Promiscuous plasmid replication in thermophiles: Use of a novel hyperthermophilic replicon for genetic manipulation of *Clostridium thermocellum* at its optimum growth temperature

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**Abstract**

*Clostridium thermocellum* is a leading candidate for the consolidated bioprocessing of lignocellulosic biomass for the production of fuels and chemicals. A limitation to the engineering of this strain is the availability of stable replicating plasmid vectors for homologous and heterologous expression of genes that provide improved and/or novel pathways for fuel production. Current vectors rely on replications from mesophilic bacteria and are not stable at the optimum growth temperature of *C. thermocellum*. To develop more thermostable genetic tools for *C. thermocellum*, we constructed vectors based on the hyperthermophilic *Caldicellulosiruptor bescii* replicon pBAS2. Autonomously replicating shuttle vectors based on pBAS2 reproducibly transformed *C. thermocellum* at 60 °C and were maintained in multiple copy. Promoters, selectable markers and plasmid replication proteins from *C. bescii* were functional in *C. thermocellum*. Phylogenetic analyses of the proteins contained on pBAS2 revealed that the replication initiation protein Repl, is unique among thermophiles. These results suggest that pBAS2 may be a broadly useful replicon for other thermophilic Firmicutes.

**Keywords:** Plasmid, Genetic tools, Transformation, Thermophile, Clostridia

**1. Introduction**

*Clostridium thermocellum* has been the subject of intense study because of its unique mechanism for solubilizing plant biomass. *C. thermocellum* secretes proteins that form an extracellular organelle, called a cellulosome (Bayer et al., 2013; Lamed and Bayer, 1988), for attachment to and digestion of complex plant biomass. Its ability to produce ethanol from cellulose has made it a leading candidate for consolidated bioprocessing (Akinosho et al., 2014; Demain et al., 2005). Recent work to engineer *C. thermocellum* for increased ethanol production has eliminated competing fermentation pathways (Argyros et al., 2011; Biswas et al., 2015; Deng et al., 2013; Papanek et al., 2015; Tripathi et al., 2010) and allowed synthesis of other fuel molecules like isobutanol (Lin et al., 2015).

Published genetic methods in *C. thermocellum* allow transformation of plasmid DNA (Guss et al., 2012; Olson and Lynd, 2012a; Tyurin et al., 2004), the generation of gene deletions (Argyros et al., 2011) and some methods for gene expression (Deng et al., 2013; Lin et al., 2015; Mearls et al., 2015; Olson et al., 2015). While gene deletions and gene expression have led to significant increases in ethanol production, the expression systems for robust high-level expression of homologous and heterologous proteins in *C. thermocellum* have limitations. The plasmid replicon from pNW33N that has been used extensively in published *C. thermocellum* transformation protocols (Tripathi et al., 2010) is not stable at elevated temperatures (Olson and Lynd, 2012b). Therefore, current methods for DNA transformation require that selection of transformants be performed at 50–51 °C (Olson and Lynd, 2012a), a suboptimal growth temperature for *C. thermocellum*, which grows optimally at 60 °C (McBee, 1954). Because this origin does not support stable replication at 60 °C, it may contribute to both
the observed plasmid instability (Olson et al., 2015) and chromosomal integration (Lin et al., 2015), even at 50–55 °C. A plasmid that could faithfully replicate at 60 °C would enable optimal in vivo expression and rapid screening of cell wall decomposition enzymes or metabolic enzymes for strain engineering.

Selective markers that have been used in C. thermocellum (Olson and Lynd, 2012a) include pyrF, that confers uracil prototrophy in a pyrF deletion strain; chloramphenicol acetyl transferase (cat), that confers thiamphenicol resistance; and neomycin acetyl transferase (neo), that confers neomycin resistance. Counter-selectable markers include pyrF, that confers sensitivity to 5-fluoroorotic acid (5-FOA); tdk, that confers sensitivity to 5-fluoro-2’-deoxyuridine (FUDR); and hpt, that confers sensitivity to toxic purine analogs such as 8-azahypoxanthine (8AZH) (Pritchett et al., 2004; Stout and Caskey, 1985). Counter-selection with tdk may be performed in a wild type strain, while use of pyrF and hpt require the use of a strain deleted for the C. thermocellum chromosomal copy of the gene.

Many plasmids have been shown to be capable of intergeneric and even interphyletic replication, raising the possibility of finding additional origins of replication for use in C. thermocellum. Early work in mesophilic Gram-positive bacteria identified several such replicons. Staphylococcus aureus plasmid pC194, that replicates by a rolling circle mechanism (Khan, 2005), as well as other Staphylococcus plasmids successfully transformed Bacillus subtilis (Dubnau, 1983; Ehrlich, 1977). Enterococcus faecalis plasmid pM61 was used to transform Lactobacillus acidophilus (Luchansky et al., 1988) and B. subtilis (Jannière et al., 1990), and has since become a broad-range host vector for members of the Firmicutes. An example of interphyletic plasmid replication is pNG2, that was isolated from Corynebacterium diphteriae, a member of the Actinobacteria phylum, but replicates in Escherichia coli, a member of the Proteobacteria phylum (Serwold-Davis et al., 1987).

Unfortunately, many of the commonly used plasmid origins were isolated from mesophilic organisms, so functionality at thermophilic temperatures is a concern. For instance, most Staphylococcal plasmids can be cured by growth at 43 °C (McNamara, 2008). The B. subtilis plasmid plM13 was used to transform S. aureus (Projan et al., 1987), and more recently the replicon was used for the construction of plKM1 for use in the thermophile Thermoanaerobacterium saccharolyticum at 48 °C, although its optimal growth temperature is 60 °C (Maì et al., 1997). The plP404 replicon (Garnier and Cole, 1988) from Clostridium perfringens, that grows optimally at 43 °C, was used to transform Thermoanaerobacter ethanolicus JW200, albeit at a temperature much lower than the recipient’s optimum growth temperature of 69 °C (Peng et al., 2006). Although the promiscuity of many plasmids has been demonstrated, there are limitations to mesophilic bacterial plasmids being used at the optimal growth temperatures of thermophiles. Issues of structural instability (Olson et al., 2015) chromosomal integration (Lin et al., 2015; Olson et al., 2015), and decreased copy number (Klapatch et al., 1996) have been reported.

An alternative approach is to identify origins of replication from thermophilic organisms as a starting point for developing more thermostable shuttle vectors. One example is plasmid pBAS2 that is native to Caldicellulosiruptor bescii (Clausen et al., 2004), a thermophilic Firmicute. C. bescii grows optimally at 75 °C (Yang et al., 2009) and plasmids using the pBAS2 origin of replication have been shown to replicate at high copy at 65 °C (Groom et al., 2014). The replication functions of pBAS2 have been used to construct vectors for transformation of C. bescii (Chung et al., 2013, 2015) and Caldicellulosiruptor hydrothermalis (Groom et al., 2014). Bioinformatic analysis revealed significant sequence identity to other Gram-positive rolling circle replication (RCR) origins, but the mechanism of pBAS2 replication is not clear. The conserved nick site is not obvious, and single stranded DNA that typically accumulates as a consequence of rolling circle replication was not observed for pBAS2 (Clausen et al., 2004). The pBAS2 replicon encodes a replication protein with homology to known Repl proteins and a Xer-like recombinase (Chung et al., 2013; Clausen et al., 2004), which is potentially responsible for the resolution of multimers of both chromosomes and plasmids that form during replication (Colloms, 2013; Subramanya et al., 1997). The open reading frame encoding Cbes2777 (the Xer-like recombinase XerD) was shown to be necessary for autonomous replication in C. bescii (Chung et al., 2013). While some replicons rely on host recombinases that work in trans to resolve multimers, pBAS2 encodes its own recombinase to resolve plasmid multimers and maintain sequence fidelity.

Based on the thermophilic source of pBAS2 and the presence of both a replication gene and a recombinase gene, we hypothesized that pBAS2-derived plasmids would be capable of independent replication in C. thermocellum at its optimal growth temperature of 60 °C. We therefore built and tested plasmids carrying the pBAS2 origin of replication with different selectable markers to explore this possibility. Here we report the transformation and stable replication in C. thermocellum of plasmid vectors based on pBAS2. We show that the pyrF gene from C. bescii functions to complement a deletion of the pyrF gene in C. thermocellum and that promoters from C. bescii function to drive expression of genes at levels sufficient for selection of transformants. Transformation of C. thermocellum DSM 1313 was performed at 60 °C, the optimal growth temperature of this strain, and plasmids derived from pBAS2 replicate stably in C. thermocellum at this temperature. Phylogenetic analysis of pBAS2 protein sequences suggests that the replication protein is novel and unique among known thermophilic proteins.

2. Methods

2.1. Bacterial media and growth conditions

C. thermocellum DSM 1313 and its derivatives were grown anaerobically in modified CTFUd medium (Olson and Lynd, 2012a) at 60 °C, under an atmosphere of 85% nitrogen, 10% CO2, and 5% Hydrogen. Defined medium for transformation and selection was CTFUd-NY (Olson and Lynd, 2012a), which contains a vitamin mix of p-aminobenzoic acid, vitamin B12, biotin, and pyridoxamine in place of the yeast extract. CTFUd-NY was supplemented with 360 μM uracil when needed. Complex medium for recovery after transformation was similar to CTFUd but contained casein (0.2 w/v) and less yeast extract (0.1% w/v), which is referred to as CTFUd+C. E. coli was grown in Luria-Bertani broth supplemented with 50 μg/mL apramycin when selecting for the presence of a plasmid.

2.2. Plasmid vector construction

All PCR reactions for cloning were performed with Q5 polymerase (New England Biolabs, Ipswich, MA) according to the manufacturer’s instructions (98 °C duplex denaturation, 60 °C annealing temperature, 30 s per kb at 72 °C for elongation). Plasmid pDCW196 was constructed by ligating C. bescii shuttle vector pDCW89 (Chung et al., 2013) to the cat gene from C. thermocellum vector pMU612 (Tripathi et al., 2010). Primers X013 and X014 amplified a 7.69 kb fragment from pDCW89, and primers X015 and X016 amplified a 1.053 kb fragment from pMU612. These fragments were digested with BamHI and Nde1 (New England Biolabs) twice in succession for 45 min at 37 °C and ligated with the FastLiT e ligation kit (Epigencentre, Madison, WI) according to the manufacturer’s instructions. Plasmid pJGW37 was constructed by

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amplifying a 7.467 kb fragment from pDCW196 using primers JG024 and JG099, digesting with XbaI (New England Biolabs) and ligating as described above to circularize the linear fragment. All plasmid sequences were verified by Sanger DNA sequencing (Genewiz). All primers are listed in Supplementary Table S1.

2.3. Transformation of C. thermocellum ΔpyrF

A 20 mL starter culture of C. thermocellum DSM 1313 ΔpyrF was grown at 60 °C to mid-exponential phase in defined CTFUD medium (Olson and Lynd, 2012a) supplemented with uracil. 15 mL of this culture was transferred to 150 mL of the same medium and grown to OD600 = 0.6. Cells were cooled to room temperature for 25 min, harvested aerobically at 7500 × g for 10 min, and washed twice with pre-chilled 10% sucrose. Competent cells were divided into 30 μL aliquots, and those not used immediately were frozen in a dry ice–ethanol slurry and stored at −80 °C. 500 ng of plasmid DNA pDCWB9 was incubated with each aliquot for 15 min in pre-chilled 1 mm cuvettes. Cells were electrotransformed with a single exponential pulse with a Gene Pulser (BioRad, Hercules, CA) set at 1.8 kV, 25 μF, and 350 Ω, and placed immediately into 60 °C CTFUD+C medium (Olson and Lynd, 2012a) supplemented with 360 μM uracil for recovery. A 0.25% inoculum was transferred to defined liquid CTFUD-NY medium lacking uracil every three hours over the period of recovery to select for transformants.

2.4. Transformation of C. thermocellum Δhpt

A 10 mL starter culture of C. thermocellum Δhpt was grown at 60 °C to mid-exponential phase in rich CTFUD medium (Olson and Lynd, 2012a). This culture was transferred to 150 mL of the same medium and grown to OD600 = 0.9. Cells were cooled to room temperature for 25 min, harvested aerobically at 7500 × g for 8 min, and washed twice with pre-chilled 10% glycerol, 260 mM sucrose. Competent cells were divided into 30 μL aliquots, and those not used immediately were frozen in a dry ice–ethanol slurry and stored at −80 °C. 350 ng of plasmid DNA pGW37 was incubated with each aliquot for 15 min in pre-chilled 1 mm cuvettes. Cells were electrotransformed with a single exponential pulse with a BioRad Gene Pulser (1.8 kV, 25 μF, 350 Ω), and placed immediately into warm CTFUD+C medium for 6 h at 60 °C, after which they were serially diluted onto CTFUD plates containing yeast extract (0.1% w/v) and 10 μg/mL thiamphenicol (Sigma, St. Louis, MO). Colonies appeared after 4–5 days of growth at 60 °C under an atmosphere of 85% nitrogen, 10% CO2, and 5% Hydrogen. Transformation efficiency was calculated in CFU/μg of plasmid DNA, and experiments were performed in biological triplicate.

2.5. Total DNA isolation from Clostridium thermocellum

For plasmid copy number determination over the growth phase, a 0.1% (v/v) inoculum was grown without shaking at 60 °C. 2% (v/v) samples were removed for qPCR analysis at the indicated time points (Fig. 3). For plasmid maintenance experiments (Table 2), 0.25% inocula were grown in serial passages to exponential phase (OD600 = 0.4 for C. thermocellum ΔpyrF, ~0.7 for C. thermocellum Δhpt) and 5 ml of culture was harvested at 4 °C at 6100 × g. The pellet was resuspended in 200 μL 40 mM EDTA, 50 mM Tris–HCl, 25% sucrose (w/v) with 30 mg/mL lysozyme (Sigma, St. Louis, MO) and 1 μL/mL RNase A (Qiagen), and incubated for 30 min at 37 °C. Cells were frozen and thawed five times using a dry ice–ethanol slurry and a 37 °C water bath. 500 μL 6 M Guanidine–HCl pH 8.5 (Sigma, St. Louis, MO) was immediately added to the cell lysate and allowed to incubate at 75 °C for 10 min. The lysate was washed twice with Phenol:chloroform:

isoamyl alcohol and once with chloroform. To the aqueous layer, 3 M sodium acetate pH 5.2 was added, and DNA was precipitated at −80 °C for 3 h, washed with cold 70% ethanol and resuspended in 10 mM Tris–HCl. Total DNA for C. thermocellum transformants containing pGW37 was visualized on a 1% agarose gel (Sigma, St. Louis, MO) stained with ethidium bromide (Fig. 2C).

2.6. Verification of plasmid transformation and structural stability

Taq polymerase (Sigma, St. Louis, MO) was used for PCR reactions to confirm presence of the plasmid using total DNA purified from C. thermocellum transformants as template. Reactions were performed with primers DC091 and DC508 according to the manufacturer’s instructions (94 °C duplex denaturation, 56 °C annealing temperature, 1 min per kb at 72 °C for elongation). PCR products were visualized on a 1% agarose gel with an NEB 1 kb ladder for size verification (NEB, Ipswich, MA). To verify structural stability in C. thermocellum, total DNA was electrot transformed into E. coli BL21 with a BioRad Gene Pulser (Biorad, Hercules, CA) using an exponential pulse in a 2 mm cuvette (2.5 kV, 25 μF, 200 Ω). Selection in E. coli was performed with 50 μg/mL apramycin, and colonies were screened for the presence of the plasmid by performing restriction digests with EcoRI and ApaLI (NEB, Ipswich, MA).

2.7. Quantitative polymerase chain reaction (qPCR)

qPCR experiments were carried out with a LightCycler 480 Real-Time PCR instrument (Roche, Basel, Switzerland) with LightCycler 480 SYBR Green I master mix (Roche). Cycle thresholds resulting from amplification with two independent sets of primers specific to either the pGW07 plasmid (Q1/Q2 inside Cbes2777, Q3/Q4 inside Cbes2778) or the C. thermocellum chromosome (CTQ1/CTQ2 inside Clo1313_0092, CTQ3/CTQ4 inside Clo1313_0090) were used to compute relative copy number of the plasmid to the chromosome. The formula for this calculation was PCN = [Cλchromosome − Cλplasmid]/l. Four replicate reactions for each primer set were performed, and the average of the two primer sets on each sample was used to calculate the plasmid copy number in each serial subculture (Table S1) according to the method of Lee et al. (2006). Amplification efficiency over 10^4 fold range of DNA concentration was 102%, within the ratio of 90–110% that is considered acceptable.

2.8. Bioinformatic analysis

The National Center for Biotechnology Information (NCBI) and the Kyoto Encyclopedia of Genes and Genomes (KEGG) were used...
to search for homologous proteins to the ORFs on plasmid pBAS2. Amino acid similarity was also determined with the NCBI Basic Local Alignment Search Tool (BLAST). Clustal Omega was used for multiple sequence alignment with default settings (Goujon et al., 2010; Sievers et al., 2011). ProtTest 3.4 (Darriba et al., 2011) was used to estimate models of evolution for the multiple sequence alignments. To determine posterior probabilities, MRBAYES version 3.2.5 was run for 1,000,000 generations with the WAG + I + G + F model of evolution (Whelan and Goldman, 2001) for Cbes2777 and the LG + I + G + F model (Le and Gascuel, 2008) for Cbes2780. Figtree version 1.4.2 (http://tree.bio.ed.ac.uk/software/figtree/) was used to visualize the phylogenetic trees generated by MRBAYES. MEME was used for motif analysis (Bailey and Elkan, 1994), searching for 6 motifs for Cbes2777 and 3 motifs for Cbes2780. MAST was used for motif searches with default settings (Bailey and Elkan, 1994). CodonO software (Angellotti et al., 2007) was used with default settings on genome files uploaded from NCBI.

3. Results and discussion

3.1. The XerD protein encoded by the C. bescii native plasmid pBAS2 has homology to known thermophilic proteins but the replication initiation protein, RepL, does not

As noted previously (Clausen et al., 2004), the protein sequences encoded by the pBAS2 open reading frames Cbes2777 and Cbes2780 (Fig. 1) are homologous to Xer-like recombinases and the RepL family of replication initiation proteins, respectively. Open reading frames Cbes2778 and Cbes2779 are short ORFs that would encode proteins of 109 aa and 73 aa respectively and are annotated as hypothetical proteins of unknown function (Tables S2 and S3).

Xer recombinases are known to resolve multimers of both chromosomes and plasmids that form during replication (Colloms, 2013; Subramanya et al., 1997). To better understand the replication machinery of the pBAS2 origin, we performed phylogenetic analysis of Cbes2777 and Cbes2780 with more recently acquired...
sequence data (Geer et al., 2010; Kanehisa and Goto, 2000; Kanehisa et al., 2014). That analysis revealed that Cbes2777 is homologous to XerC and XerD proteins in other Firmicutes, and it weakly clusters with a group of proteins from the thermophilic, anaerobic genera *Thermoanaerobacter* and *Thermoanaerobacterium* (Fig. 2A). This is not surprising, as *Caldicellulosiruptor* is known to be closely related to these genera (Rainey et al., 1993). Motif analysis revealed that all of these Xer-like recombinases contain a phage integrase-like domain and a DNA breaking-rejoining domain (Fig. 2B). The members of the cluster containing Cbes2777 have strong motif signatures for each of these protein domains (Bailey and Elkan, 1994; Marchler-Bauer et al., 2015), in particular the RHRY conserved catalytic active site (Subramanya et al., 1997) (Fig. 2B).

Xer recombination sites for multimer resolution like *cer* on CoIE1 and *psi* on pSC101 require extra sequences for the binding of accessory proteins (Cornet et al., 1994; Summers and Sherratt, 1988). One such protein for *cer* is the arginine repressor ArgR (Stirling et al., 1988), which binds to specific sites on the plasmid to orient the recombination sites during multimer resolution. We were unable to identify canonical XerC- and XerD-binding sites (Blake et al., 1997) in the pBAS2 sequence, but a 1.4 kb region of non-coding sequence on plasmid pBAS2 contains predicted ArgR binding sites (Fig. 1) existing as an approximate direct repeat for...
regulatory protein binding. This site has striking sequence similarity to known ArgR binding sites, particularly those from the roc and car operons from *B. subtilis* (Makarova et al., 2001). Thus, one possibility is that this sequence behaves like cer and psi for the resolution of plasmid multimers using the XerD recombinase encoded in cis by plasmid pBAS2. This would require that the host chromosome encodes an ArgR gene which is, in fact, present in both *C. bescii* and *C. hydrothermalis*, where pBAS2 has previously been shown to replicate. Further, it is present in *C. thermocellum* and other thermophiles including *T. saccharolyticum* and *T. ethanolicus* (Marchler-Bauer et al., 2015), suggesting that pBAS2 may have a broader host range than the *Caldicellulosiruptor* genus.

In contrast to Cbes2777, homology searches for Cbes2780 revealed a limited number of homologs with relatively low sequence identity (< 30%). There were no plasmid-encoded thermophilic homologs, and very few Firmicute homologs (Fig. 3A). Cbes2780 clusters with RepL proteins from diverse bacterial phyla including Proteobacteria, Cyanobacteria, and Firmicutes (Fig. 3A and S1). Among all the replication protein homologs there is a single strong consensus sequence ΦNPΣG in a helix-turn-helix DNA binding domain (Fig. 3B), although the *C. bescii* protein has many more arginine residues at this site. It is possible that these residues play a role in thermostability, as arginine residues have been associated with protein thermostability, particularly by increasing hydrophilicity when they replace lysine residues (Barton, 2005; Nadir et al., 1992), and when they exist in clusters (Phillips et al., 2013). These findings, in light of the lack of thermophilic homologs and the fact that Cbes2780 resides on a plasmid, suggest that this protein might have been co-opted by *C. bescii* from a distantly-related mesophilic organism, or alternatively from a currently undiscovered thermophilic organism. Importantly, unique Rep proteins provide for unique incompatibility groups (Shintani et al., 2015) making these plasmids potentially compatible with other known replicons.

It is noteworthy that *Cyanothecae* sp. PCC 8802 MarR2/RepL and *SoDallas glossinidius* RepL are encoded on plasmids that encode Xer-like recombinases, reminiscent of the arrangement of RepL and XerD on pBAS2. In fact, the use of portions of pBAS2 to construct other vectors revealed that the XerD recombinase is required for plasmid replication in *C. bescii* (Chung et al., 2013). The organization of pBAS2 may contribute to plasmid promiscuity with both a replication initiation protein and a multimer-resolving recombinase on the same plasmid.

3.2. A thermostable replicon from the *C. bescii* plasmid pBAS2 transforms *C. thermocellum* at 60 °C

Two pBAS2-derived plasmids were used for testing transformation of *C. thermocellum*. Plasmid pDCW89 (Fig. 4A) was previously constructed from the native *C. bescii* plasmid pBAS2 (Fig. 1) (Clausen et al., 2004) for use as an *E. coli/Caldicellulosiruptor* shuttle vector (Chung et al., 2013). It contains the pBAS2 origin of replication, the *E. coli* plasmid pSC101 origin of replication, an apramycin resistance cassette for selection in *E. coli*, and the *C. bescii* *pyrF* wild type allele used to select uracil prototrophy in strains containing a deletion of the *pyrF* gene. Previously, a *C. thermocellum* *pyrF* deletion was constructed (Tripathi et al., 2010), allowing transformation of pDCW89 to be tested (see below) using uracil prototrophy as the selection. Because thiamphenicol selection is also commonly used for *C. thermocellum* transformations, we constructed plasmid pJGW37 (Fig. 4B). This plasmid was based on pDCW89 (Fig. 4A) but with the *pyrF* gene replaced with a chloramphenicol acetyltransferase (cat) gene for selection of transformants in *C. thermocellum* strains that are wild type at the *pyrF* locus.

Using a method developed for *C. bescii* (Chung et al., 2012), pDCW89 was electrotetroformed into a *C. thermocellum* strain containing a deletion of the *pyrF* gene (Tripathi et al., 2010). As with the *C. bescii* *pyrF* deletion, this strain is a uracil auxotroph resistant to 5-fluoroorotic acid (5-FOA) (Table 1), allowing selection and counter-selection of the *pyrF* wild type allele. Transformants of pDCW89 were successfully selected for uracil prototrophy in defined liquid medium at 60 °C. Further, plasmid pJGW37 was successfully transformed into *C. thermocellum* containing a deletion of the hypoxanthine phosphoribosyl transferase (hpt) (*Argyros et al., 2011*) selecting thiamphenicol resistance, again at 60 °C. Hpt forms nucleotide monophosphates from purines, and can lead to the incorporation of toxic purine analogs such as 8-azahypoxanthine (AZH) into DNA and RNA (Stout and Caskey,
The Δhpt strain is resistant to AZH allowing selection of transformants that are thiampenicol resistant and subsequent counter selection for AZH resistance.

Total DNA isolated from C. thermocellum transformants containing pDCW89 or C. thermocellum transformants containing pJGW37 was used to back-transform E. coli. Two different restriction endonuclease digests performed on plasmid DNA purified from nine independent E. coli colonies (3 shown) resulted in identical banding patterns relative to the original plasmid (Fig. S2). This proves that the plasmids were successfully transformed into C. thermocellum and suggests that there was no major structural instability of the plasmid during transformation and replication in C. thermocellum and back-transformation into E. coli. Plasmid pJGW37 was also purified directly from C. thermocellum and could be visualized on an agarose gel of total DNA (Fig. 4C).

### 3.4. Heterologous proteins and promoters from C. bescii function in C. thermocellum

Plasmid pDCW89 contains the C. bescii pyrF wild type allele and this gene functions to complement the pyrF deletion in C. thermocellum. The PyrF proteins have 45% amino acid sequence identity. The G+C content of the C. bescii pyrF gene is 35.7%, and that of the C. thermocellum pyrF gene is 39.8%. In fact, CodonO software (Angelotti et al., 2007) indicates that codon usage bias between the entire C. bescii and C. thermocellum genomes is not statistically significant with a p value of > 0.33. Both pDCW89 and pJGW37 rely on a C. bescii ribosomal protein promoter (Chung et al., 2013) to direct transcription of the selectable marker gene (Fig. 1), the pyrF gene in pDCW89 and the cat gene in pJGW37. While we have no direct evidence for the efficiency of these promoters at this time, they clearly function at a level sufficient to allow selection. No promoter from C. bescii has been characterized experimentally so the transcription start site and RNA polymerase binding site for this promoter is unknown, but the sequence does contain a prototypical RNA polymerase binding site sequence for growth phase dependent transcription in Gram-positive bacteria.

### 4. Conclusions

While many plasmid origins of replication are derived from mesophilic sources, genetic tools for thermophiles must function at elevated temperatures. The use of thermophilic sources offers a solution to this issue, and we have demonstrated that plasmid origin pBAS2 from thermophile C. bescii can replicate in C. thermocellum at its optimal growth temperature of 60 °C. This expansion of the genetic tools available for C. thermocellum will facilitate more rapid genetic engineering toward the goal of developing an organism for the efficient conversion of lignocellulosic biomass to fuels and chemicals. Further, the demonstration of stable, autonomous replication of the C. bescii pBAS2 replicon in C. thermocellum suggests that this replicon might serve as a new tool for plasmid-based expression in other thermophiles, as well.

### Conflict of interest

LL has an equity interest in Enchi Corporation, a biofuel start-up company with a financial interest in C. thermocellum.
Determination of plasmid copy number.

| Plasmid   | Passages with selection | Passages without selection |
|-----------|-------------------------|---------------------------|
| pDCW89    | 3.0 ± 0.4               | 9.2 ± 0.9                 |
| pJGW37    | 6.7 ± 0.6               | 7.5 ± 1.7                 |

Plasmid copy number was determined at mid-log phase after the indicated number of passages. Maintenance experiments were performed in biological duplicate. Uracil prototrophy was used to select for pDCW89. Resistance to thiamphenicol was used to select for pJGW37.

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Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at http://dx.doi.org/10.1016/j.meteno.2016.01.004.

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