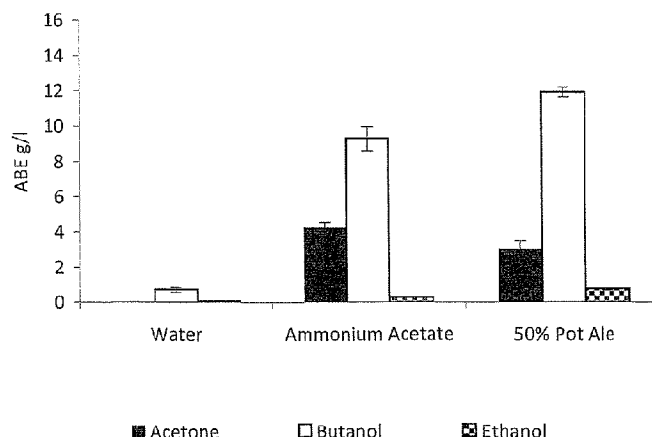




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(54) **Titre : PROCÉDE POUR LA FABRICATION DE BUTANOL OU D'ACETONE**
(54) **Title: PROCESS FOR THE MANUFACTURE OF BUTANOL OR ACETONE**



(57) **Abrégé/Abstract:**

A process for the manufacture of butanol, acetone and/or other renewable chemicals is provided wherein the process utilises one or more of the group comprising by-products of the manufacture of malt whisky, such as pot ale and/or spent lees, biomass substrates, such as paper, sludge from paper manufacture and spent grains from distillers and brewers, and diluents, such as water and spent liquid from other fermentations. The process comprises treating a substrate to hydrolyse it and fermenting the treated. Also provided is a biofuel comprising butanol manufactured according to the process of the invention.

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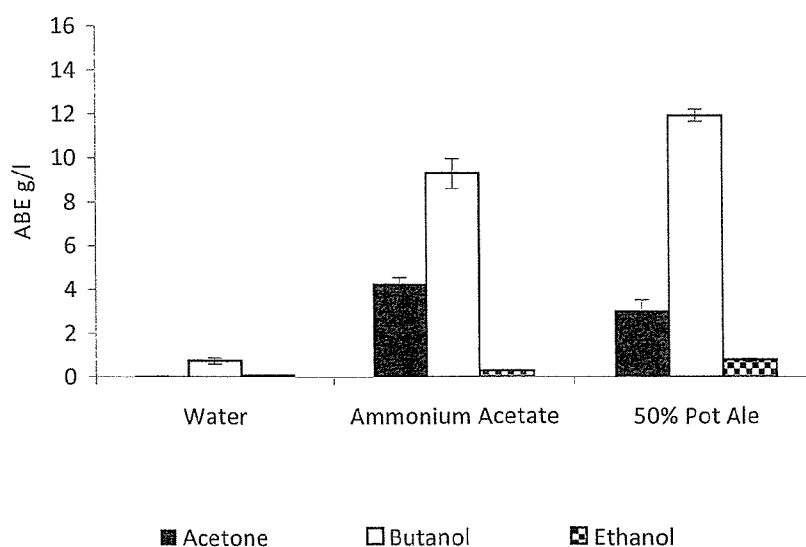
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(54) Title: PROCESS FOR THE MANUFACTURE OF BUTANOL OR ACETONE

Figure 3

(57) Abstract: A process for the
manufacture of butanol, acetone and/
or other renewable chemicals is pro-
vided wherein the process utilises
one or more of the group comprising
by-products of the manufacture of
malt whisky, such as pot ale and/or
spent lees, biomass substrates, such
as paper, sludge from paper manufac-
ture and spent grains from distillers
and brewers, and diluents, such as
water and spent liquid from other fer-
mentations. The process comprises
treating a substrate to hydrolyse it
and fermenting the treated. Also pro-
vided is a biofuel comprising butanol
manufactured according to the pro-
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PROCESS FOR THE MANUFACTURE OF BUTANOL OR ACETONE

Field of the Invention

The present invention relates to a process for the manufacture of biofuels and renewable chemicals. More particularly, the invention relates to a process for the manufacture of butanol. The invention further relates to a process for the manufacture of acetone.

Background to the Invention

In recent years, higher oil prices, depleting fuel supplies and environmental concerns have led to a renewed interest in the production of fuels from biomass ("biofuels"). Biobutanol is produced by fermentation of biomass using bacteria, typically of the genus *Clostridium*. In addition to butanol, these organisms also produce acetone, which is an important solvent, and ethanol. Such a fermentation reaction is often referred to as the "ABE process" (Acetone/Butanol/Ethanol process). Currently used feedstocks or substrates include energy crops, such as sugar beets, sugar cane, corn grain and wheat, as well as agricultural by-products, such as straw and corn stalks. Such substrates provide a balance of nutrients necessary to promote the growth of the organism during fermentation. The use of biobutanol as a fuel has several advantages over the use of ethanol. However, as biobutanol production is currently more expensive than ethanol production it has not been commercialized on a large scale.

Many current substrates for the manufacture of butanol and/or acetone by fermentation utilize the energy crops discussed above, which are already in high demand as ingredients for foodstuffs rather than as substrates for biofuels. A need therefore exists to provide a process for the manufacture of butanol and/or acetone utilizing alternative substrates.

As well as energy crops and agricultural by-products, other organic materials, such as organic waste products and/or the organic by-products of manufacturing processes, may be potential substrate components. However, many prove to be unsuitable. For instance, some organic waste or by-products may not contain the balance of nutrients suitable to promote the growth of the organism, may be harmful to the organism and/or may not contain sufficient essential nutrients in a form accessible to the organism. For this reason, many organic waste or by-products are of limited utility as substrates and consequently have little economic value.

The present invention provides a process for the manufacture of butanol and/or acetone utilizing a substrate comprising pot ale, a low value organic by-product produced as part of the manufacture of whisky, particularly malt whisky.

Malt whisky refers to whisky which has been produced from no grain other than malted barley. Production of malt whisky begins with malting of barley by steeping the barley in water. Malting releases enzymes that break down starches in the grain and convert them into sugars. When the desired state of germination is reached, the malted barley is dried. The dried malted barley is mashed in a mash-tun. In mashing, the enzymes that were developed during the malting process are allowed to convert or hydrolyse the barley starch into sugar. The resulting liquid which contains the sugars is referred to as wort. This is transferred to a large vessel called a washback where it is cooled and allowed to ferment to form the "wash". The residue remaining after extraction of the soluble sugars or wort is known as draff. This comprises spent barley solids or spent grains.

The wash is distilled in a copper distillation vessel or pot still known as a wash still to produce an alcohol-containing liquid distillate, known as low

wines. The distillation residue or liquor remaining in the pot still after the first distillation of spirit is known as pot ale or burnt ale. The low wines are distilled for a second and sometimes a third time in spirit stills to produce raw spirit, which is matured in oak casks to produce malt whisky. The remaining liquor in the second and subsequent distillations is called spent
5 lees.

The by-products of the manufacture of malt whisky therefore comprise draff, pot ale and spent lees. Pot ale has low total solids content and
10 contains dead yeast cells, yeast residue, soluble protein, soluble nutrients, carbohydrates and other material from the fermentation and mashing steps. It can also contain a significant amount of copper from the stills themselves. Pot ale is low in nutrients and may have high copper content, making it a difficulty for disposal within the industry. Significant quantities
15 are disposed of in the sea at a cost to the industry.

Summary of the Invention

The inventors of the present application have developed a process for the
20 manufacture of butanol and/or acetone and optionally other chemicals which utilises pot ale, a low or negative economic value by-product obtainable from the manufacture of malt whisky, and a carbohydrate source, such as one obtained from an organic waste product and/or an organic by-product.

25

According to a first aspect of the present invention there is provided a process for the manufacture of butanol and/or acetone, comprising at least the steps of:

- treating a substrate comprising a carbohydrate source and pot ale to solubilise at least a portion of the carbohydrate source to provide a treated substrate; and

- 5 - fermenting the treated substrate in the presence of a culture of butanol- and/or acetone-forming micro-organisms at a concentration of free copper ions of less than 20 μM to provide a fermented product containing butanol and/or acetone.

10 In one embodiment, the pot ale is the by-product of the manufacture of malt whisky, preferably Scotch malt whisky.

15 In a further embodiment, the carbohydrate source may include, but is not limited to, municipal organic waste, industrial organic waste, agricultural crops and crop residues, wood and forestry waste, marine biomass and bio-energy crops. Thus, the carbohydrate source may be selected from one or more of the group comprising paper, sludge from paper manufacture, spent grains such as those derived from grain distillers and brewers, fruit and vegetable waste, waste from the baking industry, seaweed and seaweed extracts, wood chip and other forestry derivatives, food crops, grain and crop residues, chocolate, algae (macro and micro algae), non-edible crops (and residues) and energy crops, such as switchgrass. According to a preferred embodiment the carbohydrate source is not malt whisky draff.

25 In particularly preferred embodiments, the substrate is paper, such as waste paper. Alternatively the substrate may be food waste and/or wheat or wheat-derived substrates such as bread.

In a further embodiment, the fermented product may further comprise one or more of the compounds selected from the group comprising ethanol, carbon dioxide, hydrogen, acetate and butyrate.

- 5 In another embodiment, the substrate may further comprise a diluent. In certain embodiments, the diluent is selected from one or more of the group comprising water, spent lees and spent liquid from other fermentations. In particular, the diluent may be water.
- 10 In a still further embodiment the step of treating the substrate may further comprise:
- diluting the substrate to provide a concentration of free copper ions of less than 20 μM .
- 15 In one embodiment the step of treating the substrate may comprise:
- hydrolysing the carbohydrate source in the presence of water and hydrogen ions or water and hydroxide ions.
- In another embodiment, the step of treating the substrate may comprise:
- 20 - hydrolysing the carbohydrate source in the presence of an aqueous solution of sulphuric acid.
- In a further embodiment, the step of treating the substrate does not comprise heating the substrate at high temperature (i.e. above 120 °C, preferably above 100 °C, and more preferably above 80 °C) and/or
- 25 treating the substrate with an acid, such as sulphuric acid. This particular embodiment is particularly suitable for the treatment of paper or paper-derived products as substrates.
- 30 In yet another embodiment, the step of treating of the substrate comprises:

- treating the carbohydrate source with one or more enzymes.

5 In one embodiment, the step of fermenting the treated substrate may be carried out at a pH of 5.1 or above, more typically in a pH range of from 5.3 to 5.7, still more typically at a pH of about 5.5. In a further embodiment, the pH may be adjusted during the step of fermenting the treated substrate with a buffer. Alternatively no pH adjustment is required, in particular when the substrate is from wood or paper origin.

10 In another embodiment, the concentration of free copper ions in the fermentation step can be less than 15 μ M. In a further embodiment, the concentration of free copper ions in the fermentation step can be reduced by the addition of a diluent to the treated substrate, such as a diluent described above.

15 In a still further embodiment, the culture of butanol- and/or acetone-forming micro-organisms may comprise bacteria of the genus clostridium such as: *C. acetobutylicum* ATCC 824, *C. saccharolyticum* NCP 262, *C. beijerinckii* NCIMB 8052 and *C. saccharoperbutylacetonicum* NCIMB
20 12606 (N1-4).

In another embodiment, the fermentation step can be carried out without the removal of solids therefrom.

25 In further embodiments, the treating and fermenting steps can be carried out simultaneously or can be carried out sequentially.

In one embodiment, the pot ale may be provided by the steps of:

- adding yeast to a wort comprising water and one or more carbohydrates selected from the group comprising glucose and oligosaccharides of glucose;

5 - fermenting the wort to provide a wash comprising water and one or more alcohols; and

- distilling the wash in a copper distillation vessel to provide a low wines distillate comprising one or more alcohols and a distillation residue of pot ale.

10 In a further embodiment, the wort can be provided by the further steps of:

- grinding malted barley comprising starch to provide ground malted barley;

- mixing the ground malted barley with water to provide a mash comprising water and ground malted barley;

15 - hydrolysing at least a part of the starch in the ground malted barley of the mash to provide draff comprising spent barley solids and wort comprising water and one or more carbohydrates selected from the group comprising glucose and oligosaccharides of glucose; and

- separating the draff from the wort.

20

According to a further aspect of the present invention there is provided a biofuel comprising butanol manufactured according to the process of any of the aspects of the present invention.

25 According to a further aspect of the invention there is provided a product comprising butanol and/or acetone manufactured according to the process of any aspects of the present invention.

According to a yet further aspect of the present invention there is provided use of pot ale from the production of malt whisky in the manufacture of butanol and/or acetone by fermentation

5 Detailed Description of the Invention

The present invention utilises pot ale and a carbohydrate source as a substrate in the manufacture of butanol and/or acetone by a fermentation process. In particular, the present inventors have surprisingly discovered
10 that it is possible to carry out fermentation in the presence of pot ale. It was expected that the high copper content in the pot ale from the copper pot stills would inhibit butanol- and/or acetone-forming micro-organisms, such as bacteria of the genus *Clostridium*. However, the present inventors have shown that when the substrate has concentration of free copper ions
15 to below 20 μM , there is no inhibitory effect.

Pot ale provided direct from the first distillation residue from the copper pot generally has a concentration of free copper ions at levels which are detrimental to butanol- and/or acetone-forming micro-organisms, such as
20 copper concentrations in excess of 20 μM . Consequently, unmodified pot ale may be an unsuitable medium for forming the biomass substrate. However, free copper ion concentrations which are not inhibitory for the butanol- and/or acetone-forming micro-organisms can be achieved by reducing the free copper ion concentration, for instance by diluting the pot
25 ale, either before it is combined with the carbohydrate source to provide the substrate, after it has been combined with the carbohydrate source to provide the substrate, or after the treatment of the substrate to provide the treated substrate. However, it will be apparent that whenever the reduction in free copper ion concentration is carried out, this should
30 preferably be before combination of the treated substrate comprising the

pot ale with the butanol- and/or acetone-forming micro-organisms in order to prevent the micro-organisms being harmed.

5 The use of pot ale in the manufacture of butanol, acetone and/or other renewable chemicals has several associated advantages. Pot ale is currently categorised as being of low or negative economic value. The use of pot ale in the present invention allows the economic value of pot ale to be increased. Furthermore, the pot ale acts as a diluent to hydrate the carbohydrate source. Thus, the amount of water or other diluent required
10 is reduced when pot ale is used. In addition, pot ale provides essential nutrients to the microorganisms, particularly providing a source of nitrogen which thereby improves the fermentation and overall conversion of substrate to products.

15 The use of organic waste or by-products as the carbohydrate source in the substrate for the manufacture of butanol and/or acetone and optionally other chemicals is also advantageous as it provides a solution to the disposal of these substances. In certain aspects, the present invention utilises paper, and in particular waste paper, as a carbohydrate source.
20 The present invention therefore further provides a solution to the disposal of waste paper, for example, old newspapers or used photocopier paper. It should be noted that the use of pot ale has been found necessary to create a suitable fermentation medium for the production of butanol and/or acetone from paper based substrates.

25 The substrate must be treated to solubilise at least a part of the carbohydrate source. The solubilisation may take the form of hydrolysis, thus breaking down the carbohydrate source into a form suitable for fermentation. Accordingly, in certain embodiments the substrate is
30 subjected to one or more treatment steps to solubilise and particularly

hydrolyse the carbohydrate source, for example, mashing, heating, addition of acid or alkali, addition of enzymes or a combination thereof. In certain embodiments, the treating of the substrate to solubilise the carbohydrate source comprises the step of hydrolysing the carbohydrate source in the presence of water and hydrogen ions or water and hydroxide ions. In certain embodiments, the treating of the carbohydrate source to solubilise it is carried out in the presence of any suitable acid which is capable of hydrolysing the carbohydrate source. Examples of suitable acids include sulphuric acid and nitric acid. Sulphuric acid is a preferred example of an acid for use in the process described herein. When the substrate is paper based, or paper derived, no temperature or acid treatment is required.

In certain embodiments, the treating of the substrate to hydrolyse the carbohydrate source comprises addition of one or more enzymes, such as cellulase and hemicellulase. In certain embodiments, a combination of treatments may be utilised, for example, addition of both acid and enzymes, to provide a treated substrate in a form suitable for fermentation. The combination of treatments may be applied simultaneously or sequentially.

In another embodiment wherein the substrate comprises at least one of paper, paper waste, wheat and wheat based products such as bread, the treatment may comprise the addition of enzymes. This enzyme treatment may occur at a temperature lower than that which is optimal for enzyme activity (i.e. about 50°C). More specifically the temperature may be the same as the fermentation temperature (e.g. 33°C), allowing both processes to proceed simultaneously. Suitable temperature ranges can be 25 to 40 °C, preferably 30 to 35°C, and more preferably 32 to 34°C. Only enzyme treatment may be required because steps earlier in the manufacture of

products such as bread and/or paper waste can concentrate the carbohydrate content and remove inhibitory materials (e.g. lignin from paper based products).

5 Fermentation of the treated substrate is carried out at pH in the range of 5.1 or above, more preferably in the range of from 5.2 to 6.2, still more preferably in the range of from 5.3 to 5.7 and yet more preferably at about 5.5. The use of the pH range of from 5.3 to 5.7 has been shown to provide high yields of butanol and/or acetone. Furthermore, this pH range
10 allows fermentation to be carried out without the need to remove solids therefrom, thus reducing costs and avoiding any technical problems caused by the requirement to remove solids. This pH range prevents any potential toxicity from the treated substrate while maximising butanol and/or acetone production.

15 Fermentation is carried out in the presence of a culture of butanol- and/or acetone-forming micro-organisms. The butanol- and/or acetone-forming micro-organisms may be selected from any solvent producing micro-organisms which are capable of fermenting the substrate to form butanol and/or acetone. Suitable micro-organisms include micro-organisms
20 engineered to produce solvents. Examples of suitable micro-organisms include those currently used in ABE (Acetone/Butanol/Ethanol) manufacture, and, in particular, bacteria of the genus *Clostridium* such as *C. acetobutylicum*, *C. beijerinckii*, *C. saccharoperbutylacetonicum* and *C. saccharobutylicum*. In some embodiments, the butanol- and/or acetone forming micro-organisms comprise *C. acetobutylicum*. In other
25 embodiments the micro-organisms comprises *C. Saccharoperbutylacetonicum*. This particular strain is particularly effective on paper-based substrates.

30

Fermentation is carried out at a concentration of free copper ions of less than 20 μM . This ensures that the presence of the copper ions have no/minimal negative effect. In certain embodiments, water or another aqueous solution may be added to lower the concentration of free copper ions to below 20 μM free copper ions. In certain embodiments, the concentration of free copper ions is less than 19, 18, 17, 16, 15, 14, 13, 12, 11, 10, 9, 8, 7, 6 or 5 μM free copper ions during at least the fermentation step. In certain embodiments, the concentration of free copper ions is less than 15 μM . In certain embodiments, the concentration of free copper ions is less than 10 μM .

In certain embodiments, the treating and fermenting steps are carried out simultaneously. This reduces the amount of time required, the number of steps involved and the associated cost of manufacture.

In alternative embodiments, the treating and fermenting steps are carried out sequentially. For example, the substrate may be pre-treated in two steps, first with acid and subsequently with enzyme, prior to fermentation.

In certain embodiments, the fermented product further comprises one or more of the compounds selected from the group comprising ethanol, carbon dioxide, hydrogen, acetate and butyrate. Butanol and/or acetone may be separated out of the fermented product using conventional separation techniques. Alternatively, the fermented product may be used as a fuel or otherwise without further purification.

The term "biobutanol" as used herein refers to butanol made from biomass.

The term “pot ale” as used herein refers to the liquor remaining in the wash (copper pot) still after the first distillation in the manufacture of malt whisky. It is the residue of the wash after extraction by distillation of the low wines.

5

The term “spent lees” as used herein refers to the liquor remaining in the distillation vessel after second and subsequent distillations in the manufacture of malt whisky. It is the residue of the low wines after extraction by distillation of raw spirit.

10

The term “concentration of free copper ions” refers to the concentration of copper ions which is not bound to solids, that is, the concentration of copper ions in the supernatant. The total concentration of copper in the pot ale will be higher than the concentration of free copper ions as some copper remains bound to solids, such as dead yeast cells.

15

The term “Scotch whisky” as used herein refers to whisky made in Scotland. In alternative embodiments, the malt whisky is a malt whisky manufactured in other countries, such as Ireland or India, where the process for manufacture of malt whisky in that country is similar or identical to the process used in Scotland for the manufacture of Scotch malt whisky.

20

The present invention will now be described with reference to the following examples which are provided for the purpose of illustration and are not intended to be construed as being limiting on the present invention.

25

Brief Description of the Figures

Figure 1 shows the influence of initial pH on fermentation of acid and enzyme pre-treated draff in pot ale by *C. acetobutylicum* ATCC 824. Draff was pre-treated with 0.08 M H₂SO₄ and the pH adjusted to between pH 5.0 – 6.0 prior to enzyme addition. After enzyme hydrolysis, the pH was adjusted to 4.5, 4.8, 5.0, 5.5, 6.0 or 6.5 for fermentation. Figure 1 (a) shows sugars resulting from acid and enzyme treatment, Figure 1 (b) shows residual sugars after fermentation, Figure 1 (c) shows the ABE products from fermentation and Figure 1 (d) shows yield of butanol and ABE from draff;

Figure 2 shows ABE production by *C. saccharoperbutylacetonicum* NCIMB 12606 from (a) white office paper and (b) newspaper dissolved in either water or 50% pot ale.

Figure 3 shows the production of acetone, butanol and ethanol from fermentations (150 ml) carried out with white paper solubilised with water, ammonium acetate or 50% pot ale.

Figures 4a to 4c show the production of acetone, butanol and ethanol from fermentations (150 ml) carried out with four strains of clostridia on a) white Paper, b) newsprint and c) leaflet Paper. Substrates were diluted in 50% pot ale, buffered to pH 5.5 and treated with industrial enzymes.

Figure 5 shows the production of acetone, butanol and ethanol from fermentations (150 ml) carried out with four strains of clostridia on bread in pot ale.

Figure 6 shows the production of acetone, butanol and ethanol from fermentations (150 ml) carried out with four strains of clostridia on wheat in pot ale.

5 Figure 7 shows the production of acetone, butanol and ethanol from fermentations (150 ml) carried out with two strains of clostridia on food waste in pot ale.

10 Figure 8 shows the production of acetone, butanol and ethanol from fermentations (150 ml) carried out with four strains of clostridia in pot ale with wheat that had been subjected to acid treatment.

15 Figure 9 shows the production of acetone, butanol and ethanol from fermentations (1 litre) carried out with newsprint, white paper and leaflet paper without buffering.

EXAMPLES

20 Reference Example 1 and Example 2 and 3 show the effect of pH and free copper ion concentration of the substrate upon the fermentation process. In both of Reference Example 1 and Example 3, glucose was used as the carbohydrate source. Reference Example 1 is designated as a reference test because no pot ale was present in the substrate.

25 Reference Example 1 shows that ABE can be produced from the fermentation of glucose in a pH range of above 4.5 to 6.5. At a pH of 4.5 no glucose was consumed or ABE produced. At a pH of 6.5, only acids were produced, with no ABE. Example 2 shows a preferred initial range pH of about 5.5 for the fermentation of a substrate comprising pot ale and
30 a substrate (draff).

Example 3 shows that undiluted pot ale has a concentration of free copper ions which is detrimental to fermentation, resulting in a reduction in ABE yield compared to a substrate without added free copper ions.

- 5 Furthermore as the concentration of free copper ions was reduced from 20 μM to 5 μM , the yield of ABE increased to that obtained from a substrate with no added free copper ions.

- 10 Example 4 shows that the addition of pot ale to a substrate results in an increase in ABE yield, compared to the addition of a diluent such as water.

- Example 5 shows that the addition of pot ale to a substrate has the same effect as the addition of ammonium sulphate, indicating the pot ale acts as a nitrogen source.

- 15 Example 6 shows that ABE can be produced using pot ale in combination with various substrates and microorganisms. The substrates exemplified are various types of paper material, bread, wheat and food waste.

- 20 Example 7 shows that certain substrates do not require acid treatment. Indeed acid treatment of wheat produces lower yields of ABE than untreated wheat.

- 25 Example 8 shows that pre-fermentation buffering is not always required on paper based substrates.

General Methods

- 30 The following organisms were used: *C. acetobutylicum* ATCC 824, *C. saccharolyticum* NCP 262, *C. beijerinckii* NCIMB 8052 and *C.*

saccharoperbutylacetonicum NCIMB 12606 (N1-4). Clostridia were maintained as spore suspensions at 4°C. Spores were heat shocked at 80°C for 10 minutes and inoculated into reinforced clostridia media (RCM, Oxoid Ltd, Cambridge, UK).

5

In examples 1 to 3 the cultures were incubated for 24 hours and then subcultured into tryptone-yeast extract-ammonium acetate media (TYA) media containing glucose before being used as a starting culture (at 5% v/v) for all experiments. TYA consisted of (g/l) tryptone, 6; yeast extract, 2; ammonium acetate, 3; KH₂PO₄, 0.5; MgSO₄·7H₂O, 0.3; FeSO₄·7H₂O, 0.01 supplemented with 5% glucose. All clostridia cultures were incubated in an anaerobic workstation under an N₂-H₂-CO₂ (80:10:10) atmosphere at 33°C. In examples 4 to 8 starter cultures were incubated for 24 h and then subcultured into 50% pot ale buffered to pH 5.5. All clostridia cultures were incubated in an anaerobic workstation under an N₂-H₂-CO₂ (80:10:10) atmosphere at 33°C. Small scale fermentation (150 ml) was carried out in glass bottles in the anaerobic workstation.

For 1 L scale, fermentations were conducted in fermenters (Biostat A Plus, Sartorius Stedim Ltd, Surrey, UK). Oxygen-free conditions were achieved by sparging the media in the fermenters with oxygen-free N₂ for 1 hour prior to inoculation with clostridia. For all 1 L fermentations, agitation was set at 200 rpm and temperature at 33°C.

Pot ale was obtained from distilleries and was diluted in a 1:1 ratio by volume with water. This was sufficient to reduce the concentration of free copper ions to less than 20 µM.

Wet draff, as received from the distilleries, had a moisture content between 75-80%. Where stated, draff was dried at 80°C to a moisture content of approximately 4% and milled prior to further processing.

5 Ethanol, Acetone and Butanol were measured using a Chrompack 9001 gas chromatograph equipped with a flame ionisation detector and a CP SIL 5CB column of length 10 m and diameter 0.32 mm (all Chrompack, Middelburg, Netherlands). All samples were filtered through 0.2 µm cellulose acetate syringe filters before analysis and concentrations were
10 determined by reference to ethanol, acetone and butanol standards.

For acid (acetic and butyric) and monosaccharide (glucose, xylose and arabinose) analysis, samples were filtered through 0.2 µm syringe filters and acidified with H₂SO₄. Samples were analysed by HPLC using a
15 Varian 920 LC fitted with integrated UV-VIS dual wavelength and refractive index detectors (Varian Ltd., Oxford, UK). Components were separated at room temperature on a Rezex ROA Organic acid H⁺ 8% 300 × 7.8 mm column (Phenomenex, Cheshire, UK) with 0.005 N H₂SO₄ as the mobile phase at a flow rate of 0.5 ml/min. Acids were detected at 210 nm
20 while sugars were detected with the RI detector and concentrations were determined by reference to the corresponding standards.

Reference Example 1 – Effect of pH control on ABE production by clostridia

25 The effect of pH on fermentation of glucose in TYA media by *C. acetobutylicum* ATCC 824 was investigated. Fermentations were conducted at 1 L scale and the pH was controlled at a range of set points between pH 4.5-6.5 with automated addition of either alkali or acid. At pH 4.5, no glucose utilisation, acid or ABE production was detected. For all
30 other fermentations, glucose was completely consumed within 48 hours

and acids (butyric and acetic) and solvent (acetone, butanol and ethanol) were produced (Table 2). ABE production was highest at pH 4.8 and 5.0, corresponding to yields of 0.34 and 0.30 g ABE/g sugar, respectively. Acid production increased between pH 5.5 to 6.5, with a corresponding decrease in conversion of sugar to ABE. At pH 6.5, acids only were produced with final concentrations of 7.8 and 12.8 g/l acetic and butyric acid, respectively.

Table 1. Conversion of 5% glucose to acid and ABE by *C. acetobutylicum* ATCC 824 in TYA media controlled at either pH 4.8, 5.0, 5.5, 6.0 or 6.5. Acid (butyric and acetic) and ABE concentrations were determined after 68 hours with ABE yield expressed as g of ABE produced per g of sugar consumed.

pH	Acid (g/l)	ABE (g/l)	Yield (g ABE/g sugar)
4.8	0.7	15.2	0.34
5.0	0.9	14.3	0.30
5.5	7.9	12.3	0.25
6.0	13.6	6.7	0.13
6.5	20.5	0.6	0.01

15

Example 2 – Influence of initial pH on ABE production by clostridia

The effect of initial pH on fermentation of a pre-treated substrate comprising draff and pot ale was investigated. Dried, milled draff was pre-treated by adding 10.5% (w/v) to 250 ml duran bottles with 0.08 M H₂SO₄ in 50% pot ale and sterilised at 121°C for 15 min. After cooling, the pH was adjusted to between pH 5.0 – 6.0 by addition of 10 M NaOH and incubated with cellulase and hemicellulase enzymes at 33°C for 24 hours. For fermentation, the initial pH of the solutions was adjusted to either 4.5, 4.8, 5.0, 5.5, 6.0 or 6.5 prior to inoculation with *C. acetobutylicum* ATCC

20

824. The initial sugar concentration was monitored before fermentation and the residual sugar, ABE concentration and ABE yield were calculated after fermentation (Fig 1). The initial concentration of sugars was similar for all samples, with approximately 9.6, 11.2, and 9.9 g/l glucose, xylose and arabinose. No growth or gas production was apparent at pH 5.0 or lower and no sugars were utilised. ABE production was greatest at pH 5.5 (14.2 g/l) with a yield of 13.2 g/100 g draff. This was reduced at pH 6.0, with 9.3 g ABE/100 g draff. At pH 6.5, approximately half the sugar was utilised but there was poor conversion to ABE with a final concentration of 2.3 g/l.

Example 3 – Pot ale as a growth medium for ABE production by clostridia

Pot ale was collected from a Scottish malt distillery and analysed for copper content. The pot ale had 71.8 μM total Cu of which 21.1 μM was determined to be available as "free" Cu in the supernatant with the rest bound to the solids. To assess whether this Cu concentration was toxic to *C. acetobutylicum* ATCC 824, fermentation of 5% glucose in 100 ml TYA media supplemented with different concentrations of Cu was compared (Table 3). Cu had no effect on ABE production at 5 and 10 μM with ABE concentrations of approximately 12 g/l being similar to that of the control without Cu. At the higher Cu concentration, ABE concentration was reduced to 8.6 g/l, indicating that at this concentration Cu was inhibitory to clostridia. As the pot ale had a "free" Cu content of 21.1 μM , it was decided to test clostridia fermentation in half strength pot ale in order to reduce the Cu concentration below inhibitory levels. Half-strength pot ale supplemented with glucose provided enough nutrients for growth of 824 with ABE production similar to the TYA control (Table 3).

Table 2. Conversion of 5% glucose to ABE by *C. acetobutylicum* ATCC 824 in either TYA, TYA containing 5, 10 or 20 μ M Cu or 50% pot ale.

Media	ABE (g/l)
TYA	12.4 $\pm$ 0.3
TYA, 5 μ M Cu	12.3 $\pm$ 0.3
TYA, 10 μ M Cu	11.6 $\pm$ 0.1
TYA 20 μ M Cu	8.6 $\pm$ 2.0
50% pot ale	12.0 $\pm$ 1.7

5

Example 4 – Process for conversion of waste paper to butanol and acetone

White office paper and newspaper were shredded to 5 mm wide strips and 6.7% (w/v) was mixed with either water or 50% pot ale in 250 ml duran bottles and the pH adjusted to pH 5.5. After sterilisation, the bottles were cooled and cellulase and *C. saccharoperbutylacetonicum* NCIMB 12606 added. After fermentation, the ABE concentrations were determined (Fig 2). There was poor conversion of paper to ABE in water compared to pot ale, demonstrating that pot ale was required to provide additional nutrients. In pot ale, the ABE yields after fermentation with *C. saccharoperbutylacetonicum* were 24.8 g ABE per 100 g office paper and 16.8 g ABE per 100 g newspaper.

Example 5 – Effect of pot ale as a nitrogen source on ABE production by clostridia

Certain substances which have had their carbohydrate content increased by processing may be deficient in nitrogen. A substrate comprising paper

as a carbohydrate source and 50% pot ale was compared to a substrate comprising paper and water and a substrate comprising paper and aqueous ammonium acetate. The modus operandi used was the same than the one use for the previous example.

5

As shown in figure 3, fermentation of the substrate comprising paper and water produced little ABE. Supplementing a substrate comprising paper and water with ammonium acetate produced good results, indicating that a paper carbohydrate source was deficient in nitrogen. The substrate
10 comprising paper and pot ale produced even better results as the substrate supplemented with ammonium acetate. Figure 3 shows that fermentations using either ammonium sulphate or pot ale produced more ABE than those using water. The strain used was N1-4.

15 Example 6 – Production of ABE from a range of substrates comprising pot ale

A range of substrates comprising a carbohydrate source were selected from the group comprising white paper, newsprint, leaflet paper, bread (white from Kingsmill), wheat and food waste. The first five carbohydrate
20 sources represent readily available waste materials, while wheat represents an alternative agricultural crop. The carbohydrate sources were prepared as follows: paper samples were shredded using a commercial paper shredder, the bread, wheat and food waste samples were shredded in a domestic food processor. The carbohydrate sources were solubilised
25 in 50% pot ale, buffered at pH 5.5 with sodium hydroxide, treated with industrial enzymes (C-Tech and H-Tech, commercially available cellulase and hemicellulase from Novozymes) and fermented with the four strains of clostridia bacteria previously mentioned. All the fermentations produced ABE (shown in Figures 4 – 7). The N1-4 strain was particularly efficient

with respect to the paper based substrates as the ABE concentration produced was consistently above 7 g/l and can even be above 12 g/l.

Example 7 – Pre-treatment of different carbohydrate sources

- 5 Malt whisky draff requires treatment with acid at high temperature to render its structure accessible to the enzymes used to release sugars for fermentation. In contrast, carbohydrate sources such as bread and paper wastes do not require treatment as processes carried out earlier in the manufacturing process have removed inhibitory materials such as lignin
10 and concentrated the carbohydrate content. Thus, such carbohydrate sources require only enzyme treatment at low temperature before fermentation.

- Figure 8 shows the concentration of ABE obtained after wheat is subjected
15 to an acid treatment. As it can be seen when compared with the results shown in figure 6, when used on wheat, the treatment does not lead to any improvements and can even show severe detrimental effects with respect to particular strains of clostridia (i.e. P262 and N1-4).

Example 10 – Pre-fermentation buffering

- 20 Those substrates that have not undergone acid treatment have higher pH values than those that have undergone acid treatment. Thus, buffering to the fermentation pH in those substrates which have not undergone acid treatment may be less intensive and in some cases is not required. As
25 shown in figure 9, when paper is used as a carbohydrate source, buffering was not required because the optimal strain, N1-4, can ferment at a higher pH such that satisfactory performance was achieved without buffering before inoculation.

Although the invention has been described in connection with specific preferred embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments.

Indeed, various modifications of the described modes of carrying out the
5 invention which are obvious to those skilled in the art are intended to be covered by the present invention.

Claims

1. A process for the manufacture of butanol and/or acetone, comprising at least the steps of:
 - treating pot ale and a biomass substrate other than draff, pot ale being a by-product of the first spirit distillation from a copper distillation vessel in the manufacture of malt whisky, to solubilise at least a portion of the biomass substrate to provide a treated substrate; and
 - fermenting the treated substrate in the presence of a culture of butanol- and/or acetone-forming micro-organisms at a concentration of free copper ions of less than 20 μM to provide a fermented product containing butanol and/or acetone.
2. The process as claimed in claim 1 wherein the step of treating the substrate comprises:
 - hydrolysing the biomass substrate in the presence of water and hydrogen ions or water and hydroxide ions.
3. The process as claimed in claim 1 or claim 2 wherein the step of treating the substrate comprises:
 - hydrolysing the biomass substrate in the presence of an aqueous solution of sulphuric acid.
4. The process as claimed in any one of claims 1 to 3 wherein the step of treating of the substrate comprises:
 - treating the biomass substrate with one or more enzymes.
5. The process as claimed in any one of claims 1 to 4 wherein the step of treating the substrate further comprises:
 - diluting the substrate to provide a concentration of free copper ions of less than 20 μM .
6. The process as claimed in any one of claims 1 to 5 wherein the step of fermenting the treated substrate is carried out at a pH of 5.1 or above.
7. The process as claimed in any one of claims 1 to 6 wherein the concentration of free copper ions in the fermentation step is less than 15 μM .
8. The process as claimed in any one of claims 1 to 7 wherein the culture of butanol- and/or acetone-forming micro-organisms comprises bacteria of the genus *Clostridium*.

9. The process as claimed in any one of claims 1 to 8 wherein the fermentation step is carried out without the removal of solids therefrom.

10. The process as claimed in any one of claims 1 to 9 in which the treating and fermenting steps are carried out simultaneously.

11. The process as claimed in any one of claims 1 to 9 in which the treating and fermenting steps are carried out sequentially.

12. The process as claimed in any one of claims 1 to 11 wherein the pot ale is provided by the steps of:

- adding yeast to a wort comprising water and one or more carbohydrates selected from the group comprising glucose and oligosaccharides of glucose;
- fermenting the wort to provide a wash comprising water and one or more alcohols; and
- distilling the wash in a copper distillation vessel to provide a low wines distillate comprising one or more alcohols and a distillation residue of pot ale.

13. The process as claimed in claim 12 wherein the wort is provided by the further steps of:

- grinding malted barley comprising starch to provide ground malted barley;
- mixing the ground malted barley with water to provide a mash comprising water and ground malted barley;
- hydrolysing at least a part of the starch in the ground malted barley of the mash to provide draff comprising spent barley solids and wort comprising water and one or more carbohydrates selected from the group comprising glucose and oligosaccharides of glucose; and
- separating the draff from the wort.

14. The process as claimed in any one of claims 1 to 13 wherein the pot ale is the by-product of the manufacture of Scotch malt whisky.

15. The process as claimed in any one of claims 1 to 14 wherein the biomass substrate is selected from one or more of the group comprising paper, sludge from paper manufacture, fruit and vegetable waste, waste from the baking industry, seaweed and seaweed extracts, forestry derivatives, food crops, grain and crop residues, spent grains, chocolate, algae, non-edible crops and their residues and energy crops.
16. The process as claimed in any one of claims 1 to 15 wherein the fermented product further comprises one or more of the compounds selected from the group comprising ethanol, carbon dioxide, hydrogen, acetate and butyrate.
17. A process for the manufacture of butanol and/or acetone, comprising at least the steps of:
- treating a substrate comprising draff, being a by-product of the manufacture of malt whisky, and pot ale, being a by-product of the manufacture of malt whisky from a copper distillation vessel, to hydrolyse the substrate to provide a treated substrate, said draff comprising spent grain consisting essentially of malted barley; and
 - fermenting the treated substrate in the presence of a culture of butanol- and/or acetone-forming micro-organisms at an initial pH in the range of 5.0 to 6.0 and at a concentration of free copper ions of less than 20 mM to provide a fermented product containing butanol and/or acetone.
18. The process as claimed in claim 17 wherein the initial pH is in the range of 5.3 to 5.7.
19. The process as claimed in claim 17 or 18 wherein the fermentation step is carried out without the removal of solids therefrom.
20. The process as claimed in any one of claims 17 to 19 wherein the spent grain consists of 100% malted barley.
21. The process as claimed in any one of claims 17 to 20 in which the treating and fermenting steps are carried out simultaneously.
22. The process as claimed in any one of claims 17 to 20 in which the treating and fermenting steps are carried out sequentially.

23. The process as claimed in any one of claims 17 to 22 wherein the substrate is diluted to lower the concentration of free copper ions to below 15 mM during at least the fermentation step.

24. The process as claimed in any one of claims 17 to 23 wherein the draff is provided by the further steps of:

- grinding malted barley comprising starch to provide ground malted barley;
- mixing the ground malted barley with water to provide a mash comprising water and ground malted barley;
- hydrolysing at least a part of the starch in the ground malted barley of the mash to provide draff comprising spent barley solids and wort comprising water and one or more carbohydrates selected from the group comprising glucose and oligosaccharides of glucose; and
- separating the draff from the wort.

25. The process as claimed in claim 24 wherein the pot ale is provided by the further steps of:

- adding yeast to the wort;
- fermenting the wort to provide a wash comprising water and one or more alcohols; and
- distilling the wash in a copper distillation vessel to provide a low wines distillate comprising one or more alcohols and a distillation residue of pot ale.

26. The process as claimed in any one of claims 17 to 25 wherein the treating of the substrate to hydrolyse it comprises the step of:

- hydrolysing the substrate in the presence of water and hydrogen ions or water and hydroxide ions.

27. The process as claimed in any one of claims 17 to 26 wherein the treating of the substrate to hydrolyse it comprises the step of:

- hydrolysing the substrate in the presence of an aqueous solution of sulphuric acid.

28. The process as claimed in any one of claims 17 to 27 wherein the treating of the substrate to hydrolyse it comprises the step of:

- treating the substrate with one or more enzymes.

29. The process as claimed in any one of claims 17 to 28 wherein the culture of butanol- and/or acetone-forming micro-organisms comprises bacteria of the genus *Clostridium*.

30. The process as claimed in any one of claims 17 to 29 wherein the fermented product further comprises one or more of the compounds selected from the group comprising ethanol, carbon dioxide, hydrogen, acetate and butyrate.

31. The process according to any one of claims 19 to 30 wherein the malt whisky is Scotch malt whisky.

Figure 1

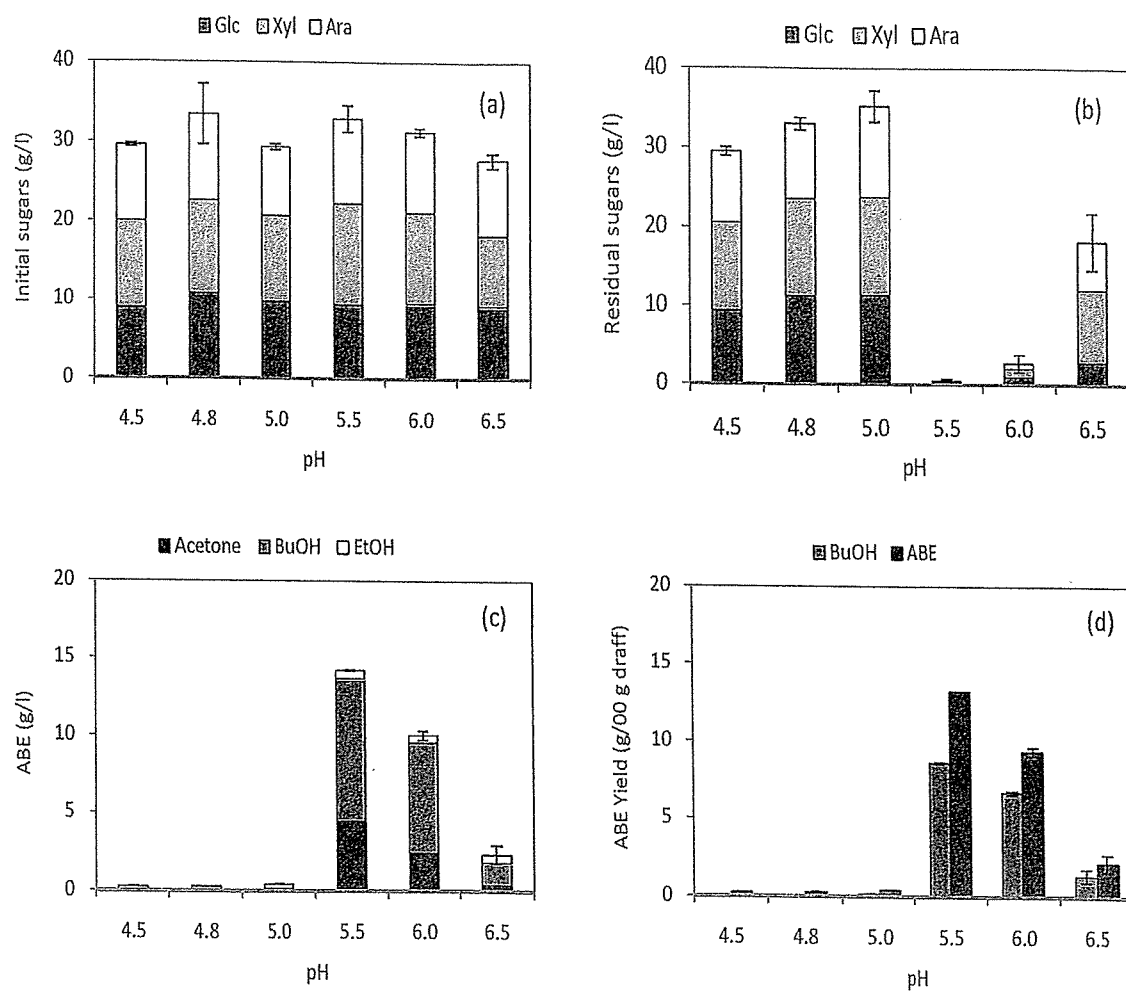


Figure 2

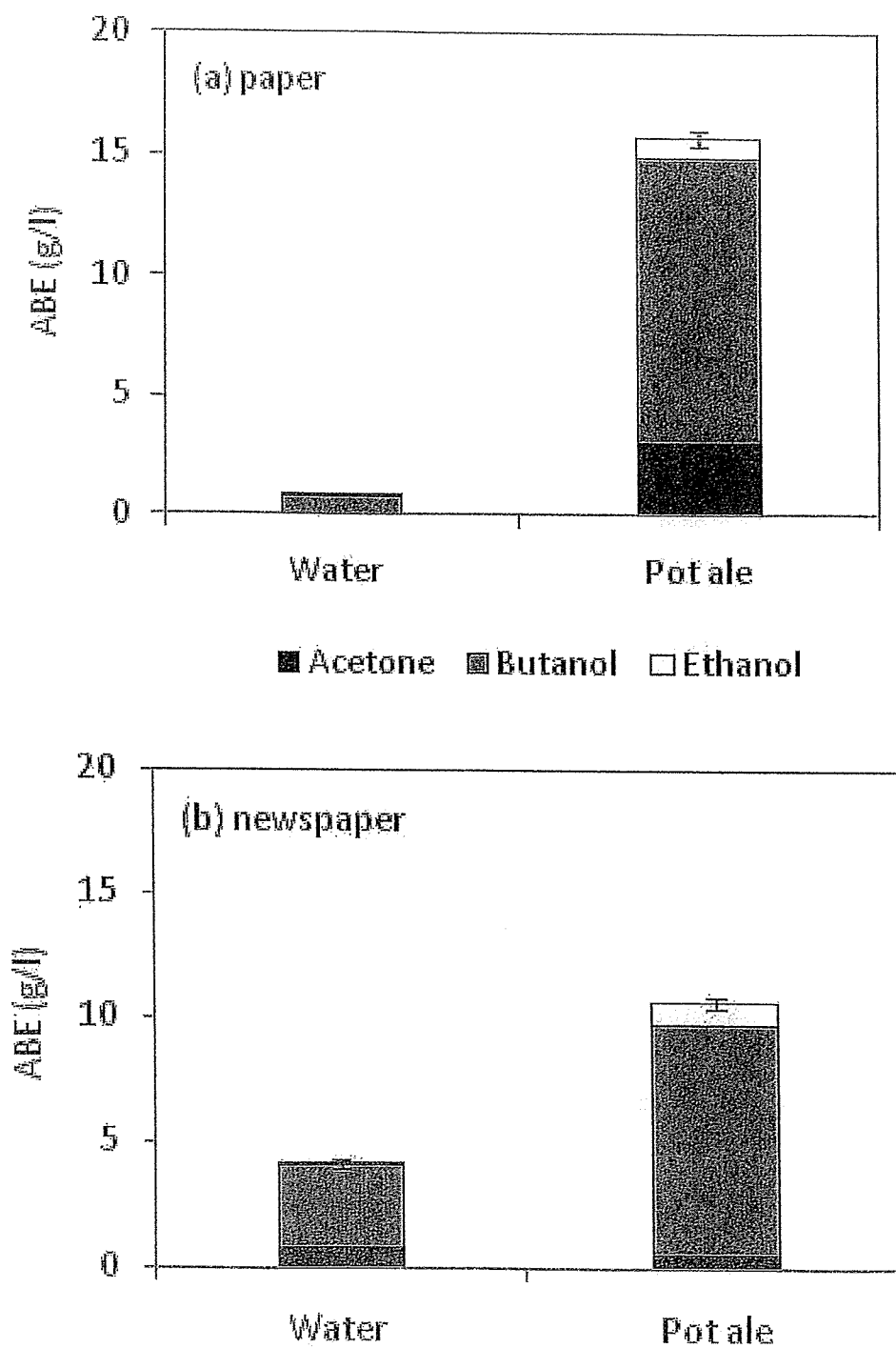


Figure 3

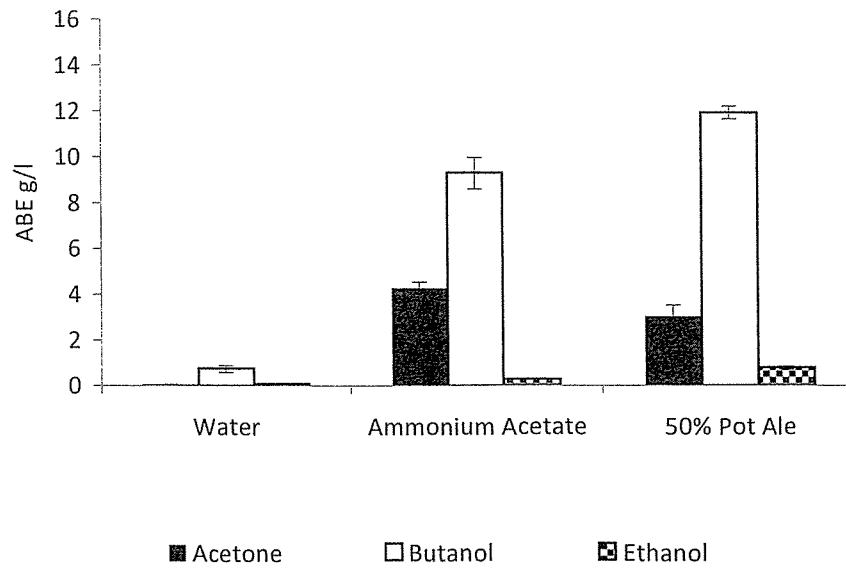


Figure 4

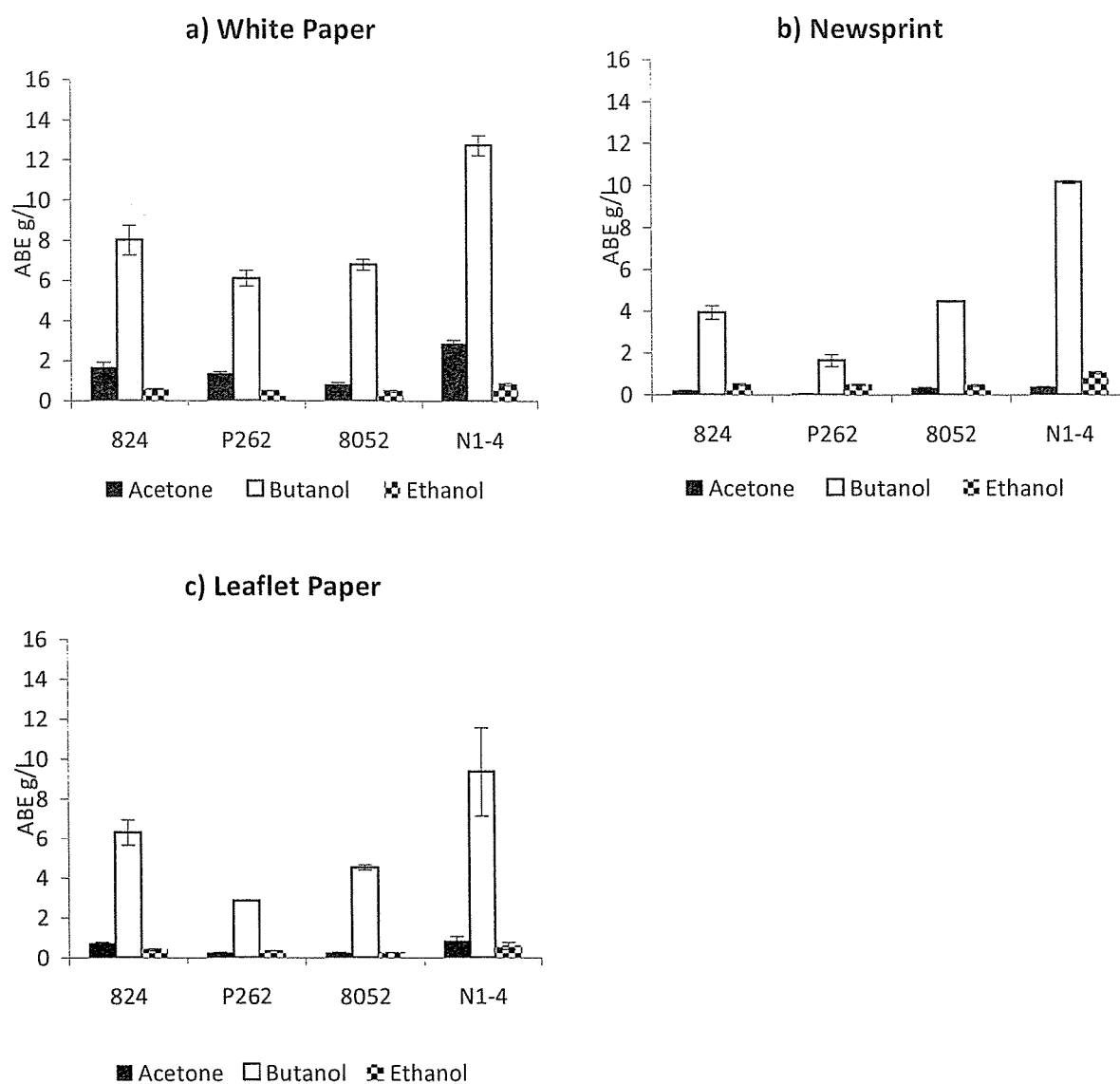


Figure 5

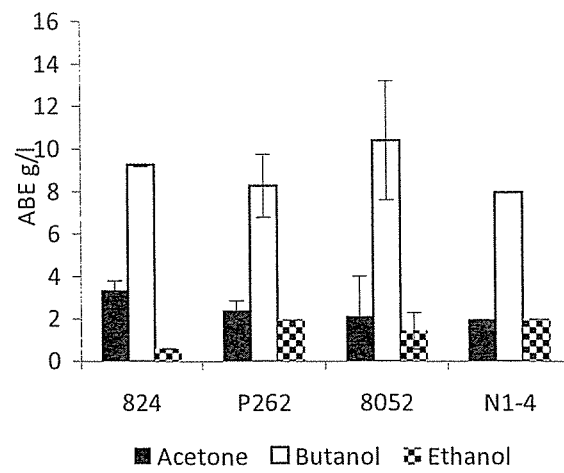


Figure 6

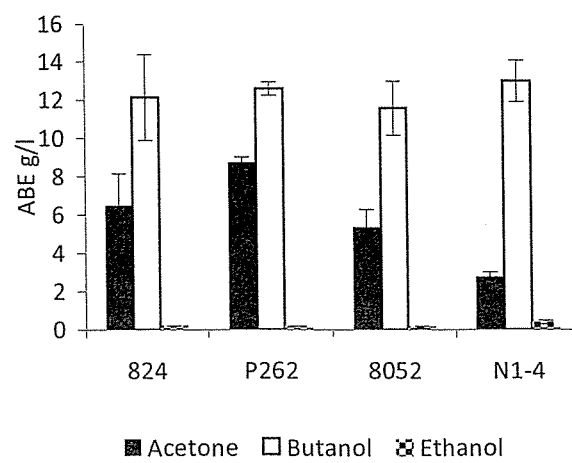


Figure 7

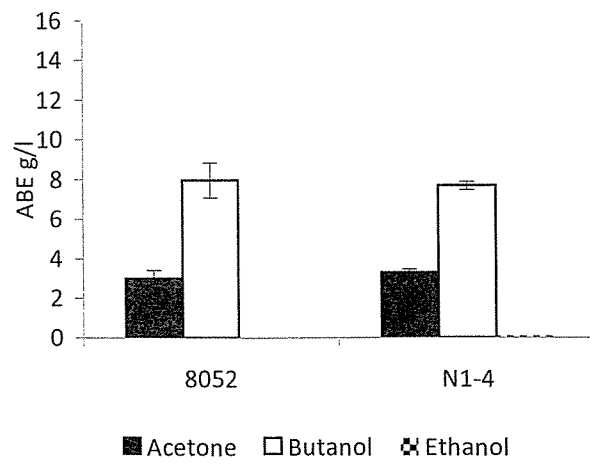


Figure 8

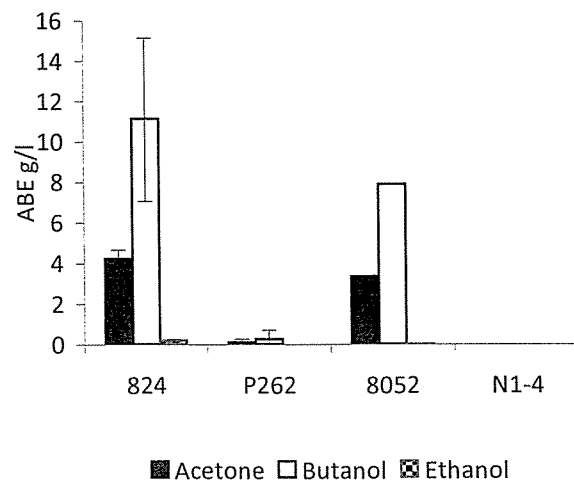


Figure 9

