

Residue Leu-641 of Acetyl-CoA Synthetase is Critical for the Acetylation of Residue Lys-609 by the Protein Acetyltransferase Enzyme of *Salmonella enterica**

Received for publication, May 3, 2005, and in revised form, May 16, 2005
Published, JBC Papers in Press, May 17, 2005, DOI 10.1074/jbc.M504863200

Vincent J. Starai‡, Jeffrey G. Gardner, and Jorge C. Escalante-Semerena§

From the Department of Bacteriology, University of Wisconsin, Madison, Wisconsin 53726-4087

Posttranslational regulation of protein function by acetylation is present throughout nature. Regulation of protein function by Sir2 protein (sirtuin) deacetylases is conserved in all domains of life. In the prokaryote *Salmonella enterica*, the metabolic enzyme acetyl-coenzyme A synthetase (Acs) is regulated by a Sir2-dependent protein acetylation/deacetylation system (SDPADS). The recent identification of the acetyltransferase enzyme responsible for the acetylation of Acs defined the SDPADS in prokaryotes. This report identifies one residue in Acs, Leu-641, which is critical for the acetylation of Acs by the protein acetyltransferase enzyme. *In vivo* and *in vitro* evidence shows that mutations at Leu-641 prevent the acetylation of Acs by protein acetyltransferase, maintain the Acs enzyme in its active state, and bypass the need for sirtuin deacetylase activity during growth on acetate.

Sir2 proteins (sirtuins) are biologically conserved NAD⁺-dependent protein deacetylases involved in the posttranslational modification of a wide variety of protein substrates including histones (1–5), tumor suppressor protein p53 (6, 7), and microtubule protein α -tubulin (8). Evidence that archaeal sirtuins regulate gene expression by modulating the acetylation state of the chromatin protein Alba has been reported (9, 10). The interactions of Sir2 family members with their protein substrates have been characterized through structural work (11–16), providing insights into the mechanisms of substrate binding and deacetylation chemistry.

Studies of Gcn5-related N-acetyltransferases (GNATs) identified a superfamily of protein acetyltransferases involved in the acetylation of Sir2 protein targets (17–21). Members of the GNAT superfamily of proteins are widely distributed throughout all domains of life, yet show strikingly limited sequence similarity at the primary amino acid level (21). Factors determining the specificity of these systems in regard to their substrates and molecular partners are only now beginning to emerge (14, 17).

Recent work from our laboratory identified acetyl-coenzyme

A (Ac-CoA) synthetase (Acs¹; EC 6.2.1.1) as a substrate of the sirtuin-dependent protein acetylation/deacetylation system (SDPADS) (22). The role of Acs, an enzyme central to the metabolism of all cells, is to activate acetate to Ac-CoA (23). The Acs reaction proceeds via an Ac-AMP intermediate (Fig. 1) before yielding Ac-CoA, a high energy metabolite involved in many anabolic and catabolic processes (23). In *Salmonella enterica*, Acs activity is posttranslationally regulated via acetylation of residue Lys-609. Acetylation of the latter effectively blocks the conversion of acetate to Ac-AMP (22). Removal of the acetyl moiety from acetylated Acs protein (Acs^{Ac}) is catalyzed by the *S. enterica* Sir2 ortholog encoded by the *cobB* gene (24). As reported earlier, sirtuin-dependent deacetylation of Acs^{Ac} consumes NAD⁺ and restores full activity of the enzyme (22). We recently showed that the *pat* (formerly *yfiQ*) gene of *S. enterica* encodes a member of the GNAT family of acetyltransferase enzymes responsible for the acetylation of Acs (25). The discovery of the protein acetyltransferase (Pat) defined the SDPADS in this bacterium (Fig. 2) (25).

Here we report results from studies aimed at dissecting the molecular interactions between Acs and the *S. enterica* protein acetyltransferase, Pat. Using genetic selections, we identified a specific residue in Acs, Leu-641, as a critical amino acid for the acetylation of Acs by Pat. We propose that changes at this position alter the protein-protein interactions between Acs and Pat. Residue Leu-641 is hypothesized to be part of the Pat-binding domain of Acs. *In vitro* and *in vivo* evidence is presented in support of this conclusion. This work begins defining the mechanism of protein substrate recognition by the SDPADS of *S. enterica*.

EXPERIMENTAL PROCEDURES

Bacterial Strains, Culture Media, and Growth Conditions

All bacterial strains used in this study were derivatives of the *S. enterica* serovar Typhimurium LT2. The genotypes of strains used in this work are listed in Table I. Bacterial strains were grown on no-carbon E minimal medium (NCE) (26), supplemented with potassium acetate as the source of carbon and energy, MgSO₄ (1 mM), and L-methionine (0.5 mM). Luria-Bertani broth (LB) was used as rich medium. Growth behavior was analyzed in 96-well microtiter dishes (BD Biosciences) using a computer-controlled BioTek EL808-Ultra microplate reader (BioTek Instruments Inc.) with the incubation chamber set at 37 °C. A 2- μ l inoculum of an overnight culture of *S. enterica* was used to seed 198 μ l of freshly prepared minimal medium in each well; the medium was supplemented with 10 mM potassium acetate as the carbon and energy source. When used, L-(+)-arabinose was added to a final concentration of 100 μ M. Data points were collected every 10 min, with shaking at the highest intensity for 540 s between readings. All chem-

* This work was supported in part by National Institutes of Health Grant GM62203 (to J. C. E.-S.). The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

‡ Supported by a Pfizer predoctoral fellowship and by the Jerome Stefaniak Fellowship awarded by the Dept. of Bacteriology/University of Wisconsin-Madison. Present address: Dept. of Biochemistry, Dartmouth Medical School, Hanover, NH 03755.

§ To whom correspondence should be addressed: 264 Enzyme Institute, 1710 University Ave., Madison, WI 53726-4087. Tel.: 608-262-7379; Fax: 608-265-7909; E-mail: escalante@bact.wisc.edu.

¹ The abbreviations used are: Acs, Ac-CoA synthetase; Acs^{Ac}, acetylated Acs protein; SDPADS, sirtuin-dependent protein acetylation/deacetylation system; Pat, protein acetyltransferase; Sir2, silent information regulator 2; NCE, no-carbon E medium; GST, glutathione S-transferase.

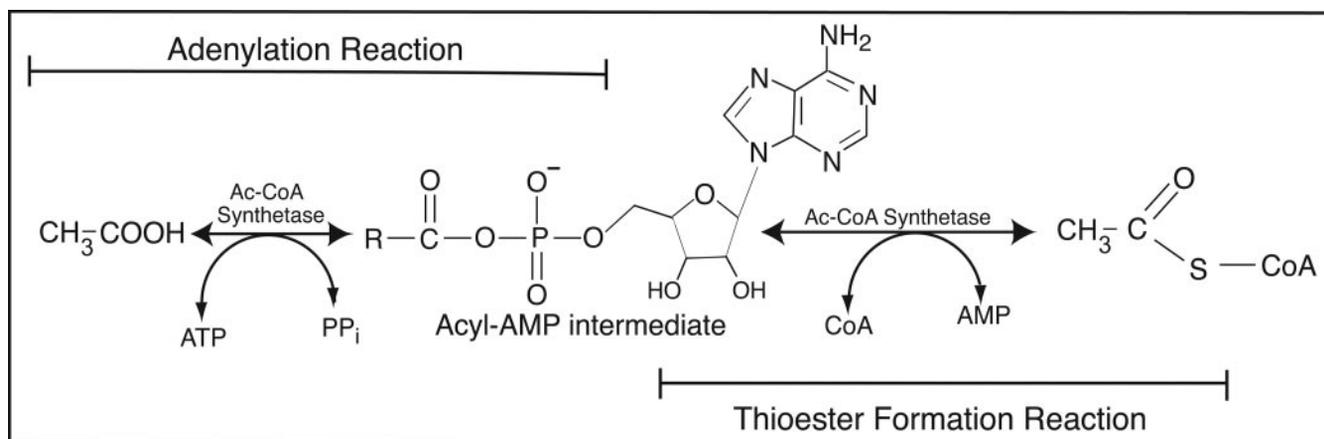


FIG. 1. Reaction catalyzed by Ac-CoA synthetase (EC 6.2.1.1).

TABLE I
Strains and plasmids

Unless otherwise stated, the strains listed were constructed during the course of this work.

Strain or plasmid	Relevant genotype	Source or ref. no. ^b
<i>S. enterica</i> strains ^a		
LT2	Wild type	Laboratory collection
LT2 derivatives		
TR6583	<i>metE205 ara-9</i>	K. Sanderson via J. Roth
TR6583 derivatives		
JE4175	pBAD30	Laboratory collection
JE6668	$\Delta 1231(acs) pta::cat^+ cobB1206::MudJ1734^b$	Laboratory collection
JE6858	$\Delta 1231(acs) pta::cat^+ cobB1206::MudJ / pBAD30$	
JE6859	$\Delta 1231(acs) pta::cat^+ cobB1206::MudJ / pACS7$	
JE7070	$\Delta 1231(acs) pta::cat^+ cobB1206::MudJ / pACS13$	
JE7462	<i>cobB1206::MudJ pat1::Tn10d(Tc) / pTara</i>	
JE7672	$\Delta 1231(acs) pta::cat^+ cobB1206::MudJ / pACS15$	
Plasmids		
pBAD30	Expression vector, $P_{araBAD} bla^+$	40
pACS7	<i>acs</i> ⁺ cloned into pBAD30	Laboratory collection
pACS10	<i>acs</i> ⁺ cloned into pTYB1 <i>bla</i> ⁺ (New England Biolabs)	22
pACS12	<i>acs4</i> (<i>Acs</i> ^{L641P}) cloned into pTYB1 <i>bla</i> ⁺	
pACS13	<i>acs4</i> (<i>Acs</i> ^{L641P}) cloned into pBAD30	
pACS15	<i>acs5</i> (<i>Acs</i> ^{L641Q}) cloned into pBAD30	
pACS16	<i>acs6</i> (<i>Acs</i> ^{L641A}) cloned into pBAD30	
pTara	Arabinose-inducible T7 RNAP ⁺	42

^a All *S. enterica* strains used in this study were derivatives of *S. enterica* serovar Typhimurium LT2.^b In the text, MudJ1734 (43) is referred to as MudJ (44).

icals were purchased from Sigma unless otherwise stated. Ampicillin was used at 100 μ g/ml and kanamycin at 50 μ g/ml.

Genetic Procedures

Phage P22 Transductions—All transductional crosses were performed as described previously (27) using phage P22 HT105/1 *int-210* (28, 29). Transductants were freed of phage as described (30).

Random Mutagenesis of *Acs*—Plasmid pACS7 (*acs*⁺) was introduced into the mutagenic *Escherichia coli* strain XL-1 Red (Stratagene) as per the manufacturer's instructions. The resulting transformants were resuspended into 5 ml of LB medium containing ampicillin and grown overnight at 37 °C with shaking. This culture was subcultured 1:100 (v:v) into 5 ml of fresh ampicillin-containing LB medium and allowed to continue growth at 37 °C with shaking. After the second overnight outgrowth, plasmid was reisolated using the Wizard Plus SV Miniprep kit (Promega).

Isolation of a Plasmid Encoding an *Acs* Enzyme Insensitive to Acetylation by *Pat*—The pool of mutagenized pACS7 plasmid was electroporated (31, 32) into *S. enterica* strain JE6668 (*acs pta cobB*), which was unable to grow on any concentration of acetate because of the lack of *Acs* and phosphotransacetylase (*Pta*) functions. The *cobB* mutation ensured that any *Acs* encoded by the plasmid introduced into the strain would remain acetylated; hence growth on acetate could only occur if the variant *Acs* protein could escape acetylation by the *Pat* enzyme. Transformed cells were plated onto NCE minimal agar supplemented with 1 mM MgSO₄, 0.5 mM L-methionine, 100 μ M L-(+)-arabinose (for induction of *acs* expression), ampicillin, and 10 mM acetate as the sole carbon and energy source. Plates were incubated at 37 °C for 48 h. Clones arising on these plates were reisolated on the same solid medium and further

characterized. The sequence of the *acs* gene encoded by plasmids that allowed strain JE6668 to grow on low acetate concentrations was determined using BigDye™ protocols (ABI PRISM, University of Wisconsin-Madison Biotechnology Center).

Construction of Plasmid pACS12 (*Acs*^{L641P})—Plasmid pACS10 encoding wild-type *Acs* protein fused at its C terminus to a chitin-binding domain was used as a template in the PCR-mediated QuikChange® XL site-directed mutagenesis kit (Stratagene). We used mutagenic primers 5'-GGTGGAGAAACCGCTCGAAGAGAA-3' and 5'-CTTCTCTTCGAGCGGTTTCTCCACC-3' to introduce the L641P mutation (underlined residues identify the mutagenic nucleotide). PCRs were performed as per the manufacturer's protocol; DNA sequencing confirmed the single base substitution. The resulting plasmid was named pACS12, which carried allele *acs4* that encoded *Acs*^{L641P} protein fused to a chitin-binding domain at its C terminus.

Construction of Plasmid pACS13 (*Acs*^{L641P})—Plasmid pACS7 carrying the wild-type *acs*⁺ gene under the control of an arabinose-inducible promoter was used as template in the PCR-mediated QuikChange® XL site-directed mutagenesis kit (Stratagene). To introduce the L641P mutation, the mutagenic primers used in the construction of plasmid pACS12 were used. The reactions were performed as per the manufacturer's protocol. DNA sequencing confirmed the single base substitution. The resulting plasmid was named pACS13, which directed the synthesis of *Acs*^{L641P} protein in response to arabinose in the medium.

Construction of Plasmid pACS15 (*Acs*^{L641Q}) and Plasmid pACS16 (*Acs*^{L641A})—Plasmid pACS15 and pACS16 were constructed as described for plasmid pACS13, except that mutagenic primers 5'-GGTGGAGAAACGCTCGAAGAGAA-3' and 5'-CTTCTCTTCGAGCTGTT-

FIG. 2. Model of posttranslational control of Acs by the SDPADS. Acs proteins are acetylated by the *S. enterica* Pat in an acetyl-CoA-dependent manner. These acetylated Acs proteins are inactive for the adenylation of acetate (left). CobB, the *S. enterica* sirtuin deacetylates Acs in a NAD⁺-dependent manner, releasing 2'-O-acetyl-ADP-ribose and nicotinamide. Unacetylated Acs is competent for the conversion of acetate to its adenyated intermediate (acetyl-AMP) and acetyl coenzyme A (right).

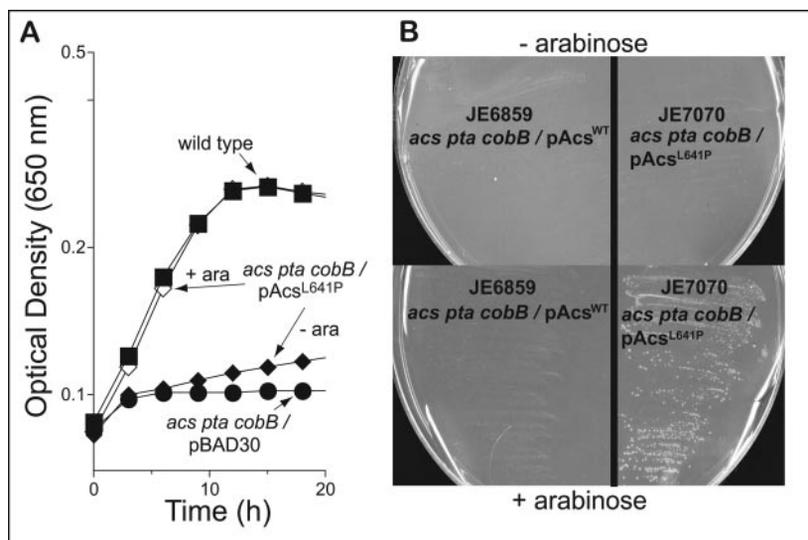
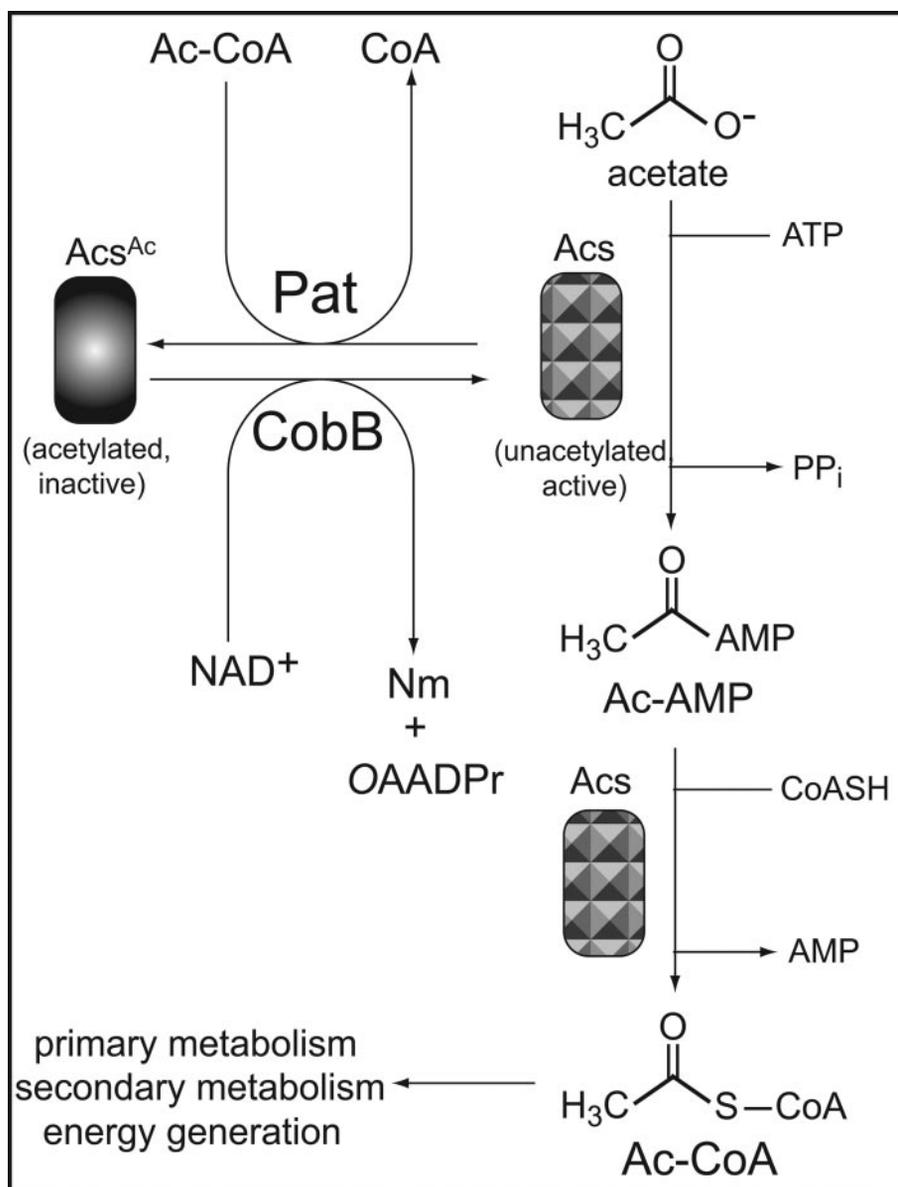


FIG. 3. The L641P mutation in Acs bypasses the need for sirtuin deacetylase activity *in vivo*. *A*, growth responses of *S. enterica* strains during growth on acetate (10 mM) as carbon and energy source. *Solid squares*, strain JE4175 (*acs*⁺; wild type); *solid circles*, JE6858 (*acs pta cobB* / pBAD30 empty vector control); *open diamonds*, JE7070 + arabinose (*acs pta cobB* / pAcs^{L641P}); *solid diamonds*, JE7070 without arabinose (*acs pta cobB* / pAcs^{L641P}). *B*, growth on minimal acetate medium with or without the addition of 100 μM L-(+)-arabinose for plasmid expression. pAcs^{WT}, pACS7; pAcs^{L641P}, pACS13.

TCTCCACC-3' were used to introduce the L641Q change, and the primers 5'-GTGGTGGAGAAAGCGCTCCGAAGAAGAAG-3' and 5'-CTTCTCTTCGAGCGCTTCTCCACCAC-3' were used to introduce the

L641A change. Plasmid pACS15 directed the synthesis of Acs^{L641Q} protein, whereas pACS16 directed the synthesis of Acs^{L641A} protein, both in response to arabinose in the medium.

TABLE II
Acs^{wt} and *Acs*^{L641P} activities purified from *cobB*⁺ and *cobB* strains

<i>acs</i> expression host ^a	Additions ^b	<i>Acs</i> ^{wt} -specific activity ^c	<i>Acs</i> ^{L641P} -specific activity ^c
<i>CobB</i> ⁺	None	3.9 ± 0.01	4.7 ± 0.05
	5 μg of CobB, 1 mM NAD ⁺	4.4 ± 0.1	4.3 ± 0.3
<i>CobB</i>	None	0.1 ± 0.01	3.2 ± 0.09
	5 μg of CobB, 1 mM NAD ⁺	4.2 ± 0.04	4.4 ± 0.1

^a The relevant genotype of the strain from which the *Acs* protein was purified. All strains also carried *metE205 ara-9* mutations in the genome.

^b The basic reaction mixture contained: coenzyme A, Mg(II)/ATP, bovine serum albumin, and purified *Acs* proteins.

^c Specific activity is defined as μmol of Ac-CoA min⁻¹ mg⁻¹. Results are the average of three independent determinations.

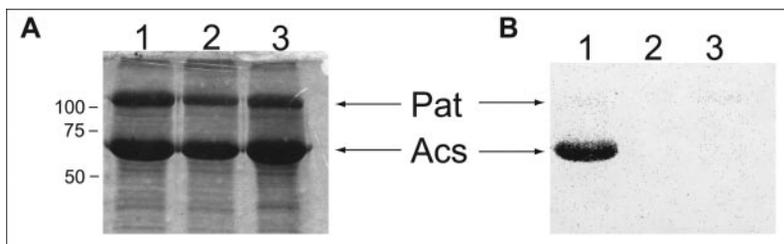


FIG. 4. *Acs*^{L641P} is not a substrate for Pat *in vitro*. A, Coomassie Blue-stained SDS-polyacrylamide gel, detecting the acetylated *Acs*. Lanes 1, *Acs* incubated with GST-Pat; lanes 2, heat-inactivated GST-Pat; lanes 3, *Acs*^{L641P} incubated with GST-Pat. Other components of the reaction mixture were as described above. Numbers to the left of the SDS-polyacrylamide gel indicate molecular mass in kilodaltons.

Biochemical Procedures

Purification of *Acs*^{L641P} Protein—*Acs* protein carrying the L641P mutation was obtained by expressing the *acs* allele carried by plasmid pACS12. The variant protein was purified from *S. enterica cobB*⁺ and *cobB* genetic backgrounds as described for wild-type *Acs* (22).

Purification of GST-Pat—GST-Pat was purified as described (25). Purified GST-Pat protein was dialyzed against 0.05 M Tris-HCl buffer, pH 7.5, containing 0.1 M KCl and 25% (v/v) glycerol. The protein was drop frozen in liquid nitrogen and stored at -80 °C.

In Vitro AMP-forming Ac-CoA Synthetase Reactions—Ac-CoA synthetase activity of *Acs* was monitored as described (22, 33, 34). *Acs*-dependent synthesis and isolation of radiolabeled Ac-CoA using [¹⁴C,C-1]acetate, ATP, and CoA with thin-layer chromatography were performed as described (22), except that 1 μg of unacetylated bovine serum albumin was added to each reaction to increase *Acs* stability.

Pat-mediated Acetylation of *Acs*—Protein acetylation reactions using [¹⁴C,C-1]Ac-CoA were performed as described (25). Briefly, 10 μg of *Acs* protein was added to 5 μg of GST-Pat protein in 0.05 M HEPES buffer, pH 7.5, containing 20 μM [¹⁴C,C-1]Ac-CoA (specific activity, 47 mCi/mmol) (Moravsek) and 200 μM Tris(2-carboxyethyl)phosphine hydrochloride in a final volume of 100 μl. Reactions were incubated for 2 h at 37 °C after which proteins were precipitated by the addition of trichloroacetic acid to a final concentration of 5% (w/v). Precipitated proteins were washed twice with 0.5 ml of ice-cold ethanol and allowed to air-dry. Proteins were resuspended in sample buffer and analyzed by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) (35) followed by Coomassie staining (36). Polyacrylamide gels were dried onto a piece of Whatman® paper and placed in a sealed film cassette under a Cyclone™ Storage Phosphor Screen (Packard Instruments). Signal intensity was analyzed after 8 h of exposure using the Cyclone™ Phosphor Detection System as per the manufacturer's directions.

Pat-mediated Inhibition of Ac-CoA Synthetase Activity—*Acs* proteins (*Acs*^{wt} and *Acs*^{L641P}) were isolated from *S. enterica* as described above, except that strain JE7462 (*pat cobB*/pTara) was used as the overexpression host. *Acs* proteins were acetylated by purified Pat protein in the following manner. A 150-μl total volume reaction containing 0.05 M HEPES buffer, pH 7.5, 200 μM Tris(2-carboxyethyl)phosphine, 40 μM Ac-CoA, and 10% glycerol (v/v) contained either active or heat-inactivated GST-Pat at a concentration of 0.95 μM (18.5 μg); *Acs* proteins were added to 1.9 μM (20 μg). Reactions were incubated at 37 °C for 2 h. GST-Pat was removed from the reactions by 80 μl of GST-Mag beads (50% slurry (v/v) (Novagen)) pre-equilibrated with the acetyltransferase reaction buffer. Incubation and removal of the magnetic beads was performed as per the manufacturer's instructions. Reaction supernatants were filtered through 0.45-μm Spin-X® centrifuge tube filters (Costar). The protein concentration of the filtered reaction was determined and tested for Ac-CoA synthetase activity as described (33, 34),

TABLE III
Pat-mediated inhibition of *Acs*^{wt} and *Acs*^{L641P} proteins

Conditions ^a	<i>Acs</i> ^{wt} -specific activity ^b	<i>Acs</i> ^{L641P} -specific activity ^b
+ Heat-inactivated GST-Pat	30.2 ± 1.1	11.8 ± 0.6
+ GST-Pat	1.9 ± 0.1	10.0 ± 0.5

^a Acetyltransferase reaction components are listed under "Experimental Procedures."

^b Specific activity is defined as μmol of acetyl-CoA formed min⁻¹ mg⁻¹. Results are the average of three independent determinations.

except that a total of 0.5 mg of *Acs* protein was used in the Ac-CoA synthetase reactions.

Protein Determination—Protein concentration in the samples was determined using the Bradford protein assay protocol (Bio-Rad Laboratories) with bovine serum albumin as standard per the manufacturer's instructions.

RESULTS AND DISCUSSION

The *in vitro* and *in vivo* evidence discussed below indicate that residue Leu-641 of *Acs* is critical for the acetylation of the active site residue Lys-609 by the Pat enzyme. Because residue Leu-641 lies close to the site of acetylation in *Acs* (Lys-609), we postulate that Leu-641 plays a critical role in the interaction of *Acs* with Pat.

The Variant *Acs*^{L641P} Enzyme Is No Longer Posttranslationally Controlled by the SDPADS—We used a genetic approach to identify derivatives of *Acs* that were no longer acetylated by Pat. *cobB pat*⁺ strains of *S. enterica* do not grow at low acetate concentrations (≤10 mM), because in the absence of CobB sirtuin deacetylase *Acs* remains acetylated (*i.e.* inactive, Fig. 2) (37). The rationale for the search was centered on the premise that if Pat failed to acetylate *Acs* in the *pat*⁺ *cobB* background, *Acs* would remain active and cells would grow on acetate despite the lack of CobB sirtuin deacetylase. To isolate variants of *Acs* that were no longer acetylated by Pat, we mutagenized plasmid pACS7 *acs*⁺ (*Acs*^{wt}) and selected for plasmids that would restore growth of the strain JE6668 (*acs pta cobB*) on minimal medium supplemented with 10 mM acetate as carbon and energy source and L(+)-arabinose (100 μM) as inducer of plasmid-borne *acs*. It was important to inactivate the chromosomal copies of *acs* and phosphotransacetylase (*pta*) to completely block acetate conversion to Ac-

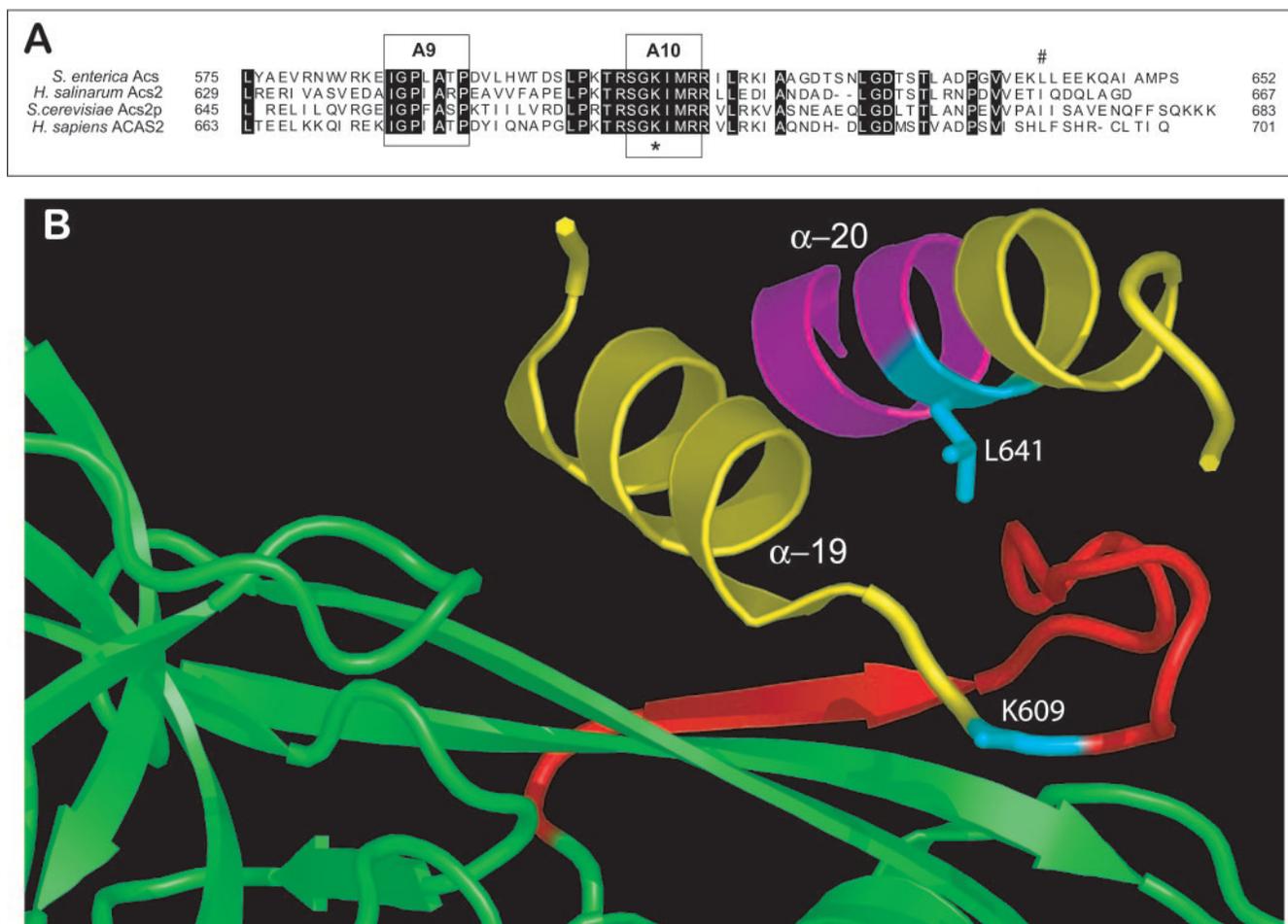


FIG. 5. Structural location of leucine 641P in relation to lysine 609. A, alignment of the C-terminal ~80 aa of Acs from prokaryotic, eukaryotic, and human Acs enzymes showing the location of the acetylation site (asterisk, Lys-609 on *S. enterica* Acs) and Leu-641 of the *S. enterica* enzyme (pound symbol). Panel B was generated using PyMol v. 97 software (pymol.sourceforge.net) (Protein Data Bank code 1PG4).

CoA (37–39) and eliminate background growth on acetate (Fig. 3B).

Sequencing of the *acs* allele encoded by a plasmid that supported robust growth of strain JE6668 on acetate revealed a single TA to CG transition at nucleotide 1922. This base substitution resulted in a mutant Acs protein with a replacement of a leucyl residue with a prolyl residue at position 641.

The L641P mutation was reconstructed using site-directed PCR mutagenesis to confirm the phenotype associated with this mutation. Acs protein encoded by the reconstructed plasmid (pACS13 *acs4* (Acs^{L641P})) restored the ability of strain JE6668 (*acs pta cobB*) to grow on acetate when arabinose was added to the medium to induce the synthesis of Acs^{L641P} (Fig. 3A, *open diamonds versus solid diamonds*). A strain carrying a plasmid encoding wild-type Acs protein failed to grow even when arabinose was included in the medium to induce the synthesis of wild-type Acs protein (Fig. 3B, *lower left panel*).

To determine whether the substitution of residue Leu-641 by Pro was specific, we replaced Leu-641 with Gln or Ala; the presence of the mutations was verified by DNA sequencing. Growth analyses involving strains carrying plasmid pACS15 *acs5* (Acs^{L641Q}) or plasmid pACS16 *acs6* (Acs^{L641A}) were performed using a strain carrying plasmid pACS13 *acs4* (Acs^{L641P}) as control. Strain JE6668 (*acs pta cobB*) harboring the plasmid pACS15 *acs5* (Acs^{L641Q}) or pACS16 *acs6* (Acs^{L641A}), as well as the strain carrying plasmid pACS13 *acs4* (Acs^{L641P}) (data not shown), grew on 10 mM acetate, indicating that the effect of a

change on Leu-641 was not specific to the side chain of the prolyl residue and emphasizing the importance of the side chain of the leucyl residue at that position.

Purified Acs^{L641P} Is Active in the Absence of CobB-Sirtuin Deacetylase Activity—Acs^{wt} and Acs^{L641P} Acs enzymes were purified from *S. enterica cobB*⁺ and *cobB* genetic background to gain insights into the mechanism by which the L641P mutation afforded CobB-independent Acs activity *in vivo*. Specific activities of these proteins were calculated, measuring the formation of [¹⁴C,C-1]Ac-CoA from [¹⁴C,C-1]acetate, Mg(II)/ATP, and HS-CoA (Table II). As expected, Acs^{wt} purified from the sirtuin-deficient *cobB* background was substantially less active than the identical enzyme purified from the sirtuin-proficient *cobB*⁺ background (~36-fold reduction in activity, Table II). Incubation of inactive Acs^{wt} with CobB sirtuin and NAD⁺ restored activity to wild-type levels (from 0.1 ± 0.01 to 4.2 ± 0.04 μmol/min/mg protein; Table II). In striking contrast to these results, the Acs^{L641P} enzyme purified from the *cobB* background was nearly as active as the Acs^{wt} enzyme purified from the *cobB*⁺ background (3.2 ± 0.09 versus 3.9 ± 0.01 μmol/min/mg protein, respectively). Incubation of CobB and NAD⁺ with the Acs^{L641P} enzyme purified from the sirtuin-deficient background did not change the activity of the Acs^{L641P} enzyme. In addition, Acs^{L641P} enzyme isolated from the *cobB*⁺ background showed little difference (~1.5-fold) in specific activity when compared with the same enzyme purified from the *cobB* background (4.7 versus 3.2 mmol of Ac-CoA min⁻¹ mg⁻¹), in-

dicating that the mutant protein was no longer under SDPADS control. These results could be explained by a lack of acetylation of the Acs^{L641P} enzyme by Pat. We also considered the unlikely alternative explanation would be that Acs^{L641P} enzyme gained the ability to activate acetate in the presence of acetylated Lys-609.

Pat Does Not Acetylate Acs^{L641P} Protein and Does Not Inhibit Its Activity—To address the possibility that the L641P mutation blocked Acs acetylation by Pat, purified Acs^{L641P} enzyme was incubated with Pat and [¹⁴C,C-1]Ac-CoA, and the presence of labeled Acs protein was analyzed by SDS-PAGE and phosphorimaging. Control experiments showed that Pat efficiently transferred the [¹⁴C,C-1]acetyl moiety of [¹⁴C,C-1]Ac-CoA to wild-type Acs protein (Fig. 4, lanes 1) and that heat-inactivated Pat did not (Fig. 4, lanes 2). In contrast to Acs^{wt}, Acs^{L641P} enzyme was not a substrate for Pat under the *in vitro* conditions tested (Fig. 4, lanes 3). These results were consistent with the idea that the Acs^{L641P} enzyme was not acetylated by Pat *in vivo*, allowing the CobB siruoin deacetylase-deficient strain, JE6668, to grow on acetate.

To show that purified Pat did not acetylate Acs^{L641P} at Lys-609, we monitored Acs activity after incubation with Pat. As expected, Pat-mediated inhibition of Acs^{wt} activity was observed (~16-fold reduction). Strikingly, the activity of the Acs^{L641P} enzyme was not significantly affected (10 versus 12 μmol of Ac-CoA/min/mg of protein; Table III). However, the Acs^{L641P} enzyme activity was about one-third that of the Acs^{wt} enzyme, suggesting that the L641P mutation may affect the stability of the enzyme. The effect of the L641P mutation on Acs activity or stability has not been further characterized.

Conclusions—Results from these studies suggest that the structural contributions of residue Leu-641 of Acs are critical for its interaction with the Pat protein acetyltransferase. This conclusion is based on the robust negative effects that changes at position 641 had on the ability of Pat to acetylate Acs, despite drastic changes in the side chain introduced at that position. Although the crystal structures of unacetylated and acetylated Acs from *S. enterica* were solved in the presence of ligands (41), the region of Acs that contains residue Lys-609 was disordered in both models, making it difficult to determine the spatial relationship of Leu-641 to Lys-609. Leu-641 is located in α-helix 20 (residues 637–645), the last α-helix of the Acs protein (Fig. 5), which could suggest that this helix represents a possible binding site on Acs for the Pat protein. Further studies of Pat-Acs interactions are under way.

REFERENCES

- Brachmann, C. B., Sherman, J. M., Devine, S. E., Cameron, E. E., Pillus, L., and Boeke, J. D. (1995) *Genes Dev.* **9**, 2888–2902
- Imai, S., Armstrong, C. M., Kaeberlein, M., and Guarente, L. (2000) *Nature* **403**, 795–800
- Landry, J., Sutton, A., Tafrov, S. T., Heller, R. C., Stebbins, J., Pillus, L., and Sternglanz, R. (2000) *Proc. Natl. Acad. Sci. U. S. A.* **97**, 5807–5811
- Smith, J. S., Brachmann, C. B., Celic, I., Kenna, M. A., Muhammad, S., Starai, V. J., Avalos, J. L., Escalante-Semerena, J. C., Grubmeyer, C., Wolberger, C., and Boeke, J. D. (2000) *Proc. Natl. Acad. Sci. U. S. A.* **97**, 6658–6663
- Rine, J., and Herskowitz, I. (1987) *Genetics* **116**, 9–22
- Luo, J., Nikolaev, A. Y., Imai, S., Chen, D., Su, F., Shiloh, A., Guarente, L., and Gu, W. (2001) *Cell* **107**, 137–148
- Vaziri, H., Dessain, S. K., Ng Eaton, E., Imai, S. I., Frye, R. A., Pandita, T. K., Guarente, L., and Weinberg, R. A. (2001) *Cell* **107**, 149–159
- North, B. J., Marshall, B. L., Borra, M. T., Denu, J. M., and Verdin, E. (2003) *Mol. Cell* **11**, 437–444
- Bell, S. D., Botting, C. H., Wardleworth, B. N., Jackson, S. P., and White, M. F. (2002) *Science* **296**, 148–151
- Zhao, K., Chai, X., and Marmorstein, R. (2003) *J. Biol. Chem.* **278**, 26071–26077
- Min, J., Landry, J., Sternglanz, R., and Xu, R. M. (2001) *Cell* **105**, 269–279
- Jackson, M. D., and Denu, J. M. (2002) *J. Biol. Chem.* **277**, 18535–18544
- Chang, J. H., Kim, H. C., Hwang, K. Y., Lee, J. W., Jackson, S. P., Bell, S. D., and Cho, Y. (2002) *J. Biol. Chem.* **277**, 34489–344998
- Avalos, J. L., Celic, I., Muhammad, S., Cosgrove, M. S., Boeke, J. D., and Wolberger, C. (2002) *Mol. Cell* **10**, 523–535
- Zhao, K., Harshaw, R., Chai, X., and Marmorstein, R. (2004) *Proc. Natl. Acad. Sci. U. S. A.* **101**, 8563–8568
- Zhao, K., Chai, X., and Marmorstein, R. (2004) *J. Mol. Biol.* **337**, 731–741
- Poux, A. N., and Marmorstein, R. (2003) *Biochemistry* **42**, 14366–14374
- Stern, D. E., and Berger, S. L. (2000) *Microbiol. Mol. Biol. Rev.* **64**, 435–459
- Rojas, J. R., Trievel, R. C., Zhou, J., Mo, Y., Li, X., Berger, S. L., Allis, C. D., and Marmorstein, R. (1999) *Nature* **401**, 93–98
- Grant, P. A., Duggan, L., Cote, J., Roberts, S. M., Brownell, J. E., Candau, R., Ohba, R., Owen-Hughes, T., Allis, C. D., Winston, F., Berger, S. L., and Workman, J. L. (1997) *Genes Dev.* **11**, 1640–1650
- Neuwald, A. F., and Landsman, D. (1997) *Trends Biochem. Sci.* **22**, 154–155
- Starai, V. J., Celic, I., Cole, R. N., Boeke, J. D., and Escalante-Semerena, J. C. (2002) *Science* **298**, 2390–2392
- Starai, V. J., and Escalante-Semerena, J. C. (2004) *Cell Mol. Life Sci.* **61**, 2020–2030
- Tsang, A. W., and Escalante-Semerena, J. C. (1998) *J. Biol. Chem.* **273**, 31788–31794
- Starai, V. J., and Escalante-Semerena, J. C. (2004) *J. Mol. Biol.* **340**, 1005–1012
- Berkowitz, D., Hushon, J. M., Whitfield, H. J., Roth, J., and Ames, B. N. (1968) *J. Bacteriol.* **96**, 215–220
- Davis, R. W., Botstein, D., and Roth, J. R. (1980) *A Manual for Genetic Engineering: Advanced Bacterial Genetics*, pp. 78–79, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY
- Schmieger, H. (1971) *Mol. Gen. Genet.* **100**, 378–381
- Schmieger, H., and Bakhaus, H. (1973) *Mol. Gen. Genet.* **120**, 181–190
- Chan, R. K., Botstein, D., Watanabe, T., and Ogata, Y. (1972) *Virology* **50**, 883–898
- Seidman, C. G., Struhl, K., Sheen, J., and Jessen, T. (1997) in *Current Protocols in Molecular Biology* (Ausubel, F. M., Brent, R., Kingston, R. E., Moore, D. D., Seidman, J. G., Smith, J. A., and Struhl, K., eds) Vol. 1, pp. 1.8–1.8.10, Wiley Interscience, New York
- Wong, T. K., and Nuemann, E. (1982) *Biochem. Biophys. Res. Comm.* **107**, 584
- Williamson, J. R., and Corkey, B. E. (1969) in *Methods in Enzymology* (Lowenstein, J. M., ed) Vol. XIII, pp. 494–497, Academic Press, New York
- Horswill, A. R., and Escalante-Semerena, J. C. (2002) *Biochemistry* **41**, 2379–2387
- Laemmli, U. K. (1970) *Nature* **227**, 680–685
- Sasse, J. (1991) in *Current Protocols in Molecular Biology* (Ausubel, F. A., Brent, R., Kingston, R. E., Moore, D. D., Seidman, J. G., Smith, J. A., and Struhl, K., eds) Vol. 1, pp. 10.6.1–10.6.8, Wiley Interscience, New York
- Starai, V. J., Takahashi, H., Boeke, J. D., and Escalante-Semerena, J. C. (2003) *Genetics* **163**, 545–555
- Brown, T. D., Jones-Mortimer, M. C., and Kornberg, H. L. (1977) *J. Gen. Microbiol.* **102**, 327–336
- Fox, D. K., and Roseman, S. (1986) *J. Biol. Chem.* **261**, 13487–13497
- Guzman, L.-M., Belin, D., Carson, M. J., and Beckwith, J. (1995) *J. Bacteriol.* **177**, 4121–4130
- Gulick, A. M., Starai, V. J., Horswill, A. R., Homick, K. M., and Escalante-Semerena, J. C. (2003) *Biochemistry* **42**, 2866–2873
- Wycuff, D. R., and Matthews, K. S. (2000) *Anal. Biochem.* **277**, 67–73
- Castilho, B. A., Olfson, P., and Casadaban, M. J. (1984) *J. Bacteriol.* **158**, 488–495
- Hughes, K. T., and Roth, J. R. (1988) *Genetics* **119**, 9–12