



Article Preliminary Studies on the Use of Reactive Distillation in the Production of Beverage Spirits

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Abstract: Distilled alcoholic beverages have been produced through fermentation and distillation for centuries but have not purposefully involved a chemical reaction to produce a flavoring. Introducing a microorganism to produce butyric acid along with the typical yeast ethanol fermentation sets up a reactive distillation system to flavor a spirit with ethyl butyrate and butyric acid. The ternary interactions of water, ethanol, and butyric acid allow all three to vaporize in the stripping distillation, thus they are concentrated in the low wines and give a large excess of ethanol compared to butyric acid for better reaction completion. The stripping distillation has also been modeled on Aspen Plus[®] V9 software (by Aspen Technology, Inc. Bedford, MA, USA) and coincides well with a test stripping distillation at the bench scale. Amberlyst[®] 15 wet catalyst was added to a subsequent distillation, resulting in the production of the desired ethyl butyrate in the distillate, measured by gas chromatography. Primary sensory evaluation has determined that this process has a profound effect on the smell of the spirit with the main flavor being similar to fruity bubble gum. The current results will provide a pathway for creating spirits with a desired flavor on demand without acquiring a heavy capital cost if a beverage distillation column is already purchased.

Keywords: reactive distillation; esterification; spirits; ethyl butyrate; butyric acid; ethanol

1. Introduction

The basic concept of fermentation and distillation to create an alcoholic beverage has remained largely unchanged since its inception, mainly due to tradition. There have been advancements made as knowledge of the process grows. For instance, it is now known why the use of copper is important in a still, as it has a catalytic effect of removing sulfur compounds during distillation. Studies have determined that the main compound being removed by the copper is dimethyl trisulfide which has a low odor detection threshold (33 parts per trillion in 20% ethanol) [1].

Reactive distillation combines the two unit operations of a chemical reaction and a distillation into one [2–4]. This technique is useful for reactions that need a special condition that can be achieved through distillation, such as a large excess of one of the reactants, or for removing a product to drive an equilibrium reaction to completion [2,4,5].

Combining two processes into one decreases time and energy to produce a desired result. Initial capital cost and maintenance costs are also lowered because the reactor and column is the same piece of equipment. A common use for reactive distillation is esterification. Other than sulfur adsorption onto copper, the alcoholic beverage industry does not purposefully use this technique in the production of spirits.

Introducing esterification into the production of beverages' spirits will create a food grade, flavored spirit while adhering to the standards of identity for distilled alcoholic

beverages in the United States (27 CFR Section 5.22 https://www.ecfr.gov/cgi-bin/text-idx? SID=39303a4811a9732df1b66270946fee1d&mc=true&node=pt27.1.5&rgn=div5#se27. 1.5_122). This paper will discuss the materials and unit operations that will be combined to develop this process and create the proposed spirit.

Esters

An ester is formed when an acid and alkoxy group join together though a condensation reaction. It can be any acid and any alkoxy group, but is usually an organic acid and an alcohol. As this is a slow reaction without the presence of a catalyst, a classic method is to use the Fischer-Speier esterification with a sulfuric acid catalyst. Figure 1 below shows the condensation of butyric acid and ethanol to create ethyl butyrate; this is the reaction that is desired in this beverage research.



Figure 1. The chemical equation for the dehydration reaction of ethanol and butyric acid to form ethyl butyrate and water.

Esters are commonly used in flavorings and fragrances because of their pleasant odor. Ethyl butyrate is commonly used because it has the smell of pineapples or fresh orange juice [6].

2. Modeling

Modeling techniques were used to determine if this study was possible. A ternary residue curve map (RCM) was made with the three main compounds at the corners: water, ethanol, and butyric acid. Aspen Plus[®] was also used to design a column that shows this system would work for the stripping portion of the beverage distillation process.

2.1. Ternary Residue Curve Map

Figure 2 shows a ternary RCM of water, ethanol, and butyric acid in weight fraction. Along with the ethanol/water azeotrope, a binary azeotrope is present between water and butyric acid which boils at 99.5 °C. A separatrix (or simple batch distillation boundary) is drawn in green which divides the diagram into two distillation regions. Residue curves are drawn in light blue with arrows indicating increasing temperature in the pot (or time during the distillation).

The concentrations of water, ethanol, and butyric acid in the pot at the beginning of a run determine the starting point on Figure 2 [2,7]. The residue curves show the concentration of the components in the pot as the distillation proceeds. As an example, if the concentrations of water, ethanol, and butyric acid are 75% w/w, 20% w/w, and 5% w/w respectively, the pot concentration will be 100 % w/w water when the distillation is finished with all of the ethanol and butyric acid from the system in the distillate. If the starting point was to the left of the distillation boundary, the pot would eventually consist of pure butyric acid.

The presence of these binary azeotropes in this system makes it possible for the high boiling butyric acid to be carried over in the distillate stream by the water during the first distillation of the spirit. This then sets up the system for the second distillation over a catalyst to produce the desired ethyl butyrate in the finished spirit.



Figure 2. A ternary residue curve map of water, ethanol, and butyric acid with separatrix. The separatrix is drawn in green which starts at the ethanol/water azeotrope and ends at the butyric acid/water azeotrope. Residue curves are drawn in light blue. This diagram is measured in weight fraction.

2.2. Aspen Plus[®] Column Design

A column was designed in Aspen Plus[®] to distill a whiskey mash with a requirement to have butyric acid present in the distillate of the column. The column was designed to have 17 bubble cap trays, stage seven as the feed stage, and a reflux ratio of five. The following Figures 3–6 are screenshots from the Aspen Plus[®] software used for the simulation.



Figure 3. Aspen Plus[®] screenshots of the flowsheet and overall stream table. (**a**) Flowsheet. (**b**) Stream Table.

condensely, rop stage performance					
	Name	Value	Units		
	Temperature	82.1776	с		
	Subcooled temperature				
	Heat duty	-143114	cal/sec		
	Subcooled duty				
	Distillate rate	8.71435	kmol/hr		
	Reflux rate	43.5718	kmol/hr		
	Reflux ratio	5			
(a)					

Figure 4. Aspen Plus[®] screenshots of the performance of the top and bottom stages. (**a**) Top Stage Performance. (**b**) Bottom Stage Performance.

		Specifications Design	Results Profiles	
B1 Sizing and Rating Tray Sizing - 1 × B1 (RadFrac) - F	esults ×] +	Section starting stage: Section ending stage:	2	
CTraved section		Stage with maximum diameter:	7	
Starting stage: 2 Endi	ng stage: 16	Column diameter:	0.476058	meter 🔻
Tray type: Bubble Cap Nun	iber of passes: 1	Downcomer area / Column area:	0.1	
Tray geometry		Side downcomer velocity:	0.0403392	m/sec 🔹
Tray spacing: 0.6096	meter •	Flow path length:	0.327075	meter 🔹
Cap slot area to active area ratio: 0.12	meter	Side downcomer width:	0.0744914	meter 🔹
Sieve hole area to active area ratio: 0.12		Side weir length:	0.345909	meter 🔹



(b)

B1 Sizing and Rating Tray Sizing - 1 × B1 (RadFrac) - Results × +

Figure 5. Aspen Plus[®] screenshots for the sizing of the column. (a) Tray Sizing Input. (b) Tray Sizing Results.

Block B1: Composition Profiles 0.95 0.90 Liquid mass fraction WATER 0.85 Liquid mass fraction ETHAN-01 Liquid mass fraction N-BUT-01 0.80 0.75 0.70 0.65 0.60 Mass fraction 0.55 0. 0.45 0.4 0.35 0.30 0.25 0.20 0.15 0.10 0.0 0.00 Stage

Figure 6. A graph of the column tray compositions as produced by Aspen ${\rm Plus}^{\circledast}$.

As seen in Figure 3b, the distillate stream of the column contains 94.7 mol % of the butyric acid in the system. This simulation shows that the main components will be present in the distillate of the stripping run. The low wines produced from this can then be distilled again using a catalyst to convert the butyric acid into the desired ethyl butyrate.

2.3. Lab-scale Fractional Distillation

A lab fractional distillation was performed in conjunction with the Aspen Plus[®] simulation to support the results. The experiment used a 500 mL round-bottom flask, Vigreux column, and a straight condenser setup in a fractional distillation configuration. The flask started with 250 mL of an aqueous solution containing 79.2 g/L ethanol and 4 g/L butyric acid. Figure 7 shows the concentrations of ethanol and butyric acid of the experiment as a function of distillate volume collected. This ternary distillation was performed to determine at what point the butyric acid is present in the distillate.

A final distillate of 70 mL was collected containing 230 g/L ethanol and 6.2 g/L butyric acid. This experiment coincides well with the Aspen Plus[®] simulation in Section 2.2.



Figure 7. Distillate concentrations of ethanol and butyric acid during the lab-scale fractional distillation vs. the volume of distillate collected.

3. Materials and Methods

This research was preformed on the bench scale. The mash was stripped 1 L at a time in a round-bottom flask. The flask was connected to a 36 cm vigreux column, a 75° distillation adapter, and finally a 20 cm straight Liebig condenser. All connections are \$24/40. Six strips of 14 AWG copper wire (1.63 mm diameter) were put into the vigreux column to represent the copper in a beverage distillation column.

For each stripping run, the distillate was collected until the ethanol fell below 10% ABV. These low wines were then distilled a second time over Amberlyst[®] 15 wet catalyst with varying loadings and positions within the column. All samples were analyzed on a Shimadzu GC-17A with FID. The column used was a Zebron ZB-WAXplus 30 m \times 0.25 mm \times 0.25 µm. The GC oven was ramped up from 40 °C to 140 °C to analyze everything including butyric acid. The column needed to be shortly reconditioned by quickly ramping up to 220 °C at the end of every run.

4. Results

4.1. Reactive Distillation

Figure 8 shows component concentrations in a reactive spirit distillation as a function of distillate volume collected. The distillation consisted of 150 mL of low wine diluted to 30% ABV. Butyric acid was added to the pot to give a 4.0 g/L starting concentration and 15 g of Amberlyst[®] 15 wet catalyst was contained in a mesh bag and added to the pot during the distillation; no ethyl butyrate was added prior to the experiment, as seen in Table 1.



Table 1. Compounds of interest in the reactive distillation system. Concentrations in [g/L].

Figure 8. Component concentrations in the distillate of a spirit run with 150 mL of low wines at 30% ABV, 4.0 g of butyric acid, and 15 g of Amberlyst[®] 15 wet catalyst in the pot.

As seen in Figure 8, the ethanol and ethyl butyrate concentration curves have the same shape and are inverse of the butyric acid concentration curve. During the distillation, butyric acid was reacted with ethanol over the catalyst to form the desired ethyl butyrate. This product was then carried over in the distillate stream by the ethanol.

Note: If creating a spirit, the heads would have been cut after collecting 5 mL, and the tails cut after collecting 40 mL. These cuts would have resulted in the final hearts product being 35 mL in volume and having low ethyl acetate and butyric acid concentrations.

4.2. Sensory Evaluation

Four variations of the reactive distillation from Section 4.1 were distilled and 35 mL of hearts were collected from each as discussed above. Table 2 shows the differences in the four samples.

Table 2. Difference in the samples used for the sensory evaluation to determine if this process has an effect on the final spirit.

Sample	Butyric Acid [g/L]	Catalyst [g]		
1	0	0		
2	0	7.5		
3	5	0		
4	5	7.5		

Each sample was proofed to 40% ABV with deionized water before being analyzed. The participant was asked to smell each sample and give a description of what they smelled (Table 3). They were also asked if they could tell a difference between samples (Table 4).

Participant	Sample 1	Sample 2	Sample 3	Sample 4
1	whiskey	softer than 1	N/A	juicy fruit gum
2	corn	corn	cheese	licorice/fruity
3	alcohol	alcohol	same as 2	apples
4	corn	corn	corn	bubble gum
5	corn whiskey	corn whiskey	buttered popcorn	buttered popcorn
6	Popsicle stick	less strong, but same as 1	Olay bar soap	minty
7	whiskey	bubble yum	soap	sour candy, apple
8	plastic, organic, sweet	less sweet	N/Â	popcorn

Table 3. Sample notes from individual sensory evaluations.

Table 4. Could the participant tell the difference between samples of the sensory evaluation

Participant	Sample					
	1v2	1v3	2v3	1v4	2v4	3v4
1	yes	no	yes	yes	yes	yes
2	no	yes	yes	yes	yes	yes
3	yes	yes	no	yes	yes	yes
4	yes	yes	yes	yes	yes	yes
5	no	yes	yes	yes	yes	yes
6	yes	yes	yes	yes	yes	yes
7	yes	yes	yes	yes	yes	yes
8	yes	no	yes	yes	yes	yes

The evaluation showed that samples 1 and 2 were similar with a smell of whiskey and they could be told apart; sample 1 had a stronger whiskey smell than sample 2. Sample 3 had notes of licorice and popcorn, which is expected as there was butyric acid present. Sample 4 was overwhelmingly different than the others having a fruity smell similar to that of banana candy or bubble gum.

5. Discussion

For research purposes, food grade butyric acid was added to the fermented mash before the first distillation. Future work will develop a co-fermentation between *Saccharomyces cerevisiae* and *Clostridium tyrobutyricum* to produce ethanol and butyric acid, respectively.

Introduction of ester production in beverage spirit distillations will create a new series of products that have unique smells and tastes. These qualities will give the spirit an increased value and marketability, thus mitigating the reduced ethanol yield from the production of the organic acid in the fermentation.

This research will provide a pathway for creating spirits with a desired flavor on demand without acquiring a heavy capital cost if a beverage distillation column is already purchased. Manufacturers would be able to produce a uniquely flavored spirit of their own with only the purchase of the correct catalyst and microorganism.

The code of federal regulations (CFR) has set rules of what can and cannot be done when it comes to distilled spirits, specifically (27 CFR Section 5 https://www.ecfr.gov/cgi-bin/text-idx?SID= 39303a4811a9732df1b66270946fee1d&mc=true&node=pt27.1.5&rgn=div5).

The reaction of ethanol and butyric acid to ethyl butyrate proceeds naturally. The catalyst that is put into the column speeds the reaction rate and drives the reaction more toward the product, i.e., the catalyst is not adding a new reaction, only accelerating an already existing reaction.

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Abbreviations

The following abbreviations are used in this manuscript:

- RCM Residue Curve Map
- g/L grams per liter
- GC Gas Chromatograph
- AWG American Wire Gauge
- CFR Code of Federal Regulations
- ABV Alcohol By Volume

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