.

Detailed Description

15 In contrast with the teaching in the prior art the present invention allows stable solutions of acrolein polymers to be formed without the requirement for grinding or heating of the solid polymer in an air stream in order to oxidize the solid. This potentially reduces the expense and time required for preparation of these highly active biocides.

20 The acrolein polymer is formed by polymerisation of acrolein in the presence of a base such as sodium hydroxide. Polymerisation in the presence of radical initiators may also be used in preparing acrolein polymers as described in some of the examples of US patent 5290894 but such polymers are not generally used in the process of the present invention.

25 The composition is dissolved in an alcohol optionally with addition of water. Dissolution of the polymer may take place as a discrete step or may occur as part of the heating process. Generally the dissolution will occur within 15 minutes of heating a PEG solution of the polymer at 35 to 65°C.

30 The heating of the polymer of acrolein in the alcohol solution is preferably followed by mixing of the heated solution with aqueous base. The step of heating in the alcohol is important for achieving stability of the acrolein polymer in aqueous solutions following the mixing with aqueous base. The temperature and period of heating will depend on the polymer, the type of alcohol used and the extent and

term of stability required. Generally we have found that good results are obtained if the polymer of acrolein is heated for sufficient time so the when mixed with aqueous base to provide an alkaline solution the polymer will not precipitate. Preferably the polymer will not precipitate even when further diluted by a factor of one in ten (parts
5 by volume) with water. Having regard to this type of testing a skilled person will, without undue experimentation, be able to determine appropriate heating conditions.

Following the heating step a base is mixed with the composition preferably in the
10 form of an aqueous base composition.

The polyacrolein used in the method of the invention is formed by polymerisation of acrolein monomer conducted in an alkaline solution and the acrolein polymer may be collected as a precipitate. It will be apparent to those skilled in the art that a co-
15 monomer, especially a water-soluble or latently water-soluble co-monomer may be used in step (a). Typically when a co-monomer is used it will constitute less than 10% by weight of the total monomer composition. We prefer that the acrolein polymer is a homopolymer.

20 The precipitate may, and preferably will be dissolved in the alcohol without oxidation of the solid by heating in air or oxygen, to form poly(2-propenal, 2-propenoic acid). The acrolein polymer will generally be isolated from the polymerisation reaction and is preferably heated in the alcohol with a solution of pH of no more than 7. The precipitate formed in the preferred aspect of the invention
25 may be dissolved in the alcohol without needing to further process it.

In the process of the invention, the acrolein homopolymer is dissolved in the alcohol. This process would generally involve heating the polyacrolein in the alcohol to a temperature in the range of from 40 to 90°C. The preferred alcohol is a
30 polyalkylene glycol and preferably has a molecular weight in the range of from 200 to 20,000. More preferably the molecular weight is in the range of from 200 to 10000 and most preferably from 200 to 2000. The acrolein polymer is heated in the alcohol to form an acetal derivative. Typically, the alcohol solution will be heated for a period of time in the range of from fifteen minutes to five hours with the alcohol

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and at a temperature in the range from 50 to 90°C, more preferably from 60 to 90°C.

Generally, the polyacrolein formed in step (a) used in the process of the invention
5 will have a low acid content typically of less than 1 mole of carboxyl groups per kg
of polymer and most preferably less than 0.5 mole acid groups per kilogram of
polymer. Despite the low content of carboxyl groups, we have found that when the
polymer is heated in the alcohol for a sufficient time and the alkali is added, the
alkaline solution resists precipitation when diluted with water, which would not have
10 been expected for an unoxidized polymer.

The process of the invention includes a step of adding base to the composition; the
base is generally added to the alcohol solution following the heating step. The pH
of the resulting solution of is preferably in the range of from 7 to 9.5 and more
15 preferably is from 7.5 to 8.5. The preferred base for addition to the alcohol solution
of polyacrolein is an alkali metal carbonate particularly sodium carbonate or
potassium carbonate. Alkaline metal hydroxide such as sodium hydroxide or
potassium hydroxide may also be used but are less preferred. Typically the alkali is
added as an aqueous solution. Preferably the solution is cooled to room
20 temperature before adding the base in the above step.

The concentration of polymer of acrolein used in the step comprising heating in that
alcohol is generally from 0.5 to 50% by weight and more preferably from 0.5 to 40%
by weight. In the case of polyalkylene glycols the polyol polymer content will
25 depend on the molecular weight of the polyol polymer. For lower molecular weight
polyol polymers the content may be as high as from 50 to 90% by weight whereas
for higher molecular weight polyol polymers (eg. 1500 or more) dilute compositions
of the polyol polymer (eg. 2 to 50 %) may be preferred.

30 The acrolein polymer prepared by the method of the invention is useful in a range of
applications, especially antimicrobial applications. The preferred applications
include antiseptic compositions, disinfectant compositions and compositions for use
in treatment of gastrointestinal disease. The compositions formed in accordance
with the invention generally have a good long-term antimicrobial activity. Typically,

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the acrolein polymers provided by the method of manufacture described above will provide a minimum kill concentration after storage at 40°C for no less than twenty days of less than 150 ppm against a range of bacteria, e.g. *E. coli*, at 10^4 - 10^9 cfu/mL.

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The composition prepared by the method of the invention is particularly suited to use in administration of animals for treatment or prophylaxis of gastrointestinal disease, particularly gastrointestinal microbial infection. The composition prepared by the method of the invention may be administered to animals via drinking water, via food or other suitable means such as tablets, syrups and the like.

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The invention will now be described with reference to the following examples. It is to be understood that the examples are provided by way of illustration of the invention and that they are in no way limited to the scope of the invention.

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Examples

Aqueous Solution Stability Test

We found the following test ("herein referred to as the aqueous solution stability test") to be useful in determining the temperature and period of heating for providing good long term solution stability and activity.

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The heated alcohol solution of acrolein polymer was allowed to cool and mixed with dilute (0.4% w/w) aqueous sodium carbonate to provide an alkaline pH. If the resulting composition showed no sign of polymer precipitate the resulting composition was diluted with water to provide a one in ten dilution composition which was again examined for a precipitate at room temperature. After a sufficient period of heating of the acrolein polymer in the alcohol solution in accordance with step (c), preferably at a temperature in the range of from 60 to 105°C, the diluted composition was clear.

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Example 1 – Preparation of Polyacrolein

Water (720 mL at ambient temperature, about 20°C) and acrolein (60g; freshly distilled, plus optionally hydroquinone added to 0.25% w/w) were placed in an open beaker, within a fume cupboard, and very vigorously stirred, mechanically. Then,

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0.2 M aqueous sodium hydroxide (21.4 mL) was added to bring the pH to 10.5 - 11.0. The solution immediately turned a yellow typical of the hydroquinone anion and within a minute, the colour had disappeared and the clear solution became milky. About 1 minute later, precipitation of a white, flocculent polymer began, and
5 appeared complete within 15-30 minutes. The precipitate was filtered and washed with water.

Example 2

Polyacrolein (5.0 g) was added to hot PEG-200 (64.0 g, 65 °C) and the mixture
10 stirred until the solid dissolved (20 min). $\text{Na}_2\text{CO}_{3(\text{aq})}$ solution) (31 g of 1.29% w/w) was then added and the mixture heated at 65 °C for 10 min. The mixture was then allowed to cool and the sample made up with water to 100 g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g total mass, was
15 tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*. The composition was found to have poor stability and actually deteriorated rapidly.

Example 3

20 50.1 grams of the solution from Example 2 was then heated at 90 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water to 50 g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was
25 tested against *E.coli*.

Example 4

Polyacrolein (5.0 g) was added to hot PEG-200 (64.0 g, 65 °C) and the mixture stirred until the solid dissolved (5 min). Water (25.0 g) was then added and the
30 mixture heated at 105 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water (a portion of which contained Na_2CO_3 (0.40 g)) to 100 g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g

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total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

Example 5

- 5 Polyacrolein (5.0 g) was added to hot PEG-200 (64.0 g, 65 °C) and the mixture stirred until the solid dissolved (10 min). Water (26.0 g) was then added and the mixture heated at 90 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water (a portion of which contained Na₂CO₃ (0.40 g)) to 100 g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested
10 with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

Example 6

- 15 Polyacrolein (5.0 g) was added to hot PEG-200 (64.1 g, 65 °C) and the mixture stirred until the solid dissolved (5 min). Water (24.6 g) was then added and the mixture heated at 105 °C for 4 hours. The mixture was then allowed to cool and the sample made up with water (a portion of which contained Na₂CO₃ (0.40 g)) to 100 g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested
20 with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

Example 7

- 25 Polyacrolein (1.0 g) was added to hot PEG-200 (12.8 g, 65 °C) and the mixture stirred until the solid dissolved (5 min). Water (5.2 g) was then added and the mixture heated at 90 °C for 0.5 hours then at 105 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water (a portion of which contained Na₂CO₃ (0.04 g)) to 20 g to give a solution 5% w/w in polyacrolein. The
30 pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

Example 8

Polyacrolein (5.0 g) was added to hot PEG-200 (64.0 g, 65 °C) and the mixture stirred until the solid dissolved (10 min). Water (20.0 g) was then added and the mixture heated at 90 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water (a portion of which contained Na₂CO₃ (0.40 g)) to 100 g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

10 Example 9

Polyacrolein (5.0 g) was added to hot PEG-200 (64.0 g, 65 °C) and the mixture stirred until the solid dissolved (10 min). Water (20.0 g) was then added and the mixture heated at 90 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water (a portion of which contained Na₂CO₃ (0.40 g)) to 100 g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10 g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

20 Example 10

Polyacrolein (1.25g) was added to hot PEG-2000 (16.0g) at 65 °C and the mixture stirred until the solid dissolved (5min). The mixture was heated at 105 °C for two hours, before 0.1g of Na₂CO₃ dissolved in 7.65g of water was added to give a solution 5% w/w polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 3.0g sample diluted with water to 30g total mass was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

Example 11

30 Polyacrolein (1.00g) was added to hot PEG-200 (12.81g) at 65 °C and the mixture stirred until the solid dissolved (5min). The mixture was heated at 105 °C for seven hours, before cooling to room temperature. Once at room temperature 0.081g of Na₂CO₃ dissolved in 6.14g of water was added to give a solution 5% w/w

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polyacrolein. 1.0g of the sample diluted with water to 10g total mass was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

Example 12

- 5 Polyacrolein (5.003g) was added to hot PEG-200 (63.990g) at 65 °C and 0.403g of sodium carbonate dissolved in 30.641g of water was added. The solution was maintained at 65 °C before the temperature was increased to 90 °C and held for two hours. The sample was stored for 2 weeks at 40 °C before the MKC was tested against *E.coli* and determined to have a MKC of 500ppm.

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Example 13

- Polyacrolein (1.0g) was added to hot PEG-200 (12.8g, 65 °C) and the mixture stirred until the solid dissolved (5 min). Water (6.2g) was then added and the mixture heated at 90 °C for 2 hours. The mixture was then allowed to cool and the sample
15 made up with water to 20g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and the pH of a 1.0 g sample, diluted with water to 10g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

20 Example 14

- Polyacrolein (5.0g) was added to hot PEG-200 (64.0g, 65 °C) and the mixture stirred until the solid dissolved (10 min). The mixture was then allowed to cool and the sample made up with water to (a portion of which contained Na₂CO₃ (0.04g)) to give 100g to give a solution 5% w/w in polyacrolein. The pH of the neat solution
25 was tested with PANPEHA pH sticks and the pH of a 3.0g sample diluted with water to 10g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against *E.coli*.

Example 15

- 30 Polyacrolein (5.0g) was added to hot PEG-200 (64.1g, 65 °C) and the mixture stirred until the solid dissolved (10 min). The mixture was heated at 90 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water to (a portion of which contained Na₂CO₃ (0.20g)) to give 100g to give a solution 5% w/w in polyacrolein. The pH of the neat solution was tested with PANPEHA pH sticks and

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the pH of a 1.0g sample diluted with water to 10g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against E.coli.

Example 16

- 5 Polyacrolein (1.0g) was added to hot PEG-200 (12.8g, 65 °C) and the mixture stirred until the solid dissolved (5 min). Water (5.2g) was then added and the mixture heated at 90 °C for 0.5 hours then at 105 °C for 2 hours. The mixture was then allowed to cool and the sample made up with water to (a portion of which contained Na_2CO_3 (0.040g)) to 20g to give a solution 5% w/w in polyacrolein. The pH of the
- 10 neat solution was tested with PANPEHA pH sticks and the pH of a 1.0g sample diluted with water to 10g total mass, was tested using a pH probe. The minimum-kill-concentration (MKC) was tested against E.coli

Table 1

Sample (or)	Temp of dissolution	Add water	Add Na_2CO_3 (aq)	Time & Temp of 2 nd Heating	Add Na_2CO_3 (aq)	Colour	pH of neat solution	pH of 1-in-10 dilution	MKC	PPT
Example 2	65 °C	No	(0.4%)	10 min @ 65 °C	No	yellow/orange	8.5	8.85	31 ppm	No but poor stability
Example 3	65 °C	No	(0.4%)	2 h @ 90 °C	No	orange	7.0	6.62	62 ppm	No
Example 4	65 °C	Yes	No	2 h @ 105 °C	(0.4%)	brown	8.0	7.12	31 ppm	No
Example 5	65 °C	Yes	No	2 h @ 90 °C	(0.4%)	brown	8.0	7.11	62 ppm	No
Example 6	65 °C	Yes	No	4 h @ 105 °C	(0.4%)	brown	8.0	7.23	62 ppm	No
Example 7	65 °C	Yes	No	2 h @ 105 °C	(0.2%)	brown	5.0	5.06	125 ppm	No
Example 8	65 °C	Yes	No	2 h @ 90 °C	(0.4%)	brown	8.0	7.51	62 ppm	No
Example 9	65 °C	Yes	No	2 h @ 90 °C	(0.4%)	brown	8.6	8.69	125 ppm	No
Example 10	65 °C	No	No	2 h @ 105 °C	(0.4%)	brown	7.5	7.55	31 ppm	No
Example 11	65 °C	No	No	7 h @ 105 °C	(0.4%)	brown	-	8.31	125 ppm	No
Example 12	65 °C	No	(0.4%)	2 h @ 90 °C	No	-	-	-	500 ppm	No
Example 13	65 °C	Yes	No	2 h @ 65 °C	(0.4%)	Orange	8.0	7.13	62 ppm	ppt
Example 14	65 °C	No	No	2 h @ 90 °C	(0.04%)	Brown	5.0	3.37	62 ppm	ppt
Example 15	65 °C	No	No	2 h @ 80 °C	(0.2%)	brown	6.0	5.27	62 ppm	ppt
Example 16	65 °C	Yes	No	2 h @ 105 °C	(0.2%)	brown	5.0	5.06	125 ppm	No

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Table 2 Stability Data for Example 5

Sample	MKC	days at r.t.	MKC	days at 40 °C	Biocidal	days at r.t.	MKC	days at 40 °C	Biocidal	days at 40 °C
Example 5	62 ppm	2	31 ppm	17	Pass	31	62	52	Pass	59

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The Examples show that by heating the polymer in acrolein for a sufficient time at an elevated temperature, preferably above 65 C, the stability is significantly increased.

- 5 The addition of aqueous base to provide a pH of at least 7 also provides a further significant increase in stability and/or activity is demonstrated in examples 7, 14 and 16.

- 10 The aqueous base is preferably added after heating in alcohol as demonstrated by comparison of Examples 8 and 12.

CLAIMS

1. A method of manufacture of a soluble, microbiologically active and stable acrolein polymer comprising the following steps in sequence:
 - (a) polymerising acrolein in the presence of base to form a polymer of acrolein;
 - (b) dissolving the polymer of acrolein in an alcohol selected from monoalcohols and polyols optionally with addition of water to form an alcohol solution of the polymer of acrolein and providing a pH of no more than 7;
 - (c) heating the alcohol solution of the polymer of acrolein of pH of no more than 7 to react the polymer of acrolein with the alcohol; and
 - (d) mixing base with the polymer of acrolein.
2. A method according to claim 1 wherein the acrolein polymer comprises a comonomer in an amount of up to 10% by weight of the total monomer composition.
3. A method according to claim 1 wherein the acrolein polymer is a homopolymer.
4. A method according to claim 1 wherein the acrolein polymer is collected from the polymerisation reaction as a precipitate, and dissolved in the alcohol.
5. A method according to claim 1 wherein the acrolein polymer is not subject to oxidation by heating of the acrolein polymer solid in air at a temperature of at least 60°C before dissolving in the alcohol.
6. A method according to claim 1 wherein the acrolein polymer is isolated as a solid from the step of polymerisation in the presence of base and dissolved in the alcohol without a step of oxidising the isolated solid by heating in air.
7. A method according to claim 1 wherein the acrolein polymer is dissolved in the alcohol by heating the acrolein polymer in the alcohol to a temperature in the range of from 40 to 105°C.
8. A method according to claim 1 wherein alcohol is a polyalkylene glycol.

9. A method according to claim 1 wherein the heating of the polymer of acrolein in alcohol is continued for a time sufficient so that it does not precipitate when subject to the hereinafter defined aqueous solution stability test.
- 5 10. A method according to claim 1 wherein the acrolein polymer is heated in the alcohol at a temperature in the range from 50 to 105°C, for a period in the range of from fifteen minutes to five hours.
- 10 11. A method according to claim 1 wherein the acrolein polymer dissolved in the alcohol in step (b) has an acid content of less than 1 mole of carboxyl groups per kilogram of polymer.
12. A method according to claim 11 wherein said acid content is less than 0.5 mole acid groups per kilogram of polymer.
- 15 13. A method according to claim 1 wherein the base is added to the alcohol solution following dissolution of the alcohol.
14. A method according to claim 13 wherein the pH of the resulting solution is in the range of from 7 to 9.5.
- 20 15. A method according to claim 13 wherein the pH of the resulting solution is in the range of from 7.5 to 8.5.
- 25 16. A method according to claim 1 wherein the base comprises a compound selected from the group consisting of alkali metal carbonate alkali metal hydroxide such as sodium hydroxide and mixtures thereof.
17. A method according to claim 16 wherein the base comprises sodium carbonate and/or potassium carbonate.
- 30 18. A method according to claim 1 wherein the polymer of acrolein used in the step of heating in the alcohol is in a concentration in the alcohol of from 0.5 to 50% by weight.

19. A method according to claim 18 wherein the concentration is from 0.5 to 40% by weight.
- 5 20. A method according to claim 1 wherein the alcohol is polyethylene glycol and in present at a concentration in the range of from 5 to 90% by weight.
21. A method according to claim 1 wherein the alcohol is a polyethylene glycol of molecular weight in the range of from 200 to 20,000.
- 10 22. A composition prepared according to the method of any one of claims 1 to 21.
23. Use of the acrolein polymer prepared according to any one of claims 1 to 15 as an antimicrobial.
- 15 24. A use according to claim 23 wherein the acrolein polymer provides a minimum kill concentration after storage at 40°C for no less than twenty days of less than 150 ppm against *E. coli*, at 10^4 - 10^9 cfu/mL.
- 20 25. A use according to claim 23 in administration to animals for treatment or prophylaxis of gastrointestinal microbiological infection.
26. A use of an acrolein polymer prepared according to claim 1 in preparation of a medicament for oral administration to animals in treatment or prevention of
- 25 gastrointestinal infection.
27. A use according to claim 26 wherein the composition is in a form for administered to animals via drinking water or via food.