Comparison of various organic acids for xylo-oligosaccharide productions in terms of pKa values and combined severity

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Abstract

Background: Methods to produce XOS have been intensively investigated, including enzymatic hydrolysis, steam explosion, and acid hydrolysis. Acid hydrolysis is currently the most widely used method to produce XOS due to its advantages of fewer processing steps, stronger raw material adaptability, higher yield, and better reproducibility. Especially, organic acids such as acetic acid, formic acid and xylenic acid work better as compared with mineral acids. However, the catalytic mechanism of different organic acids has been little studied. In this paper, four different organic acids, including formic acid, glycolic acid, lactic acid, and acetic acid were selected to compare their hydrolytic effects.

Results: Using pKa values as the benchmark, the yield of XOS increased with the increasing value of pKa. The yield of XOS was 37% when hydrolyzed by 5% acetic acid (pKa=4.75) at 170°C for 20 min. Combined severity (CS), a parameter associated with temperature and reaction time was proposed, was proposed to evaluate the hydrolysis effect. The results of CS were consistent with that of pKa values on both the yield of XOS and the inhibitor.

Conclusion: The results based on pKa values and combined severity, a parameter associated with temperature and reaction time, concluded that acetic acid is a preferred catalyst. Combining the techno-economic analysis and environmental benefits, acetic acid hydrolysis process has lower factory production costs, and it is also an important metabolite and a carbon source for wastewater anaerobic biological treatment. In
conclusion, production of xylo-oligosaccharides by acetic acid is an inexpensive, environment-friendly, and sustainable processing technique.

**Keywords:** xylo-oligosaccharide; acid hydrolysis; combined severity; technoeconomic analysis; pKa.
Background

Agricultural and forestry waste is an important biomass and renewable resource [1]. Discussions regarding the conversion and utilization of agricultural and forestry waste, such as straw, chaff, and hardwood, have dominated research in recent years. China is a large agricultural country, producing about 900 million tons of agricultural waste and 140 million tons of forestry waste every year. The utilization of waste is tied to the development of the country’s economy and environment. The most common and traditional methods for this include composting and incineration. Composting is simple in operation and low in cost [2], nevertheless, the accumulated wastes tend to cause pollution and harm the environment to a large extent over a period of time. Incineration is the most widely used treatment method that effectively reduces the accumulation of wastes and generates electricity to produce heat. However, the process produces a large number of harmful gases which not only cause environmental pollution but also causes the waste of resources [3]. The generation of greater economic and environmental benefits requires the application potential of agricultural and forestry wastes to be further explored. This was demonstrated in a number of studies that used lignocellulose found in rich amounts in agricultural and forestry waste, and is a valuable resource for industrial application. Lignocellulose, the most abundant renewable resource on earth, is a complex of cellulose, hemicellulose, lignin, extractives, and inorganic components. It is well established that lignocellulose is too recalcitrant to be utilized by microorganisms or enzymes due to the presence of lignin seals and hemicellulose sheaths,
and due to the crystallinity of cellulose. Consequently, lignocellulosic biomass pretreatment, a method that unfolds the structure of lignocellulose, plays a vital role in the lignocellulosic feedstock based biorefinery. Most widely studied pretreatment methods include physical, chemical, and biological methods. After pretreatment, hemicellulose can be converted into a mixture of oligosaccharides called xylo-oligosaccharides (XOS), an extremely valuable degradation product [4]. XOS are functional oligomers of sugars with straight as well as branched chains formed by 2-10 xylose molecules through β-1,4-glycosidic bonds [5]. At present, XOS are mainly used in food, feed, medicine, and health care products. According to the XOS consumption market share survey, the global turnover of XOS was about RMB 403 million. From this, XOS used in feed accounted for 41.54% of the market, while its use in medicine and health care products accounted for 29.01% and food and beverage accounted for 27.2%. As a type of functional oligosaccharide, XOS has attracted much attention due to its unique physiological properties. It was demonstrated in a number of studies that XOS can promote the proliferation of beneficial bacteria [5], strengthen immunity [7], prevent constipation, and lower serum cholesterol [8] among other benefits. In 2018, the European Union officially approved XOS as a new food additive.

In recent years, methods to produce XOS have been intensively investigated, including enzymatic hydrolysis [9,10], steam explosion [11], and acid hydrolysis. Acid hydrolysis is currently the most widely used method to produce XOS due to its advantages of fewer processing steps, stronger raw material adaptability, higher yield,
and better reproducibility. Especially, organic acids such as acetic acid, formic acid and xylonic acid work better as compared with mineral acids. In the current work’s preliminary studies, the preparation of XOS by organic acid has resulted in promising output. Zhang et al [12] reported a yield of 45.91% XOSs by acetic acid at pH 2.7 and at a temperature of 150 °C. The study used 0.64M xylonic acid to produce 44.5% XOS at 154 °C. In terms of the yield of XOS, these studies achieved significant results. The effect of different organic acid catalysts in the industrial production of XOS from wheat straw was reported in this study. Using formic acid, glycolic acid, lactic acid, and acetic acid as examples, a comparative selection method for hydrolyzed organic acids is proposed in this study.

Acidity coefficient (pKa) is an important benchmark of acids in the field of biochemistry. It is an equilibrium constant which represents the ability of an acid to dissociate hydrogen ions [13]. The pKa values affect the activity, water solubility, and spectral properties of chemicals. The four organic acids used in this study had different pKa values, and the conclusion is based on the yield of XOS and by-products. For more accurate visualization analysis, combined severity (CS) a function of reaction time, pH, and temperature was established to evaluate the acid pretreatment process. The comparison using CS avoids the influence of temperature on pKa. The conclusion at this stage was also based on the conversion rate of xylan. Finally, the whole industrial production process was projected and designed. In addition, techno-economic analysis was carried out during acid hydrolysis, and the cost of producing one-ton XOS by
different acids was calculated and demonstrated. The combination of these three means of comparison makes the selection of organic acids more convenient, effective, and convincing.

Results and discussion

The kinetics of acidolysis by various acids and degradation of wheat straw

Four small molecular weight organic acids at 0.8mol/L concentration were analyzed in this study. To compare their hydrolytic ability, their pKa values were taken as the benchmark (Table S1).

As observed in Figure 1, the yield of XOSs increased with higher pKa value. When the pKa value was 4.75, the yield reached 37%. Another interesting finding was that each component of XOS (from xylo-disaccharides to xylo-hexose) increased with the increase in the value of pKa. In addition, Figure 2 shows the yield of several by-products such as, xylose, furfural, and HMF. The yield of xylose shows a downward trend accompanied by the value of pKa ranging from 3.75 to 4.75. Notably, the ratio of XOS to the total by-products is also shown in Figure 2. This indicates a good agreement between the pKa value and the hydrolytic efficiency of different acids.

Acidic hydrolysis of lignocellulose is a homogeneous process and can be described with the kinetic model of Arrhenius equation (Gooch, 2011). The Arrhenius equation is shown in equation (2):

\[ k = k^0 \times e^{-Ea/RT} \]  

where, k is kinetic constant, s-1; k0 is a pre-factor, s-1; Ea is activation energy,
kJ/g·mol·K; R is gas constant - 8.314×10^{-3}, KJ/mol·K and T is the temperature, K.

To concretely describe the acid hydrolytic process, the Arrhenius equation was deformed using ionization constant. The equation can then be used for assessing the effect of different pKa values. The reformed equation is as followed:

\[ y = y^0 \times (ka)^m \times e^{-\frac{Ea}{RT}} \]  

(3)

In this equation, y represents the yield, %; y0 represents pre-factor, %; m is index factor; ka represents the ionization constant. The pKa value is directly related to the ionization constant as follows:

\[ pKa = -log(ka) \]  

(4)

According to Eq. 3, the acid hydrolysis of hemicellulose is related to the ionization constant and temperature. The kinetics of hemicellulose hydrolysis at different temperatures and ka values can be determined. In this study, the temperature is kept at 170 °C. Then use formula (4) to fit the yield of xylose in Figure 2, and the activation energy could be calculated in Table 1. It was observed that when the hydrolysis was catalyzed by an acid with a higher pKa value, lower activation energy was required to produce xylose. The activation energy corresponds to the complexity level of the reaction. This is also supported by Figure 1 and Figure 2 which represent that during hydrolysis with a low pKa value acid, XOS are more easily degraded to xylose, thereby reducing the yield of XOS.

To compare the hydrolytic effects of various acids on subsequent enzymatic hydrolysis reactions, 25 FPIU/g of cellulose was used as the hydrolytic enzyme. The
result obtained after 108 h of treatment is shown in Figure 3.

When the residue was pretreated with acids of smaller pKa value and used as the substrate for enzymatic hydrolysis, the enzymatic hydrolysis rate was higher and reached 98%. However, the maximum enzymatic hydrolysis rate of the residues was 85% when hydrolyzed with acids of 4.75 pKa value, but had the highest XOS yield. On comparing the yield of XOS with enzymatic hydrolysis (Figure S1), acetic acid with a pKa value of 4.75 was found to be significantly superior in terms of XOS yield as well as effectiveness. Therefore, acetic acid was the preferred catalyst for the pretreatment process.

Establishment and application of combined severity for the evaluation of pretreatment process

The catalytic effect of acids changed with their pKa value, but the calculation function of pKa and temperature has not been determined. For more accurate visualization analysis, CS- a function of reaction time, pH, and temperature- was established to evaluate the acid pretreatment [14]. The CS is defined as follows[15]:

\[
CS = \log\{t \times \exp\left[\left(\frac{T_H - T_R}{\omega}\right)\right]\} - \text{pH} \tag{6}
\]

where, \(t\) is reaction time in min, \(T_H\) is the hydrolysis temperature in °C, \(T_R\) is a base temperature set at 100 °C, and \(\omega\) is a fitting parameter which in most studies is assigned with the value of 14.75.

The pH of a weak acid decreases with an increase in temperature because of its heated ionization process, represented by the Henderson–Hasselbalch equation. Thus,
a higher pH than the original value was used to calculate the CS of weak acids. In the present study, four organic acids, including formic acid, glycolic acid, lactic acid, and acetic acid were used for the pretreatment at different temperatures. The conditions and constants are shown in Table 2.

In order to evaluate the effect of pretreatment with different acids, CS was employed to describe the pretreatment condition. The value of CS has been plotted against XOS yield, Xylose yield, and the ratio of XOS and by-products (Figure 4).

The trend of XOS or Xylose (by-products) yield by different acids was inconsistent and irregular, because CS is a parameter comprising of temperature, pH, and time, but not acid type. Although the trend is irregular, there is no doubt that each graph shows an evident peak value. In Figure 4 (a), the peak was observed with acetic acid with a value of 37%. The order of the effect of different acids on XOS yield was acetic acid > lactic acid > glycolic acid > formic acid. In contrast, the effect of acids on the peak values of xylose yield was in the opposite order - formic acid > glycolic acid > lactic acid > acetic acid. From the results of CS evaluation, it can thus be concluded that acetic acid should be preferred for either higher XOS yield or xylose (by-products) yield. Moreover, it’s an excellent discovery that the results of CS evaluation are consistent with pKa values of different acids.

**Tech-noeconomic analysis**

Techno-economic analysis is extremely important in research and development of lignocellulosic biofuels production processes at NREL from over two decades [16],
These include both biochemical and thermochemical approaches. In order to analyze the techno-economic aspects, the whole process of industrial application is depicted in Figure 5.

Baled wheat straw was delivered to the processing plant where they were broken, washed, and milled. Straw was then loaded for pretreatment at a certain (such as 1:10) solid-to-liquid ratio. After solid-liquid separation, the liquid portion was treated to obtain the target product XOS. The solid residue portion, on the other hand, goes through enzymatic degradation and fermentation process. An approximate cost composition analysis of the whole process is shown in Figure 6.

In general, the maximum cost of the process is associated with the raw materials, which is about 38% of the production cost for one ton XOS. It can be observed from Figure 6 that the techno-economic difference between four different acid hydrolysis processes lies in the cost of pretreatment and conditioning, in which the ratio of capital recovery charge is 11%, raw material is 2%, process electricity is 0.8% and the fixed cost is 3%. Among these, capital recovery charge and fixed cost do not change with the type of acids. In order to compare the cost of four different acid hydrolysis processes, the price of acid is the main concern and next is the process electricity (take the best advantage of each acid in this experiment as an example). The detailed cost accounting is shown in the following table (Table 3).

For each ton of XOS produced, the order of the acids in terms of the combined cost of raw materials and process electricity was as follows: Glycolic acid > Lactic
acid > Formic acid > Acetic acid. Considering the production cost of the factory, acetic acid should be preferred as catalyst. Furthermore, besides the lower production costs, acetic acid has other advantages. Previous laboratory studies have shown that about 75% of acetic acid can be recovered by distillation [17]. Anaerobic biological treatment plays an important role in industrial wastewater treatment, where acetic acid [18] is a metabolite in the second stage of wastewater treatment and a carbon source in the third stage (Figure S2). This has important implications in the industrial production.

4. Conclusion

The hydrolytic effect of four different organic acids on wheat straw was relatively obvious. Using pKa values as the benchmark, the yield of XOS increased with the increasing value of pKa. The yield of XOS was 37% when hydrolyzed by 5% acetic acid (pKa=4.75) at 170°C for 20 min. To eliminate the impact of temperature on the dissociation constant, combined severity (CS), a parameter associated with temperature and reaction time was proposed. The results of CS were consistent with that of pKa values on both the yield of XOS and the inhibitor. In terms of technoeconomic analysis, the overall cost of acetic acid was lower. For environmental benefits, acetic acid can be recovered by more than 75% and it is also an important metabolite and carbon source for wastewater anaerobic biological treatment.

Materials and Methods

Materials

The lignocellulosic material, wheat straw, was obtained from Lianyungang city
(Jiangsu Province, China). It was grouped into 20 mesh powder and was air dried to reduce the moisture under 8%. The chemical composition (wt %, based on the oven-dried weight) of the wheat straw was: 23.53 % xylan, 35.34 % glucan, 30.63 % acid-insoluble lignin and 3.10 % acid-insoluble lignin.

**Acidolysis of wheat straw**

This study used four different acids (formic acid, acetic acid, lactic acid, and glycolic acid) as catalysts and the concentration of each acid was 5 % (w/v). The pretreatments were carried out in 35 mL stainless-steel reactors, to which 15 mL mixture, containing the substrate and the acid catalyst at a 1:10 solid-to-liquid ratio, was added. The reaction mixture was heated in a 170 ℃ oil bath for 20 min. After the reaction was complete, the supernatant was measured and the slag was used for enzymatic hydrolysis.

**Analytical methods**

The composition of the corncob was determined according to the standard protocol utilized by the National Renewable Energy Laboratory (NREL). XOS (xylobiose, xylotriose, xylotetraose, xylopentaose, and xylohexasose) were analyzed using high-performance anion exchange chromatography (HPAEC-PAD), coupled with pulsed amperometry detector (Dionex ICS-5000). The mobile phase used was 0.1 M NaOH and 0.5 M NaOAc (containing 0.1 M NaOH) at a flow rate of 0.3 mL/min and the anion-exchange column was CarboPac™ PA200.

The monosaccharide and other by-products were detected by high-performance
liquid chromatography (HPLC, Agilent 1260, Agilent Technologies, Santa Clara, CA, USA) using an Aminex Bio-Rad HPX-87H column and a refractive index detector at 55 °C, with 5 mmol/L sulfuric acid (H₂SO₄) as the mobile phase at a flow rate of 0.6 mL/min.

The yield of XOS was calculated according to the equation (1):

\[
\text{XOS yield} = \frac{\text{XOS}}{\text{initial xylan content in substrate}} \times 100\% \quad (1)
\]
Declarations

Abbreviations

XOS: xylooligosaccharide; CS: combined severity; HPLC: high performance liquid chromatography; HPAEC: high performance anion-exchange chromatography; NREL: National Renewable Energy Laboratory; CRM: cost of raw material; CPE: cost of process electricity.

Authors’ contributions

RC and XLL developed the idea for the study. RC and JMG performed the research. RC conducted the data analysis and prepared the manuscript. YX helped to revise the manuscript. All authors read and approved the final manuscript.

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Competing interests

The authors declare that they have no competing interests.

Availability of data and materials

All data generated and analyzed in this study are included in this published article.

Consent for publication

Not applicable
Ethics approval and consent to participate

Not applicable

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Supplementary material

Supplementary material relate to this article can be found in text.
References:

[1] Xie H, Zhang D, Mao G, Wang F, Song A. Availability of lignocellulose from forestry waste for use as a biofuel in China. Biotech. 2018; 8:234.

[2] Jia T, Xu Y, Luan Y. Agricultural and Forestry Waste Composts as Substitutes for Peat in Potting Media: Effects on Root Growth and Fractal Features of New Guinea impatiens (Impatiens hawkeri). Agricultural Research. 2016; 5:269-76.

[3] Joseph S, Xu CY, Wallace HM, Farrar M, Nguyen TT, Bai SH, et al. Biochar Production From Agricultural and Forestry Wastes and Microbial Interactions. 2017:443-73.

[4] Guo J, Huang K, Zhang S, Xu Y. Optimization of selective acidolysis pretreatment for the valorization of wheat straw by a combined chemical and enzymatic process. Journal of Chemical Technology & Biotechnology. 2019; 95:694-701.

[5] Carvalho AFA, Neto PDO, Silva DFD, Pastore GM. Xylo-oligosaccharides from lignocellulosic materials: Chemical structure, health benefits and production by chemical and enzymatic hydrolysis. Food Res Int. 2013;51 :75-85.

[6] Carlson JL, Erickson JM, Hess JM, Gould TJ, Slavin JL. Prebiotic Dietary Fiber and Gut Health: Comparing the in Vitro Fermentations of Beta-Glucan, Inulin and Xylooligosaccharide. Nutrients. 2017;9.

[7] Hansen CHF, Frøkiaer H, Christensen AG, Bergström A, Licht TR, Hansen AK, et al. Dietary Xylooligosaccharide Downregulates IFN-γ and the Low-Grade Inflammatory Cytokine IL-1β Systemically in Mice. The Journal of Nutrition.
[8] Kyung M, Choe HS, Jung S, Lee K, Jo SE, Seo S, et al. Effects of xylooligosaccharide-sugar mixture on glycemic index (GI) and blood glucose response in healthy adults. Journal of Nutrition & Health. 2014;47.

[9] Peter, Immerzeel, Peter, Falck, Mats, Galbe, et al. Extraction of water-soluble xylan from wheat bran and utilization of enzymatically produced xylooligosaccharides by Lactobacillus, Bifidobacterium and Weissella spp. Lwt Food Science & Technology. 2014.

[10] Lu J, Liu H, Xia F, et al. The hydrothermal-alkaline/oxygen two-step pretreatment combined with the addition of surfactants reduced the amount of cellulase for enzymatic hydrolysis of reed[J]. Bioresource Technology, 2020, 308:123324.

[11] Azevedo, Carvalho, Ana, Flavia, Marcondes, Wilian, et al. The potential of tailoring the conditions of steam explosion to produce xylo-oligosaccharides from sugarcane bagasse. Bioresource Technology Biomass Bioenergy Biowastes Conversion Technologies Biotransformations Production Technologies. 2018.

[12] Zhang H, Xu Y, Yu S. Co-production of functional xylooligosaccharides and fermentable sugars from corncob with effective acetic acid prehydrolysis. Bioresource Technol. 2017; 234:343-9.

[13] Lim C, Bashford D, Karplus M. Absolute pKA calculations with continuum dielectric methods. Journal of Physical Chemistry. 1991;95.

[14] Kabel MA, Bos G, Zeevalking J, Voragen AGJ, Schols HA. Effect of pretreatment
severity on xylan solubility and enzymatic breakdown of the remaining cellulose from wheat straw. Bioresource Technol. 2007; 98:2034-42.

[15] Chum HL, Johnson DK, Black SK. Organosolv pretreatment for enzymic hydrolysis of poplars. 2. Catalyst effects and the combined severity parameter. Ind.eng.chem.res. 1990;29.

[16] Wright JD. Economics of enzymatic hydrolysis processes. American Institute of Chemical Engineers Spring National Meeting1988.

[17] Huang K, Luo J. β-Factor Based Separation Characteristics of Bio-derived Chemicals Present in Lignocellulosic Hydrolysates Using Vacuum Distillation. Acs Sustain Chem Eng. 2019; 7:2406-13.

[18] Ye C, Cheng JJ, Creamer KS. Inhibition of anaerobic digestion process: A review. Bioresour Technol. 2008; 99:4044-64.
Figure Captions

Fig. 1. The concentration and yield of XOS with different pKa values

Fig. 2. The yield of by-products with different pKa values

Fig. 3. Enzymic yield of acidolysis residue with different acids

Fig. 4. Combined severity versus XOS yield (a), Xylose yield (b), XOS/By-products (c)

Fig. 5. Overall industrial process

Fig. 6. The cost composition analysis of the whole process
Fig. 1
Fig. 2
Fig. 3
Fig. 4

(a) XOS yield (%) vs. CS

(b) Xylose yield (%) vs. CS

(c) XOS/By-products vs. CS

Symbols:
- ▲ Formic acid
- □ Glycolic acid
- △ Lactic acid
- □ Acetic acid
Fig. 5
Fig. 6

- Storage
- Utilities
- Boiler/Turbogenerator
- Wastewater Treatment
- Enzymolysis/Fermentation
- Pretreatment/Conditioning
- Feed Handing
- Biomass

Cost composition (%)

- Capital Recovery Charge
- Raw Materials
- Process Electricity
- Grid Electricity
- Fixed Costs

(Included in price of Purchased Feedstock)
Table 1. The activation energy value of degradation to xylose

Table 2. Conditions and constants of four organic acids during the acidolysis

Table 3. The detailed cost accounting
| pKa | $ka \times 10^{-5}$ | $y^0$ (%) | $E_a$ (kJ/g • mol • k) | Adjusted $R^2$ |
|-----|-----------------|----------|------------------------|----------------|
| 3.75 | 17.78           | 0.52     | 107.12                 |                |
| 3.83 | 14.79           | 0.76     | 109.21                 | 0.973          |
| 4.14 | 7.24            | 1.53     | 115.52                 |                |
| 4.75 | 1.78            | 5.65     | 131.98                 |                |
| Temperature (°C)/Time(min) | Acid type  | pH\(^a\) | CS  |
|---------------------------|------------|----------|-----|
| 130/20                    | Formic acid| 1.63     | 0.55|
|                           | Glycolic acid| 1.64    | 0.54|
|                           | Lactic acid| 1.82     | 0.36|
|                           | Acetic acid| 2.12     | 0.06|
| 130/45                    | Formic acid| 1.51     | 1.03|
|                           | Glycolic acid| 1.56   | 0.98|
|                           | Lactic acid| 1.75     | 0.79|
|                           | Acetic acid| 2.04     | 0.5 |
| 130/75                    | Formic acid| 1.36     | 1.4 |
|                           | Glycolic acid| 1.41  | 1.35|
|                           | Lactic acid| 1.54     | 1.22|
|                           | Acetic acid| 1.89     | 0.87|
| 150/20                    | Formic acid| 1.45     | 0.38|
|                           | Glycolic acid| 1.51  | 0.32|
|                           | Lactic acid| 1.70     | 0.13|
|                           | Acetic acid| 1.98     | -0.15|
| 150/45                    | Formic acid| 1.32     | 0.86|
|                           | Glycolic acid| 1.42  | 0.76|
|                           | Lactic acid| 1.58     | 0.6 |
|                           | Acetic acid| 1.87     | 0.31|
| 150/75                    | Formic acid| 1.12     | 1.29|
|                           | Glycolic acid| 1.24  | 1.17|
|                           | Lactic acid| 1.31     | 1.1 |
|                           | Acetic acid| 1.61     | 0.8 |
| 170/20                    | Formic acid| 1.26     | 0.72|
|                           | Glycolic acid| 1.35  | 0.63|
|                           | Lactic acid| 1.58     | 0.4 |
|                           | Acetic acid| 1.73     | 0.25|
| 170/45                    | Formic acid| 1.12     | 1.20|
|                           | Glycolic acid| 1.24  | 1.08|
|                           | Lactic acid| 1.40     | 0.92|
|                           | Acetic acid| 1.61     | 0.71|
| 170/75                    | Formic acid| 0.98     | 1.57|
|                           | Glycolic acid| 1.11  | 1.44|
|                           | Lactic acid| 1.17     | 1.38|
|                           | Acetic acid| 1.45     | 1.10|
Table 3

| Organic acid | CRM (RMB/t) | CPE (RMB/t XOS) | Costs |
|--------------|-------------|----------------|-------|
| Formic acid  | 2800        | 4017.75        | 592.142 |
| Glycollic acid | 8000     | 2203.25        | 1617.624 |
| Lactic acid  | 7800        | 3978.75        | 1591.83 |
| Acetic acid  | 2260        | 1214.25        | 461.714 |

Cost of raw material; Cost of process electricity; Cost = CRM × 8% + CPE × 0.2%