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System-level cost evaluation for economic viability of cellulosic biofuel manufacturing



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- A system-level cost model for cellulosic biofuel manufacturing is established.
- The relationships between individual process characteristics are studied.
- Two numerical cases are conducted to illustrate the effectiveness of the proposed model.
- The result shows that 12.8% of total cost can be reduced from baseline case.

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ABSTRACT

Biofuel is a clean and renewable energy source and is considered a promising alternative to traditional fossil fuels. The economic viability is crucial in promoting large-scale adoption and long-term sustainability of biofuel. Most of the current literature on biofuel economics assumes the individual biofuel manufacturing processes are independent of each other. Consequently, the interrelationships between parameters within and across processes regarding manufacturing cost and biofuel yield are not well investigated. In this paper, a system-level cost model for cellulosic biofuel manufacturing is established across multiple production processes to investigate the relationships between the individual process characteristics and the system performance to reduce the overall cost under the constraint of biofuel yield. Two numerical case studies are conducted to illustrate the effectiveness of the proposed model. Compared with the baseline case, the cost-effective case shows that 12.8% of the total cost is reduced without ethanol yield loss.

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1. Introduction

The continuously increasing energy consumption coupled with limited fossil fuel reserves is one of the most serious challenges to the sustainable development of human society. Although significant research progress has been made to investigate renewable energy resources as sustainable energy alternatives, fossil fuels accounts for the largest portion of the worldwide energy supply. The usage of fossil fuels has become the main source of carbon dioxide (CO₂) emission, which is one of the major greenhouse gases (GHGs) that lead to global warming. The global temperature increase associated with GHGs emissions from burning fossil fuels is expected to be 3.6 °C by 2040 [1]. The severe environmental burdens caused by fossil fuels make it critical to highly prioritize sustainable low-carbon fuels.

As a promising alternative, biofuel gains its popularity as a renewable energy source that can be sustainably developed [2]. Biofuels, especially bio-ethanol produced from cellulosic feedstock, appear to be environmentally friendly with no net CO_2 and very low sulfur emissions. In addition, biofuels can help decrease a country's energy dependence on imported oil, and have positive impacts on both the economy and the environment [3]. Given these advantages, biofuels have been receiving growing attention from countries around the world. As a result, global biofuel production in 2013 has increased seven times compared to 2000 [4]. Moreover, it is estimated that 30% of the U.S. liquid transportation fuels will be replaced by biofuels by 2022 [5].

Based on different biomass feedstocks and associated manufacturing systems, biofuels are divided into several categories such as corn-based biofuel, cellulosic biofuel, and algae biofuel, etc. Cellulosic biofuel has been considered as the most promising biofuel due to its capability of satisfying the fuel demands using agricultural wastes such as corn stover. As shown in Fig. 1, a typical



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Nomenclature

Bold face		M_{gl}	molecular weight of glucose (g/mol)	
R	reaction diagonal matrix	M _x	molecular weight of xylan (g/mol)	
F	formation diagonal matrix	M _{xl}	molecular weight of xylose (g/mol)	
S	state variable set	M _{xo}	molecular weight of xylose oligomer (g/mol)	
		No	amount of substance of accessible cellobiose lattices/g	
Upper case		N	cellulose	
A	pre-exponential factor (1/s)	N _A	Avogadro constant	
Aca	area of the cellobiose lattice (m^2)	$P_{i,g}$	threshold ethanol concentration of glucose (g/mol)	
Amay	maximum enzyme adsorption per g cellulose	$P_{i,x}$	threshold ethanol concentration of xylose (g/mol)	
AS	total surface area accessible to enzyme (m^2)	$P_{m,g}$	maximum ethanol concentration of glucose (g/mol)	
A.	reactor surface area (m^2)	$P_{m,x}$	maximum ethanol concentration of xylose (g/mol)	
Casid	active acid concentration (w/w%)	R	gas constant	
	initial acid concentration (w/w%)	Т	temperature (K)	
C^D	process-dependent cost (\$)			
C^{ID}	process-independent cost (\$)	Lower ca	se	
E Fa	activation energy	Α	ratio of the liquid volume and solid spheres volume	
E_{L}	bound enzyme concentration (g/kg)	Ce	concentration of ethanol (kg/L)	
	bound concentration of endo- β -1 4-glucanase and exo-	Cg	concentration of glucose (kg/L)	
LDI	β -1 4-glucanase (g/kg)	C_{gb}	concentration of cellobiose (kg/L)	
Fra	bound concentration of β -glucosidase (g/kg)	C_{xl}	concentration of xylose (kg/L)	
	free enzyme concentration ($\sigma/k\sigma$)	C _{xo}	concentration of xylose oligomer (kg/L)	
E _f Em	concentration of free β -glucosidase (g/kg)	C_Z	concentration of recombinant (kg/L)	
L _{f2} F,	energy consumption by heat transfer (kl)	d	feedstock particle diameter (m)	
En F	energy consumption by reaction (kl)	d _r	thickness of reactor (m)	
L _r F	energy consumption by heating up steam (kl)	h _{cellubiose}	entropy of cellobiose (kJ/kg)	
L _S F	total energy consumption (kI)	h _{cond}	conductivity of pretreatment reactor (W/m)	
K.	inhibition constant of glucose when glucan transforms	h _{conv}	convection coefficient (W/m^2)	
K I,gl	to glucose (g/I)	h _{ethanol}	entropy of ethanol (kJ/kg)	
К	inhibition constant of cellobiose when glucan trans-	hglucan	entropy of glucan (kJ/kg)	
K1,gb	forms to glucose (g/L)	hglucose	entropy of glucose (kJ/kg)	
K.	inhibition constant of vulose when glucan transforms to	h _{xvlan}	entropy of xylan (kJ/kg)	
$\mathbf{K}_{1,xl}$	alucose (a/L)	h _{xvlose}	entropy of xylose (kJ/kg)	
K	inhibition constants of glucose when glucan transforms	h _{xvlose-olig}	-oligomer entropy of xylose oligomer (kJ/kg)	
R _{2,gl}	to cellobiose (α/I)	k	reaction rate (1/s)	
K	inhibition constants of cellobiose when glucan trans-	ġ	heat flux (W/m^2)	
R _{2,gb}	forms to cellobiose (q/I)	$q_{emax,g}$	overall maximum specific ethanol production rate by	
<i>K</i>	inhibition constants of vulose when glucan transforms	118	glucose (g)	
$\mathbf{K}_{2,xl}$	to cellobiose (q/I)	q _{emax x}	overall maximum specific ethanol production rate by	
K	inhibition constants of glucose when cellobiose trans		xylose (g)	
K3,gl	forms to glucose (g/L)	$q_{smax,g}$	overall maximum specific glucose utilization rate (g)	
K.	cellobiose saturation constants when cellobiose trans	$q_{smax,x}$	overall maximum specific xylose utilization rate (g)	
K _{3m}	forms to glucose (g/L)	p_{σ}	mass of glucan in feedstock (kg)	
V	inhibition constants of vulose when collopiose trans	p_x	mass of xylan in feedstock (kg)	
к _{3,xl}	forms to glucose (g/L)	1.0		
V	inhibition constant of glucose when glucose transforms	Creek		
к _{i,g}	the other of (a/L)	a cicck	weight factor of glucose consumption	
V	to ethanol (g/L)	ß	initial water volume (I)	
κ _{i,x}	to otherpol (g/L)	γ λ	ratio of cellohiose lattices occupied to bound enzyme	
V	to entation constant in terms of L/π collulated	10	molecule	
К _р И	dissociation constant in terms of L/g cellulose	0	density of the solid (ka/m^3)	
К _{s,g} И	limitation constant of glucose (g/L)	Psolid	maximum overall specific growth rate of glucose $(1/s)$	
K _{S,X}	minitation constant of xylose (g/L)	µmax,g	maximum overall specific growth rate of vulces (1/s)	
IVIg	molecular weight of glucan (g/mol)	µmax,x	maximum overan specific growth fate of xylose (1/5)	

cellulosic biofuel manufacturing system consists of three main individual processes: pretreatment, enzymatic hydrolysis, and fermentation [6]. In the pretreatment process, the resistance of biomass cell walls is reduced by the overheated steam and acid. In enzymatic hydrolysis, the exposed cellulose chains are decomposed into fermentable sugars. These fermentable sugars are subsequently converted into biofuel (e.g., ethanol) in the fermentation process. Most research efforts regarding the biofuel manufacturing system are devoted to improving these processes from a biochemical perspective, such as pretreatment technology development [7–9] and fermentation bacterial cultivation [10–12]. The outcomes of these research efforts have built a theoretical foundation for the biofuel manufacturing industry.

The U.S. government has given a mandate to produce 16 billion gallons of cellulosic biofuel annually by 2022 [13]. To achieve this production target, many additional large-scale biofuel manufacturing plants are being constructed. Economic viability is one of the most critical aspects affecting the evaluation of large-scale implementation and long-term sustainability of cellulosic biofuel manufacturing system. Consequently, studies regarding the cost-benefit analysis of cellulosic biofuel have emerged [14–17]. However, in the current literature, most cost evaluation studies on cellulosic

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