Abstract: The global demand for acrylic acid (AA) is increasing due to its wide range of applications. Due to this growing demand, alternative AA production strategies must be explored to avoid the exacerbation of prevailing climate and global warming issues since current AA production strategies involve AA production using fossil resources. Investigations on alternative strategies for AA production therefore constitute an important research interest. The present study therefore assesses waste apple pomace (WAP) as a feedstock for the sustainable AA production. To undertake this assessment, process models, based on two production pathways were designed, modelled and simulated in ASPEN plus® software. The two competing production pathways investigated include a process incorporating WAP conversion to lactic acid (LA), prior to LA dehydration to generate AA (denoted as the FD pathway) and another process involving WAP conversion to propylene, prior to propylene oxidation to generate AA (denoted as the TFO pathway). Economic and environmental performances of the FD and TFO pathways were assessed via the minimum selling price (MSP) and potential environmental impacts per h (PEI/h) metrics. The study was able to show that the FD pathway presented an improved economic performance (MSP of AA: US $1.17 per kg) performance compared to the economic performance (MSP of AA: US $1.56 per kg) of the TFO pathway. Crucially, the TFO process was shown to present an improved environmental performance (2.07 kPEI/h) compared to the environmental performance of the FD process (8.72 kPEI/h). These observations suggest that the selection of the preferred AA production will require a trade-off between the performance measures, and the integration of a multi-criteria decision assessment in future work.

Keywords: acrylic acid; economic assessment; waste valorization; waste apple pomace; bio-refinery
1. Introduction

The increasing threat of catastrophic climate effects [1] due to anthropogenic greenhouse gas (GHG) emissions and natural resource depletion have, in recent times, motivated research into the production of ‘green’ products. Some of these green products, produced via bio-refinery of biomass, are capable of partly (or completely) replacing their fossil fuel alternatives [2,3]. Bio-refinery technologies constitute sustainable progressions that incorporate biomass conversion technologies to facilitate the conversion of biomass to high-value products, and thus, promote the circular bio-economy paradigm [4]. Given that the implementation of such bio-refineries has thus far been limited by low profitability and poor commercialization concerns [3], the use of waste biomass streams was proposed as one strategy that may enhance overall economic performance given that feedstock cost constitutes an important factor that influences bio-refinery economic performance [3][4]. The present study therefore proposes the utilization of the waste biomass resources of waste apple pomace (WAP) as a useful bio-refinery feedstock. WAP constitutes an abundant biomass resource, with ~18 ×10^9 kg of pomace estimated to be generated globally per year [5]. The poor management of this WAP stream may lead to unfavorable environmental outcomes such as uncontrolled GHG emissions from the disposal of pomace in landfills, unwanted acidification of agricultural soils via leachate formation and also the pollution of underground waters [6]. The utilization of WAP as a biomass resource therefore presents an opportunity for value extraction while also circumventing waste management concerns [7]. The current study therefore seeks to investigate the production of AA from WAP. Acrylic acid (AA; CH₂=CHCOOH) is the simplest of the unsaturated acids [8-10]. AA is an important product that is characterized by several significant end uses such as supper absorbent polymers, artificial leather and textiles (Figure 1). Figure 1 also shows that 42% of acrylic acid generated is employed in the manufacturing of dispersions i.e. detergents. In addition, AA may have other applications ‘others’ including in the production of acrylic-based polymers for fabricating corneal prosthesis, contact lenses, and in tissue engineering (these applications are shown as ‘Other’ in Figure 1) [11,12].

![Figure 1: Acrylic Acid by End-Use](image)

The wide range of AA application is, partly due to its ability to achieve rapid polymerization and its efficiency in delivering carboxylate functionality in commercial products [14]. The importance of AA is demonstrated by the annual global market size and market demand, predicted to reach US$ 22.55 billion and 8750 kton respectively by 2022 [15,16]. Currently, AA is commercially obtained via the catalytic oxidation of olefins [8-10] which are derived from the refining of fossil crude [8,9]. Such a commercial use of fossil crude exacerbates GHG emissions and natural resource depletion concerns. AA can also be produced from the non-fossil feedstocks of glycerol or biomass [17-24]. Notably, since
glycerol typically requires additional production operations (i.e. as a by-product of the biodiesel production process), the use of freely available biomass as a feedstock may constitute a more sustainable strategy. The literature reveals that (non-fossil) AA may be produced from biomass, via stepwise processes, involving an initial LA acid production step (via sugar fermentation) followed by LA dehydration [25-27]. However, in the current study ‘syngas pathway’ that incorporates the conversions of biomass to (→) propylene → AA (i.e. TFO pathway) has been suggested. We hypothesize that this pathway may be more favorable because unlike the biomass→ LA → AA pathway that only utilizes polysaccharides present in biomass for AA production (i.e. via polysaccharide fermentation), all fractions of the biomass feedstock including the lignin fraction are employed in AA production in the proposed pathway via a preliminary syngas production step (i.e. thermochemical transformation). Therefore, to evaluate our hypothesis, in the present study we undertake an economic and environmental assessment of the two pathways that can transform WAP to AA (i.e. via the TFO and FD pathways). In spite of the importance of AA to our best of knowledge, no comparative economic assessment of AA production processes from WAP has been reported in the literature. Additionally, recognizing the importance of addressing environmental performance concerns, the current study also seeks to comparatively explore the environmental performances of the alternative AA production pathways. Since the feedstock collection and transportation to the AA production site is assumed to be similar for both pathways, only environmental impacts of the stand-alone AA production facilities will be assessed. Thus the assessment of impacts associated with environmental risk, waste minimization opportunities and raw materials [28-30] acquisition are beyond the scope of the present study and are not presented here. The comprehensive approach to the comparative study in the present work therefore emphasizes its novelty.

2. Materials and Methods

In this study, the commercial software ASPEN Plus® version 11 (Aspen Technology Inc., Cambridge, Massachusetts, USA) was used to design, model and simulate AA production via the TFO and FD pathways. Calculations, involving mass balance, energy balance, chemical and phase equilibrium were undertaken in ASPEN Plus®. The WAP was modelled according to its reported chemical composition [31], proximate and elemental composition [32]. These data are presented in Tables S1 and S2 in the supplementary file A.

Modelling of other chemical inputs was achieved using the data library in ASPEN Plus. The non-random two-liquid (NRTL) thermodynamic property method is sufficient to model the properties (i.e. activities, fugacities, fugacity coefficients etc.) of chemical species in different phases and in complex systems [33-36]. We therefore, employed NRTL in the modeling process. In the present study, the reactors were modelled based on experimental results obtained from literature as discussed below (section 2.1), using the stoichiometric reactor models (RSTOIC). The RSTOIC reactors are based on the knowledge of fractional conversion of reacting species. For complex, reactions such as those involved in gasification processes, it was assumed that thermodynamic equilibrium is attained since the rates of associated chemical and physical-chemical reactions are infinitely high [37]. The gasification reaction was therefore modelled using the two equilibrium reactor models (RGIBBS) reactors in ASPEN Plus®. In the RGIBBS model, ASPEN plus determines the concentration of species by minimizing the Gibbs free energy ($G_{total}$) of the system which is defined as follows [38-40]:

$$G_{total} = \sum_{i=1}^{N} n_i G_{f.i} + \sum_{i=1}^{N} n_i RT \ln \left( \frac{f_i}{f_i^0} \right)$$ (1)

where $f^0$ for gas phase species, is equal to 1 bar such that,

$$f_i = \theta_i y_i P$$ (2)

while for liquid phase species,

$$\frac{f_i}{f_i^0} = a_i$$ (3)
where \( f_i \), \( G_i^{\text{fug}} \), and \( \theta_i \) represent the standard molar fugacity, the Gibbs free energy of formation at 1 bar and the fugacity coefficient of the \( i \)th specie respectively. \( T \), \( R \) and \( y_i \) represent the temperature in K, the universal gas constant (8.314 J/mol·K), the mole fraction vapour of the \( i \)th specie respectively.

Furthermore, all pumps, turbines and compressors were modelled as isentropic units [41]. Finally, distillation processes were modelled using the RADFRAC model Aspen Plus®, which facilitates rigorous energy and mass balance calculations and can be used to facilitate the separation of highly non-ideal liquid mixtures [42]. To determine the column design specifications, the design parameters (i.e. number of stages, reboiler duty, reflux ratio etc.) were estimated using the Fenske-Underwood-Gilliland correlation model in Aspen Plus® [41,43].

### 2.1 Process description

Initially, the FD process that employs LA as a precursor to AA via its dehydration was investigated, according to the simplified process flow diagram presented in Figure 2 and is based on similar techniques which were used for LA production as described in [44]. Figure 2 shows that initially, the water content of the WAP is reduced to 50 wt.% from 67 wt.% (Materne company, Brussels Belgium) via mechanical dewatering operations. The dewatered WAP is then homogenized at 1 atm and at a temperature of 25 °C. The duties required for the dewatering and homogenisation operations are specified as 2 kW per 1 m³ of feed and 1.5 kW per ton of feed, respectively and are incorporated in the simulation using Fortran codes [45,46]. Pre-treatment of the WAP is subsequently undertaken, for lignin separation, via hot water treatment to minimize the generation of inhibitors (i.e furfural) that may limit the efficiency of the downstream fermentation operations [6,47,48]. This hot water treatment approach was also established to limit polysaccharide losses during biomass pre-treatment as discussed in the literature [6,47,48]. According to the well-established experimental approach employed in the literature [49], the hot water pre-treatment is undertaken at the temperature of 142.4 °C and pressure 1 atm with a 65 % sugar (C6 and C5) recovery achieved according to the reaction equations presented in Table S3 in the supplementary file A. To enhance process efficiency and energy self-sufficiency, onsite steam generation is achieved using the lignin fraction as a solid fuel [41].

The lignin separated is employed as a fuel for heat and electricity generation via a combined heat and power (CHP) plant, in accordance with the Rankine cycle as described in the literature [6]. In the CHP plant, pressurization of water to 20 atm is initially achieved using a pump operating at 0.9 efficiency. The pressurised water is then heated using heat generated from the boiler via the combustion of the solid fuel (in this case lignin) to facilitate the production of superheated steam. Electrical energy is then generated via the expansion of the pressurised superheated steam to 1 atm, in an isentropic turbine. The steam is then condensed and is made available for reuse within the system. Figure 2 then shows that the lignin-free stream (after hot water treatment) is subjected to a fermentation operation under the action of Lactobacillus casei sub sp. rhamnous (ATCC 10863) at 45 °C and pressure of 1 atm [50]. This microbe is used in the present study due to its capability to metabolise C5 (~80 % conversion) and C6 (~100 % conversion) sugars to produce LA according to the reactions summarised in Table S3 (supplementary file A). At the conclusion of the fermentation process, the broth is subjected to precipitation treatment followed by acid treatment using Ca(OH)₂ at the temperature of 95 °C to facilitate the production of C₆H₁₀CaO₆ for its separation from the solution [51-53]. The C₆H₁₀CaO₆ is then subjected to acid treatment to enable the recovery of LA, with the waste stream of gypsum generated.

Stage-wise separation of the C₆H₁₀CaO₆ and the gypsum is assumed to be sufficient and is achieved using filtration at 25 °C and 1 atm [51-53]. Since the dehydration of the LA to AA requires that the LA stream has a LA content of 50 wt.% [54], a vaporization operation is imposed at the temperature of 100 °C to increase the concentration of LA and achieve 50 wt.% concentration [51-53]. In accordance with the work of Ghantani et al., [54] LA (50 wt.% moisture) dehydration to produce AA is achieved under the action of Ca/P (1.3 molar ratio), with LA presenting a 100 % conversion when subjected to a temperature condition of 375 °C. The dominant dehydration reaction facilitates the production of AA, water and a side decarboxylation reaction facilitates the production of CO and acetaldehyde (C₃H₄O), with the AA selectivity specified as 60% [54]. The exit stream is then initially degassed for the recovery of CO, which is utilized as a gaseous fuel in the CHP plant. The CO gas has a higher heating value (HHV) of 4.368 MJ/kg [55]. Since it is not possible to separate AA from water via conventional distillation or rectification because of the formation of a minimum boiling azeotrope [56], an extractive distillation approach is employed [57]. Several organic solvents have been reported in the literature for AA...
extraction from an aqueous solution. These solvents include diisopropyl ether, isopropylacetate, methylisobutyl ketone and toluene [57]. Song et al., [57] however recommended toluene as the preferred solvent for the extraction when the combined effects of extraction capacity and solvent cost are considered. In the present simulation, toluene is therefore employed to facilitate AA extraction. In this approach the stream exiting the dehydration reactor is rapidly cooled to 25 °C using a quencher to reduce the risk of polymerization of AA, after which the mixture is fed to a liquid–liquid extractor, with toluene as the solvent to facilitate the separation of aqueous components from AA [57]. The toluene–AA steam is then distilled to facilitate solvent recovery with associated AA generated as the bottom stream [56]. The recovered toluene is cooled and condensed and made available for further extractions. Finally, the aqueous stream is distilled to facilitate acetaldehyde recovery, as a by-product of the process. The alternative AA production process (simplified) is presented in Figure 3 which shows the TFO pathway. The gasification operation described in Figure 3 is based on generic processes previously described in the literature [58-60]. After the initial dewatering step, the WAP stream is then subjected to a drying operation. The dried WAP is then transferred to the reactor where the partial oxidation of the WAP is achieved under the action of steam (Figure 3).
Figure 2: Simplified process flow diagram for acrylic acid production from waste apple pomace using the fermentation-dehydration (FD) pathway involving LA production as a precursor to AA production.
Figure 3: Simplified process flow diagram for acrylic acid production from waste apple pomace using the thermo-chemical-fermentation-oxidation pathway involving the propylene production as a precursor.
The typical gasification reactions are summarised in Table S3. Herein, we used steam gasification as steam facilitates the production of syngas (CO and H₂) characterized by higher H₂ and reduced CO₂ contents [60,61]. To further decrease in the CO₂ generation a steam-biomass ratio of 0.5 is imposed as detailed before [62]. The gasification operation is specified as being undertaken at the temperature of 1000 °C as the applying temperatures higher than 800 °C minimizes tar formation in gasification systems [58]. Notably, at gasification temperatures ranging from 900 °C-1000 °C, tar formation is significantly mitigated due to tar cracking [58,62]. This statement is consistent with other process models developed in the literature where tar formation was not considered [63,64]. Additionally such high temperatures reduce the risk of the production of heavy hydrocarbons (i.e > C4) in the gasification process [65]. At the conclusion of the gasification reaction, the exit stream is cooled to 37 °C and the solids (char) present in the stream, removed using a cyclone. The current study assumes that the char is composed of carbon and ash only [62]. The syngas product is then transferred to a fermenter, operating at the mesophilic temperature of 37 °C and pressure of 1 atm. Fermentation of the syngas is achieved under the action of acetogenic microorganisms such as Clostridium ljungdahlii and Clostridium carboxidivorans [66,67] according to equations presented in Table S3. CO and H₂ conversions to ethanol are specified as 90 % and 70 % respectively, as previously reported by Phillips et al., [68]. According to Klasson et al.’s work, [68], the aforementioned conversions are achieved when fermentation of syngas occurs in the presence of ~0.8 g of Clostridium ljungdahlii per litre of fermentation broth. Having concluded the fermentation operation, the fermentation broth containing the ethanol is purified via a stage-wise distillation process as described in the literature [69]. Briefly, the fermentation broth is transferred to a ‘beer’ column (B) where stripping of ethanol from the stream is achieved with the resulting vapour phlegm composed of ~35 wt.% of ethanol and ~65 wt.% of water. Further concentration of the vapour phlegm is achieved when the stream is transferred to the rectification column (R) such that a distillate containing ~91 wt.% of ethanol is produced. To facilitate further concentration of the ethanol containing stream, the distillate is dehydrated using molecular sieves, to achieve an ethanol product with a purity of ~ 99 wt.%. Having generated the ethanol product, the stream is transferred to a rectification reactor where dehydration is achieved at 350 °C and 1 atm for ethylene production in accordance with the work of Arvidsson and Lundin [70] (reactions in Table S3). In accordance to literature the conversion of ethanol to ethylene is specified as 99.9 % while the side reaction involves the conversion of the residual ethanol to diethyl ether [69,70]. Given that acetic acid is produced as a by-product of the syngas fermentation process and unreacted ethanol is retained in the stream after the dehydration reaction, it is assumed that the acetic acid and ethanol are available to undergo esterification reaction for ethyl acacetate production. Both streams are therefore transferred to an esterification reactor (reaction in Table S3) operating at 80 °C and 1 atm with a reaction conversion of 84% achieved (ethanol as the limiting reactant) [71].

Cleaning of the ethylene product is achieved using activated carbon. To produce propylene from ethylene, the methods highlighted in the novel and recent study by Beucher et al., [72] are employed. To this regard, the ethylene product is subjected to a cascade of reactions involving two reactors for propylene formation. In the first reactor, ethylene is converted (~85 wt. % conversion) to 2-butenes under the action of a Nickel based catalyst at temperature and pressure conditions of 120 °C and 29.6 atm respectively. The exit stream is then transferred to the second reactor, where the 2-butenes in the stream react with the previously unconverted ethylene at temperature and pressure conditions of 450 °C and 1 atm respectively to produce propylene, under the action of Tungsten oxide based catalyst. To produce AA from propylene, a well-established method were used [73]. According to Sanchez et al., [73], the stream containing the propylene is fed to a reactor together with air compressed at the pressure of 5.95 atm and low pressure steam at the pressure and temperature of 5.95 atm and 159 °C respectively. The propylene, air and steam feed fractions are maintained at 1/7.75/3.75 (mass basis) [73]. In the first reactor, the propylene is then oxidized to facilitate its initial conversion to acrolein, AA, acetic acid and CO₂ with conversions of 79%, 6%, 12% and 3% respectively achieved, under the action of Bismuth molybdate catalyst at the temperature of 350 °C [73]. The initial oxidation of propylene reactions in reactor 1 are presented in the Table S3. Figure 3 subsequently shows that the hot gaseous mixture, containing mostly acrolein, is transferred to the next reactor 2, where acrolein conversion to AA is achieved under the action of vanadium-molybdenum oxide, at the temperature of 300 °C. An AA production yield of 85%, is attained in reactor 2 [73]. The gaseous product is then cooled
from 300 °C to 25 °C with the liquid–liquid extraction module discussed in FD pathway above executed to facilitate AA recovery. In the TFO pathway, the heat from all exothermic reactor units in the bio-refinery (i.e. propylene oxidation reactors [74]), is recovered in form of high pressure steam at 20 atm and employed in electricity generation (via the CHP). The specifications of major operations discussed above are presented in Table S3 in the supplementary file A. Having designed and modelled the AA production processes, energy balance calculations were undertaken, using the ASPEN energy analyser® V.11 [69,75].

2.2 Economic assessments

Plant availability was assumed to be 7200 h per year. To assess the economic performances of the AA productions processes, the minimum selling prices (MSPs) of the AA product generated from both processes were determined. The MSP was specified as the price of AA for which the net present value (NPV) of the project was equal to zero, with the NPV determined using a discounted flow table and calculated as follows [60]:

\[
NPV = -TCI + \sum_{n=1}^{30} \frac{N_R}{(1+i)^n} + \frac{S_n}{(1+i)^n}
\]  

(4)

where TCI, i and n denotes the total capital investment cost in US$, discount rate (10%), and project lifespan (30 y) respectively. \( S_n \) denotes the salvage value specified as zero, with straight-line depreciation assumed. \( N_R \) denotes the net cash flow (annual cash flow less the assets) in US$.

The total operating costs (TOC) of the FD and TFO processes were also determined as a summation of the fixed production cost components (i.e costs incurred irrespective of the plant productivity), and the variable production costs (i.e. cost proportional to the plant outputs) [60]. The costing relations employed in estimating these cost components are presented in supplementary file A. The TCI was determined from equipment purchase costs provided by the ASPEN process economic analyzer V11 (APEA). These purchase costs were determined based on mapping algorithms that employ the mass and energy balance simulation results for equipment sizing calculations. Based on the equipment purchase costs, the TCI costing approach is summarized in Figure 4 [76].

![Figure 4: Total capital investment cost components [76]](image-url)

The full description of each of the cost components are beyond the scope of the current study and may be found elsewhere [77]. The associated TCI cost correlations are presented in the supplementary file A. In the present study the purchase cost of general equipment i.e distillation columns, heater, cooler, tanks and pumps were obtained from APEA. In line with previous studies [78-80], the cost of specialized equipment such as the gasification reactor, cyclone, reactors and molecular sieves were obtained from literature and commercial vendors [81,82] and the purchase costs of these specialized equipment
summarized in the supplementary file A. To account for inflationary effects from equipment purchase costs obtained in different years, Chemical Engineering Plant Cost Index (CEPCI) values were employed according to standard methods [60], with the CEPCI for the year of 2020 specified as 607.5 [6]. The purchase cost $p_i$ of these specialized, $i$th equipment, in US$ was subsequently adjusted for the desired characteristic factor as follows [60,83]:

$$p_i = p_{i,ref} \left( \frac{F}{F_{ref}} \right)^{0.65}$$

where $F$ and $F_{ref}$ denote the characteristic factors such as the volume of the equipment in the present study and reference volume of the equipment; $p_{i,ref}$ denotes the purchase cost for the equipment with the reference characteristic factor and 0.65 [84] denotes the scaling factor employed in the present study.

The economic assessment technique described thus far is the study estimate type (i.e. class 4 estimate) and is typically employed in facilitating conceptual studies [85]. The economic assessment technique was therefore specified as sufficient for the current comparative investigation. Finally, a summary of the economic assumptions employed in the present study are shown in Table 1.

| Parameter                  | Value   |
|----------------------------|---------|
| Base year                  | 2020    |
| Project lifetime (y)       | 30      |
| Plant availability (h/y)   | 7200    |
| Tax rate (%)               | 30      |
| Discount rate (%)          | 10      |
| Salvage value (US$)        | 0       |
| Depreciation               | Straight line |

2.3 Environmental performance assessment

Since the current study is interested in the comparative assessment of the stand-alone alternative AA production pathways, the potential environmental impact (PEI) of the alternative processes were investigated was selected as a sufficient metric in this regard [86,87]. The PEI was measured using the WAR algorithm since it constitutes a powerful tool that facilitates rapid assessments of environmental impacts during the design stages of manufacturing processes [87,88]. PEI employs energy and mass balance results of the simulated process as inputs for the calculation of its constituent impact categories [69] highlighted in Table 2. The PEI out of a process is calculated as follows [89]:

$$i_{out, ep} = \sum_i I_i = \sum_i M_i \sum_k x_k \phi_k$$

(6)

$$i_{out, ep} = \sum_i M_i \sum_k x_k \phi_k$$

(7)

where $i_{out, ep}$ denotes the rate of PEI out of the process from chemical interactions; $i_{out, ep}$ denotes the rate of PEI out of a system from energy generation processes; $M_i$ is the mass flow rate of exit streams from the process $i$, $x_k$ is the mass fraction of component $k$ in the exit stream $i$ and $\phi_k$ is the potential environmental impact due to component $k$. The parameter, $\phi_k$, denotes the sum of the specific PEI due to disposal of component $k$ to the environment leading to different impact categories $I$ as follows [89]:

$$\psi_k = \sum_i \alpha_i \phi_{ki}$$

(8)
where, $a_i$ denotes the relative weighting factor of impact categories, $J$ (Table 2), with all impact categories considered equally significant in the present study. In the present study the environmental impact of the processes were reported in terms of the PEI per unit mass of product stream. For simplicity, PEI/h from the biomass (WAP) have been considered negligible. Further descriptions of the WAR algorithm and the governing Equations are outside the scope of the present study and are presented elsewhere [87,89,90].

Table 2: Impact categories that constitute the basis of the potential environmental impact [87,90]

| General Impact Category | Impact Category | Measure of Impact Category |
|-------------------------|-----------------|---------------------------|
| Human toxicity          | Ingestion (HTPI) | LD50                      |
|                         | Inhalation/dermal (HTPE) | OSHA PEL                |
| Ecological toxicity     | Aquatic toxicity (ATP) | Fathead Minnow LC50     |
|                         | Terrestrial toxicity (TTP) | LD50                    |
| Global atmospheric impacts | Global warming potential (GWP) | GWP                    |
|                         | Ozone depletion potential (ODP) | ODP                    |
| Regional atmospheric impacts | Acidification Potential (AP) | AP                     |
|                         | Photochemical oxidation potential (PCOP) | PCOP                  |

3. Results

Based on the processes described in section 2, the ASPEN plus developed models are presented in Figures 5 and 6, with the mass balance results of major streams presented in Table 3. Figure 5 represents the FD process which integrates WAP handling and homogenization, with pre-treatment (for sugar release), fermentation (for LA production) and dehydration (for AA production). The model developed based on TFO process is shown in Figure 6 and this includes WAP handling and gasification (for syngas production), syngas fermentation (for ethanol production), ethanol purification, ethanol dehydration (for ethylene production), ethylene to propylene conversions and propylene oxidation (for AA production) operations. The result in Table 3 shows that the FD process will facilitate the production of 1.94 tons/h (14 ktons/y) of AA (97 wt.% purity) and 0.34 tons/h (2.45 ktons/y) of acetaldehyde as by-product. It is estimated that for the equal mass of feedstock, the resulting mass of AA generated via the FD process is approximately 6 times (1.94 tons/h, 97 wt. % purity) than that the TFO process (0.313 tons/h, 97 wt. % purity). A by-product of AC of 1.2 tons/h was also generated.
Figure 5: ASPEN plus output for AA production from WAP via the fermentation dehydration (FD) pathway (black continuous lines, red dash lines and blue dashed line denote material, heat and work flows).

Figure 6: ASPEN plus output for AA production from WAP via the thermochemical-fermentation-oxidation (TFO) pathway (black continuous lines, red dash lines and blue dashed line denote material, heat and work flows).
Figure 7: Potential environmental impacts per kg of product (PEI/kg) for the fermentation dehydration (FD) and thermochemical fermentation oxidation (TFO) pathways.

Table 3: Mass balance results for major streams in the AA production processes

| Stream name  | Fermentation dehydration (FD) process | Thermochemical-fermentation-oxidation (TFO) process |
|--------------|---------------------------------------|--------------------------------------------------|
|              | Hydrolysate   | Stream LA | AA | AC | Syngas | Ethyl acetate | Stream P | AA |
| 2-Butene     | 0            | 0         | 0  | 0  | 0      | 0.550        | 0        | 0  |
| AC           | 0            | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| AA           | 0            | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| Carbon dioxide| 0         | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| Carbon monoxide| 0       | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| Ethanol      | 0            | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| Fructose     | 0.018        | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| Glucose      | 0.024        | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| Hydrogen     | 0            | 0         | 0  | 0  | 0      | 0            | 0        | 0  |
| LA           | 0            | 0.500     | 0  | 0  | 0      | 0            | 0        | 0  |
Methane 0 0 0 0 0 0 0 0 0
Propylene 0 0 0 0 0 0 0 0.450 0
Toluene 0 0 0.030 0 0 0 0 0 0.030
Ethyl acetate 0 0 0 0 0 0 1 0 0
Water 0.955 0.500 0 0.020 0 0 0 0 0
Xylose 0.006 0 0 0 0 0 0 0 0

Mass Flows (tons/h) 99.30 7.83 1.94 0.33 0.483 1.166 0.478 0.313

LA and P denote the streams containing crucial intermediate compounds of Lactic acid and propylene respectively. AC denotes Acetaldehyde.

Table 4: External utility requirements of the conversion processes, assuming full heat integration.

| Process result                          | (FD) process | (TFO) process |
|-----------------------------------------|--------------|---------------|
| Externally required heating utility (MW)| 59.66        | -             |
| Externally required cooling utility (MW)| 27.48        | 15.87         |
| Net electricity generated (MW)          | 11.45        | 0.83          |

FD denotes the Fermentation dehydration process and TFO denotes the Thermochemical-fermentation-oxidation process.

Table 5: Cost distribution for total capital investment for acrylic acid production via two pathways

| Costing components                  | Fermentation-dehydration pathway (FD) | Thermochemical-fermentation-oxidation (TFO) |
|-------------------------------------|---------------------------------------|---------------------------------------------|
| Warehouse cost (US $)               | 342807.9                              | 536738.2                                    |
| Equipment installation cost (US $)  | 771317.8                              | 1207661.0                                   |
| Home office and construction fee (US $) | 385658.9                              | 603830.5                                    |
| Site development (US $)              | 8570186.2                             | 13418455.1                                  |
| Prorateable expenses (US $)          | 1006998.3                             | 1576668.5                                   |
| Project contingency (US $)           | 1006998.3                             | 1576668.5                                   |
| Additional piping (US $)             | 2013996.6                             | 3153337.0                                   |
| Field expenses (US $)                | 1006998.3                             | 1576668.5                                   |
| Other costs (start up , permits etc) (US $) | 1006998.3                             | 1576668.5                                   |
| Working capital (US $)               | 805598.6                              | 1261334.8                                   |
| Total capital investment (US $)      | 16917571.2                            | 26488030.6                                  |
Table 6: Cost distribution for total operating cost for acrylic acid production via two pathways

| Costing components       | Fermentation-dehydration pathway (FD) | Thermochemical-fermentation-oxidation (TFO) |
|--------------------------|---------------------------------------|---------------------------------------------|
| Labour cost (US $)       | 1504296                               | 1504296                                    |
| Maintenance cost (US $)  | 1353866.4                             | 1353866.4                                  |
| Labour burden (US $)     | 113616.9                               | 214100                                     |
| Property insurance (US $)| 1127838.1                             | 1765868.7                                  |
| Total variable cost (US $)| 14834442.6                            | 8791658.6                                  |
| Total operating cost (US $)| 18934059.9                           | 13629789.7                                 |
Table 7: The minimum selling price (MSP) of acrylic acid

| Process                                         | MSP (US$/kg) |
|------------------------------------------------|--------------|
| Fermentation-dehydration pathway (FD)          | 1.17         |
| Thermochemical-fermentation-oxidation (TFO)    | 1.56         |

Employing ASPEN energy analyzer® V.11 discussed above in section 2.1, the externally required heating and cooling utilities required are highlighted in Table 4. Employing methods discussed in section 2.3, the economic performances of FD and TFO processes were assessed. The TCI and the total operating cost (TOC) components results are presented in Tables 5 and 6 respectively. Employing the TOC and TCI values, the economic performance parameters were determined and summarized in Table 7. Employing the WAR software and using the methods discussed earlier above in section 2.3 above, the PEI/kg for both processes are summarized in Figure 7.

4. Discussions

Table 3 shows that based on the AA purity of 97 wt. % achieved in the present study, the AA product can be designated as technical grade (i.e. 94 wt.% ≤ AA purity < 99.5 wt.%) [91,92]. Such technical grade AA can be readily used in the synthesis of acrylate esters that can be transformed into co-monomers [93]. These co-monomers can be polymerized to produce valuable products such as surface coatings, adhesives, textiles, paints, polymeric materials etc.[94]. Table 3 also shows that the composition of syngas generated on a mass basis is 8%, 31%, and 61% for H₂, CO, and CO₂, which translates to mole basis equivalents of 62%, 17%, and 21% respectively. This mole basis yields of H₂, CO, and CO₂ is comparable to the experimentally determined mole composition of syngas of ~60%, ~22%, and ~18% reported in the literature [94]. The mass balance results for major streams in the AA production processes are summarized in Table 3. The table shows that only 0.478 tons/h of propylene is generated from 0.786 tons/h of ethanol, translating to a yield of 27 wt.%. This observation is consistent with experimentally reported yield of propylene from ethanol, which was shown to range from 24.5-28.6 wt.% [95]. In addition, considering the mass flow rate of sugars (C₆+C₅) in the hydrolysate (4.77 ton/h) and the mass flow rate of LA (i.e. 3.915 ton/h) presented in Table 3, the LA yield per unit mass of sugars can be calculated to be 82.1 wt.%. This yield is consistent with the yield of LA of 88 kg of LA per 100 kg sugars (from apple pomace) and based on experimental work [96]. As expected, in both cases, the actual yield of LA is less than the theoretical yield of 1 kg of lactic per 1 kg of sugar [96]. Table 4 shows that while the external heating utility requirement of the FD pathway for AA production is 59.66 MW the external heating utility requirement of the TFO pathway is determined to be negligible. These results are based on the assumption that full heat integration is achieved. This observation is expected and consistent with processes that incorporate exothermic processes [97-99]. In the TFO process, the oxidation of propylene for the generation of acrolein, AA, acetic acid and CO₂ as discussed in section 2.1 is highly exothermic [74]. For such exothermic systems, the need for addition external heating utility is redundant. Similarly, since the exit hot streams (from the exothermic processes) are cooled by being employed as heat sources for steam generation the externally required cooling duty calculated (15.87 MW) is shown to be lower than the externally required cooling duty required in the FD process (27.48 MW). Table 4 also shows that the FD process has the potential for higher net electricity generation potential of 11.45 MW, compared to the TFO process with the net electricity production potential of 0.83 MW. This observation is expected since in FD, the lignin fraction (HHV = ~ 25.5 MJ/kg (100)), employed as a solid fuel in the boiler for steam generation has a higher energy density compared to the TFO process where CO (HHV = 4.368 MJ/kg[55]) is employed as the gaseous fuel. Clearly, the combustion of the lignin will lead to the generation of more boiler heat leading to more energetic steam generation, compared to the combustion of CO. Table 5 shows that the total capital investment (TCI) calculated for the TFO process of US$ 26.5 million is 56 % more (~) than the value calculated for the FD process of US$ 16.9 million. This observation could be justified by the differences in processing complexity, with the TFO process clearly shown to incorporate more processing steps than the FD process. A higher number of processing steps employed typically translates to a higher capital cost. This statement is consistent with the fundamental basis of the ‘functional unit’ approach in capital cost estimation [101,102]. Table 5 also shows that in both FD and TFO processes, the equipment installation cost is the most crucial costing component and constitutes 51% of the total TCI. This observation is consistent with the class 4 estimation approach that considers the major equipment cost as the most crucial in calculating initial project cost estimates. Table 6 on the other hand, shows that, the total operating cost of the FD process (US$ 18.9 million) exceeds the total operating cost of the TFO process (US$ 13.6 million) with the variable cost component accounting for 78 % and 65 % respectively. This observation is expected since based on model descriptions in section 2.1, the FD process will involve a higher variable cost (i.e. chemical inputs of Ca(OH)₂, H₂SO₄, catalyst, toluene required and waste gypsum management cost) compared to the TFO process (catalyst, toluene). Table 7 shows that the MSPs of the AA produced via the TFO and FD processes are US$ 1.56 per kg and US$ 1.17 per kg respectively. This observation is due to the generation of more AA product in FD process, as discussed above and also the lower TCI requirement of the FD process as discussed above.
Figure 7 shows that the FD pathway has a higher environmental impact (8.72 kPEI/h) compared to the TFO pathway (2.07 kPEI/h). In both pathways the photochemical oxidation potential (PCOP) index constitutes the highest impact category with values of 6.06 kPEI/h and 1.48 kPEI/h for the FD and TFO pathways respectively. This observation may be due to the larger mass of toluene required in the FD process (642.2 kg/h) for AA recovery compared the TFO process (102.7 kg/h) and employed in AA recovery. Volatile organic compounds such as toluene have the potential of undergoing photochemical reactions with oxides such as nitrous oxides (NO\textsubscript{x}), leading to unwanted smog formation, implying that higher masses of toluene will exacerbate the PCOP pollution risk [87]. Figure 7 also shows that for the impact categories of HTPI, HTPE and TTP, the impacts associated with the FD process exceed the TFO process. These observations may be a consequence of the higher mass of toluene employed in FD compared to the TFO process. Notably the wastewater stream generated via the FD process is characterized by residual sulphuric acid and calcium sulfate which are consequences of the lactic acid purification process discussed in 2.1 above. Sulphuric acid and calcium sulfate have a potential of exacerbating unfavorable environmental impacts via an increase in indexes of ATP (aquatic toxicity) and TTP (Terrestrial toxicity) when improperly managed. The presence of these pollutants in the waste stream from the FD process may explain the higher ATP (0.047 kPEI/h) and TTP (1.08 kPEI/h) indexes of the FD process compared to the AP (0.0085 kPEI/h) and TTP (0.278 kPEI/h) values of the TFO process. The higher impact AP (acidification potential) index value (0.213 kPEI/h) for the FD process relative to the AP index value (0.213 kPEI/h) of the TFO process is due to need for external fossil energy to satisfy the heating energy needs of the FD process. This is because the use of these fuels will lead to the unwanted production of oxides such as CO\textsubscript{2} that are capable of forming weak acids (i.e. H\textsubscript{2}CO\textsubscript{3} for acid rain production. The study therefore suggests the preference for the TFO process compared to the FD process if only the environmental performance metric is considered. This observation contradicts the conclusions based on the economic performance metric since the FD process was shown to be preferable to the TFO process as illustrated by its lower MSP of AA. Such a scenario therefore presents a conundrum, with the selection of the preferred AA production pathway dependent on tradeoffs between the need for favorable economic and environmental outcomes by decision makers.

5. Conclusion
The study presents an initial comparative analysis of the two acrylic acid (AA) production pathways while employing waste apple pomace (WAP) as a renewable resource. In this study, we assessed FD and TFO processes as alternative approaches. The FD process incorporates an initial pretreatment operation for the production of a hydrolysate stream containing useful sugars followed by fermentation of these sugars for lactic acid (LA) production prior to LA dehydration for AA production. The TFO process on the other hand integrates the gasification of WAP followed by the fermentation of the syngas for alcohol production, which is employed as a precursor for propylene gas production, prior to a catalytic oxidation process for AA production. Considering both processes, although the designed processes are speculative, the FD process was shown to present an improved economic performance relative to the TFO process. This is highlighted by the lower MSP of the AA product from the FD process of US$ 1.17 per kg relative to the MSP of US$ 1.56 per kg from the AA product of the TFO process. Crucially, the TFO process presented an improved environmental performance compared to the FD process thus presenting a decision challenge for policy makers. The need for further multi-criteria decision assessment is therefore proposed.

| Abbreviations | Full words |
|---------------|------------|
| FD            | Fermentation-dehydration |
| TFO           | Thermochemical-fermentation-oxidation |
| WAP           | Waste apple pomace |
| AA            | Acrylic acid |
| LA            | Lactic acid |
| GHG           | Greenhouse gas |
| NRTL          | Non-random two-liquid |
| RSTOIC        | Stoichiometric reactor model |
| RGIBBS        | Equilibrium reactor model |
| RADFRAC       | Rigorous distillation column model |
| CHP           | Combined heat and power |
| HHV           | higher heating value |
| TCI           | total capital investment |
| TOC           | total operating costs |
| FCI           | Fixed capital investment |
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