



(51) International Patent Classification:

C07C 29/16 (2006.01) C07C 31/125 (2006.01)
C07C 29/80 (2006.01)

(21) International Application Number:

PCT/IB2018/058925

(22) International Filing Date:

13 November 2018 (13.11.2018)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/588,571 20 November 2017 (20.11.2017) US

(71) Applicant: **SABIC GLOBAL TECHNOLOGIES B.V.**
[NL/NL]; Plasticslaan 1, 4612 PX Bergen Op Zoom (NL).

(72) Inventors: **RALLAPALLI, Jagan Mohan**; SABIC T&I,
PO Box 42503, Riyadh, 11551 (SA). **GHAMDI-AL,
Ameen**; SABIC T&I, PO Box 42503, Riyadh, 11551 (SA).
PAUL, Somak; SABIC T&I, PO Box 42503, Riyadh,
11551 (SA).

(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM,
AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ,
CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO,
DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN,
HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP,

KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME,
MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ,
OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA,
SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN,
TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ,
UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ,
TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK,
EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV,
MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM,
TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,
KM, ML, MR, NE, SN, TD, TG).

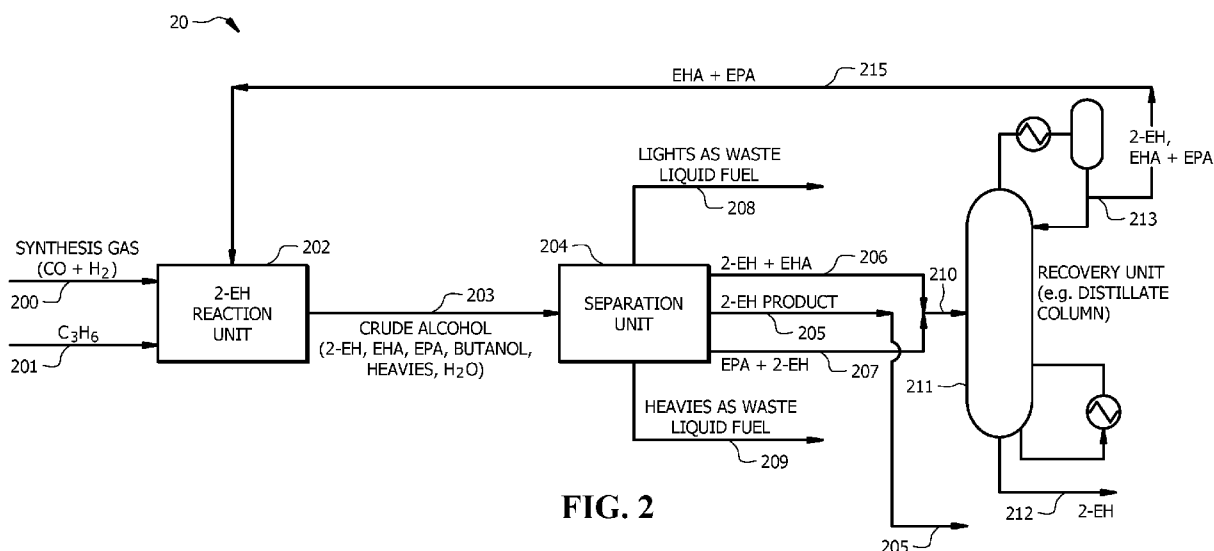
Declarations under Rule 4.17:

- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))

Published:

- with international search report (Art. 21(3))

(54) Title: RECOVERY OF ETHYL HEXANOL FROM RECYCLE STREAMS IN 2-ETHYL HEXANOL PROCESS



(57) Abstract: A method of recovering 2-ethyl hexanol (2-EH) from crude alcohol that includes the 2-ethyl hexanol, 2-ethyl, 3-propyl acrolein (EPA), and 2-ethylhexanal (EHA) is disclosed. The crude alcohol stream is first separated into a first stream of primarily 2-EH, a second stream of primarily EPA and 2-EH collectively, and a third stream of primarily 2-EH and EHA collectively. The second stream and the third stream are combined to form a combined stream. The combined stream is processed further to form a first product stream comprising primarily 2-EH and a second product stream comprising primarily, 2-EH, EHA, and EPA collectively.



RECOVERY OF ETHYL HEXANOL FROM RECYCLE STREAMS IN 2-ETHYL HEXANOL PROCESS

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims the benefit of priority of U.S. Provisional Patent
5 Application No. 62/588,571, filed November 20, 2017, which is hereby incorporated by
reference in its entirety.

FIELD OF INVENTION

[0002] The present invention generally relates to the production of 2-ethyl hexanol.
Specifically, the present invention relates to the production of crude alcohol and recovery of
10 2-ethyl hexanol from the crude alcohol.

BACKGROUND OF THE INVENTION

[0003] 2-ethyl hexanol (2-EH) is a high boiling point, low volatility liquid used as a
solvent, a fragrance, and precursor material for the production of plasticizers, lubricants,
paints, and coatings. 2-EH can be produced by a series of reactions. First, synthesis gas and
15 propylene are reacted with each other to produce butyraldehyde. Second, the butyraldehyde
is then converted to 2-ethyl, 3-propyl acrolein (EPA) by simultaneous aldolisation and
dehydration. Third, 2-EH is produced by hydrogenation of EPA in two steps. In the first
hydrogenation step, EPA is partially hydrogenated to produce 2-ethylhexanal (EHA). In the
second hydrogenation step, the EHA is further hydrogenated to produce 2-EH in a crude
20 alcohol product. It should be noted that, typically, not all of the EPA is converted to EHA
and not all the EHA is converted to 2-EH.

[0004] Thus, the crude alcohol product contains mainly 2-EH, but also contains EHA,
EPA, butanol, heavies, and water. Thus, in order to prepare the 2-EH for market, a product
purification unit purifies the crude alcohol. The purification unit can contain one or more
25 distillation columns for separating the 2-EH from the various other components in the crude
alcohol. The purification unit produces a first stream comprising primarily 2-EH product, a
second stream comprising primarily 2-EH and EHA, a third stream comprising primarily
EPA and 2-EH, a fourth stream comprising primarily lights discharged as waste liquid fuel,
and a fifth stream comprising heavies discharged as waste liquid fuel.

[0005] FIG. 1 shows prior art system 10 for producing 2-EH. System 10 includes 2-EH reaction unit 102 and separation unit 104. 2-EH reaction unit 102 includes one or more aldolisation reactor(s), dehydration reactor(s), and dehydrogenation reactor(s). Separation unit 104 includes distillation units.

5 [0006] To produce 2-EH involves reacting, in 2-EH reaction unit 102, synthesis gas 100 with propylene 101 to produce intermediary products such as EHA and EPA, at least some of which is ultimately converted to 2-EH, as described above. The 2-EH is flowed from 2-EH reaction unit 102 as a part of crude alcohol 103. Based on the process carried out in 2-EH reaction unit 102, crude alcohol 103 may comprise 2-EH, EHA, EPA, butanol, 10 heavies, and water. In system 10, separation unit 104 refines crude alcohol 103 by removing EHA, EPA, butanol, heavies, and water to produce 2-EH product 105, which typically has 99.7 mol. % 2-EH. Byproducts of the separation process in separation unit 104 include 2-EH & EHA stream 106, EPA & 2-EH stream 107, lights stream 108, and heavies stream 109. 2-EH & EHA stream 106 and EPA & 2-EH stream 107 are recycled to 2-EH reaction unit 102.

15 [0007] In system 10, a considerable quantity of 2-EH product that is produced is recycled back to 2-EH reaction unit 102. Because of this, there is high-energy consumption in 2-EH reaction unit 102 to control the reaction conditions. Improvements in this and other aspects of the production of 2-EH are desired.

BRIEF SUMMARY OF THE INVENTION

20 [0008] A method has been discovered for recovering 2-EH from crude alcohol. The discovered method involves refining a crude alcohol stream to form a high concentration 2-EH stream and byproduct streams. In the discovered method, the byproduct streams of the refining process are combined. Then the combined stream is refined to form additional 2-EH product.

25 [0009] Embodiments of the invention include a method of recovering 2-EH from a crude alcohol stream comprising (1) primarily 2-EH, (2) EPA, and (3) EHA. The method includes separating the crude alcohol stream into a first stream comprising primarily 2-EH, a second stream comprising primarily EPA and 2-EH collectively, and a third stream comprising primarily 2-EH and EHA collectively. The method further includes combining 30 the second stream and the third stream to form a combined stream. Further yet, the method

includes separating the combined stream to form a first product stream comprising primarily 2-EH and a second product stream comprising primarily 2-EH, EHA, and EPA collectively.

[0010] Embodiments of the invention include a method of recovering 2-EH from a crude alcohol stream comprising (1) primarily 2-EH, (2) EPA, and (3) EHA. The method
5 includes reacting, in a reaction unit, synthesis gas and propylene to form the crude alcohol stream. The method also includes separating the crude alcohol stream into a first stream comprising primarily 2-EH, a second stream comprising primarily EPA and EH collectively, and a third stream comprising primarily EH and EHA collectively. The method further includes combining the second stream and the third stream to form a combined stream.
10 Further yet, the method includes flowing the combined stream to a packed recovery distillation column and separating the combined stream, by the packed recovery distillation column, to form a bottom product stream comprising primarily 2-EH and a distillate product stream comprising primarily, 2-EH, EHA, and EPA collectively. The method also includes recycling the distillate product stream to the reaction unit and processing, in the reaction unit,
15 EHA and EPA of the distillate product stream to produce 2-EH.

[0011] The following includes definitions of various terms and phrases used throughout this specification.

[0012] The terms “about” or “approximately” are defined as being close to as understood by one of ordinary skill in the art. In one non-limiting embodiment the terms are
20 defined to be within 10%, preferably, within 5%, more preferably, within 1%, and most preferably, within 0.5%.

[0013] The terms “wt.%”, “vol.%” or “mol.%” refer to a weight, volume, or molar percentage of a component, respectively, based on the total weight, the total volume, or the total moles of material that includes the component. In a non-limiting example, 10 moles of
25 component in 100 moles of the material is 10 mol.% of component.

[0014] The term “substantially” and its variations are defined to include ranges within 10%, within 5%, within 1%, or within 0.5%.

[0015] The terms “inhibiting” or “reducing” or “preventing” or “avoiding” or any variation of these terms, when used in the claims and/or the specification, includes any
30 measurable decrease or complete inhibition to achieve a desired result.

[0016] The term “effective,” as that term is used in the specification and/or claims, means adequate to accomplish a desired, expected, or intended result.

[0017] The term “primarily,” as that term is used in the specification and/or claims, means greater than any of 50 wt. %, 50 mol. %, or 50 vol. %. For example, “primarily” may include 50.1 wt. % to 100 wt. % and all values and ranges there between, 50.1 mol. % to 100 mol. % and all values and ranges there between, 50.1 vol. % to 100 vol. % and all values and ranges there between.

[0018] The use of the words “a” or “an” when used in conjunction with the term “comprising,” “including,” “containing,” or “having” in the claims or the specification may mean “one,” but it is also consistent with the meaning of “one or more,” “at least one,” and “one or more than one.”

[0019] The words “comprising” (and any form of comprising, such as “comprise” and “comprises”), “having” (and any form of having, such as “have” and “has”), “including” (and any form of including, such as “includes” and “include”) or “containing” (and any form of containing, such as “contains” and “contain”) are inclusive or open-ended and do not exclude additional, unrecited elements or method steps.

[0020] The process of the present invention can “comprise,” “consist essentially of,” or “consist of” particular ingredients, components, compositions, etc., disclosed throughout the specification.

[0021] In the context of the present invention, at least fifteen embodiments are now described. Embodiment 1 is a method of recovering 2-ethyl hexanol (2-EH) from a crude alcohol stream containing (1) primarily 2-EH, (2) 2-ethyl, 3-propyl acrolein (EPA), and (3) 2-ethylhexanal (EHA). The method includes the steps of separating the crude alcohol stream into a first stream containing primarily 2-EH, a second stream containing primarily EPA and 2-EH collectively, and a third stream containing primarily 2-EH and EHA collectively; combining the second stream and the third stream to form a combined stream; and separating the combined stream to form a first product stream containing primarily 2-EH and a second product stream containing primarily, 2-EH, EHA, and EPA collectively. Embodiment 2 is the method of embodiment 1, further including the step of recycling the second product stream to a process that uses the EHA and the EPA of the second product stream to form additional 2-EH. Embodiment 3 is the method of any of embodiments 1 and 2, further including the step

of flowing the combined stream to a distillation column, wherein the separating of the combined stream is carried out by the distillation column, wherein the first product stream is a bottom product stream and the second product stream is a distillate product stream. Embodiment 4 is the method of any of embodiments 1 to 3, wherein 70 to 95 mol. % of 2-EH in the combined stream is contained in the first product stream. Embodiment 5 is the method of any of embodiments 1 to 4, wherein the first product stream is greater than 95 mol. % 2-EH. Embodiment 6 is the method of any of embodiments 1 to 5, wherein the first product stream is greater than 99 mol. % 2-EH. Embodiment 7 is the method of any of embodiments 1 to 6, wherein the first product stream is 99.75 mol. % or greater of 2-EH. Embodiment 8 is the method of any of embodiments 3 to 7, wherein the distillation column is operated such that the first product stream is at a temperature in a range of 150 to 160 °C and a pressure in a range of 0.2 to 0.6 bars. Embodiment 9 is the method of any of embodiments 3 to 8, wherein the distillation column is operated such that the second product stream is at a temperature in a range of 205 to 215 °C and a pressure of 1.8 to 2.2 bars. Embodiment 10 is the method of any of embodiments 1 to 9, wherein the first product stream contains more 2-EH than the second product stream. Embodiment 11 is the method of any of embodiments 1 to 9, wherein a mass ratio of 2-EH in the first product stream to 2-EH in the second product stream is in a range of 1.2/1 to 1.6/1. Embodiment 12 is the method of any of embodiments 1 to 11, wherein the mass of 2-EH in the first product stream is 55% to 65% of the mass of 2-EH in the combined stream.

[0022] Embodiment 13 is a method of recovering 2-ethyl hexanol (2-EH) from a crude alcohol stream containing (1) primarily 2-EH, (2) 2-ethyl, 3-propyl acrolein (EPA), and (3) 2-ethylhexanal (EHA). The method includes the steps of reacting, in a reaction unit, synthesis gas and propylene to form the crude alcohol stream; separating the crude alcohol stream into a first stream containing primarily 2-EH, a second stream containing primarily EPA and EH collectively, and a third stream containing primarily EH and EHA collectively; combining the second stream and the third stream to form a combined stream; flowing the combined stream to a packed recovery distillation column; separating the combined stream, by the packed recovery distillation column, to form a bottom product stream containing primarily 2-EH and a distillate product stream containing primarily, 2-EH, EHA, and EPA collectively; and recycling the distillate product stream to the reaction unit; and processing, in the reaction unit, EHA and EPA of the distillate product stream to produce 2-EH. Embodiment 14 is the method of embodiment 13, wherein the packed recovery distillation

column is operated such that the bottom product stream is at a temperature in a range of 150 to 160 °C and a pressure in a range of 0.2 to 0.6 bars. Embodiment 15 is the method of any of embodiments 13 and 14, wherein the packed recovery distillation column is operated such that the distillate product stream is at a temperature in a range of 205 to 215 °C and a pressure of 1.8 to 2.2 bars.

[0023] Other objects, features and advantages of the present invention will become apparent from the following figures, detailed description, and examples. It should be understood, however, that the figures, detailed description, and examples, while indicating specific embodiments of the invention, are given by way of illustration only and are not meant to be limiting. Additionally, it is contemplated that changes and modifications within the spirit and scope of the invention will become apparent to those skilled in the art from this detailed description. In further embodiments, features from specific embodiments may be combined with features from other embodiments. For example, features from one embodiment may be combined with features from any of the other embodiments. In further embodiments, additional features may be added to the specific embodiments described herein.

BRIEF DESCRIPTION OF THE DRAWINGS

[0024] For a more complete understanding, reference is now made to the following descriptions taken in conjunction with the accompanying drawings, in which:

[0025] FIG. 1 shows a prior art system for producing 2-EH;

[0026] FIG. 2 shows a system of recovering 2-EH from a crude alcohol stream, according to embodiments of the invention; and

[0027] FIG. 3 shows a method of recovering 2-EH from a crude alcohol stream, according to embodiments of the invention.

DETAILED DESCRIPTION OF THE INVENTION

[0028] A method has been discovered for recovering 2-EH from a crude alcohol stream. The discovered method involves refining the crude alcohol stream to form concentrated 2-EH and byproduct streams (which also contain 2-EH, EPA and EHA), combining the byproduct streams of the refining process, and refining the combined stream to

form additional 2-EH product. In embodiments of the invention, the byproduct streams do not have to be recycled as is but can be subjected to a recovery processes to recover additional 2-EH. For example, in embodiments of the invention, 70% of the existing 2-EH in what would be recycle streams in conventional processes is recovered. The recovery can be implemented by the use of a 2-EH recovery distillation column. Recovering the 2-EH product by the recovery distillation column, according to embodiments of the invention, helps improve the productivity and reduce the energy requirement in the reaction system.

[0029] FIG. 2 shows system 20 for recovering 2-EH from a crude alcohol stream, according to embodiments of the invention. FIG. 3 shows method 30 for recovering 2-EH from a crude alcohol stream, according to embodiments of the invention. Method 30 may be implemented by system 20.

[0030] System 20 includes 2-EH reaction unit 202, separation unit 204, and recovery unit 211. 2-EH reaction unit 202 may include one or more aldolisation reactor(s), dehydration reactor(s), and dehydrogenation reactor(s). Separation unit 204 may include one or more distillation unit(s). Recovery unit 211 may be a distillation column adapted to separate a stream that comprises 2-EH, EHA, and EPA to form a first stream comprising primarily 2-EH product and a second stream comprising primarily 2-EH, EHA, and EPA. In embodiments of the invention, recovery unit 211 is a packed recovery distillation column.

[0031] According to embodiments of the invention, method 30 begins at block 300, which may include producing 2-EH by reacting, in 2-EH reaction unit 202, synthesis gas 200 with propylene 201 to produce intermediary products and eventually 2-EH. Block 300 may first involve reacting synthesis gas and propylene to produce butyraldehyde. Second, the butyraldehyde may then be converted to 2-ethyl, 3-propyl acrolein (EPA) by simultaneous aldolisation and dehydration. Third, 2-EH can be produced by hydrogenation of EPA in two steps. In the first hydrogenation step, EPA can be partially hydrogenated to produce 2-ethylhexanal (EHA). In the second hydrogenation step, the EHA can be further hydrogenated to produce 2-EH in a crude alcohol product. The 2-EH may be flowed from 2-EH reaction unit 202 as a component of crude alcohol 203. Based on the process carried out in 2-EH reaction unit 202, crude alcohol 203 may comprise 2-EH, EHA, EPA, butanol, heavies, and water. Typically, crude alcohol 203 comprises 86 to 92 mol. % 2-EH, 0.2 to 1.2 mol. % EHA, 0.05 to 0.2 mol. % EPA, 1.5 to 4 mol. % butanol, 0.1 to 0.8 mol. % heavies, and 5 to 8 mol. % water. Method 30 may involve, at block 301, separation unit 204 refining crude

alcohol 203 by removing EHA, EPA, butanol, heavies, and water to produce 2-EH product 205. Typically, 2-EH product 205 comprises 99.6 to 99.8 mol. % 2-EH, 0.002 to 0.006 mol. % EHA, and 0.0001 to 0.0002 mol. % EPA.

[0032] Byproduct streams of the separation of block 301 may include 2-EH & EHA stream 206, EPA & 2-EH stream 207, lights stream 208, and heavies stream 209. 2-EH & EHA stream 206 may comprise primarily 2-EH and EHA. Typically, 2-EH & EHA stream 206 comprises 90 to 94 mol. % 2-EH, and 5 to 9 mol. % EHA. EPA & 2-EH stream 207 may comprise primarily EPA and 2-EH. Typically, EPA & 2-EH stream 207 comprises 88 to 92 mol. % 2-EH and 8 to 12 mol. % EPA. According to embodiments of the invention, EH & EHA stream 206 and EPA & 2-EH stream 207 are combined to form combined stream 210, at block 302. Combined stream 210, according to embodiments of the invention, comprises 2-EH, EHA, and EPA.

[0033] Method 30 may involve, at block 303, separating combined stream 210 to form a bottom stream (first product stream 212 comprising primarily 2-EH) and a distillate stream (second product stream 213) comprising primarily 2-EH, EHA, and EPA collectively. In embodiments of the invention, a distillation column of recovery unit 211 is operated such that first product stream 212 is at a temperature in a range of 150 to 160 °C and a pressure in a range of 0.2 to 0.6 bars. In embodiments of the invention, a distillation column of recovery unit 211 is operated such that second product stream 213 is at a temperature in a range of 205 to 215 °C and a pressure of 1.8 to 2.2 bars.

[0034] According to embodiments of the invention, 70 to 95 mol. % of 2-EH in combined stream 210 is recovered and is comprised in first product stream 212. According to embodiments of the invention, the bottom stream (first product stream 212) is greater than 95 mol. % 2-EH. In embodiments of the invention, first product stream 212 comprises 95.0 to 99.9 mol. % 2-EH, including all values therein; *e.g.*, 95, 96, 97, 98, 99, and ranges therein; *e.g.*, 95.0 to 95.9 mol. %, 96.0 to 96.9 mol. %, 97.0 to 97.9 mol. %, 98.0 to 98.9 mol. %, 99.0 to 99.9 mol. %. Further, in particular embodiments of the invention, the bottom stream (first product stream 212) is greater than 99 mol. % 2-EH. In embodiments of the invention, first product stream 212 comprises 99.0 to 99.9 mol. % 2-EH and 1.0 to 0.1 mol. % EPA. In embodiments of the invention, second product stream 213 comprises 78 to 82 mol. % 2-EH and 4 to 8 mol. % EHA. Second product stream 213 may be recycled to 2-EH reaction unit 202 to form additional 2-EH.

[0035] In embodiments of the invention, first product stream 212 comprises more 2-EH than second product stream 213. In embodiments of the invention, a mass ratio of 2-EH in first product stream 212 to 2-EH in second product stream 213 is in a range of 1.2/1 to 1.6/1. Further, in embodiments of the invention, the mass of 2-EH in the first product stream is 55% to 65% of the mass of 2-EH in the combined stream.

[0036] Although embodiments of the present invention have been described with reference to blocks of FIG. 3, it should be appreciated that operation of the present invention is not limited to the particular blocks and/or the particular order of the blocks illustrated in FIG. 3. Accordingly, embodiments of the invention may provide functionality as described herein using various blocks in a sequence different than that of FIG. 3.

[0037] In sum, embodiments of the invention can do one or more of the following: improve 2-EH product quality, reduce propylene specific consumption, and reduce the energy consumed in the reaction system and for product separation.

[0038] As part of the disclosure of the present invention, a specific example is included below. The example is for illustrative purposes only and is not intended to limit the invention. Those of ordinary skill in the art will readily recognize parameters that can be changed or modified to yield essentially the same results.

EXAMPLE**Simulation of 2-EH Recovery System of FIG. 2**

Table 1 below shows the results of a simulation of 2-EH recovery using system 20 of FIG. 2.

The simulation was carried out using the simulation package ASPEN PLUS®.

5

Table 1

Equipment	Units	Separation Unit 204	Separation Unit 204	Recovery Column	
		2-EH & EHA stream 206	EPA & 2-EH stream 207	second product stream 213 (distillate)	first product stream 212 (2-EH recovery)
Description					
Temperature	C	196.448655	170	209.4110662	154.1154088
Pressure	bar	1.44325	2	2.01325	0.41325
Mole Flows	kmol/hr	6.476274779	9.441228072	7.470583675	8.446919175
Mole Fractions					
H ₂		0.00	0	0	0
N ₂		0.00	0	0	0
WATER		0.00	0	0	0
N-BAL		0.00	0	0	0
BUTANOL		0.00	0	0	0.00
EPA		0.00	0.103354189	0.129126293	0.001319025
EHA		0.071514111	0	0.061991413	3.91E-06
EMPOH		0.006340111	0.000957201	0.005273409	0.001266972
2-EH		0.922145687	0.89568861	0.803608808	0.997410096
C ₁₆ RESID		0	0	0	0
I-BAL		0.00	0	0	0
Mass Flows	kg/hr	0.00	1225.6	968.07366	1100
H ₂	kg/hr	0.00	0	0	0
N ₂	kg/hr	0.00	0	0	0
WATER	kg/hr	0.00	0	0	0
N-BAL	kg/hr	0.00	0	0	0
BUTANOL	kg/hr	4.22E-05	0	4.22E-05	0.00
EPA	kg/hr	1.90E-06	123.1433519	121.737287	1.406066549
EHA	kg/hr	59.38188106	0	59.3776508	0.004230135
EMPOH	kg/hr	5.347296356	1.176911492	5.130481188	1.393726652
2-EH	kg/hr	777.7444385	1101.279737	781.8281988	1097.195977
C ₁₆ RESID	kg/hr	0	0	0	0
I-BAL	kg/hr	1.78E-15	0	0	0

[0039] As evident in Table 1, the results show that recovery of a 2-EH stream of 99.7 mol. % is achievable in embodiments of the invention. Further, Table 1 shows that in embodiments of the invention, first product stream 212 comprises more 2-EH than second product stream 213. Further yet, Table 1 shows that, in embodiments of the invention, a mass ratio of 2-EH in first product stream 212 to 2-EH in second product stream 213 is approximately 1.4/1. Also, Table 1 shows that, in embodiments of the invention, the mass of 2-EH in first product stream 212 is 55% to 65% of the mass of 2-EH in combined stream 210 (2-EH & EHA stream 206 and EPA & 2-EH stream 207).

[0040] Although embodiments of the present application and their advantages have been described in detail, it should be understood that various changes, substitutions and alterations can be made herein without departing from the spirit and scope of the embodiments as defined by the appended claims. Moreover, the scope of the present application is not intended to be limited to the particular embodiments of the process, machine, manufacture, composition of matter, means, methods and steps described in the specification. As one of ordinary skill in the art will readily appreciate from the above disclosure, processes, machines, manufacture, compositions of matter, means, methods, or steps, presently existing or later to be developed that perform substantially the same function or achieve substantially the same result as the corresponding embodiments described herein may be utilized. Accordingly, the appended claims are intended to include within their scope such processes, machines, manufacture, compositions of matter, means, methods, or steps.

CLAIMS

1. A method of recovering 2-ethyl hexanol (2-EH) from a crude alcohol stream comprising (1) primarily 2-EH, (2) 2-ethyl, 3-propyl acrolein (EPA), and (3) 2-ethylhexanal (EHA), the method comprising:
separating the crude alcohol stream into a first stream comprising primarily 2-EH, a second stream comprising primarily EPA and 2-EH collectively, and a third stream comprising primarily 2-EH and EHA collectively;
combining the second stream and the third stream to form a combined stream;
and
separating the combined stream to form a first product stream comprising primarily 2-EH and a second product stream comprising primarily, 2-EH, EHA, and EPA collectively.
2. The method of claim 1, further comprising:
recycling the second product stream to a process that uses the EHA and the EPA of the second product stream to form additional 2-EH.
3. The method of any of claims 1 or 2, further comprising:
flowing the combined stream to a distillation column, wherein the separating of the combined stream is carried out by the distillation column, wherein the first product stream is a bottom product stream and the second product stream is a distillate product stream.
4. The method of any of claims 1 or 2, wherein 70 to 95 mol. % of 2-EH in the combined stream is comprised in the first product stream.
5. The method of any of claims 1 or 2, wherein the first product stream is greater than 95 mol. % 2-EH.
6. The method of any of claims 1 or 2, wherein the first product stream is greater than 99 mol. % 2-EH.
7. The method of any of claims 1 or 2, wherein the first product stream is 99.75 mol. % or greater of 2-EH.

8. The method of claim 3, wherein the distillation column is operated such that the first product stream is at a temperature in a range of 150 to 160 °C and a pressure in a range of 0.2 to 0.6 bars.
9. The method of claim 3, wherein the distillation column is operated such that the second product stream is at a temperature in a range of 205 to 215 °C and a pressure of 1.8 to 2.2 bars.
10. The method of any of claims 1 or 2, wherein the first product stream comprises more 2-EH than the second product stream.
11. The method of any of claims 1 or 2, wherein a mass ratio of 2-EH in the first product stream to 2-EH in the second product stream is in a range of 1.2/1 to 1.6/1.
12. The method of any of claims 1 or 2, wherein the mass of 2-EH in the first product stream is 55% to 65% of the mass of 2-EH in the combined stream.
13. A method of recovering 2-ethyl hexanol (2-EH) from a crude alcohol stream comprising (1) primarily 2-EH, (2) 2-ethyl, 3-propyl acrolein (EPA), and (3) 2-ethylhexanal (EHA), the method comprising:
 - reacting, in a reaction unit, synthesis gas and propylene to form the crude alcohol stream;
 - separating the crude alcohol stream into a first stream comprising primarily 2-EH, a second stream comprising primarily EPA and EH collectively, and a third stream comprising primarily EH and EHA collectively;
 - combining the second stream and the third stream to form a combined stream;
 - flowing the combined stream to a packed recovery distillation column;
 - separating the combined stream, by the packed recovery distillation column, to form a bottom product stream comprising primarily 2-EH and a distillate product stream comprising primarily, 2-EH, EHA, and EPA collectively; and
 - recycling the distillate product stream to the reaction unit; and
 - processing, in the reaction unit, EHA and EPA of the distillate product stream to produce 2-EH.

14. The method of claim 13, wherein the packed recovery distillation column is operated such that the bottom product stream is at a temperature in a range of 150 to 160 °C and a pressure in a range of 0.2 to 0.6 bars.
15. The method of any of claims 13 and 14, wherein the packed recovery distillation column is operated such that the distillate product stream is at a temperature in a range of 205 to 215 °C and a pressure of 1.8 to 2.2 bars.

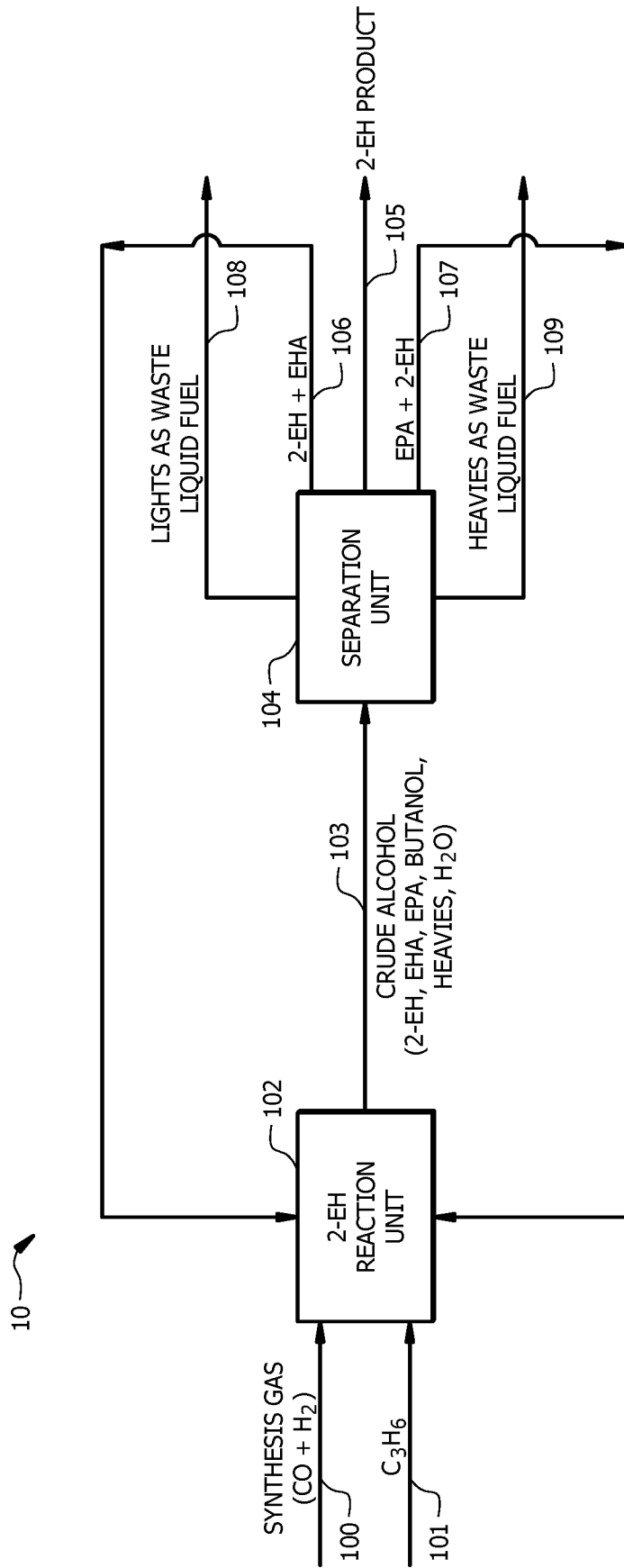


FIG. 1
(Prior Art)

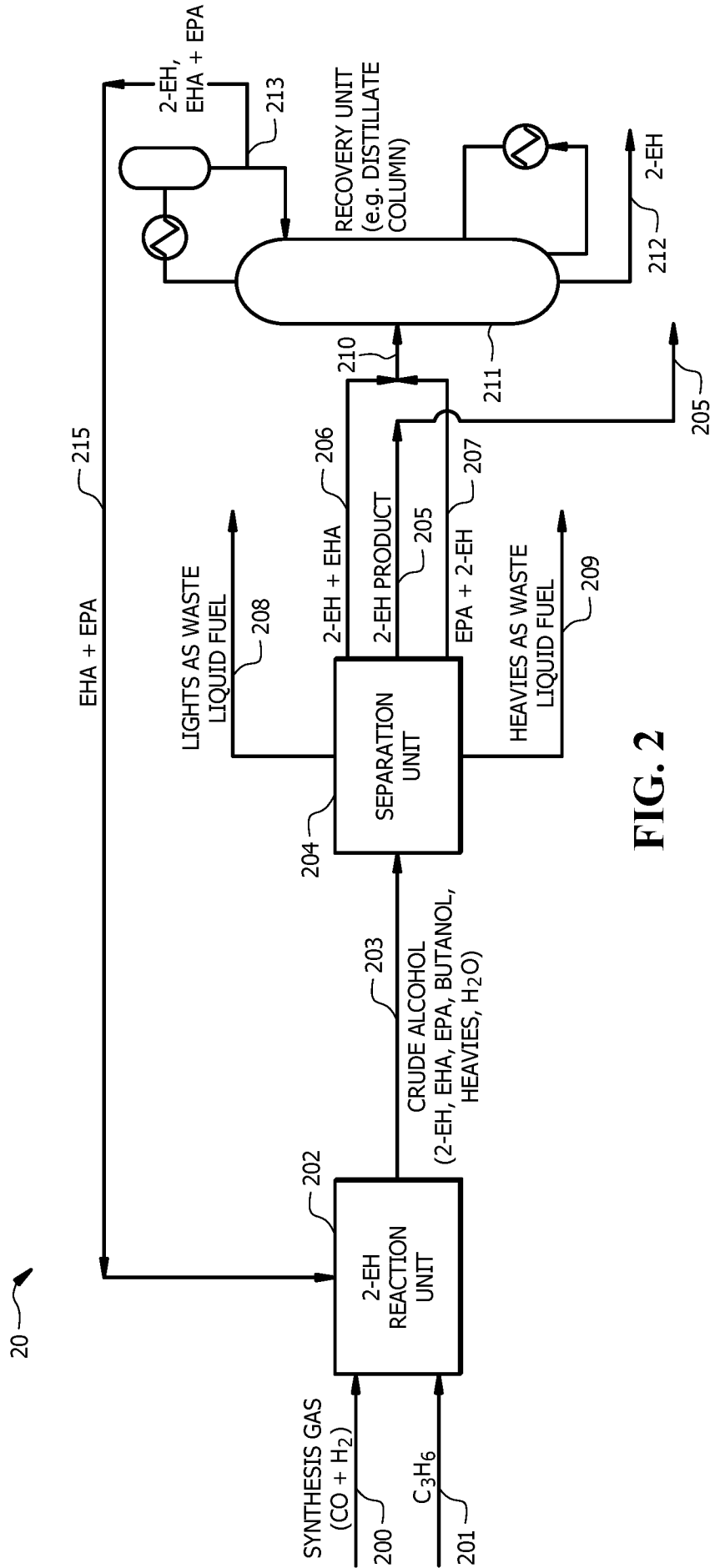
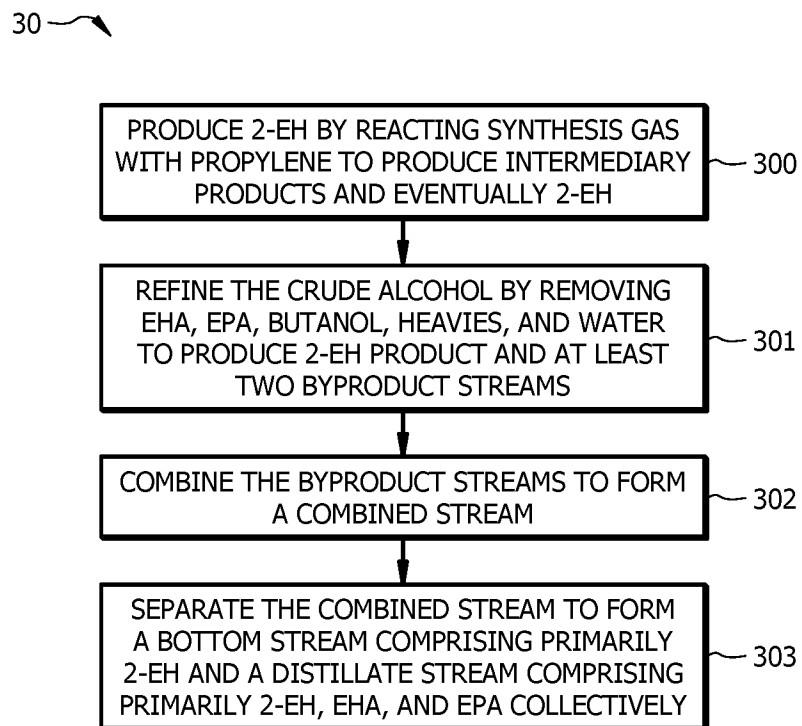


FIG. 2

**FIG. 3**

INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2018/058925

A. CLASSIFICATION OF SUBJECT MATTER
 INV. C07C29/16 C07C29/80 C07C31/125
 ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED
 Minimum documentation searched (classification system followed by classification symbols)
 C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
 EPO-Internal, CHEM ABS Data, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2016/075621 A1 (HASHMI SYED AZHAR [SA] ET AL) 17 March 2016 (2016-03-17) paragraph [0111]; claim 1 -----	1-15
A	US 4 138 588 A (TUMMES HANS ET AL) 6 February 1979 (1979-02-06) claim 1 -----	1-15

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search 13 February 2019	Date of mailing of the international search report 21/02/2019
--	---

Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Lacombe, Céline
--	--

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/IB2018/058925

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2016075621	A1	17-03-2016	CN 105189426 A 23-12-2015
			EP 2991956 A1 09-03-2016
			US 2016075621 A1 17-03-2016
			WO 2014178031 A1 06-11-2014

US 4138588	A	06-02-1979	AT 358548 B 10-09-1980
			AU 519741 B2 17-12-1981
			BE 865146 A 21-09-1978
			BR 7801564 A 31-10-1978
			CA 1088103 A 21-10-1980
			DE 2713434 A1 28-09-1978
			ES 468005 A1 16-11-1978
			FR 2384735 A1 20-10-1978
			GB 1581898 A 31-12-1980
			HU 179297 B 28-09-1982
			IT 1104185 B 21-10-1985
			JP S566405 B2 10-02-1981
			JP S53119804 A 19-10-1978
			NL 7704631 A 28-09-1978
			PL 205451 A1 26-03-1979
			RO 74430 A 01-02-1982
SE 436351 B 03-12-1984			
US 4138588 A 06-02-1979			
