



Integrated supply chain design for commodity chemicals production via woody biomass fast pyrolysis and upgrading



Yanan Zhang^a, Guiping Hu^{b,c,*}, Robert C. Brown^{a,b}

^a Department of Mechanical Engineering, Iowa State University, Ames, IA 50011, United States

^b Bioeconomy Institute, Iowa State University, Ames, IA 50011, United States

^c Industrial and Manufacturing Systems Engineering, Iowa State University, Ames, IA 50011, United States

HIGHLIGHTS

- Economic feasibility and the optimal production planning for commodity chemicals.
- Trade-off between economic and environmental metrics for chemicals is analyzed.
- Biomass availability and facility capital costs are the most important factors.

ARTICLE INFO

Article history:

Received 7 November 2013

Received in revised form 11 January 2014

Accepted 14 January 2014

Available online 29 January 2014

Keywords:

Supply chain optimization

Commodity chemicals

Fast pyrolysis

Woody biomass

ABSTRACT

This study investigates the optimal supply chain design for commodity chemicals (BTX, etc.) production via woody biomass fast pyrolysis and hydroprocessing pathway. The locations and capacities of distributed preprocessing hubs and integrated biorefinery facilities are optimized with a mixed integer linear programming model. In this integrated supply chain system, decisions on the biomass chipping methods (roadside chipping vs. facility chipping) are also explored. The economic objective of the supply chain model is to maximize the profit for a 20-year chemicals production system. In addition to the economic objective, the model also incorporates an environmental objective of minimizing life cycle greenhouse gas emissions, analyzing the trade-off between the economic and environmental considerations. The capital cost, operating cost, and revenues for the biorefinery facilities are based on techno-economic analysis, and the proposed approach is illustrated through a case study of Minnesota, with Minneapolis-St. Paul serving as the chemicals distribution hub.

© 2014 Elsevier Ltd. All rights reserved.

1. Introduction

The growing interest in biofuels production has generated much related research in economic analysis, environmental assessment, and supply chain system design (An et al., 2011; Blottnitz and Curran, 2007; Bowling et al., 2011; Giarola et al., 2012; Hamelinck et al., 2005; Hess et al., 2007; Larson, 2006; Stephen et al., 2010; You et al., 2012; Zhang et al., 2013a,b,d). Biomass logistics are complicated by the bulky, distributed nature of biomass and by the high volumes of low energy density materials to be collected and transported to the conversion facilities (Tallaksen, 2011). The unique nature of biomass feedstock provides great impetus for the exploration of sustainable and robust supply chain systems.

* Corresponding author. Address: Industrial and Manufacturing Systems Engineering, 3014 Black Engineering Building, Iowa State University, Ames, IA 50011, United States. Tel.: +1 5152948638.

E-mail address: gphu@iastate.edu (G. Hu).

Numerous studies have been devoted to optimal design and operational planning of the bioethanol supply chain. You et al. (2012) developed a multi-objective mixed integer linear programming (MILP) model which addressed the optimal design and planning of the cellulosic ethanol supply chain under economic, environmental, and social objectives. Dunnett et al. (2008) proposed a system model to optimize the lignocellulosic bioethanol supply chain under assumptions of energy integration. Bai et al. (2011) optimized biofuel refinery location and supply chain planning for bioethanol production, taking into account of traffic congestion issues. Giarola et al. (2012) developed a stochastic modeling framework adopting a scenario-based approach to assess the effects of trading greenhouse gas (GHG) emissions allowances under market uncertainty for bioethanol production.

Researchers have also been aggressively exploring the supply chain design for biomass-derived transportation fuels (gasoline and diesel fuel). You and Wang (2011) presented the optimal design and planning of a biomass-to-liquids (BTL) supply chain under

Vispute et al. (2010) proposed a novel integrated catalytic thermochemical pathway to convert woody biomass to commodity chemicals, such as benzene, toluene, and xylene aromatic hydrocarbons (BTX). In this pathway, the bio-oil produced from woody biomass fast pyrolysis undergoes two-stage hydrotreatment followed by fluid catalytic cracking (FCC). Due to the high selectivity of commodity chemicals products attainable using this production pathway, the pathway has garnered significant attention and has inspired further examination of its economic feasibility and environmental effects. A techno-economic study has been conducted to examine the five commodity chemicals production scenarios, one of which was Vispute's two-stage hydrotreating followed by FCC. Vispute's pathway is found to be the most profitable among the five scenarios (Brown et al., 2012). Another techno-economic study concluded that this chemicals production pathway is

Although there have been many studies of supply chain design and optimization for biofuel production, there have been few papers addressing supply chain design and optimization for renewable chemicals production from woody biomass via the thermochemical pathway. In this paper, a supply chain network is designed and optimized for the biobased chemical production pathway, using MILP modeling to optimize the locations and capacities of distributed preprocessing hubs and centralized biorefinery facilities. This paper examines both economic and environmental criteria in a multi-objective framework that allows analysis of trade-offs between economic feasibility and environmental impact. A case study for the state of Minnesota is presented to illustrate the integrated supply chain network design model.

2.1. Integrated catalytic processing pathway

Vispute et al. (2010) has proposed an integrated catalytic processing pathway for commodity chemicals production via woody biomass (Fig. 1). In this pathway, woody biomass is preprocessed (chopped, dried, and grinded) and then sent to a pyrolyzer to produce bio-oil. The bio-oil undergoes phase separations through a liquid-liquid extractor, resulting in separate water insoluble and aqueous phases. The water insoluble phase consists mainly of pyrolytic lignin, which is treated as a co-product. The aqueous phase is sent to a low temperature hydrotreating process (125 °C, 100 bar). Then the hydrotreated bio-oil is sent to a high temperature hydro-treating process for further hydrodeoxygenation (200 °C, 100 bar) over catalysts. After the two-stage hydrotreating process, FCC is performed on the hydrotreated aqueous phase to produce commodity chemicals. In addition to the primary raw material the woody biomass, hydrogen is needed for the two-stage hydrotreating process.

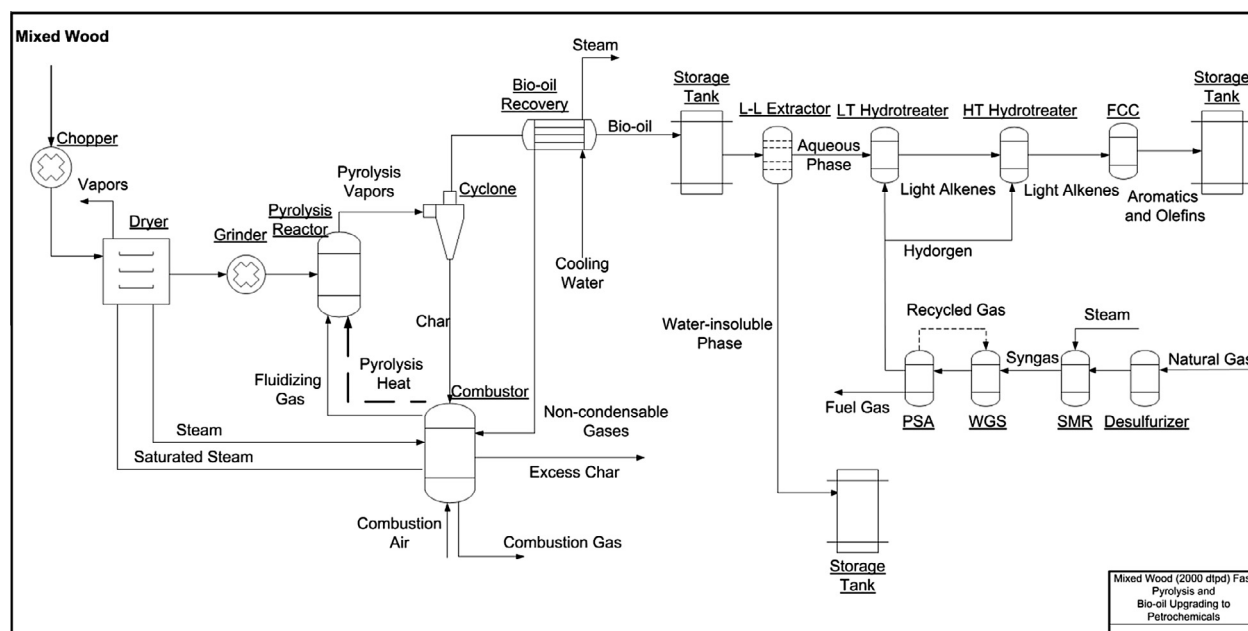


Fig. 1. Process diagram for mixed wood fast pyrolysis and bio-oil upgrading to commodity chemicals (Adapted from Zhang et al. (2013b)).

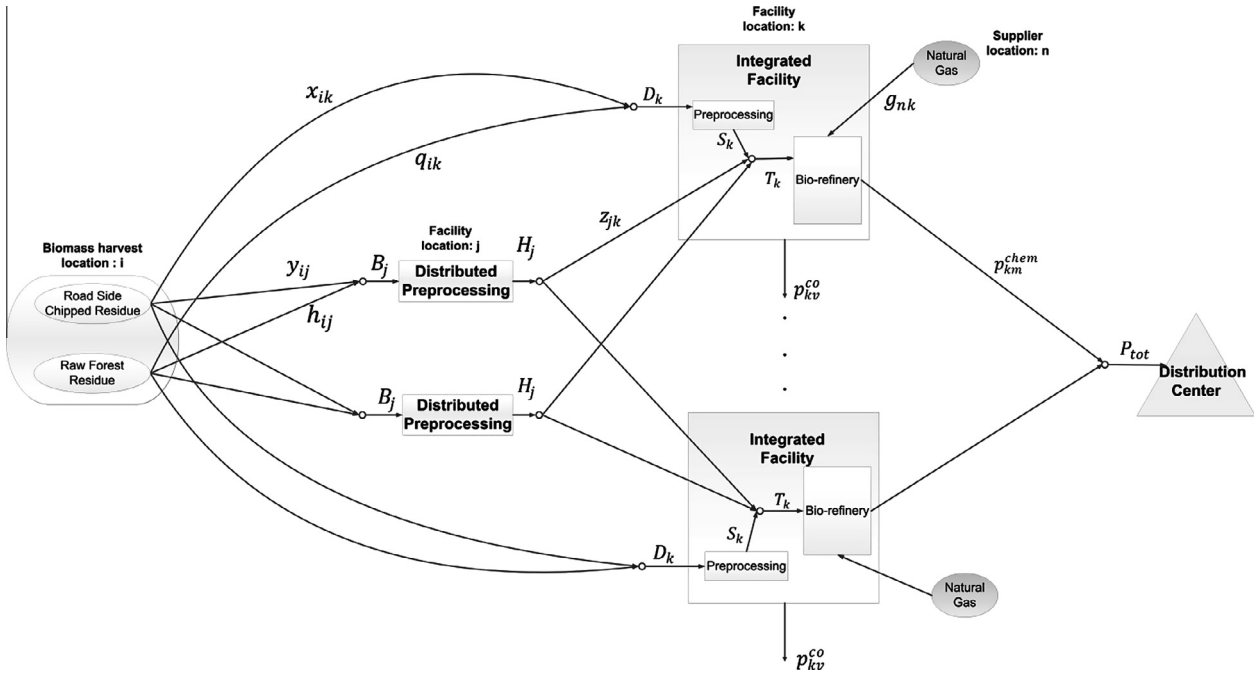


Fig. 2. Supply chain schematic for chemicals production via woody biomass fast pyrolysis. Note: This figure is a schematic diagram of the biomass flows; it does not represent an actual number of facilities.

Hydrogen is produced through the steam reforming of natural gas. Natural gas usually contains sulfur, so the gas goes through a desulfurizer for purification before entering the steam methane reformers and water gas shift reactors. The produced hydrogen is then separated from the syngas and sent to the hydrotreaters.

2.2. Supply chain model description

In this paper, the optimal plant sizes, locations, biomass and product flows are considered as the decision variables for the integrated supply chain design. Table A1 (in the appendix) shows descriptions for decision variables and the parameters for the economic and environmental objectives.

Fig. 2 illustrates the supply chain network schematics for chemicals production via woody biomass fast pyrolysis and upgrading. First, the woody biomass is harvested and collected from location i . Two types of woody biomass are considered: raw forest residue and the residue chipped with a road-side chipping method. Both woody biomass types need to be preprocessed for size and moisture reduction before conversion. For biomass preprocessing, two methods are considered. One method is distributed preprocessing, where multiple preprocessing centers are located close to biomass sources. The other method is integrated preprocessing, where the biomass is gathered into one integrated facility. The integrated facility has a preprocessing facility and the biorefinery facility. All the preprocessing facilities are to chop, dry, and grind the biomass to reduce the moisture and sizes. Then the preprocessed biomass is sent to the biorefinery facilities. Chemicals and co-products are produced at the integrated facility location. The co-products are char and lignin which are left at the local location and the chemicals are transported to the distribution center.

2.3. Model formulation

2.3.1. Economic objective

The economic objective is to maximize the net present profit for a 20-year project producing commodity chemicals via woody biomass fast pyrolysis and upgrading:

$$\max \text{PROFIT} = \sum_{t=1}^{ny} \varphi_t (\text{COST}_t^{\text{Revenue}} - \text{COST}^{\text{VarOper}} - \text{COST}^{\text{FixOper}} - \text{COST}^{\text{Collect}} - \text{COST}^{\text{Trans}}) - \text{COST}^{\text{Capital}} \quad (1)$$

The PROFIT is a function of the annual revenue ($\text{COST}_t^{\text{Revenue}}$), annual variable operating cost ($\text{COST}^{\text{VarOper}}$), annual fixed operating cost ($\text{COST}^{\text{FixOper}}$), annual biomass collection cost ($\text{COST}^{\text{Collect}}$), annual biomass transportation cost ($\text{COST}^{\text{Trans}}$), the plant capital cost ($\text{COST}^{\text{Capital}}$) and the discount factor (φ_t).

The discounted factor φ_t is used to calculate the net present. Annual interest r is assumed to be 10% for the 20-year project (2011–2032). The discount factor is shown below:

$$\varphi_t = \frac{1}{(1+r)^{t-1}} \quad (2)$$

The annual revenue $\text{COST}_t^{\text{Revenue}}$ is the sum of the revenue from chemicals product and the revenue from the co-products at individual plant location in year t as described in Eq. (3). The annual revenue is not same for every year since the selling price of the chemical product m in year t (c_{mt}^{chem}) is assumed to be changing every year. The prices of chemicals throughout the years are predicted based on EIA petroleum price prediction.

$$\text{COST}_t^{\text{Revenue}} = \sum_{k=1}^K \sum_{v=1}^V p_{kv}^{\text{co}} c_{kv}^{\text{co}} + \sum_{k=1}^K \sum_{m=1}^M p_{km}^{\text{chem}} c_{km}^{\text{chem}} \quad (3)$$

The annual variable operating cost $\text{COST}^{\text{VarOper}}$ is a sum of variable operating costs for the distributed preprocessing facilities, integrated preprocessing facilities, and integrated biorefinery facilities, which is shown in Eq. (4). The variable operating costs include the costs for plant operation, such as electricity, process water, and catalysts.

$$\text{COST}^{\text{VarOper}} = \sum_{k=1}^K T_k C_k^{\text{VCenB}} + \sum_{i=1}^I \sum_{j=1}^J h_{ij} C_{ij}^{\text{VPre}} + \sum_{i=1}^I \sum_{j=1}^J y_{ij} C_{ij}^{\text{VPre2}} + \sum_{i=1}^I \sum_{k=1}^K q_{ik} C_{ik}^{\text{VPre}} + \sum_{i=1}^I \sum_{k=1}^K x_{ik} C_{ik}^{\text{VPre2}} \quad (4)$$

The annual fixed operating cost $COST^{FixOper}$ is defined by Eq. (5). The fixed operating cost includes the salaries, overhead, and maintenance costs for the distributed preprocessing facilities and integrated facilities.

$$COST^{FixOper} = \sum_{j=1}^J \sum_{l=1}^L r_{ij} C_l^{FD} + \sum_{k=1}^K \sum_{l=1}^L u_{lk} C_l^{FC} \quad (5)$$

The annual biomass collection cost $COST^{Collect}$ is the sum of collection costs for raw biomass and roadside chipped biomass given in Eq. (6).

$$COST^{Collect} = \sum_{i=1}^I \sum_{j=1}^J h_{ij} C_i^{CR} + \sum_{i=1}^I \sum_{k=1}^K q_{ik} C_i^{CR} + \sum_{i=1}^I \sum_{j=1}^J y_{ij} C_i^{CC} + \sum_{i=1}^I \sum_{k=1}^K x_{ik} C_i^{CC} \quad (6)$$

The annual biomass transportation cost $COST^{Trans}$ includes the transportation costs of all of the materials (biomass, chemicals, and natural gas), as shown in Eq. (7).

$$COST^{Trans} = \sum_{i=1}^I \sum_{j=1}^J h_{ij} C^{TR} \tau_{dij} + \sum_{i=1}^I \sum_{k=1}^K q_{ik} C^{TR} \tau_{dik} + \sum_{i=1}^I \sum_{j=1}^J y_{ij} C^{TC} \tau_{dij} + \sum_{i=1}^I \sum_{k=1}^K x_{ik} C^{TC} \tau_{dik} + \sum_{j=1}^J \sum_{k=1}^K z_{jk} (C^{TPre} \tau_{djk} + C^{DFC}) + \sum_{n=1}^N \sum_{k=1}^K g_{nk} C^{TN} \tau_{dnk} + \sum_{k=1}^K P_k C^{TP} \tau_{dk}^{DC} \quad (7)$$

The plant capital cost, the sum of capital investment for all of the facilities, is assumed to be invested in the current year, so the discount factor is not applied (see Eq. (8)).

$$COST^{Capital} = \left(\sum_{j=1}^J \sum_{l=1}^L r_{ij} C_l^{CapID} + \sum_{k=1}^K \sum_{l=1}^L u_{lk} C_l^{CapIC} \right) \quad (8)$$

2.3.2. Environmental objective

The environmental objective for GHG-emissions minimization is defined as follows:

$$\min E_{TOT} = E^{Collection} + E^{Transport} + E^{Production}$$

where:

$$E^{Collect} = \sum_{i=1}^I \sum_{j=1}^J h_{ij} e_i^{CR} + \sum_{i=1}^I \sum_{k=1}^K q_{ik} e_i^{CR} + \sum_{i=1}^I \sum_{j=1}^J y_{ij} e_i^{CC} + \sum_{i=1}^I \sum_{k=1}^K x_{ik} e_i^{CC} \quad (9)$$

$$E^{Trans} = \sum_{i=1}^I \sum_{j=1}^J h_{ij} e^{TR} \tau_{dij} + \sum_{i=1}^I \sum_{k=1}^K q_{ik} e^{TR} \tau_{dik} + \sum_{i=1}^I \sum_{j=1}^J y_{ij} e^{TC} \tau_{dij} + \sum_{i=1}^I \sum_{k=1}^K x_{ik} e^{TC} \tau_{dik} + \sum_{j=1}^J \sum_{k=1}^K z_{jk} e^{TPre} \tau_{djk} + \sum_{k=1}^K P_k e^{TP} \tau_{dk}^{DC} + \sum_{n=1}^N \sum_{k=1}^K g_{nk} e^{TN} \tau_{dnk} \quad (10)$$

$$E^{Production} = \sum_{k=1}^K T_k e^{CenB} + \sum_{i=1}^I \sum_{j=1}^J h_{ij} e^{Pre} + \sum_{i=1}^I \sum_{j=1}^J y_{ij} e^{Pre2} + \sum_{i=1}^I \sum_{k=1}^K q_{ik} e^{Pre} + \sum_{i=1}^I \sum_{k=1}^K x_{ik} e^{Pre2} \quad (11)$$

In Eq. (9), $E^{Collect}$ is the CO₂-equivalent GHG emissions associated with the biomass collection processes. Here e_i^{CR} is the emission of a collection-unit amount of raw biomass from harvest site i , and e_i^{CC}

is the emission of a collection-unit amount of roadside chipped biomass. In Eq. (10), E^{Trans} is the CO₂-equivalent GHG emissions associated with the materials transportation processes. The term e^{TR} is the emission of a transporting-unit amount of raw biomass, e^{TC} is the emission of a transporting-unit amount of roadside chipped biomass, and e^{TPre} is the emission of a transporting-unit amount of preprocessed biomass. The term e^{TN} is the emission of a transporting unit amount of natural gas, and e^{TP} is the emission of a transporting unit amount of chemicals. In Eq. (11), $E^{Production}$ is the emissions associated with the biomass conversion processes. Here e^{Pre} is the emission of raw biomass preprocessing process, e^{Pre2} is the emission of roadside chipped biomass preprocessing process and e^{CenB} is the emission of a converting unit amount of preprocessed biomass at biorefinery facility location k .

2.3.3. Biomass supply constraints

In this section, the mass balance of biomass flows and facility capacities constraints are included. The total collected biomass F_i should not exceed the total biomass allowed for collection in harvesting location i . In Eq. (12), α is the sustainability factor, which illustrates the allowed collection percentage of the available biomass.

$$F_i \leq \alpha A_i, \forall i \quad (12)$$

The total collected biomass F_i can be categorized into two types: raw biomass and roadside chipped biomass. They both can be transported to either the distributed preprocessing facility location j or the integrated facility location k . In Eqs. (13)–(15), y_{ij} and h_{ij} are the amount of transported raw biomass and roadside chipped biomass from harvest location i to distributed preprocessing location j . x_{ik} and q_{ik} are the amount of raw biomass and roadside chipped biomass transported from harvest location i to integrated facility location k . ϵ is the loss factor for the biomass transportation process. B_j is the total received biomass (raw biomass and roadside chipped biomass) in distributed preprocessing facility location j , and D_k is the total biomass (raw biomass and roadside chipped biomass) received in integrated preprocessing facility location k .

$$\sum_{j=1}^J (y_{ij} + h_{ij}) + \sum_{k=1}^K (x_{ik} + q_{ik}) = F_i (1 - \epsilon), \forall i \quad (13)$$

$$\sum_{i=1}^I (y_{ij} + h_{ij}) = B_j, \forall j \quad (14)$$

$$\sum_{i=1}^I (x_{ik} + q_{ik}) = D_k, \forall k \quad (15)$$

2.3.4. Distributed preprocessing facility constraints

The distributed preprocessing facility constraints are shown in Eqs. (16)–(20).

$$B_j \leq \sum_{l=1}^L Cap_l^{Dis} r_{ij}, \forall j \quad (16)$$

For each candidate location j , there is at most one facility with capacity level l .

$$\sum_{l=1}^L r_{ij} \leq 1, \forall j \quad (17)$$

The total number of distributed preprocessing facilities at location j with capacity level l should not exceed the maximum number Num^{Dis} .

$$\sum_{l=1}^L \sum_{j=1}^J f_{lj} \leq \text{Num}^{\text{Dis}}, \forall j \quad (18)$$

For Eqs. (8) and (9), the received biomass B_j is preprocessed with a yield β of H_j at distributed preprocessing facility location j and then H_j is transported to the integrated biorefinery location k .

$$B_j \beta = H_j, \forall j \quad (19)$$

$$\sum_{k=1}^K z_{jk} = H_j, \forall j \quad (20)$$

2.3.5. Integrated facility constraints

The total biomass (raw biomass and roadside chipped biomass) received in integrated preprocessing facility location k is presented as D_k . As indicated in Eq. (21), the received biomass is preprocessed to dry biomass with a yield β of S_k at location k . The total preprocessed biomass T_k is the sum of preprocessed biomass from integrated preprocessing facility S_k and that from distributed preprocessing facility $\sum_{j=1}^J z_{jk}$, as described in Eq. (22). In the integrated biorefinery facility location k , the preprocessed biomass T_k is converted to various chemicals, as shown in Eq. (23). μ_m is the conversion rate for specific chemical m , and p_{km}^{chem} is the production quantity of chemical m at location k . In addition to the chemicals, the co-products pyrolytic lignin and biochar are produced at integrated biorefinery facility. In Eq. (24), γ_v is the conversion rate for co-products and p_{kv}^{co} is the production of co-product v at location k .

$$D_k \beta = S_k, \forall k \quad (21)$$

$$\sum_{j=1}^J z_{jk} + S_k = T_k, \forall k \quad (22)$$

$$T_k \mu_m = p_{km}^{\text{chem}}, \forall k \quad (23)$$

$$T_k \gamma_v = p_{kv}^{\text{co}}, \forall k, \forall v \quad (24)$$

The total preprocessed biomass for the integrated biorefinery facility at location k should not exceed the capacity of the integrated biorefinery $\text{Cap}_l^{\text{Cen}}$ if facility level l is selected (Eq. (25)). At most one facility can exist in one location as indicated in Eq. (26). In Eq. (27), the total number of facilities should not exceed the maximum facility number Num^{Cen} due to budget constraints.

$$T_k \leq \sum_l u_{lk} \text{Cap}_l^{\text{Cen}}, \forall k \quad (25)$$

$$\sum_{l=1}^L u_{lk} \leq 1, \forall k \quad (26)$$

$$\sum_l \sum_k u_{lk} \leq \text{Num}^{\text{Cen}} \quad (27)$$

2.3.6. Natural gas and chemicals constraints

In Eq. (28), the total natural gas demand at the biorefinery locations is the sum of natural gas flows from various natural gas suppliers. The supplied natural gas from location n to all biorefinery facilities should not exceed the available natural gas in supply location n as indicated in Eq. (29). In Eq. (30), natural gas demand is calculated as a factor θ of the preprocessed total biomass T_k at biorefinery location k . The total chemicals production P_{tot} (Eq. (31)) is the sum of all types of chemical m produced from all of the integrated facility locations. In Eq. (32), the total chemicals production should not exceed the maximum chemicals demand.

$$\sum_{n=1}^N g_{nk} = Q_k, \forall k \quad (28)$$

$$\sum_{k=1}^K g_{nk} \leq \text{NG}_n, \forall n \quad (29)$$

$$Q_k = \theta T_k, \forall k \quad (30)$$

$$\sum_{k=1}^K \sum_{m=1}^M p_{km}^{\text{chem}} = P_{\text{tot}} \quad (31)$$

$$P_{\text{tot}} \leq T^{\text{max}} \quad (32)$$

$$x_{ik}, q_{ik}, y_{ij}, h_{ij}, z_{jk}, g_{nk}, \geq 0, r_{ij}, u_{lk} \in \{0, 1\}, \forall i, j, k, n, l, m, v, t \quad (33)$$

3. Results and discussion

3.1. Data sources

In this case study, forest residue is the feedstock and the state of Minnesota is employed. It should be noted that this case study is a hypothetical scenario. The facilities are not being built or proposed. We chose the state of Minnesota due to the forest residue availability. This decision making framework can be applied to other region of interests with minor modifications. The amount of available forest residue is obtained from the National Renewable Energy Laboratory (NREL, 2013). Each county in Minnesota is considered as a candidate harvesting site, a potential distributed preprocessing facility location, and potential integrated facility location. The Minneapolis-St. Paul metro area has the most convenient transportation resources; therefore, Minneapolis-St. Paul is selected to be the distribution center. The chemicals demand data are based on the commodity flow survey for Minnesota (BTS, 2007). All of the chemicals are assumed to be transported to the distribution center in Minneapolis-St. Paul. The information about the natural gas suppliers and their gas availability is obtained from the U.S. Energy Information Administration (EIA, 2011).

Five capacity levels ($L1, L2, L3, L4$, and $L5$) are considered for distributed preprocessing and integrated facilities; $L1, L2, L3, L4$, and $L5$ correspond to 100, 200, 500, 1000, and 2000 metric ton/day dry biomass processing capacities. The 2000 metric ton/day capacity plant is selected as the reference plant and the bio-oil conversion rate is assumed to be 52 wt.% of dry biomass. The capital costs for the distributed and integrated facilities are based on the techno-economic analysis (Brown et al., 2012; Zhang et al., 2013b). A scale factor of $n = 0.6$ is employed to estimate the capital costs. In Eq. (34), S_{new} and S_0 represent the new plant size and the reference plant size, and C_{new} and C_0 are the capital costs for the new plant and the reference plant.

$$C_{\text{new}} = C_0 \left(\frac{S_{\text{new}}}{S_0} \right)^n \quad (34)$$

For biomass preprocessing, two methods are considered. One is to preprocess biomass in distributed preprocessing facilities and the other is to preprocess biomass in integrated preprocessing facilities. Table A2 (in the Appendix) details the capital costs and the fixed operating cost for the distributed preprocessing facility and for the integrated preprocessing and biorefinery facility at various levels (Zhang et al., 2013b). The fixed operating cost includes salaries, overhead, maintenance, and insurance. The maintenance fees are assumed to be 6% of the facility capital cost. The overhead and insurance are assumed to be 2% and 1.5% of the total salaries, respectively.

The variable costs for the distributed preprocessing facility and integrated preprocessing facility for same biomass are assumed to be the same. But for raw biomass and roadside chipped biomass preprocessing, variable operating costs are different. For roadside chipped biomass preprocessing process, the variable operating cost does not include chopping cost in Table A3 (in the Appendix). For the environmental impact assessment, all GHG emissions related to biomass collection, materials transportation, and production processes are based on the Aspen Plus, SimaPro and GREET models (Zhang et al., 2013b,c). The emission for the distributed preprocessing facility and integrated preprocessing facility for same biomass are assumed to be the same. But raw biomass preprocessing and roadside chipped biomass preprocessing have different emissions. The variable operating costs for facilities and emissions data are derived from the reference plant data (Table A3) (Zhang et al., 2013b,c).

The 2012–2035 chemicals prices are based on the technoeconomic analysis (Zhang et al., 2013b). The correlations between each chemical species and petroleum price are used to calculate the prices for the next 20 years. The chemical yield and market prices for the next 20 years are shown in Table A4 (in the Appendix). The co-products yields include char, pyrolytic lignin, and fuel gas. The prices of the co-products are \$18.21, \$22.05, and \$200 per metric ton for char, pyrolytic, and fuel gas respectively.

The collection costs for raw biomass and roadside chipped biomass are based on Leinonen (2004). Forest haulage cost is \$9.8/ton for raw forest residue. The stumpage price for the forest residue is assumed to be \$5/metric ton. So the collection cost for raw biomass is \$15.8 /metric ton. For roadside chipped forest residue, there is a \$9.8/ton haulage cost, \$9.8/ton chipping cost, and stumpage cost of \$5/metric ton. Therefore, the collection cost for roadside chipped forest residue is \$26.6/metric ton.

The costs of the harvesting methods of forest residues also have been reported by Leinonen (2004). The four harvest methods include bundle, terrain chip, road chip, and plant chip. The road transportation costs for raw forest residue and roadside chipped forest residue are \$12.8/ton and \$18.3/ton for 80 km. As calculated,

the variable transportation costs for raw forest residue and roadside chipped forest residue are assumed to be \$0.41/metric-ton-mile and \$0.28/metric-ton-mile. The preprocessed forest residue is transported by the trucks with a fixed transportation cost of \$3.32/metric ton for wood chips loading and unloading and a variable transportation cost of \$0.124/metric-ton-mile (Searcy et al., 2007). The transportation cost of commodity chemicals is assumed to be same as the national average truck shipping cost of \$0.286/metric-ton-mile (BTS, 2012). The transportation cost of natural gas via pipeline is assumed to be same as the national average oil pipeline cost of \$0.0297/metric-ton-mile (BTS, 2012). The distances between counties are based on the great circle distances calculated based on the latitudes and longitudes. Circuitry factors are incorporated to estimate the actual transportation distances. The circuitry factors are assumed to be 1.27 (Rogers and Brammer, 2009) and 1.1 for truck and pipeline, respectively (CBO, 1982).

This model employs MATLAB to collect the data and uses geographic information system (GIS) software to map the biomass availability and locations. The mathematical model is coded in GUSEK and solved with Gurobi.

3.2. Results and analysis for economic objective model

The economic objective model is developed to determine the economic feasibility and optimal capacities and locations of the distributed preprocessing facilities and integrated facilities in Minnesota by maximizing the net present profit for a 20-year project. The northern Minnesota has the most abundant forest residue sources, especially in Lake, Itasca, St. Louis, Koochiching, Cass, Aitkin, Hubbard, Clearwater, and Beltrami Counties. The forest residue in those nine counties represents 70% of the total forest residue in Minnesota. Among these counties, St. Louis County has the largest amount of forest residue, representing approximately 19% of the total forest residue in Minnesota. The optimal locations for the distributed preprocessing, integrated facilities, and natural gas suppliers locations are illustrated in Fig. 3b. The results predict that three integrated facilities (include the preprocess facility and biorefinery

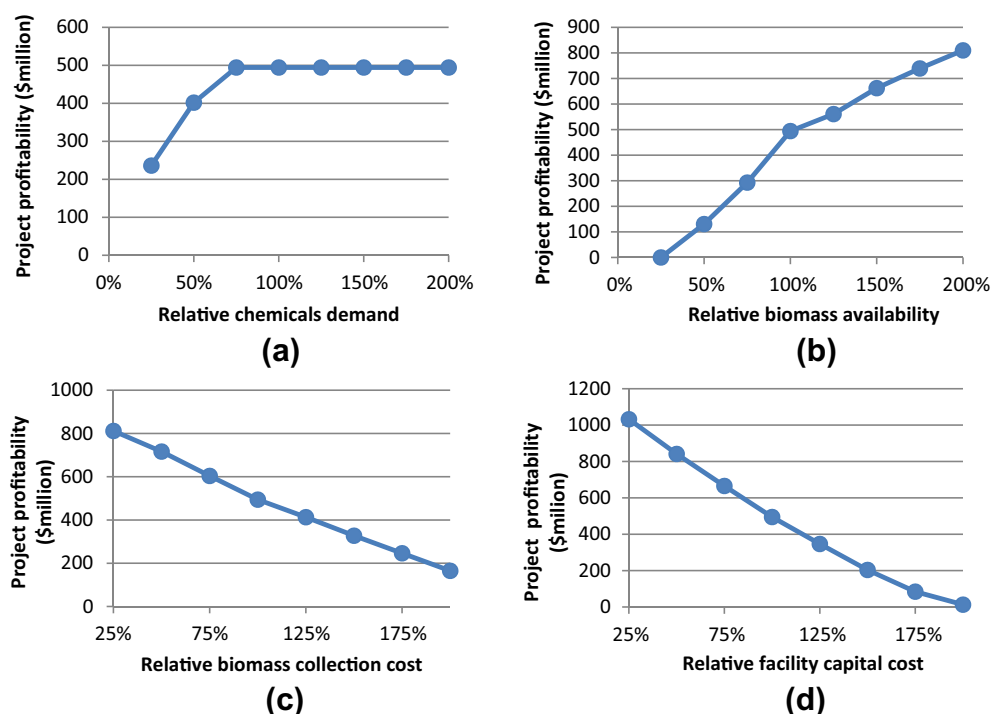


Fig. 3. Effects of chemicals demand, biomass availability, biomass collection cost and facility capital cost on project profitability.

Table 1
Mass flows for the distributed preprocessing facilities.

Biomass	Location1	Location2	Mass flow (metric ton/year)
h	Aitkin	Aitkin	113965.8
h	Carlton	Aitkin	51866.1
h	Crow wing	Aitkin	50287.87
h	Morrison	Aitkin	11498.38
h	Pine	Aitkin	46521.15
h	Beltrami	Beltrami	171182.7
h	Clearwater	Beltrami	123937.1
y	Lake of the Woods	Beltrami	47419.13
h	Mahnomen	Beltrami	8135.4
y	Marshall	Beltrami	19046.63
h	Pennington	Beltrami	1570.725
y	Polk	Beltrami	39340.28
h	Red lake	Beltrami	810.225
y	Roseau	Beltrami	30625.72
h	Itasca	Itasca	248718.6
h	Koochiching	Itasca	8882.725
h	Koochiching	Koochiching	274139.3
h	Lake	Lake	274139.3

Note: *h* represents the raw biomass transported from harvest site to distributed preprocessing location; *y* represents the roadside chipped biomass transported from harvest site to distributed preprocessing location.

facility) and five distributed preprocessing facilities would be built in the state of Minnesota.

Based on the optimization model, two of the integrated facilities with the highest capacity level (*L5*, 2000 metric ton/day) should be built in St. Louis County and Cass County. Although Dakota County contains only 6% of the forest residue in Minnesota, it is located very near Minneapolis-St Paul and thus reduces the transportation costs significantly. Therefore based on the model, an integrated facility is to be built in Dakota with capacity *L4* (1000 metric ton/day). The five distributed preprocessing facilities are modeled to be built in Aitkin (*L3*), Beltrami (*L4*), Itasca (*L3*), Koochiching (*L3*), and Lake (*L3*) Counties. These facilities are in the counties rich in forest residue for convenient collection of biomass. The biomass mass flows for the distributed preprocessing facilities and their locations are shown in Table 1.

The main biomass mass flows to the three integrated facilities (integrated preprocessing and biorefinery) are shown in Table 2. Most of the biomass arriving at the St. Louis County or Cass County integrated preprocessing facilities is raw biomass or preprocessed biomass from distributed preprocessing facilities. The third integrated facility located in Dakota County, receives raw biomass and roadside chipped biomass from nearby biomass harvest sites and preprocessing biomass from Aitkin County. The raw biomass is preprocessed and converted to commodity chemicals at the integrated facility. Each integrated facility has a natural gas supplier nearby.

The facility capital cost is the largest expenditure, representing 33% of the total cost. The production cost accounts for 30% of the total cost, which includes the fixed operating cost (19.4%) and the variable operating cost (10.3%). The remainder of the cost comes from the biomass collection and transportation cost, which are 18.9% and 18%, respectively. The transportation cost includes the costs of transporting the biomass, commodity chemicals, and the natural gas. The biomass transportation is the largest among them, representing 13.9% of the total cost.

3.3. Factors influencing project profitability

Fig. 3 illustrate the impacts of variable factors on the project profitability. The commodity chemicals demand is directly related to project revenues. Fig. 3(a) shows the impact of chemicals demand on project profitability. Here the relative chemicals demand in x-axis represents ratio of chemical demand to baseline demand. The profitability increases directly with increasing chemicals

Table 2
Main mass flows for the integrated facilities.

Biomass	Location 1	Location 2	Mass flow (metric ton/year)
x	Becker	Cass	16231.6
q	Cass	Cass	114896
q	Crow wing	Cass	10833.9
q	Hubbard	Cass	108658.9
q	Wadena	Cass	23519
z	Aitkin	Cass	80671.5
z	Beltrami	Cass	285133.8
z	Cass	Cass	176819.8
z	Itasca	Cass	164724.8
q	Anoka	Dakota	2674.4
x	Blue Earth	Dakota	3860
q	Chisago	Dakota	2499.9
q	Dakota	Dakota	164281.6
x	Fillmore	Dakota	29177.9
q	Goodhue	Dakota	72581.9
q	Hennepin	Dakota	2771
x	Houston	Dakota	22204.7
x	Kanabec	Dakota	11933
x	Mille lacs	Dakota	11931
x	Morrison	Dakota	4305.4
q	Olmsted	Dakota	8146.1
x	Renville	Dakota	12757.9
x	Sherburne	Dakota	3118.05
q	Wabasha	Dakota	17722.6
x	Winona	Dakota	12728.6
z	Aitkin	Dakota	96148.3
z	Dakota	Dakota	257526.7
q	Lake	St. Louis	6049.4
q	St. Louis	St. Louis	540124.7
z	Koochiching	St. Louis	176819.8
z	Lake	St. Louis	176819.8
z	St. Louis	St. Louis	352282.2

Note: *q* means the transported raw biomass from harvest site to integrated preprocessing location; *x* means the road side chipped biomass from harvest site to integrated preprocessing location; *z* means the preprocessed biomass from distributed preprocessed location to the integrated biorefinery location.

demand from 25% to 75% of the current production. Profitability increases to \$494 million when chemicals demand reaches 75% of the baseline. After that, profitability stays constant even as the chemicals demand increases. This is because the forest residue is not sufficient to achieve the largest profitability possible when the chemicals demand is 75% of the baseline. So even when the chemicals demand increases, the biomass supply is not sufficient to meet the demand.

Based on the analysis of impact of chemicals demand variation on total profitability, it is illustrated that the biomass availability plays a significant role in the total profitability. Seasonal and other factors (competition of biomass etc.) cause variation in biomass availability and thus lead to different optimal solutions. Fig. 3(b) illustrates the effect of variation of biomass availability on the project profitability. Here the relative biomass availability in the x-axis means ratio of biomass availability to baseline availability. The project profitability increases as the biomass availability increases. As discussed, the project profitability is limited by insufficient biomass. When there is an increase in biomass availability, the project profitability will increase significantly.

The competition for the feedstock will lead to increasing forest residue price. Biomass collection cost is an important parameter for the project profitability, representing 18.9% of the total capital cost. The effect of variation in biomass collection cost on project profitability is analyzed in Fig. 3(c). Here the relative biomass collection cost in x-axis is ratio of biomass of collection cost with respect to the baseline biomass collection cost. It is illustrated that when the biomass collection cost is reduced to 25% of the baseline, the maximum profitability for the project increases to \$812 million. The profitability decreases to just \$165 million when the biomass collection cost is twice the baseline cost.

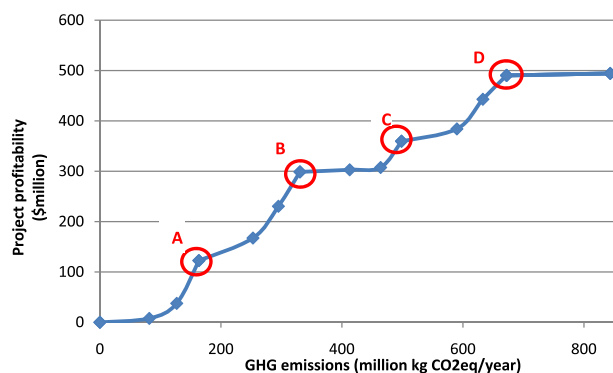


Fig. 4. Pareto curve for the economic–environmental multi-objective optimization for supply chain of commodity chemicals production via woody biomass fast pyrolysis.

Facility capital cost is the largest contributor to project profitability. As indicated in Fig. 3(d), if the facility cost double from the baseline, the profit will drop to zero. The relative facility capital cost in x-axis is ratio of facility capital cost with respect to the baseline.

3.4. Results and analysis for the economic–environmental multi-objective model

The multi-objective model is formulated to analyze the trade-off between minimizing GHG emissions and maximizing project profits. The ϵ -constraint method is used to solve this multi-objective problem. The Pareto curve generated by all the optimal solutions is shown in Fig. 4. The GHG emissions reduce from 843 million kg CO₂eq per year to zero while the total 20-year profitability decreases from 494 million dollars to zero.

In the Pareto curve, there is one integrated facility with capability L4 (1000 metric ton/day) in St. Louis for Point A. For point B, two distributed preprocessed facilities with capacity L3 are proposed in Koochiching and Lake. One integrated facility is proposed with capacity of L5 level (2000 metric ton/day) in St. Louis. The emissions for Point B are 331 million kg CO₂ eq/year and the profitability is \$300 million. From point A to point B, the optimal solution includes only one integrated facility (built in St. Louis County). After point B, other integrated facilities are proposed in addition to the integrated facility in St. Louis County. For points C, two integrated facilities are proposed which are located in St. Louis (L5, 2000 metric ton/day) and Beltrami (L4, 1000 metric ton/day) and the distributed preprocessing facilities are proposed in Itasca, Lake and Koochiching. Point D is a good point where two integrated facilities and five distributed preprocessing facilities are proposed. The two integrated facilities are located in St. Louis (L5, 2000 metric ton/day) and Case (L5, 2000 metric ton/day). The five distributed preprocessing facilities are located in Itasca, Lake, Koochiching, Aitkin, and Beltrami.

The Pareto curve illustrates the trade-offs between economics and environmental effects. When the production capacity is comparatively small, the profitability grows fast with a small increase of GHG emissions. After a certain production capacity (point B), however, the profitability grows much slowly. From point A to point D, the optimal integrated facility locations always include St. Louis County, which indicates that St. Louis County is the most favorable location to build the integrated chemicals plant.

4. Conclusion

This work investigates the economic feasibility and the optimal production planning and facility locations for commodity

chemicals production via woody biomass fast pyrolysis. The economic objective model results show that the distributed facilities biomass chipping is preferable to the roadside chipping method for forest residue. The harvest sites rich in biomass resources are the preferable locations for building biorefinery facilities. Influences of parameters on economic objective model show that the biomass availability and facility capital costs are the most important factors for the project profitability. The economic–environmental multi-objective model results illustrate the trade-off between economic and environmental considerations.

Acknowledgements

The authors would like to acknowledge the financial support of the Bioeconomy Institute and the Biobased Industry Center of Iowa State University and the National Science Foundation under Grant Number EPS-1101284.

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.biortech.2014.01.049>.

References

- An, H., Wilhelm, W.E., Searcy, S.W., 2011. A mathematical model to design a lignocellulosic biofuel supply chain system with a case study based on a region in Central Texas. *Bioresour. Technol.* 102, 7860–7870.
- Bai, Y., Hwang, T., Kang, S., Ouyang, Y., 2011. Biofuel refinery location and supply chain planning under traffic congestion. *Transp. Res. B Method.* 45, 162–175.
- Blottnitz, H.V., Curran, M.A., 2007. A review of assessments conducted on bio-ethanol as a transportation fuel from a net energy, greenhouse gas, and environmental life cycle perspective. *J. Cleaner Prod.* 15, 607–619.
- Bowling, I.M., Ponce-Ortega, J.M., El-Halwagi, M.M., 2011. Facility location and supply chain optimization for a biorefinery. *Ind. Eng. Chem. Res.* 50, 6276–6286.
- Brehmer, B., Boom, R.M., Sanders, J., 2009. Maximum fossil fuel feedstock replacement potential of petrochemicals via biorefineries. *Chem. Eng. Res. Des.* 87, 1103–1119.
- Brown, T.R., Zhang, Y., Hu, G., Brown, R.C., 2012. Techno-economic analysis of biobased chemicals production via integrated catalytic processing. *Biofuels, Bioprod. Biorefin.* 6, 73–87.
- BTS, 2012. Average Freight Revenue per Ton-mile. Bureau of Transportation Statistics. Available from: <http://www.bts.gov/publications/national_transportation_statistics/html/table_03_21.html> (accessed November, 2012).
- BTS, 2007. Commodity Flow Survey 2007. Table 12. Shipment Characteristics by NAICS and Commodity for Metropolitan Area of Origin: 2007. Available from: <http://www.rita.dot.gov/bts/sites/rita.dot.gov/bts/files/publications/commodity_flow_survey/2007/metropolitan_areas/minneapolis_st_paul_st_cloud_mn_wi_csa_mn_part/index.html>.
- CBO, 1982. Energy Use in Freight Transportation. Congressional Budget Office.
- Christensen, C.H., Rass-Hansen, J., Marsden, C.C., Taarning, E., Egeblad, K., 2008. The renewable chemicals industry. *ChemSusChem* 1, 283–289.
- Dale, B.E., 2003. ‘Greening’ the chemical industry: research and development priorities for biobased industrial products. *J. Chem. Technol. Biotechnol.* 78, 1093–1103.
- Dunnett, A., Adjiman, C., Shah, N., 2008. A spatially explicit whole-system model of the lignocellulosic bioethanol supply chain: an assessment of decentralised processing potential. *Biotechnol. Biofuels* 1, 1–17.
- EIA, 2011. Natural Gas Annual Respondent Query System. Available from: <http://www.eia.gov/cfapps/ngqs/ngqs.cfm?f_report=RP7>.
- Elia, J.A., Baliban, R.C., Floudas, C.A., Gurau, B., Weingarten, M.B., Klotz, S.D., 2013. Hardwood biomass to gasoline, diesel, and jet fuel: 2. supply chain optimization framework for a network of thermochemical refineries. *Energy Fuels* 27, 4325–4352.
- Gavrilescu, M., Chisti, Y., 2005. Biotechnology—a sustainable alternative for chemical industry. *Biotechnol. Adv.* 23, 471–499.
- Giarola, S., Shah, N., Bezzo, F., 2012. A comprehensive approach to the design of ethanol supply chains including carbon trading effects. *Bioresour. Technol.* 107, 175–185.
- Hamelinck, C.N., Hooijdonk, G.v., Faaij, A.P.C., 2005. Ethanol from lignocellulosic biomass: techno-economic performance in short-, middle- and long-term. *Biomass Bioenergy* 28, 384–410.
- Hess, J.R., Wright, C.T., Kenney, K.L., 2007. Cellulosic biomass feedstocks and logistics for ethanol production. *Biofuels, Bioprod. Biorefin.* 1, 181–190.

- Kim, J., Realff, M.J., Lee, J.H., 2011. Optimal design and global sensitivity analysis of biomass supply chain networks for biofuels under uncertainty. *Comput. Chem. Eng.* 35, 1738–1751.
- Larson, E.D., 2006. A review of life-cycle analysis studies on liquid biofuel systems for the transport sector. *Energy Sustainable Dev.* 10, 109–126.
- Leinonen, A., 2004. Harvesting Technology of Forest residues for fuel in the USA and Finland. Available from: <<http://www.vtt.fi/inf/pdf/tiedotteet/2004/T2229.pdf>>.
- NREL, 2013. Forest and Primary Mill Residues, 2013. (Available at: http://www.nrel.gov/gis/data_biomass.html).
- Oak Ridge National Laboratory (ORNL), 2007. Survey of Alternative Feedstocks for Commodity Chemical Manufacturing. Available from: <http://cepac.chemecmu.edu/pasi2011/library/cremaschi/Survey_of_alternative_feedstocks_for_the_chemical_industry.pdf>.
- Rogers, J.G., Brammer, J.G., 2009. Analysis of transport costs for energy crops for use in biomass pyrolysis plant networks. *Biomass Bioenergy* 33, 1367–1375.
- Schilling, L.B., 1995. Chemicals from alternative feedstocks in the United States. *FEMS Microbiol. Rev.* 16, 101–110.
- Searcy, E., Flynn, P., Ghafoori, E., Kumar, A., 2007. The relative cost of biomass energy transport. *Appl. Biochem. Biotechnol.* 137–140, 639–652.
- Stephen, J.D., Mabey, W.E., Saddler, J.N., 2010. Biomass logistics as a determinant of second-generation biofuel facility scale, location and technology selection. *Biofuels, Bioprod. Biorefin.* 4, 503–518.
- Tallaksen, J., 2011. Biomass Gasification: A Comprehensive Demonstration of a Community Scale Biomass Energy System. Final Report, USDA Rural Development, Grant 68-3A75-5-232. Available from: <http://renewables.morris.umn.edu/biomass/documents/USDA_Report/USDA_Main_Report.pdf>.
- Vispute, T.P., Zhang, H., Sanna, A., Xiao, R., Huber, G.W., 2010. Renewable chemical commodity feedstocks from integrated catalytic processing of pyrolysis oils. *Science* 330, 1222–1227.
- You, F., Tao, L., Graziano, D.J., Snyder, S.W., 2012. Optimal design of sustainable cellulosic biofuel supply chains: multiobjective optimization coupled with life cycle assessment and input–output analysis. *AIChE J.* 58, 1157–1180.
- You, F., Wang, B., 2011. Life cycle optimization of biomass-to-liquid supply chains with distributed-centralized processing networks. *Ind. Eng. Chem. Res.* 50, 10102–10127.
- Zhang, Y., Brown, T.R., Hu, G., Brown, R.C., 2013a. Comparative techno-economic analysis of biohydrogen production via bio-oil gasification and bio-oil reforming. *Biomass Bioenergy* 51, 99–108.
- Zhang, Y., Brown, T.R., Hu, G., Brown, R.C., 2013b. Techno-economic analysis of two bio-oil upgrading pathways. *Chem. Eng. J.* 225, 895–904.
- Zhang, Y., Hu, G., Brown, R.C. 2013c. Life Cycle Assessment of Commodity Chemicals Production from Forest Residue via Fast Pyrolysis. Technical Report, Iowa State University.
- Zhang, Y., Hu, G., Brown, R.C., 2013d. Life cycle assessment of the production of hydrogen and transportation fuels from corn stover via fast pyrolysis. *Environ. Res. Lett.* 8, 025001.