Requirement of the Type II Secretion System for Utilization of Cellulosic Substrates by *Cellvibrio japonicus* \(^*\)†‡

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Cellulosic biofuels represent a powerful alternative to petroleum but are currently limited by the inefficiencies of the conversion process. While Gram-positive and fungal organisms have been widely explored as sources of cellulases and hemicellulases for biomass degradation, Gram-negative organisms have received less experimental attention. We investigated the ability of *Cellvibrio japonicus*, a recently sequenced Gram-negative cellulolytic bacterium, to degrade bioenergy-related feedstocks. Using a newly developed biomass medium, we showed that *C. japonicus* is able to utilize corn stover and switchgrass as sole sources of carbon and energy for growth. We also developed tools for directed gene disruptions in *C. japonicus* and used this system to construct a mutant in the *gspD* gene, which is predicted to encode a component of the type II secretion system. The *gspD:pJGG1* mutant displayed a greater-than-2-fold decrease in endoglucanase secretion compared to wild-type *C. japonicus*. In addition, the mutant strain showed a pronounced growth defect in medium with biomass as a carbon source, yielding 100-fold fewer viable cells than the wild type. To test the potential of *C. japonicus* to undergo metabolic engineering, we constructed a strain able to produce small amounts of ethanol from biomass. Collectively, these data suggest that *C. japonicus* is a useful platform for biomass conversion and biofuel production.

The need for renewable alternatives to petroleum has stimulated interest in many areas of research, including the production of biofuels. Strategies for the production of ethanol from corn starch are well established, but enthusiasm for these approaches is tempered by concerns about cost, as well as indirect effects on global nutrition (38). Cellulosic biofuels represent a potentially useful alternative to starch-based ethanol, in that fuel could be produced from low-value agricultural waste products (10, 39). However, the economic viability of this approach requires overcoming the recalcitrance of plant cell walls to enzymatic degradation (55). This recalcitrance results from the crystalline nature of cellulose, as well as the complexity of plant cell walls, which are a composite of cellulose, hemicellulose, and lignin. As a result, efficient degradation of plant cell walls requires the addition of large quantities of a diverse collection of enzymes, which greatly increases the cost of biomass processing (23).

A recent analysis suggested that a combination of consolidated bioprocessing and pretreatment could significantly reduce production costs associated with cellulosic biofuels (48, 52). Consolidated bioprocessing involves the use of a single organism for the degradation of biomass to its component sugars and the subsequent conversion of these sugars to biofuel. Current approaches for the production of consolidated bioprocessors (CBPs) involve the introduction of the genes necessary for ethanol production into organisms capable of deconstructing biomass or the introduction of genes encoding biomass-degrading enzymes and their cognate secretion systems into ethanogenic microorganisms.

While *Saccharomyces cerevisiae* is arguably the most prominent industrial ethanologen, Gram-negative bacteria, such as *Zymomonas mobilis*, are also used in the commercial production of ethanol (1, 45). Furthermore, studies by the Ingram group have shown that the Gram-negative bacterium *Escherichia coli* can be engineered for efficient production of ethanol (32). The strong ethanologenic potential of these Gram-negative organisms necessitates the development of technologies for their conversion to consolidated bioprocessors. However, due to differences in cell surface architecture, unique strategies for enzyme display and secretion are required for the construction of consolidated bioprocessors in Gram-negative microorganisms. One approach to overcoming the challenges of secretion/display is to identify Gram-negative organisms that can efficiently degrade bioenergy-relevant biomass substrates. Due to the broad conservation of secretion strategies in Gram-negative microorganisms, the cellulolytic enzymes and secretion machinery from these bacteria would be expected to be transportable to ethanogenic organisms, such as *Z. mobilis* and *E. coli*. Consistent with this notion, previous studies have shown that the introduction of the type II secretion system (TTSS) from *Erwinia chrysanthemi* endowed *E. coli* with the ability to secrete heterologously expressed *E. chrysanthemi* pectinases (29).

*Cellvibrio japonicus*, originally referred to as *Pseudomonas fluorescens* var. *cellulosa* (culture no. 107, isolated in 1948 from...
soil in Saitama-ken, Japan), has long been known to be capable of cellulose degradation (51). The organism has been reported to produce an extracellular cellulase activity (22, 59), which is secreted into the culture supernatant and does not associate with the cell surface (28). Furthermore, the extracellular cellulases of C. japonicus are inducible, and their activities are increased in the presence of cellulose (35, 40, 60) and strongly downregulated when cellobiose is present in the medium (58).

An extensive literature involving the biochemical characterization of C. japonicus polysaccharide-degrading enzymes has been produced over the past 40 years, and in several cases, the corresponding genes have been cloned. Using C. japonicus genomic libraries expressed in E. coli, Wolff et al. described the isolation of four distinct carboxymethylcellulase genes (56). Subsequently, genes encoding endoglucanases, cellobextrinases, xylanases, mannanases, and an arabinofuranosidase have been cloned and characterized (5–8, 17–19, 21, 24). The isolation of four distinct carboxymethylcellulase genes (56).

### Materials and Methods

**Bacterial strains, media, and growth conditions.** C. japonicus sp. nov. strain Ueda107 was obtained from the National Collections of Industrial, Marine, and Food Bacteria. Table 1 lists the genotypes of the strains of E. coli and C. japonicus used in these studies. E. coli and C. japonicus strains were grown on M9 minimal medium (41) supplemented with MgSO4 (1 mM) and CaCl2 (0.1 mM). Glucose (0.2% [wt/vol]), cellobiose (0.34% [wt/vol]), carboxymethylcellulose (CMC) (1% [wt/vol]), and Avicel (1% [wt/vol]) were used as sole carbon and energy sources for cells grown in minimal medium. Lysogenic broth (LB) was used as rich medium (3, 4). All incubations in liquid medium were performed at 30°C with high aeration (225 rpm). For small-scale preparations, 5 ml of C. japonicus cells were grown in a 30-ml culture tube. For larger-scale preparations, 15 ml cells were grown in a 125-ml baffled flask. When used, antibiotics were present in the medium at the following concentrations: ampicillin, 100 µg/ml; gentamicin, 15 µg/ml; chloramphenicol, 25 µg/ml; and kanamycin, 50 µg/ml. All chemicals were purchased from Fisher.

**Preparation of feedstock medium.** A derivative of M9 medium was developed with corn stover or ammonia fiber expansion (AFEX)-treated corn stover (AFEX-CS) or switchgrass (AFEX-SG) as a sole carbon source. Briefly, the AFEX-CS was ground to a fine particle size (approximately 0.5 mm) with a porcelain mortar and pestle. The ground AFEX-CS then underwent a 15-min autoclave cycle with 30 min of drying. The sterile AFEX-CS was washed 5 times with sterile water before being used in the minimal medium. AFEX-treated corn stover, AFEX-treated switchgrass, and nonpretreated corn stover were added to M9 medium at a concentration of 1% (wt/vol).

**Mobilization of plasmids.** Plasmids were introduced into E. coli strains with calcium chloride-mediated transformation (41) or electroporation (14). The replicating plasmid pGG2, a derivative of pBRMCS, was mobilized into C. japonicus by conjugation. Briefly, an E. coli S17 λpir strain harboring pG2, an E. coli DH5α strain harboring helper plasmid pRK2013, and the C. japonicus recipient strain were streaked into a common region of an LB plate. After 2 days of incubation at 30°C, the cells were streaked on selective minimal medium containing M9 salts, 0.2% glucose, 15 µg/ml gentamicin, and 50 µg/ml valine (to counterselect against the E. coli strains). Exconjugants were visible after 3 to 5 days of incubation at 30°C. The resulting C. japonicus colonies were streaked twice more onto the same medium and then tested for the presence of the plasmid. Approximately 1/10th C. japonicus cells were found to harbor plasmids.

**Construction of targeted gene disruptions.** For gene disruptions in C. japonicus, plasmid pGG1, a derivative of plasmid pK18mobSacB, was introduced into S17 λpir by electroporation, selecting for kanamycin resistance. The E. coli S17 λpir strain harboring plasmid pGG1, an E. coli DH5α strain harboring helper plasmid pRK2013, and the C. japonicus recipient strain were streaked into a common region of an LB plate. After 48 h of incubation at 30°C, the cells from the LB plate were streaked on selective minimal medium containing M9 salts, 0.2% glucose, 15 µg/ml gentamicin, and 50 µg/ml valine (to counterselect against the E. coli strains). Exconjugants were visible after 3 to 5 days of incubation at 30°C. The resulting C. japonicus colonies were streaked twice onto the same medium and then tested for the presence of the plasmid by PCR.
Integration frequencies for pJGG1 in *C. japonicus* were generally low (approximately 1/10^6 *C. japonicus* cells were found to harbor integrated pJGG1).

PCR. All amplifications used *TripleMaster* polymerase (Eppendorf) and were performed in an Eppendorf Mastercycler gradient PCR thermocycler (Brinkmann Instruments). Primers were purchased from Integrated DNA Technologies. Template DNA was obtained from *C. japonicus* by adding 5 ml of an overnight LB culture directly to the PCR mixture.

Plasmid construction. Plasmid pJGG1 was constructed by amplifying an internal 500-bp fragment of the *gspD* gene using the primers 5'-GGTTGTTCTGATTCTGAGT-3' and 5'-GGTTGTTATATCCGTGGTTA-3'. The resulting PCR product was then digested with XhoI and SphI. Plasmid pJGG1 was digested with EcoRI and SphI to liberate the *gspD* and *adhB* genes, which were then ligated into pBBR1-MCS5 cut with the same enzymes. The ligation products were transformed into *E. coli* strain DH5α, selecting for kanamycin resistance on rich medium. Plasmid DNA was recovered from the kanamycin-resistant transformants, and the plasmid was purified.

Plasmid pJGG1 was constructed from plasmid pLO1295 (61), a ColEI derivative containing the *pdc* and *adhB* genes from *Z. mobilis*. Plasmid pLO1295 was digested with EcoRI and XbaI to liberate the *pdc* and *adhB* genes, which were then ligated into pBBR1-MCS5 cut with the same enzymes. The ligation products were transformed into *E. coli* strain DH5α, selecting for gentamycin resistance on rich medium. The resulting plasmid was conjugated into *C. japonicus* as described above, selecting for gentamycin resistance.

Acid and ethanol analysis. The concentrations of organic acids and ethanol that were secreted during growth in liquid medium were determined using commercially available detection kits. Formate and acetate were detected using kits from MegaZyme (Ireland, United Kingdom), whereas pyruvate, lactate, and ethanol were detected using kits from BioVision (Mountain View, CA), according to the manufacturer’s instructions. The pHs of culture supernatants were determined with an UltraBASIC pH probe (Denver Instruments).

Congo red staining. To determine the relative amount of endoglucanase secretion, an agar plate-based assay was used as described previously (49). Carboxymethylcellulose (1% [wt/vol]) and glucose (0.1% [wt/vol]) were used as carbon sources. After 48 h of growth, the plate was flooded with 0.1% (wt/vol) Congo red solution (Ricca Chemical) and stained for 15 min at room temperature. The dye was removed, and 5 ml of water was used to wash the plate. Finally, 5 ml of a 1 M NaCl solution was applied for 15 min, and the plates were then dried and photographed.

RESULTS

Growth of *C. japonicus* in the presence of cellulose substrates. Although *C. japonicus* strain Ueda107 can grow in the presence of laboratory cellulose substrates (51), little has been reported about its ability to utilize carbon sources relevant to bioenergy. We examined the ability of *C. japonicus* to grow in the presence of a collection of cellulose and noncellulosic carbon sources. The growth rates of *C. japonicus* in M9 minimal medium supplemented with 0.2% glucose, 0.34% cellobiose, or 1% carboxymethylcellulose were approximately the same (Fig. 1A, B, and C), with generation times of 2.7 h, 2.4 h, and 2.8 h, respectively. Similar results were observed when 1% glucose or 1% cellobiose was provided as a carbon source (J. G. Gardner and D. H. Keating, unpublished data).

We then examined the ability of *C. japonicus* to utilize the insoluble cellulose substrate Avicel. Because measurement of the optical density at 600 nm (OD_{600}) was challenging due to the insoluble Avicel present in the medium, we determined bacterial growth by viable-cell counting. Growth of *C. japonicus* in the presence of Avicel was biphasic, with rapid cell division (2.1-h generation time) occurring during the first 10 h, followed by a reduced growth rate (8.3-h generation time) (Fig. 1D). Although the reason for the biphasic growth is unknown, *C. japonicus* can clearly use soluble and insoluble cellulose substrates as sole sources of carbon and energy.

Growth of *C. japonicus* in the presence of biomass. The robust growth of *C. japonicus* in the presence of soluble and insoluble forms of cellulose led us to examine its ability to utilize feedstocks relevant to the bioenergy field. We chose two key biomass feedstocks for these studies: corn stover and switchgrass. Because current approaches to cellulose biofuels typically involve the use of biomass pretreatment, we focused on corn stover and switchgrass subjected to the AFEX process (37, 50). We chose AFEX as a pretreatment because it leaves the hemicellulose fraction intact (48, 50), which allowed us to examine the ability of *C. japonicus* to utilize both cellulose and hemicellulose.

When added to minimal medium, AFEX-treated corn stover was initially found to support the growth of diverse bacteria, including *E. coli* (Gardner and Keating, unpublished). Because our work, as well as the work of others, had previously shown that *E. coli* cannot utilize carboxymethylcellulose or Avicel and lacks any detectable cellulase activity, we hypothesized that the AFEX process results in the release of soluble sugars or peptides that support the growth of noncellulolytic bacteria. We therefore developed an alternative protocol involving grinding of the AFEX-treated material with a mortar and pestle and autoclaving, followed by extensive washing with sterile water. When added to M9 minimal medium, the washed, autoclaved, AFEX-treated corn stover (or AFEX-treated switchgrass) failed to support the growth of *E. coli* but allowed the growth of *C. japonicus* (Fig. 2A). Growth of *C. japonicus* in the presence of AFEX-treated corn stover was associated with release...
of glucose and xylose monosaccharides (Gardner and Keating, unpublished), suggesting that cellulose and hemicellulose were used as carbon sources.

When the growth rate was measured by viable-cell counting, *C. japonicus* grown in the presence of AFEX-treated corn stover or switch grass displayed growth kinetics distinct from what was observed with Avicel, glucose, carboxymethylcellulose, or cellobiose as a carbon source. No detectable increase in cell numbers was observed for approximately 10 h after addition of the cells to the medium (Fig. 2B). After the 10-hour growth lag, however, the cells displayed a growth rate similar to what was seen in other media (generation time, 1.9 h). Interestingly, the addition of 0.2% glucose to media containing AFEX-treated corn stover did not reduce the length of this lag (Fig. 2E), although it appeared to increase the growth rate after the initiation of cell division. *C. japonicus* was also able to utilize AFEX-treated switchgrass as a carbon source (Fig. 2C), with the cells displaying a growth lag similar to that seen with AFEX-treated corn stover but more rapid growth at later time points (generation time, 0.98 h). To determine if the lag was related to the AFEX pretreatment, we measured the rate of growth of *C. japonicus* with nonpretreated corn stover but more rapid growth at later time points (generation time, 1.7-h generation time) (Fig. 2D). However, the final cell yield was reduced by about 10-fold with respect to growth in the presence of AFEX-treated corn stover.

**FIG. 2.** Growth and degradation of biomass by *C. japonicus*. (A) Wild-type *C. japonicus* and *E. coli* were cultured in the presence of 1% (wt/vol) autoclaved, washed corn stover for 48 h at 30°C. The tubes containing cells and residual cellulose were then photographed. “Control” refers to mock-inoculated medium containing 1% (wt/vol) AFEX-corn stover. (B) Growth of *C. japonicus* in minimal medium supplemented with 1% (wt/vol) AFEX-CS. (C) Growth of *C. japonicus* in minimal medium supplemented with 1% (wt/vol) AFEX-SG. (D) Growth of *C. japonicus* in minimal medium supplemented with 1% (wt/vol) nonpretreated corn stover (nontreated-CS). (E) Growth of *C. japonicus* in minimal medium supplemented with 1% (wt/vol) AFEX-treated corn stover and 0.2% glucose (AFEX-CS + glucose). In panels B to E, the number of cells was determined by viable-cell counting. All experiments were performed in triplicate. The error bars (often too small to be seen) represent standard deviations.

Production of organic acids during growth in the presence of cellulosic substrates. A previous study noted that *C. japonicus* was capable of producing acid when grown with cellobiose and mannitol as carbon sources, but acid was not produced when glucose or other mono- and disaccharides were provided as carbon sources. The surprising variability in acid production during growth with different carbon sources led us to ask whether acid was produced when *C. japonicus* was grown in the presence of cellulosic substrates. Similar to what was reported previously (31), we observed a decrease in medium pH when *C. japonicus* was grown with 0.34% cellobiose as a carbon source (Fig. 1B). However, in contrast to what has been reported previously, we also observed a decrease in pH when 0.2% glucose was included as a carbon source (Fig. 1A). When *C. japonicus* was grown with carboxymethylcellulose as a carbon source, we observed a decrease in pH, but it was greatly diminished with respect to what was observed in the presence of glucose or cellobiose (Fig. 1C). A similar trend was observed with insoluble forms of cellulose, which led to a negligible pH decrease (Fig. 2B, C, and D). We do not have an explanation for the reduced amount of acid production observed when cells were cultured in the presence of cellulosic carbon sources, although it could simply reflect the reduced availability of monosaccharides for the bacterium.

The decrease in medium pH during growth with glucose or cellobiose as a carbon source led us to characterize the acids excreted by *C. japonicus*. When cells were grown with either glucose or cellobiose as a carbon source, we observed that acetate and pyruvate were the dominant acids excreted into the culture medium, in addition to a small amount of lactate. When *C. japonicus* was grown in the presence of carboxymethylcellulose, small amounts of lactate and acetate were produced. However, only negligible amounts of acids were detected when cells were grown with Avicel or AFEX-
treated corn stover as a carbon source, consistent with the observed stable pH.

A *C. japonicus* gspD mutant is deficient for cellulase secretion and the use of cellulose and biomass as carbon sources. Previous studies have shown that *C. japonicus* produces a diverse and extensive group of cellulases and hemicellulases (15). Furthermore, a substantial fraction (ca. 90%) of these activities have been reported to be extracellular (59, 60). In Gram-negative plant-pathogenic organisms, such as *Erwinia* spp., extracellular glycosylhydrolyase and pectate lyase activities are secreted via the TTSS (46). The genome of *C. japonicus* encodes a single predicted TTSS, but little has been reported about its role in cellulase secretion or growth on cellulosic substrates. To characterize the role of the TTSS in *C. japonicus*, we developed a system for construction of directed gene disruptions in *C. japonicus* strains. We then examined the ability of the mutant to utilize monosaccharides, disaccharides, and cellulosic substrates. The *gspD*:pJGG1 mutation displayed the same growth rate as the wild type on medium containing either glucose or cellulbiose as a carbon source (Fig. 5A and B). However, when carboxymethylcellulose was employed as a carbon source, the *gspD*:pJGG1 mutant displayed a lag in growth during the first 8 h and a reduced growth rate at later time points (Fig. 5C). Despite the reduced rate of growth, the *gspD*:pJGG1 mutant was able to reach the same final OD600 as the wild type. A more severe growth defect was observed in medium containing insoluble cellulose. The *gspD*:pJGG1 mutant showed a 6-h lag in growth in medium containing Avicel, which was followed by a brief period of rapid growth (Fig. 5D). However, growth appeared to cease at time points beyond 12 h, resulting in a greater-than-100-fold decrease in viable cells with respect to the wild type. When cultured in the presence of AFEX-treated corn stover, the *gspD*:pJGG1 mutant displayed a growth lag during the first 24 h (Fig. 5E), followed by a slight increase in the growth rate between 24 and 48 h and a greater-than-100-fold reduction in viable cells with respect to the wild type. Interestingly, growth of the *gspD*:pJGG1 mutant in the presence of AFEX-treated switchgrass resulted in a 1,000-fold reduction in viable cells with respect to the wild type (Fig. 5F).
Metabolic engineering of *C. japonicus*. The ability of *C. japonicus* to utilize bioenergy-relevant forms of cellulose led us to investigate its ability to be metabolically engineered for the production of bioproducts. We constructed a version of the broad-host-range plasmid pBBR-MCS5 (referred to as pJGG2) containing the *adhB* and *pdc* genes from *Z. mobilis*. The *pdc* gene encodes pyruvate decarboxylase, which catalyzes the decarboxylation of pyruvate to form acetaldehyde, and the *adhB* gene encodes alcohol dehydrogenase, which catalyzes the NADH-dependent reduction of acetaldehyde to ethanol. Studies from the Ingram laboratory have shown that expression of the *adhB* and *pdc* genes from *Z. mobilis* can enhance the production of ethanol in *E. coli* (61) and other Gram-negative bacteria (32). Plasmid pJGG2 replicated efficiently within *C. japonicus*, as judged by antibiotic resistance and recovery of intact plasmids from *C. japonicus* cells (Gardner and Keating, unpublished). When cultured under aerobic conditions in LB plus 0.2% glucose, cells harboring the pJGG2 plasmid grew at a rate approximately equivalent to that of cells containing the vector control but displayed a 15-fold increase in ethanol production (Fig. 6). Although ethanol production was modest under all conditions, the greatest ethanol production was observed in cells grown in the presence of LB supplemented with glucose, with growth in minimal glucose and minimal cellobiose media producing a slightly reduced amount of ethanol. In addition, growth in carboxymethylcellulose- and Avicel-containing media resulted in a low but detectable level of ethanol production. We were unable to detect any growth of *C. japonicus* containing the pJGG2 plasmid or vector control under anaerobic conditions with glucose, cellobiose, carboxymethylcellulose, Avicel, or AFEX-treated corn stover as a carbon source.

**FIG. 5.** Growth defect of the *C. japonicus* gspD mutant in media with insoluble cellulose. Wild-type *C. japonicus* and the gspD::pJGG1 mutant were cultured in M9 medium supplemented with 1 mM MgSO₄, 0.1 mM CaCl₂, and glucose (0.2% [wt/vol]) (A), cellobiose (0.34% [wt/vol]) (B), CMC (1% [wt/vol]) (C), Avicel (1% [wt/vol]) (D), AFEX-treated corn stover (1% [wt/vol]) (E), or AFEX-treated switchgrass (1% [wt/vol]) (F) as the sole carbon and energy sources, as described in Materials and Methods. All incubations were at 30°C with high aeration (225 rpm). Growth was determined by measurement of the OD₆₀₀ (A to C) or by viable-cell counting (D to F). All experiments were performed in triplicate. The error bars (often too small to be seen) represent standard deviations.

**DISCUSSION**

The *C. japonicus* genome is predicted to encode ca. 154 enzymes capable of cleaving glycosidic bonds. Consistent with this prediction, our studies showed that *C. japonicus* can utilize diverse cellulosic substrates as sources of carbon and energy, including the key bioenergy feedstocks corn stover and switchgrass. In contrast to a previous report (31), we observed that...
growth in the presence of glucose resulted in a reduction in medium pH, accompanied by the production of acetic acid, pyruvate, and a small amount of lactate. Using a directed gene disruption system developed as part of these studies, we demonstrated a central role for the type II secretion system in the utilization of insoluble cellulose. Collectively, our results support the notion that *C. japonicus* can serve as a useful source of biomass-degrading enzymes, and perhaps secretion systems, for the generation of Gram-negative consolidated bioprocessors.

**Growth of *C. japonicus* in the presence of insoluble cellulose.**

When glucose, cellobiose, or carboxymethylcellulose was supplied as a carbon source, *C. japonicus* showed similar generation times (Fig. 1A, B, and C). However, the kinetics of growth differed significantly when *C. japonicus* was cultured in the presence of insoluble forms of cellulose. In the presence of Avicel, growth of *C. japonicus* was biphasic, with the most rapid cell division occurring in the first 10 h, followed by a reduced growth rate (Fig. 1D). Interestingly, several groups have reported similar biphasic curves for cellulose hydrolysis in the presence of purified cellulases (2). In these cases, the initial rapid rate of degradation has been suggested to result from cleavage of amorphous cellulose, followed by a much slower hydrolysis of the crystalline cellulose (although other explanations for this phenomenon have been advanced [26, 39]). We suggest that the initial rapid growth of *C. japonicus* may also be supported by the hydrolysis of amorphous cellulose, whereas the lower rate of growth at later time points results from the reduced rate of hydrolysis of crystalline cellulose.

We also examined the ability of *C. japonicus* to use two sources of biomass relevant to the bioenergy industry: corn stover and switchgrass. In both cases, *C. japonicus* was able to degrade the substrates, as judged by an increase in cell number and a decrease in insoluble biomass in the culture (Fig. 2) and the release of glucose and xylose monosaccharides (Gardner and Keating, unpublished). Although the maximum growth rate of *C. japonicus* was comparable to that seen in the presence of purified cellulose, an approximate 8-h lag was observed when biomass was provided as a growth substrate. Interestingly, this lag in growth was observed in the absence of corn stover pretreatment (Fig. 2D) and in the presence of corn stover medium supplemented with 0.2% glucose (Fig. 2E). Because *C. japonicus* grows well with 0.2% glucose as a carbon source (Fig. 1A), these data strongly suggest that the growth delay observed during the first 10 h results from the presence of the biomass. Furthermore, since the growth delay was seen with biomass that had not undergone pretreatment (Fig. 2D), the lag does not result from compounds arising during the AFEX process. Plants have been reported to produce a significant number of compounds that could negatively affect the growth of *C. japonicus*. For example, lignin is a major constituent of plant cell walls (20 to 30% of the total) and contains phenolic substituents with known toxic effects (11, 13, 30, 42). Plants produce additional defensive compounds, such as phenylpropanoid derivatives and hydroxylated phenols with known antimicrobial activities (13). In particular, corn extracts contain 2,4-dihydroxy-7-methoxy-2H-1,4-benzoxazin-3(4H)-one, which has been shown to prolong the lag phase of *Erwinia* species, as well as that of other Gram-negative bacteria (12, 27). Interestingly, the ability to detoxify this compound has been reported in some organisms (25, 57). We are currently employing *C. japonicus* microarrays to measure the transcriptional response of *C. japonicus* to the presence of glucose, cellobiose, Avicel, and biomass. We expect that the results of these transcriptional-profiling studies will provide insight into the nature of this growth inhibition.

**Type II secretion system-dependent secretion of cellulases.**

Previous studies by several groups have shown that pectinases and cellulases produced by Gram-negative organisms can be secreted via the type II secretion system (34, 46). Although *C. japonicus* contains a predicted type II secretion system, the roles of these genes in cellulase and hemicellulase secretion, or growth on cellulolic substrates, had not been investigated. By the use of a vector integration strategy, we constructed a *gspD::pJGG1* mutant that would be expected to prevent the function of the type II secretion system. The *gspD::pJGG1* mutant displayed a marked delay in growth when the soluble cellulase substrate carboxymethylcellulose was provided as a carbon source but ultimately grew to the same final OD₅₀₀ as the wild-type strain (Fig. 5C). A similar phenotype was observed in measurements of endoglucanase activity, as detected by carboxymethylcellulose cleavage and Congo red staining. After 48 h of incubation, a greater-than-2-fold reduction in endoglucanase activity was observed (Fig. 4). However, prolonged incubation resulted in a detectable zone of clearing from the *gspD::pJGG1* mutant (Gardner and Keating, unpublished). We suggest that the initial delay in growth in the presence of carboxymethylcellulose, and the initial lack of detectable activity observed in the Congo red assay, results from an insufficient extracellular concentration of endoglucanases required for hydrolysis of carboxymethylcellulose. In both cases, prolonged incubation results in a greater release of endoglucanases (either by alternative secretion pathways or by cell lysis), which then allow efficient carboxymethylcellulose degradation and cell growth.

When the *gspD::pJGG1* mutant was cultured in the presence of insoluble cellulosic substrates, a more severe growth phenotype was observed. In the presence of Avicel, AFEX-treated corn stover, or AFEX-treated switchgrass, the growth of the *gspD::pJGG1* mutant arrested prematurely, which led to a greater-than-100-fold decrease in viable cells compared to the wild type. These data indicate that the type II-independent endoglucanase release is insufficient for growth on more recalcitrant carbon sources. A recent study in the bacterium *Saccharophagus degradans* demonstrated the presence of a processive endoglucanase activity encoded by the *cel5H* gene, which was capable of the conversion of insoluble cellulose to cellobiose, as well as an apparent lack of cellobiohydrolase activity (54). The cellulose-degrading apparatus of *C. japonicus* bears a great deal of sequence similarity to the *S. degradans* system (15) and may be similarly reliant on processive endoglucanases. If this class of enzymes is poorly released in the *gspD::pJGG1* mutant, this could explain the severe growth defects observed in the presence of insoluble cellulose. It remains possible that the reduced growth of the *gspD::pJGG1* mutant simply results from sensitivity of the cells to components of insoluble cellulose, although this possibility is rendered less likely by the normal rate of growth of the organism on glucose or cellobiose (Fig. 5A and B). Future studies will employ proteomics to characterize the secretion of extracellular enzymes of *C. japonicus*.
lar enzymes in the wild type and the gspD::pJGG1 mutant during growth in the presence of soluble and insoluble forms of cellulose. These data will not only expedite the identification of substrates for the type II secretion system, but should also identify the critical enzymes necessary for breakdown of cellulose and biomass.

A role for C. japonicus in biomass processing. C. japonicus can utilize diverse forms of cellulose, including bioenergy-relevant biomass. These data will not only expedite the identification of substrates for the type II secretion system, but should also identify the critical enzymes necessary for breakdown of cellulose and biomass.

Our demonstration that pBBR plasmids replicate within C. japonicus and development of methods for directed gene disruption, combined with previous reports of transposon-based gene knockouts (5, 17), provide the tools necessary for metabolic engineering. As a proof of concept, engineered C. japonicus was shown to produce ethanol from cellulolic substrates (Fig. 6), although at low yields. This modest amount of ethanol production may have resulted from the low expression of the pdc and adhB genes in C. japonicus, and we are currently testing alternative promoters to improve pdc and adhB expression.

Alternatively, the reduced ethanol yields may have resulted from our use of aerobic growth conditions, which would be expected to greatly reduce the intracellular concentrations of NADH required for ethanol production (9, 33, 53). Growth under anaerobic conditions has been reported previously for C. japonicus (16); however, we were unable to confirm this in our laboratory. Studies are currently under way to test the utility of C. japonicus to produce next-generation biofuels that can be produced under aerobic growth conditions.

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