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# Structural and Biochemical Studies of a Fluoroacetyl-CoA-Specific Thioesterase Reveal a Molecular Basis for Fluorine Selectivity<sup>†,,‡</sup>

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## **Abstract**

We have initiated a broad-based program aimed at understanding the molecular basis of fluorine specificity in enzymatic systems, and in this context, we report crystallographic and biochemical studies on a fluoroacetyl-coenzyme A (CoA) specific thioesterase (FIK) from Streptomyces cattleya. Our data establish that FIK is competent to protect its host from fluoroacetate toxicity in *vivo* and demonstrate a  $10^6$ -fold discrimination between fluoroacetyl-CoA( $k_{\rm cat}/K_{\rm M}$ =5×10<sup>7</sup>M<sup>-1</sup> s<sup>-1</sup>) and acetyl-CoA( $k_{cat}/K_M$ =30 M<sup>-1</sup> s<sup>-1</sup>) based on a single fluorine substitution that originates from differences in both substrate reactivity and binding. We show that Thr 42, Glu 50, and His 76 are key catalytic residues and identify several factors that influence substrate selectivity. We propose that FIK minimizes interaction with the thioester carbonyl, leading to selection against acetyl-CoA binding that can be recovered in part by new C=O interactions in the T42S and T42C mutants. We hypothesize that the loss of these interactions is compensated by the entropic driving force for fluorinated substrate binding in a hydrophobic binding pocket created by a lid structure, containing Val 23, Leu 26, Phe 33, and Phe 36, that is not found in other structurally characterized members of this superfamily. We further suggest that water plays a critical role in fluorine specificity based on biochemical and structural studies focused on the unique Phe 36 "gate" residue, which functions to exclude water from the active site. Taken together, the findings from these studies offer molecular insights into organofluorine recognition and design of fluorinespecific enzymes.

Although the identification of halogenated natural products containing chlorine, bromine, and iodine has rapidly accelerated over the last 30 years as greater biodiversity has been explored, the incidence of fluorinated natural products remains limited to a relatively small number of structurally related compounds (Chart 1) (1–6). Fluorine has unique properties

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<sup>&</sup>lt;sup>‡</sup>The atomic coordinates and structure factors for the structures reported herein have been deposited in the Protein Data Bank (accession numbers 3P2Q, 3P2R, 3P2S, 3P3F, and 3P3I).

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Sequences of oligonucleotides, SDS-PAGE, size-exclusion chromatograms, sequence and structural alignments, and nonenzymatic hydrolysis data. This material is available free of charge via the Internet at http://pubs.acs.org.

arising from the combination of its small size, high electronegativity, and high C–F bond strength that have found utility in the design and development of new pharmaceuticals (7–10). It is often used as a sterically conservative replacement for hydrogen in small molecules, allowing them to retain binding to their macromolecular targets while adjusting several molecular properties such as stability, reactivity, conformation, lipophilicity, and *in vivo* distribution and metabolism (7–9, 11). Furthermore, the ease of elimination of fluoride from fluorine-substituted cellular metabolites has allowed the rational design of several bioactive mechanism-based inhibitors, such as 2′,2′-difluoro-2′-deoxycytidine (Gemzar) (12–14) and 5-fluorouracil (Adrucil, Carac, Efudex, and Fluoroplex) (15, 16).

Fluoroacetate is a naturally occurring organofluorine that is surprisingly poisonous considering its simple structure, with toxicity rivaling some of the most potent anticancer therapeutics (17, 18). Like Gemzar and fluorouracil, it can be metabolized by the normal host machinery before generating a mechanism-based inhibitor that shuts down the target pathway upon fluoride elimination. Specifically, fluoroacetate can be activated to form fluoroacetyl-CoA, which is then converted to fluorocitrate via the TCA cycle (19–22). While the aconitase-catalyzed formation of the fluoro-*cis*-aconitate intermediate parallels the reaction pathway of citrate, addition of water in the next step leads to stoichiometric fluoride elimination rather than formation of fluoroisocitrate (23–26). The product of this reaction is a strong noncovalent inhibitor that remains bound in the aconitase active site with an extremely slow rate of displacement by citrate (26, 27).

Although fluoroacetate production has been mainly noted in the plant kingdom (28–32), our understanding of organofluorine biosynthesis has come exclusively from studies of Streptomyces cattleya, a soil microbe that is the only known genetic host for production of both fluoroacetate and fluorothreonine (2,4,5,33,34). Similar to other antibiotic-producing hosts (35–40), S. cattleya is expected to contain pathways for detoxification in order to protect itself from its own production of fluoroacetate. A fluoroacetyl-CoA-specific thioesterase (FIK) was recently identified in S. cattleya that selectively hydrolyzes fluoroacetyl-CoA over acetyl-CoA (41). The gene encoding FlK is found clustered with the C-F bond-forming fluorinase (flA), raising the possibility that FlK-catalyzed hydrolysis of fluoroacetyl-CoA plays a role in fluoroacetate resistance in S. cattleya by preventing the entrance of fluoroacetyl-CoA into the TCA cycle (34). Given the close structural similarity between fluoroacetyl-CoA and acetyl-CoA, the discrimination between these two substrates exhibited by FlK is remarkable. The recently reported crystal structure of FlK demonstrates that this enzyme belongs to the hot dog-fold family of thioesterases (42). From these studies, a catalytic triad composed of Thr 42, His 76, and an active site water was proposed. Based on a computational model of the FIK-substrate complex, Arg 120 and Gly 69 were hypothesized to serve as the main selectivity elements by orienting the substrate for hydrolysis through fluorine-specific interactions.

Our group is interested in further studying the origin of substrate specificity in FIK to develop a better understanding of the design principles that drive fluorine-based selectivity in protein–ligand interactions. FIK provides a particularly interesting system for study because evolutionary selection against hydrolysis of the nonfluorinated substrate congener, acetyl-CoA, is likely important for maintaining normal cell growth and has tuned discrimination based on a single fluorine substitution. We quantify this fluorine-specific substrate discrimination to be surprisingly high  $(10^6$ -fold) and demonstate that it is derived from differences in both substrate reactivity and binding by carrying out steady-state kinetic

<sup>&</sup>lt;sup>1</sup>Abbreviations: CoA, coenzyme A; TCA cycle, tricarboxylic acid cycle; *S. cattleya, Streptomyces cattleya*, 4-HBA-CoA, 4-hydroxybenzoyl- CoA; hTHEM2, human thioesterase superfamily member 2; PaaI, phenylacetyl-CoA thioesterase; MBP, maltose binding protein.

characterization of FIK with respect to both fluoroacetyl-CoA and acetyl-CoA. Through a combination of structural and biochemical studies of FIK and a series of mutants, we have identifed several features unique to FIK that provide a molecular basis for fluorine specificity. We propose a model in which FIK minimizes interactions with the carbonyl unit common to both substrates, which contribute to substrate recognition and catalysis in typical members of the superfamily, selecting for reaction with the more activated fluoroacetyl-CoA substrate and decreasing general acyl-CoA binding. We further hypothesize that fluorine-specific interactions are recovered through the unusual lid structure of FIK, which provides a hydrophobic environment that both maximizes dipolar interactions with the fluorine atom and provides an entropic driving force for C–F bond recognition.

## **MATERIALS AND METHODS**

#### **Commercial Materials**

Restriction enzymes, T4 DNA ligase, and Phusion polymerase were purchased from New England Biolabs (Ipswich, MA). Platinum Taq High Fidelity (HF) polymerase and One-Shot TOP10 chemically competent cells were purchased from Invitrogen (Carlsbad, CA). Oligonucleotides were synthesized by Integrated DNA Technologies (Coralville, IA). DNA sequencing was performed by Quintara Biosciences (Berkeley, CA). DNA purification kits and Ni-NTA agarose were purchased from Qiagen (Valencia, CA). Complete Mini EDTAfree protease inhibitor was purchased from Roche Applied Science (Penzberg, Germany). Spectra Multicolor low-range protein ladder was purchased from Fermentas (Glen Burnie, MD). Amicon Ultra 3000 MWCO centrifugal concentrators and 5000 MWCO regenerated cellulose ultrafiltration membranes were purchased from Millipore (Billerica, MA). Acrylamide/bisacrylamide (40%, 19:1, and 30%, 37:1) and Gel Filtration Standard were purchased from Bio-Rad Laboratories (Hercules, CA). Luria-Bertani (LB) broth Miller and LB agar Miller were purchased from EMD Biosciences (Darmstadt, Germany). Carbenicillin, isopropyl  $\beta$ -D-thiogalactopyranoside (IPTG), tris(hydroxymethyl) aminomethane hydrochloride (Tris-HCl), sodium chloride, and dithiothreitol (DTT) were purchased from Fisher Scientific (Pittsburgh, PA). Streptomycin sulfate, ammonium persulfate, oxalyl chloride, N,N-dimethylformamide (Sure-Seal), 5,5'-dithiobis(2nitrobenzoic acid) (DTNB), sodium fluoroacetate, coenzyme A trilithium salt (CoA), acetyl-CoA, N, N, N, N'-tetramethylethane-1,2-diamine (TEMED), and tricine were purchased from Sigma-Aldrich (St. Louis, MO).

# **Gene Synthesis**

A synthetic gene encoding flK was codon-optimized for Escherichia coli using Gene Designer (DNA 2.0; Menlo Park, CA) and synthesized using PCR assembly. Gene 20ligo (43) was used to convert the gene sequence into primer sets using default optimization settings. Primers for gene assembly are listed in Supporting Information Table S1, and a primer map is shown in Supporting Information Figure S1. To assemble the synthetic gene, each primer was added at a final concentration of 1  $\mu$ M to the first PCR (50  $\mu$ L) containing Platinum Taq HiFi buffer (20 mM Tris-HCl, 50 mM KCl, pH 8.4), MgSO<sub>4</sub> (1.5 mM), dNTPs (250  $\mu$ M each), and 5 units of Platinum Taq HF (Invitrogen). The following thermocycler program was used for the first assembly reaction: 95 °C for 5 min; 95 °C for 30 s; 55 °C for 2 min; 72 °C for 10 s; 40 cycles of 95 °C for 15 s, 55 °C for 30 s, 72 °C for 20 s plus 3 s/cycle; these cycles were followed by a final incubation at 72 °C for 5 min. The second assembly reaction (50  $\mu$ L) contained 16  $\mu$ L of the unpurified first PCR with standard reagents for Platinum Taq HF. The thermocycler program for the second PCR was as follows: 95 °C for 30 s; 55 °C for 2 min; 72 °C for 10 s; 40 cycles of 95 °C for 15 s, 55 °C for 30 s, 72 °C for 80 s; these cycles were followed by a final incubation at 72 °C for 5 min. The second PCR(16  $\mu$ L) was transferred again into fresh reagents and run using the same

program. Following gene construction, the DNA smear at the appropriate size was gel purified and used as a template for amplification with Platinum TaqHF and rescue primers sFlK F100 and sFlKR100 (Supporting Information Table S1) under standard conditions. The PCR product was digested with *Nde*I and *Bam*HI and ligated into the *Nde*I–*Bam*HI sites of pET16b. The sequence of the synthetic gene was verified by sequencing.

#### **Construction of Vectors for Protein Expression**

The synthetic gene encoding FlK was PCR amplified from pET16b with Platinum TaqHF using the primers sFlKF13 and sFlK R15 (Supporting Information Table S1). The PCR product was digested with *Sfo*I and *Xh*oI and ligated into the *Sfo*I–*Xho*I sites of pET23a vector modified to encode a His<sub>10</sub> tag and a tobacco etch virus (TEV) protease cleavage site (Macrolab, UC Berkeley). The resulting plasmid was verified by sequencing.

# **Site-Directed Mutagenesis**

Plasmids encoding mutant FlKs were prepared using oligonucleotide primers containing the desired mutation (Supporting Information Table S2). Each mutagenesis reaction contained forward and reverse primers (2  $\mu$ M each), pET23a-His $_{10}$ -Tev-sFlK template (50–100 ng), dNTPs (200  $\mu$ Meach), Phusion polymerase, and Phusion buffer. The reaction mixture was subjected to the following thermocycler program: 98 °C for 3 min; 16 cycles of 98 °C for 15 s, 55 °C for 15 s, 72 °C for 3 min 30 s; a final extension at 72 °C for 7 min. Reactions were digested with DpnI and transformed into One- Shot TOP10 chemically competent cells. All mutants were then verified by sequencing of the flK gene. The E50Q mutant was subcloned into vector pSV272 (Macrolab, UC Berkeley) using primers sFlK F13 and sFlK R15 and using the SfoI and XhoI restriction sites to generate a His $_{10}$ -MBP-Tev-sFlK-E50Q expression construct.

# E. coli Viability Assay

Electrocompetent *E. coli* BL21-(DE3) cells were transformed with empty pET23a-His $_{10}$ -Tev, pET23a-His $_{10}$ -Tev-FlK, and pET23a-His $_{10}$ -Tev-FlK-H76A, and cells were grown on LB agar containing 50  $\mu$ g/mL carbenicillin. Saturated cultures were diluted to OD $_{600\text{nm}}$ =0.3 in LB, and serial 10-fold dilutions were made in a 96-well plate. Dilutions (2  $\mu$ L) were spotted onto LB agar containing either 50  $\mu$ g/mL carbenicillin or 50  $\mu$ g/mL carbenicillin and 20 mM sodium fluoroacetate. Plates were incubated at 37 °C for 16 h and then photographed.

### **Expression and Purification of FIK Variants**

LB containing carbenicillin (50  $\mu$ g/mL) was inoculated to OD<sub>600nm</sub>=0.05 with an overnight LB culture of freshly transformed *E. coli* BL21(DE3) containing the overexpression plasmid. The cultures were grown at 37 °C at 200 rpmtoOD<sub>600nm</sub>=0.8, at which point they were cooled on ice for 20 min. Isopropyl  $\beta$ -D-thiogalactopyranoside (IPTG) was added to a final concentration of 1 mM, and cells were incubated at 16 °C for 12–16 h. Cells were harvested by centrifugation at 3696g at 4 °C for 15 min. The cell pellet was resuspended in 5 mL of lysis buffer (50 mM sodium phosphate, pH 8.0, 300 mM NaCl, 20 mM imidazole, 100  $\mu$ M phenylmethanesulfonyl fluoride, one Complete Mini EDTA-free tablet per 50 mL buffer) per gram of cell pellet wet weight and lysed by one pass through a French pressure cell (Thermo Fisher) at 14000 psi. In soluble material was removed by centrifugation at 15316g at 4 °C for 20 min. A 10% (w/v) solution of streptomycin sulfate was added to the supernatant to a final concentration of 0.9% (w/v), and precipitated DNA was removed by centrifugation at 15316g for 20 min. The supernatant was loaded onto aNi-NTA agarose column (1 mL resin/L of cell culture) using an ÄKTA Purifier FPLC system (GE Healthcare). The column was washed with 10 column volumes of wash buffer (50 mM

sodium phosphate, pH 8.0, 300 mM NaCl, 20 mM imidazole) and 10 column volumes of wash buffer plus 8% elution buffer (50 mM sodium phosphate, pH 8.0, 300 mM NaCl, 250 mM imidazole). The column was eluted using a gradient from 8% to 100% elution buffer over 10 column volumes. Protein-containing fractions were pooled by their absorbance at 280 nm and concentrated to 10-15 mL using a 5000 MWCO ultrafiltration membrane. TEV protease (purified as described previously (44)) was added at a 1:50 mass ratio, and the mixture was dialyzed against TEV cleavage buffer (50 mM sodium phosphate, pH 8.0, 300 mM NaCl, 1 mM DTT) for 16 h. The dialysate was passed over a Ni-NTA column to remove the His<sub>6</sub>-tagged TEV protease and the His<sub>10</sub> peptide, and the column was washed with 2 column volumes of wash buffer. The flow-through and wash fractions were concentrated to 2 mL and loaded onto a Superdex 75 16/60 pg (GE Healthcare) column equilibrated with size-exclusion buffer (20 mM Tris-HCl, pH 7.6, 50 mM NaCl). FIK eluted from the column as a dimer (Supporting Information Figure S2) and was concentrated to 1-10 mg/mL and stored at 4 °C. For long-term storage, 10% glycerol was added to the concentrated protein, and it was stored at -80 °C. Both His<sub>10</sub>-tagged and His<sub>10</sub>-tagged MBP fusion proteins were purified following this protocol (Supporting Information Figure S3).

#### **Crystallization and Structure Determination**

Purified FIK in size-exclusion buffer was concentrated to 17 mg/mL. Crystals of apo-FIK were obtained using the hanging drop vapor diffusion method by combining equal volumes of protein solution and reservoir solution containing 0.1 M Tris-HCl, pH 7.6, and 25% polyethylene glycol 3350. Clusters of plates grew within 2 days. Single crystals were obtained by gently agitating the clusters. The crystals were cryoprotected by soaking briefly in a solution containing 0.1 M Tris-HCl, pH 7.6, 25% polyethylene glycol 3350, and 25% (v/v) ethylene glycol and flash cooled in liquid nitrogen. Data were collected at beam line 8.2.2 at the Advanced Light Source (Lawrence Berkeley National Laboratory). Data were processed with XDS and scaled with XSCALE (45). Molecular replacement with Phaser (46) using a polyalanine model of *Thermus thermophilus* hypothetical protein TTHA0967 (PDB ID 2CWZ) identified two FlK monomers in the asymmetric unit arranged with typical hot dog-fold topology. Maps were improved by Phenix AutoSolve and AutoBuild (47). The resulting maps were of sufficient quality to discard the original model, and Arp/wARP (48) was used to build a near-complete chain trace. The remaining parts of the model were manually built using Coot (49). The structure was solved at 1.9 Å resolution and refined to an  $R_{\text{work}}/R_{\text{free}}$  of 20.1%/23.1% using Phenix Refine.

Crystals of the open conformation of FIK were obtained by soaking apo-FIK crystals in a solution of 1 mM fluoroacetyl-CoA in mother liquor for 10 min followed by brief cryoprotection by soaking in mother liquor supplemented with 25% (v/v) ethylene glycol. Crystals were then flash-cooled, and data were collected at beam line 8.3.1 (Advanced Light Source, Lawrence Berkeley National Laboratory). Molecular replacement performed with Phaser using the refined apo-FIK structure as a search model identified two monomers per asymmetric unit. The open FIK structure was manually rebuilt using Coot and refined using Phenix Refine to  $R_{\rm work} = 18.8\%$  and  $R_{\rm free} = 22.0\%$  at 2.0 Å resolution. Crystals of the FIK-fluoroacetate complex were obtained by soaking apo-FIK crystals in 2.5 mM fluoroacetate. The initial model of the protein was obtained as for the open conformation of FIK. Unmodeled peaks in the map were identified using Coot, and ligands corresponding to the density were modeled when possible. The structure was solved to 2.5 Å resolution and refined with Phenix Refine to  $R_{\rm work}/R_{\rm free}$  of 19.4%/24.0%

Crystals of FIK-F36A were obtained as for wild-type FIK. FIK-F36A crystals grew in a larger C2 crystal form that had a Matthews coefficient consistent with six monomers per asymmetric unit. Molecular replacement performed with Phaser using the refined apo-FIK structure as a search model unambiguously identified three dimers per asymmetric unit, in

agreement with the Matthews coefficient. The structure was solved to 2.3 Å resolution, manually rebuilt using Coot, and refined to  $R_{\rm work}/R_{\rm free}$  of 22.0%/24.7%. Crystals of the FIK-F36A product complex were obtained in the same conditions as apo-FIK-F36A by cocrystallization with 1 mM fluoroacetyl-CoA and were found to contain three dimers per asymmetric unit. Maps were obtained, and the model was built as for FIK-F36A. The structure was solved to 2.0 Å resolution and refined with Phenix Refine to  $R_{\rm work}/R_{\rm free}$  of 21.8%/25.1%.

#### Fluoroacetyl-CoA Synthesis

Fluoroacetyl-CoA was synthesized as described previously (41) with some modification. Sodium fluoroacetate (100 mg, 1 mmol) was dried under vacuum in an oven-dried roundbottom flask equipped with a stir bar and a reflux condenser. Dry tetrahydrofuran (2 mL), dry N,N-dimethylformamide (100  $\mu$ L), and oxalyl chloride (2 M in dichloromethane, 1 mmol) were added to the flask by syringe, and the reaction mixture was stirred and heated at 65 °C for 2–3 h under nitrogen atmosphere. After the flask was allowed to cool, the reaction mixture (650  $\mu$ L) was added to stirred solution of coenzyme A trilithium salt (50 mg, 0.06 mmol) dissolved in 1 mL of 10% sodium bicarbonate. The mixture was stirred vigorously for 3 min under a stream of nitrogen and then immediately injected onto an Agilent Eclipse XDB-C18 column (9.4× 250 mm, 5  $\mu$ m) and purified by reverse-phase HPLC (0–100% B over 30 min at 3 mL/min; A, H<sub>2</sub>O; B, acetonitrile) using an Agilent 1200 binary pump coupled to a diode-array detector. Fractions (1.5 mL) were flash frozen in liquid nitrogen and lyophilized. The lyophilized fluoroacetyl-CoA was dissolved in water and characterized using an Agilent 6130 single-quadrupole electrospray ionization mass spectrometer (ESI-MS)  $(m/z (MH^+), 828.1; m/z (MH^+_{calc}), 828.1)$ . The purified substrate was also assayed for contaminating coenzyme A by comparing the total coenzyme A absorbance at 260 nm  $(\varepsilon_{260 \text{ nm}}=13100 \text{ M}^{-1} \text{ cm}^{-1})$  with free thiol content measured by reaction with DTNB  $(\varepsilon_{412\text{nm}}=13600 \text{ M}^{-1} \text{ cm}^{-1} \text{ for } 2\text{-nitro-}5\text{-thiobenzoate})$ . Only fluoroacetyl-CoA of >95% purity was used for assays.

## **Kinetic Measurements**

The thioesterase activity of FIK was measured by spectrophotometrically monitoring the increase in absorbance at 412 nm due to reaction of enzymatically generated free coenzyme A with DTNB in a Beckman Coulter DU-800 spectrophotometer as described previously (41) with some modification. FIK activity was measured immediately after purification. Assays were performed at 25 °C in a total volume of 500  $\mu$ L containing 100 mM Tris-HCl, pH 7.6, 0.5 mM DTNB, the appropriate amount of fluoroacetyl-CoA or acetyl-CoA, and FIK (0.5 nM, fluoroacetyl-CoA; 10  $\mu$ M, acetyl-CoA) or FIK mutant (50 nM<sup>-1</sup>  $\mu$ M, fluoroacetyl-CoA; 10  $\mu$ M for acetyl-CoA) in an amount sufficient to detect activity. Commercial acetyl-CoA was also tested for contaminating CoA as described above. A CoA standard curve from 0 to 50  $\mu$ M was generated using the appropriate amounts of CoA in a solution containing 100 mM Tris-HCl, pH 7.6, and 0.5 mM DTNB. Kinetic parameters ( $k_{\rm cat}$  and  $k_{\rm M}$ ) were determined by fitting the data using Kaleidagraph 3.51 (Synergy Software, Reading, PA) to the equation:

$$v_0 = v_{\text{max}}[S]/K_M + [S]$$

where  $v_0$  is the initial rate and [S] is the substrate concentration.

#### RESULTS AND DISCUSSION

## Toxicity of Fluoroacetyl-CoA and Rescue by FIK

To test whether FlK is competent to confer fluoroacetate resistance in a heterologous bacterial system, we expressed the *flK* gene in *E. coli* and measured its viability by spotting 1:10 dilutions of culture onto solid media (Figure 1). In the presence of fluoroacetate, the viability of *E. coli* containing an empty plasmid or a plasmid encoding an FlK mutant with diminished activity was severely reduced, while *E. coli* harboring the plasmid encoding wild-type FlK exhibited viability similar to that of cultures spotted onto fluoroacetate-free media, which indicates that FlK is competent to confer fluoroacetate resistance in bacteria. As acetyl-CoA is an essential building block in the cell, these results are consistent with the conclusion that FlK demonstrates specificity toward fluoroacetyl-CoA over acetyl-CoA *in vivo*.

#### Competition with Acetyl-CoA

Previous measurements of FIK activity suggested that this enzyme exhibits remarkable *in vitro* selectivity for fluoroacetyl-CoA over acetyl-CoA, with no hydrolysis observed at concentrations up to 1 mM acetyl-CoA in the presence of 50 ng of FIK (41). We can envision two major modes for recognition of the fluorinated substrate over the nonfluorinated substrate. One mode of substrate specificity could rely on structural recognition of the fluorine atom, which could be discriminated based on polarity or size. A second possible component of selectivity is the differing chemical reactivities of fluoroacetyl-CoA and acetyl-CoA. The inductively electron-withdrawing fluorine atom activates the thioester C=O in fluoroacetyl-CoA toward nucleophilic attack. Indeed, the pseudo- first-order rate constant for uncatalyzed hydrolysis of fluoroacetyl-CoA was measured to be an order of magnitude faster  $(1.4 \times 10^{-4} \text{ s}^{-1} \text{ at pH 7.6})$  than the hydrolysis of acetyl-CoA  $(1.3 \times 10^{-5} \text{ s}^{-1} \text{ at pH 7.6})$  under the same conditions (Supporting Information Figure S4). Thus, the active site nucleophile of FIK could be tuned toward lower reactivity, which would limit the hydrolysis of acetyl-CoA compared to fluoroacetyl-CoA.

To clarify whether the exclusion of acetyl-CoA binding plays a role in FIK selectivity, we performed a competition experiment in which the fluoroacetyl-CoA concentration was fixed at  $20~\mu\text{M}$  and the rate of hydrolysis was measured in the presence of increasing concentrations of acetyl-CoA (Figure 2). No significant effect on the rate of free CoA release was observed except at the highest concentration (5 mM) of acetyl-CoA, suggesting that the absence of the fluorine modification severely diminishes substrate affinity. This result is surprising in light of previous studies that have demonstrated promiscuous substrate binding in other members of the hot dog-fold thioesterase superfamily, which have been shown to accept a range of diverse acyl-CoA substrates with physiologically relevant  $k_{\text{cat}}/K_{\text{M}}$  values (50–55). Because fluoroacetyl-CoA is larger than acetyl-CoA (estimated total size: C–F, 2.82 Å; C–H, 2.29 Å (11)), we might have expected that the binding pocket of FIK would also accommodate acetyl-CoA as a substrate, especially considering that several of the substrate-binding interactions with CoA-dependent enzymes involve the carbonyl group and the cofactor itself (50, 54).

Based on the results of the competition experiment, it seemed likely that the  $K_{\rm M}$  for acetyl-CoA is above 1 mM, the highest concentration tested in previous studies (41). After both enzyme and substrate concentrations were increased, acetyl-CoA turnover was detectable with a rate well above the nonenzymatic hydrolysis rate (Supporting Information Figure S4). Under these conditions, we measured a  $k_{\rm cat}$  of  $0.06 \pm 0.001~{\rm s}^{-1}$  and a  $k_{\rm cat}/K_{\rm M}$  of 30 M<sup>-1</sup> s<sup>-1</sup> for acetyl-CoA, which corresponds to a  $10^6$ -fold decrease in catalytic efficiency compared to the fluoroacetyl-CoA substrate ( $k_{\rm cat}$ =390 ± 20 s<sup>-1</sup>;  $k_{\rm cat}/K_{\rm M}$ =5 ×  $10^7$  M<sup>-1</sup> s<sup>-1</sup>) (Table 1). The

combination of the slow rate of turnover and low pseudo-second-order rate constant with acetyl-CoA indicates that the nonfluorinated substrate binds poorly and, potentially in unproductive conformations, at the FIK active site. In contrast, the kinetic parameters measured for fluoroacetyl-CoA are similar to the values observed for other members of this superfamily for their substrates ( $k_{cat}$ , 0.56–190 s<sup>-1</sup>;  $k_{cat}/K_M$ , 10<sup>4</sup>–10<sup>7</sup> M<sup>-1</sup> s<sup>-1</sup>) (50, 51, 53–58). Taken together, the kinetic data for fluoroacetyl-CoA and acetyl-CoA are consistent with selection against acetyl-CoA as a substrate. Given that the magnitude of the discrimination exhibited by FIK is outside the window expected of single C–H to C–F substitutions that do not perturb ligand conformation or local p $K_a$ s (8, 9, 11), we turned to structural studies of FIK and FIK mutants in order to further elucidate the interactions at the active site that might promote binding and hydrolysis of the fluorinated substrate while excluding its nonfluorinated congener.

#### Crystal Structure of FIK

A set of FIK structures were recently reported (42), with FIK found to be arranged as a dimer in a typical hot dog-fold thioesterase topology (50, 51, 54, 58). We have also determined the crystal structure of the wild-type enzyme at 1.9 Å resolution by molecular replacement using a polyalanine model of T. thermophilus hypothetical protein TTHA0967 (PDB ID 2CWZ) as a search model (Figure 3A, Table 2). The asymmetric unit contained two copies of the FIK monomer, in agreement with the apparent molecular weight measured by size-exclusion chromatography (Supporting Information Figure S2). A BLAST search (59) of FIK followed by multiple sequence alignment using MUSCLE (60) suggests that the FIK active site is composed of Thr 42, Glu 50, and His 76. Indeed, these residues are found at the dimer interface and arranged with a topology similar to the Arthrobacter clade of hot dog-fold thioesterases (Figure 3B) (50). Although FIK adopts the fold that defines the superfamily, the structural data reveal some unique features not observed in structures of other superfamily members. The most striking difference is a "lid" structure formed by short helical segments (a1 and a2, residues 23-26 and 31-34, respectively) connected by flexible linkers, which is positioned directly over the active site of FIK and constrains the size of the active site pocket available for substrate binding (Figure 3C). Tertiary contacts within the lid are mediated by an edge-face interaction between Phe 33 and Phe 36 and by close packing of the side chains of Val 23, Leu 26, Val 39, and Phe 36. The surface of the lid that faces the substrate-binding cavity is made up primarily of hydrophobic side chains (Val 23, Leu 26, Tyr 27, Phe 33, Phe 36, and Val 39). In combination with the side chains of Val 46 and Val 54 from helix  $\alpha$ 3 and Ile 72 from strand  $\beta$ 2, the lid creates a chemical environment within the active site that is largely hydrophobic with the exception of the residues involved in nucleophilic/general base catalysis or water activation. The hydrophobic residues of the lid and those lining the bottom of the active site are well conserved in uncharacterized FIK homologues identified by sequence alignment (Supporting Information Figure S5), although one group of less closely related homologues lacks helix a 2 and the second loop of the lid.

Structural alignment of FlK to other structurally characterized thioesterases with the same active site topology revealed the conspicuous absence of a conserved Asn/Gln whose position in the tertiary structure of FlK approximately corresponds to Val 23 (Supporting Information Figure S6). In these other thioesterases, the carboxamide of the Asn/Gln is proposed to function in orientation and polarization of the thioester C=O and to assist in departure of the thiolate leaving group (50, 51, 54). Based on the crystal structure of FlK, there does not appear to be another hydrogen bond donor to compensate for this function. The absence of the Asn/Gln–carbonyl oxygen interaction in FlK could contribute to the lowered catalytic efficiency for acetyl-CoA compared to fluoroacetyl-CoA both by decreasing binding interactions with a shared structural motif and by selecting against hydrolysis of the less reactive acetyl-CoA substrate.

# Mutagenesis Studies of the Catalytic Residues of FIK

Although the detailed catalytic mechanism for the hot dog-fold thioesterases still requires further elucidation, Ser/Thr, Asp/Glu, His catalytic triads have been proposed and characterized by mutagenesis for many members of this superfamily (50, 54, 57, 58). From their sequence conservation (Supporting Information Figure S5) and location in the active site pocket of FIK (Figure 3B), FIK appears to contain a similar catalytic grouping comprised by Thr 42, Glu 50, and His 76. A previous report has excluded Glu 50 as a catalytic residue based on its location in the active site pocket as well as an increase in  $k_{\rm cat}$  observed for the E50A mutant, leading to a proposal of an alternative catalytic triad comprised of Thr 42, His 76, and an active site water (42). However, the positioning of Glu 50 on the third turn of the hot dog helix ( $\alpha$ 3) is consistent with the position of the Asp/Glu in hTHEM2, 4-HBA-CoA thioesterase, and PaaI from *E. coli* (51) (Supporting Information Figure S6) and is within hydrogen-bonding distance of the catalytic Thr.

To test whether Glu 50, His 76, and Thr 42, are involved in FlK catalysis, we measured the kinetic constants for the E50Q, H76A, T42A, T42S, and T42C mutants (Table 1). On the basis of mutagenesis studies on several members of the superfamily, the Asp/Glu and Ser/ Thr residues have both been implicated as potentially involved in assisting water attack at the carbonyl (50, 54, 57, 58). Mutation of Glu 50 in FIK to Gln resulted in a significant 3000-fold decrease in  $k_{\text{cat}}$ , which is consistent with an important role for Glu 50 in catalysis. However, the rate of hydrolysis of fluoroacetyl-CoA catalyzed by FlK-E50Q remains over 2000-fold higher than that of the pseudo-first-order rate of the uncatalyzed reaction at pH 7.6. We should note at this time that the E50Q mutant could only be solubly expressed as aMBP fusion protein. FlK-E50Q remained soluble after removal of the MBP tag; however, a relatively large proportion (~60%) of the mutant eluted from a size-exclusion column in the void volume, suggesting the formation of a soluble aggregate. After protein concentration, we verified that the fraction corresponding to the dimer peak did not revert to soluble aggregate by size-exclusion chromatography after the steady-state kinetic studies were completed (Supporting Information Figure S7). Although we performed our kinetic measurements using only the dimeric form of the enzyme, we cannot rule out the possibility that some fraction of this sample was not correctly folded.

Replacement of His 76 with Ala resulted in a substantial  $10^5$ -fold decrease in  $k_{\text{cat}}$  to a rate acceleration of only 20-fold over the pseudo-first-order rate for the uncatalyzed reaction. In comparison, alanine replacement of His 65 in 4-HBA-CoA thioesterase (50) and of His 56 in hTHEM2 (54) resulted in only 4000-fold and 2-fold reductions in  $k_{cat}$ , respectively. The effect of this mutation on the overall rate of the reaction is much greater than the effect of the E50Q mutation and may point toward a more important role for His 76 in FlK than in other characterized members of the superfamily. Consistent with the observed defect in  $k_{\text{cat}}$ , FIK-H76A was not competent to confer fluoroacetate resistance in a heterologous expression system (Figure 1). As discussed previously, His 76 is located at the dimer interface, but it is not within hydrogen-bonding distance of Glu 50 (Figure 3B). Based on a homology model of the FlK-substrate complex (Supporting Information Figure S8), His 76 appears to be positioned closer to the thioester sulfur atom. Thus, it is possible that His 76 helps to replace the function of the conserved Asn/Gln found in other hot dog-fold thioesterases by assisting departure of the thiolate leaving group with less carbonyl polarization in FIK. Because fluoroacetyl-CoA is more activated toward nucleophilic attack, this change in mechanism could contribute to the large observed difference in the  $k_{\text{cat}}$ parameter between fluoroacetyl-CoA and acetyl-CoA.

We next turned our attention to examining the role of Thr 42 in catalysis by substitution with Ala, which led to a 900-fold reduction in  $k_{\text{cat}}$ . Although the  $k_{\text{cat}}/K_{\text{M}}$  value measured for the T42A mutant is similar to the E50Q mutant, the 15-fold increase in  $K_{\text{M}}$  may indicate that

this residue participates in both catalysis and recognition of the substrate. Due to speculation that FIK might achieve its selectivity by using a secondary alcohol nucleophile (41, 42), we also characterized the T42S and T42C mutants to further probe the role of nucleophilicity or basicity in the FIK reaction. The T42S and T42C mutants show 20- and 35-fold decreases in  $k_{\text{cat}}$ , respectively, but differ with regard to their response in  $k_{\text{cat}}/k_{\text{M}}$ . As  $k_{\text{cat}}/k_{\text{M}}$  is an order of magnitude lower for the T42C mutant, it is likely that the structural perturbations at the active site are greater in this mutant and do not allow us to conclusively determine the trend in reactivity for Thr 42 in catalysis.

To further probe the role of Thr 42, we compared the kinetic parameters with respect to both fluoroacetyl-CoA and acetyl-CoA for the T42S and T42C mutants (Table 1). In , the relative decrease in  $k_{\text{cat}}$  for acetyl-CoA was smaller than the decrease in  $k_{\text{cat}}$  observed with respect to fluoroacetyl-CoA. However, the most striking effect of these mutations is that neither the T42S nor the T42C mutant shows a substantial difference between fluoroacetyl-CoA and acetyl-CoA in the  $K_{\rm M}$  parameter. Interestingly, the  $K_{\rm M}$  value for acetyl-CoA in the T42S mutant is within range of other characterized hot dog-fold thioesterases for their substrates. If we assume that the same kinetic mechanism is valid for both fluoroacetyl-CoA and acetyl-CoA, comparison of  $K_{\rm M}$  values implies that discrimination in substrate binding has been lost in both the T42S and T42C mutants. In terms of  $k_{\text{cat}}/K_{\text{M}}$ , the discrimination between the fluorinated and nonfluorinated substrates is decreased from 10<sup>6</sup>-fold in the wildtype enzyme to 200-fold for the T42S mutant and to 250-fold for the T42C mutant, and the observed discrimination is accounted for almost entirely by the difference in  $k_{\text{cat}}$  between the two substrates. One possible explanation for this is that the removal of steric bulk at this position in the T42S and T42C mutants could allow for structural rearrangements that introduce new interactions with acetyl-CoA, enhancing FIK's affinity for this substrate. In support of this hypothesis, it is interesting to note that the recently reported structure of FIK-T42S in complex with acetyl-CoA shows increased flexibility at the active site as determined by the observed population of two different rotamers of Ser 42 in the crystal structure (42). Although no new interactions with acetyl-CoA were found in the structure, the observed flexibility raises the possibility that a second hydrogen bond donor is accessible to the acetyl-CoA carbonyl group that could account for the observed decrease in  $K_{\rm M}$ .

#### Mutagenesis Studies of the Putative Substrate-Binding Pocket of FIK

We next turned our attention to examining the role of the hydrophobic lid residues identified in the putative substrate-binding pocket (Figure 3C). Based on structural alignments with ligand-bound forms of 4-HBA-CoA thioesterase from Arthrobacter sp. (50) and hTHEM2 (54) (Supporting Information Figure S6), the fluorine atom would be located in a largely hydrophobic environment surrounded by Phe 33, Phe 36, Val 23, Leu 26, Val 54, and Pro 37 (Supporting Information Figure S8), which are all conserved or conservatively substituted with bulky, hydrophobic residues in closely related FIK homologues of unknown function (Supporting Information Figure S5). Mutation of Phe 36 to alanine led to a 100-fold decrease in  $k_{\text{cat}}/K_{\text{M}}$  with very little perturbation of  $k_{\text{cat}}$  for fluoroacetyl-CoA (Table 1). For the F33A, V23A, and L26A mutants, the magnitude in the decrease of  $k_{cat}/K_{M}$  (F33A, 2800fold; V23A, 100-fold; L26A, 900-fold) was larger than the impact on  $k_{cat}$  for fluoroacetyl-CoA (F33A, 14-fold decrease; V23A, 2.5-fold decrease; L26, 58-fold decrease). Interestingly, the kinetic constants with respect to acetyl-CoA for both the F36A and V23A mutants remained relatively unchanged, which is consistent with an important function for these residues in discrimination between the fluorinated and nonfluorinated substrates. Notably, all of these residues are part of the FlK-specific lid structure. Although the lid residues have been found to be conserved in several uncharacterized homologues, Phe 36 is unique and not found in other sequences published to date (Supporting Information Figure S5).

We next looked to characterizing the role of hydrogen-bonding and dipolar interactions in the substrate-binding pocket. Arg 120 is the only polar residue found in the active site pocket besides the catalytic triad and was previously proposed to function as a hydrogen bond donor to the fluorine atom (42). Mutation of Arg 120 to Ala, Gln, and Lys all led to formation of highly insoluble protein with no dimeric protein observed by size-exclusion chromatography, which implies that the 1-point hydrogen bond that this highly conserved residue shares with Glu 50 in apo FlK is important to the structural integrity of the dimer. Except for the conservative E500 mutation, all other mutants tested of Glu 50 and Arg 120 resulted in insoluble protein that could not be purified in dimeric form. In addition to any enthalpic contributions of  $C-F \cdot \cdot \cdot H-N$  dipolar interactions, it is also possible that only substrates that can replace the hydrogen-bonding interaction of Arg 120 with Glu 50, such as fluoroacetyl-CoA, can maintain the dimer interface and bind well in the active site pocket. Attempts to engineer a hydrogen bond donor for the carbonyl oxygen of the acyl-CoA substrate as observed in other members of the superfamily by replacement of Val 23 with As n and Gln yielded protein with little or no activity toward either fluoroacetyl-CoA or acetyl-CoA. These results suggest that the specifics of active site packing and configuration may have diverged between FIK and other superfamily members without allowing us to directly address the role of the putative missing hydrogen bond.

In light of the kinetic constants measured for fluoroacetyl-CoA and acetyl-CoA for the F36A, F33A, and V23A mutants, a trend emerges in which removal of hydrophobic groups near the substrate-binding pocket decreases the  $k_{\text{cat}}/K_{\text{M}}$  for fluoroacetyl-CoA by 2–3 orders of magnitude while leaving the  $k_{\text{cat}}/K_{\text{M}}$  for acetyl-CoA relatively unchanged. The magnitude of this decrease is consistent with both the difference in  $K_{\rm M}$  with respect to fluoroacetyl-CoA and acetyl-CoA measured for wild-type FlK (Table 1) and the results from competitive inhibition studies (Figure 2). Although these data suggest that the hydrophobic groups are involved in specific recognition of the fluorine atom by FlK, a survey of the chemical environments of the fluorine atom in protein crystal structures with a bound fluorinated ligand has previously shown that binding of a fluorine atom in a lipophilic pocket does not typically confer advantages in terms of binding affinity on this scale (8). It has been observed, however, that fluorinated compounds often have negative entropies of aqueous solvation due to the tendency of water molecules to order around the hydrophobic C–F unit (11). In particular, monofluorination at the α-carbon of carbonyl-containing compounds typically increases their lipophilicity as determined by octanol-water and nonanol-water partition coefficients (7). The hydrophobicity of the fluoroacetyl-CoA binding pocket could thus promote fluoroacetyl-CoA binding over acetyl-CoA binding by providing a more lipophilic environment into which fluoroacetyl-CoA can partition, resulting in the release of water and an entropic driving force for binding. Furthermore, any enthalpic dipolar interactions with the C-F unit would be favored in a more apolar environment because organic fluorine does not compete well with stronger hydrogen bond acceptors. As it therefore seems inconsistent with previous studies that these residues are utilized mostly to make enthalpically favorable interactions with the fluorine substituent in the FlK binding pocket, we propose that the a polar, aromatic nature of the substrate-binding pocket provides an entropic driving force for substrate binding by promoting release of ordered water molecules from the C-F unit.

#### Crystal Structure of the FIK-Fluoroacetate Complex

We solved the structure of the FIK-fluoroacetate complex at 2.3 Å resolution to further investigate fluoroacetyl-CoA binding to FIK (Table 2). Seven fluoroacetate molecules were trapped in the structure (Figure 4), all located either in the putative active site or along the channel where CoA is expected to bind based on homology (Supporting Information Figure S8). The fluoroacetate molecules bound closest to the putative catalytic residues are bound

in different orientations at the two active sites, suggesting that their binding may differ from how the fluoroacetyl unit is bound in the substrate complex. Strikingly, a large conformational change occurs in the lid region of one active site in the product complex, involving the swinging aside of the unique Phe 36 and twisting of Phe 33 to open a channel in the protein between the active site and solvent that is blocked by Phe 36 in the apo structure (Figure 4). These movements are consistent with recently reported structures of the wild-type FlK bound to fluoroacetate and pantothenate-based substrate analogues (42). Our structure is distinctive, however, in that we observe a 3.3 Å interaction between a fluoroacetate fluorine atom and Phe 33, suggesting a possible mechanism for disruption of the edge-face interaction between Phe 33 and Phe 36 and gate opening. The orientation of the fluoroacetyl group that leads to the disruption would not be possible in the substrate complex as it would force the CoA chain to occupy the same space as helix a3 from either chain A or chain B. This observation raises the possibility that the Phe 33-Phe 36 gate could be relevant not only for maintaining the hydrophobicity of the active site for fluoroacetyl-CoA binding but also potentially for release of fluoroacetate. The movement of Phe 33 and Phe 36 could facilitate fluoroacetate release either by opening up a channel though which fluoroacetate could exit or by allowing the entrance of water into the active site and the displacement and hydration of fluoroacetate. We were also able to determine the structure of a fluoroacetyl-CoA-soaked crystal at 2.0 Å resolution. Although electron density was clearly observed in the active site, it could not be unambiguously assigned to either substrate or products. The same "open gate" conformation was trapped in this structure, which supports the hypothesis that the observed conformational changes are relevant to the normal catalytic cycle of FlK.

#### Crystal Structures of FIK-F36A and of the FIK-F36A Product Complex

To explore the role of ordered water and Phe 36 in substrate discrimination, we solved the crystal structures of apo-FIK-F36A at 2.3 Å resolution and of the FIK-F36A product complex at 2.0 Å resolution (Table 3). The overall folds of both structures do not appear to be significantly perturbed by the F36A mutation and are in good agreement with the corresponding wild-type structures with a core rmsd of 0.5 Å for both apo-FIK-F36A and the product complex of FIK-F36A. Notably, the channel that is normally occluded by Phe 36 is unblocked in the apo FIK-F36A mutant structure and correspondingly appears to have a more highly aquated active site compared to wild-type enzyme (Figure 5). In both apo structures, water molecules occupy the space that would typically be occupied with the substrate based on homology. However, additional ordered water molecules were observed at the active site of FIK-F36A and found to form a continuous hydrogen-bonding network with the external waters that are usually blocked from the active site by Phe 36. One of these internal waters makes a 2.9 Å hydrogen bond with Arg 120, potentially competing with a favorable interaction with the fluorine atom. In contrast, three ordered water molecules were observed just outside of the Phe 36 "gate" in the wild-type FlK structure that were unable to interact with water molecules within the active site. We propose that the exclusion of water from the active site of the wild-type enzyme prevents solvation of residues that would otherwise make favorable interactions with the fluorine atom and promotes the release of ordered water molecules from the C-F unit, providing an entropic driving force for fluoroacetyl-CoA binding. The greater extent of the water network at the active site of FIK-F36A compared to wild type supports the hypothesis that its large  $K_{\rm M}$  defect with respect to fluoroacetyl-CoA arises from a diminished ability to exclude water from the active site.

We also solved the crystal structure of the FIK-F36A product complex in order to further examine the role of solvation of the fluorine atom itself in the increased  $k_{\text{cat}}/K_{\text{M}}$  of the F36A mutant (Figure 6). A  $2.5\sigma$  peak was observed in the  $F_0 - F_c$  solvent/ligand omit electron density map for fluoroacetate and for the first eight atoms of the CoA chain. The orientation

and position of the ligands differ significantly from the positions of pantothenate- derived fluoroacetyl-CoA analogues bound in two recently reported structures of FlK (42). Dias et al. have speculated that the ligands are bound in positions that are irrelevant to catalysis due to the absence of the nucleotide moiety of CoA, which is believed to be important in correct binding of the substrate. However, the positions of fluoroacetate and CoA in our structure are in agreement with the positioning of substrates and products in structural homologues of FIK (Supporting Information Figure S6) and confirm the location of the fluorine atom in the hydrophobic pocket, which is consistent with kinetic data implicating these lid residues in substrate recognition. The product-bound structure also reveals that the fluorine atom is indeed within 3.1 Å of the Arg 120 side chain and within 3.7 Å of the Gly 69 backbone amide, in agreement with the computational model generated by Dias et al. (42). Consistent with previously identified fluorine-arginine interactions, the plane of the C-F bond is oriented perpendicular to the plane of the guanidinium side chain in the product structure rather than in a linear hydrogen bond geometry. While hydrogen bonding with organic fluorine is considered rare or weak (8, 61, 62), these specific dipolar C-F · · · H-N interactions are abundant in protein structures with bound fluorinated ligands and are also thought to explain the "fluorophilic" character of arginine (8, 63). Indeed, 32 structures were found in a search of the Protein Data Bank for C-F interactions with the guanidinium side chain (8). As no analogous interactions would be possible for acetyl-CoA, these fluorinespecific interactions could make enthalpic contributions to the observed substrate selectivity. In addition to these interactions, the product structure reveals a 3.3 Å hydrogen bond between the backbone NH of Thr 42 and the thioester C=O and also shows that the second hydrogen bond equivalent to that provided by an Asn/Gln residue in structural homologues is indeed missing. In this structure, the fluorine atom appears to be solvated as it is within 2 Å of a chain of hydrogen-bonded water molecules that would likely be excluded from the substrate-binding pocket by Phe 36 in the closed conformation of FlK, potentially explaining the large observed defect in  $K_{\rm M}$  for this mutant. The product complex structure is thus consistent with a model in which the active site lid and Phe 36 in particular aid in discrimination between fluoroacetyl-CoA and acetyl-CoA by actively excluding water from the active site.

## **CONCLUSIONS**

We have demonstrated that FIK is a fluoroacetyl-CoA-specific thioesterase that exhibits a  $10^6$ -fold higher catalytic efficiency for fluoroacetyl-CoA compared to acetyl-CoA. Based on experiments in a heterologous *E. coli* host, the *in vitro* selectivity of FIK also appears to be relevant *in vivo* and is consistent with the estimated intracellular concentration of acetyl-CoA (64), which is lower than the measured  $K_{\rm M}$  for acetyl-CoA. Based on kinetic measurements, the remarkable discrimination between the fluorinated and nonfluorinated substrates by FIK arises from both differences in substrate recognition ( $k_{\rm cat}/K_{\rm M}$ ) and chemical reactivity ( $k_{\rm cat}$ ) related to the fluorine substituent. We hypothesize that the more significant part of substrate selectivity is derived from mechanistic differences between FIK and other members of the superfamily that serve to select against reaction with unactivated acyl-CoAs, such as acetyl-CoA, because of the larger defect in the  $k_{\rm cat}$  (~10<sup>4</sup>) compared to the  $K_{\rm M}$  (~10<sup>2</sup>) parameter. Using X-ray crystallographic and biochemical studies on wild-type and mutant FIKs, we have identified several components of this selectivity and explored their role in preferential hydrolysis of fluoroacetyl-CoA.

We propose that FIK has evolved a mode of substrate recognition that is distinct from the more promiscuous hot dog-fold thioesterases by minimizing interactions with the carbonyl unit common to all acyl-CoAs to select against acetyl-CoA while recouping the resulting decrease in binding energy through C–F specific properties of fluoroacetyl-CoA. This working hypothesis is supported by structural information that indicates that a key Asn/Gln

hydrogen bond to the thioester carbonyl is missing in conjunction with biochemical data comparing fluoroacetyl-CoA and acetyl-CoA hydrolysis in the T42S and T42C mutants. Our data suggest that the specificity of FlK in binding fluoroacetyl-CoA is aided in part by hydrophobic interactions with the lipophilic binding pocket and by C-F · · · H-N dipolar interactions between the fluorine atom and Arg 120. Because neither of these enthalpic contributions would be expected to account for the observed 10<sup>6</sup>-fold selectivity between the fluorinated and nonfluorinated substrates given the existing literature on protein/ organofluorine binding affinities, we propose that entropic contributions related to ordered water play a larger role in fluorine-based substrate discrimination. In support of this model, we have biochemically and structurally characterized a mutant of a unique phenylanine (Phe 36) on the hydrophobic "lid" of FlK and have shown that defects in water exclusion from the active site lead to a 100-fold decrease in fluoroacetyl-CoA binding affinity without affecting  $k_{\text{cat}}$ . We have also identified key active site residues and propose an alternative catalytic triad consisting of Thr 42, Glu 50, and His 76. Based on the low  $k_{\text{cat}}/k_{\text{uncat}}$ measured for the H67A mutant, we further hypothesize that general acid catalysis by His 76 may be sufficient to assist CoA departure in the more activated fluoroacetyl-CoA substrate, but not acetyl-CoA, in the absence of typical interactions that serve to polarize the thioester carbonyl.

# **Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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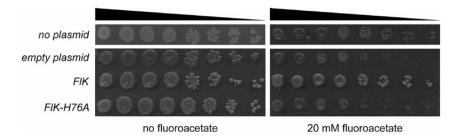
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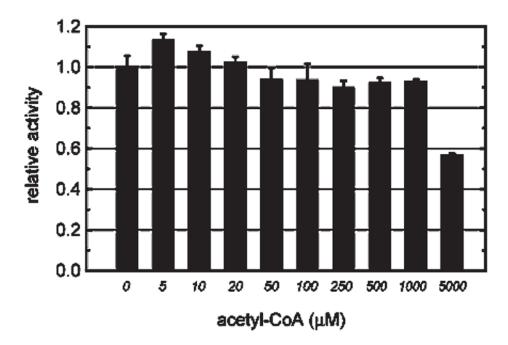
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**Figure 1.** FIK confers fluoroacetate resistance *in vivo*. The viability of *E. coli* BL21(DE3) was assayed by spotting 1:10 serial dilutions of culture onto solid media in the absence (left) or presence (right) of 20 mM fluoroacetate. *E. coli* harbored no plasmid (top row), empty pET23a-His<sub>10</sub>-Tev (second row), pET23a-His<sub>10</sub>-Tev-FIK (third row), or pET23a-His<sub>10</sub>-Tev-FIK-H76A (fourth row).



**Figure 2.** Rates of FIK hydrolysis of fluoroacetyl-CoA in the presence of varying concentrations of acetyl-CoA. The ability of acetyl-CoA to bind to FIK was tested using a competition assay in which the rate of hydrolysis of fluoroacetyl-CoA (fixed at  $20 \mu M$ ) was measured in the presence of increasing concentrations of acetyl-CoA. Data are mean  $\pm$  SE (n= 3).

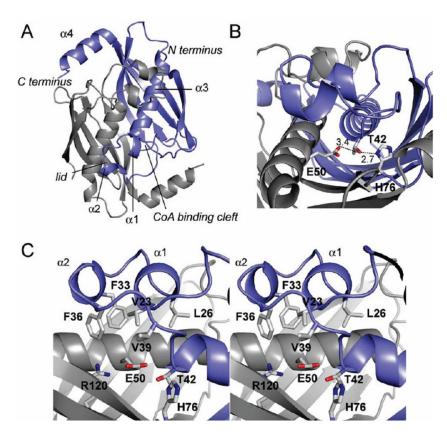


Figure 3. Crystal structure of FIK. (A) Cartoon representation of the FIK crystal structure looking down at the active site and lid structure formed by  $\alpha 1$  and  $\alpha 2$  (residues 23–26 and 31–34, repectively). Chain A is colored in blue, and chain B is colored in gray ( $\alpha 3$ , residues 42–58;  $\alpha 4$ , residues 125–135). (B) View of the hydrogen- bonding network at the putative active site showing Thr 42, His 76, and Glu 50 site water. (C) Stereoview of the active site and the hydrophobic lid (carbon, gray; nitrogen, blue; oxygen, red; sulfur, yellow).

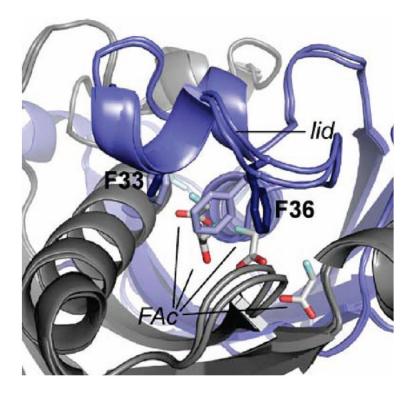
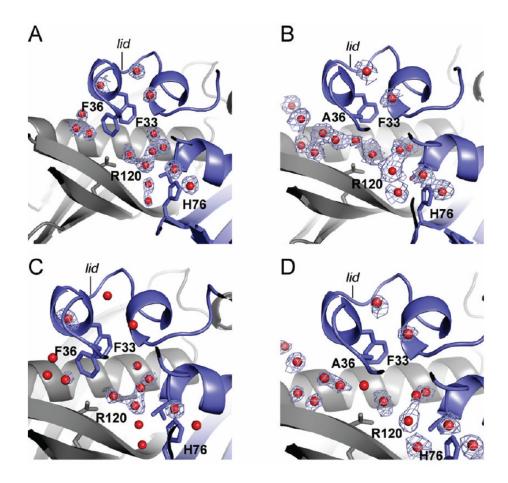
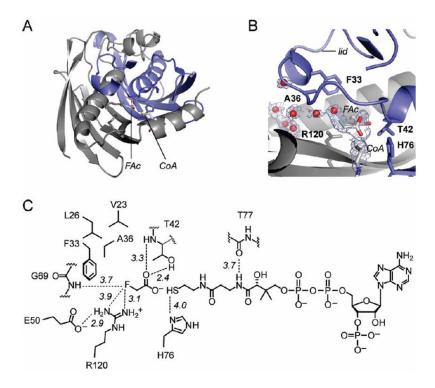


Figure 4. Conformational changes in the fluoroacetate complex of FIK. Chain A of both the unliganded and fluoroacetate-bound structures is shown in blue, and chain B of both structures is shown in gray. Phe 36 in shown in light blue for the closed conformation and dark blue for the open conformation. A total of seven fluoroacetate molecules were found in the active site (carbon, gray; oxygen, red; fluorine, light green).



**Figure 5.** Accessibility of the active site to water in FlK and FlK-F36A. Electron density maps are shown in light blue, chain A is colored in blue, chain B is colored in gray, and water molecules are shown as red spheres. (A)  $2F_0 - F_c$  map contoured at  $0.8\sigma$  for water molecules at the active site of wild-type FlK. (B)  $2F_0 - F_c$  map contoured at  $0.8\sigma$  for water molecules at the active site of FlK-F36A. (C)  $F_0 - F_c$  solvent omit map contoured at  $2.5\sigma$  for wild-type FlK overlaid with the refined FlK structure. (D)  $F_0 - F_c$  solvent omit map contoured at  $2.5\sigma$  for FlK-F36A overlaid with the refined FlK-F36A structure.



**Figure 6.** FIK-F36A product complex. Chain A is colored in blue, and chain B is colored in gray. For clarity, only the fluoroacetate and CoA ligands are colored by atom (carbon, gray; nitrogen, blue; oxygen, red; fluorine, light green; sulfur, yellow). (A) A view of the product bound at the FIK-F36A active site shown in the context of the overall fold. (B)  $F_0 - F_c$  solvent/ligand omit map contoured at  $2.5\sigma$  overlaid with the refined FIK-F36A product complex structure. (C) Chem Draw figure showing interactions observed between FIK-F36A and the products.

**Chart 1.**Biogenic Organofluorine Natural Products Found in the Environment

Table 1

Kinetic Constants Measured for Fluoroacetyl-CoA and Acetyl-CoA Hydrolysis by Wild-Type and Mutant FIKs<sup>a</sup>

Weeks et al.

		fluoroacetyl-CoA			acetyl-CoA	
enzyme	$k_{\rm cat}~({ m s}^{-1})$	$K_{\mathrm{M}}\left(\mu\mathrm{M}\right)$	$k_{\rm cat}/K_{\rm M}~({\rm M}^{-1}~{\rm s}^{-1})~k_{\rm cat}~({\rm s}^{-1})$	$k_{\rm cat}~({\rm s}^{-1})$	$K_{ m M}$ ( $\mu$ M)	$k_{\rm cat}/K_{\rm M}~({\rm M}^{-1}~{\rm s}^{-1})$
wild type	wild type $(3.9 \pm 0.2) \times 10^2 = 8 \pm 1$	8 ± 1	$(5\pm1)\times10^7$	$(6 \pm 1) \times 10^{-2}$	$(2.1 \pm 0.5) \times 10^3$ $(3 \pm 1) \times 10^1$	$(3\pm1)\times10^1$
E50Q	$(1.3\pm0.1)\times10^{-1}$	$(1.3\pm0.1)\times10^{-1}$ $(1.4\pm0.1)\times10^{1}$ $(9\pm1)\times10^{3}$	$(9\pm1)\times10^3$			
H76A	$(3.0\pm0.3)\times10^{-3}$	$(3.0\pm0.3)\times10^{-3}  (2.0\pm0.7)\times10^{1}  (1.5\pm0.5)\times10^{2}$	$(1.5\pm0.5)\times10^2$			
T42A	$(4.5\pm0.9)\times10^{-1}$	$(1.1 \pm 0.4) \times 10^2$ $(4 \pm 2) \times 10^3$	$(4\pm2)\times10^3$			
T42S	$(1.9 \pm 0.4) \times 10^1$ $(6 \pm 2) \times 10^1$	$(6\pm2)\times10^{1}$	$(3\pm1)\times10^5$	$(1.0\pm0.1)\times10^{-2}$	$(1.0\pm0.1)\times10^{-2}$ $(6.1\pm0.9)\times10^{1}$ $(1.6\pm0.3)\times10^{2}$	$(1.6\pm0.3)\times10^2$
T42C	$(1.1\pm0.2)\times10^{1}$	$(3.6\pm1)\times10^2$	$(3\pm1)\times10^4$	$(1.0\pm0.1)\times10^{-2}$	$(1.0 \pm 0.1) \times 10^{-2}$ $(3.9 \pm 0.9) \times 10^{2}$	$(1.2\pm0.6)\times10^{1}$
F33A	$(2.7\pm0.4)\times10^{1}$	$(1.5\pm0.4)\times10^3$	$(1.5 \pm 0.4) \times 10^3$ $(1.8 \pm 0.5) \times 10^4$	$^{\mathrm{hd}}b$	pu	
F36A	$(2.7\pm0.2)\times10^2$	$(7.1 \pm 0.8) \times 10^2$	$(7.1 \pm 0.8) \times 10^2$ $(3.7 \pm 0.5) \times 10^5$	$(1.0 \pm 0.2) \times 10^{-1}$ $(3 \pm 1) \times 10^3$	$(3\pm1)\times10^3$	$(2.9 \pm 0.9) \times 10^{1}$
V23A	$(1.5 \pm 0.3) \times 10^2$	$(3.0 \pm 1.5) \times 10^2$ $(5 \pm 3) \times 10^5$	$(5\pm3)\times10^5$	$(1.0 \pm 0.1) \times 10^{-2}$ $(9 \pm 1) \times 10^{2}$	$(9\pm1)\times10^2$	$(1.1\pm0.2)\times10^{1}$
L26A	$6.7\pm0.1$	$(1.2 \pm 0.2) \times 10^2$	$(1.2 \pm 0.2) \times 10^2$ $(5.7 \pm 0.8) \times 10^4$	pu	pu	

 $^{2}$ Data are mean  $\pm$  SE (n=3) as determined from nonlinear curve fitting. Error in the  $k_{\rm Cat}/K_{
m M}$  parameter was obtained from propagation of error from the individual kinetic terms.

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 $^b$  No detectable activity in the presence of up to 10  $\mu\mathrm{M}$  enzyme and up to 2.5 mM acetyl-CoA.

Table 2

Data Collection and Refinement Statistics for Wild-Type FIK Structures

	apo-FlK	FlK-FAc	FlK open
data collection			
X-ray source	ALS 8.2.2	ALS 8.3.1	ALS 8.3.1
space group	C2	C2	C2
cell dimensions			
<i>a, b, c</i> (Å)	48.4, 90.9, 62.0	62.3, 92.4, 50.3	62.8, 90.7, 52.3
$\alpha, \beta, \gamma(\deg)$	90.0, 100.1, 90.0	90.0, 102.0, 90.0	90.0, 104.5, 90.0
wavelength	0.9901	1.116	1.116
resolution $^a$	29.96–1.85 (1.90–1.85)	19.44–2.46 (2.50–2.46)	19.77–1.95 (2.00–1.95)
$R_{\mathrm{merge}}$ (%) <sup>a</sup>	9.0 (60.1)	10.6 (70.3)	8.5 (60.4)
I/σI <sup>a</sup>	12.25 (2.31)	12.94 (2.14)	14.01 (2.19)
completeness (%) <sup>a</sup>	99.7 (99.8)	99.5 (100)	99.7 (99.7)
redundancy <sup>a</sup>	4.1 (4.1)	4.1 (4.2)	3.7 (3.7)
refinement			
resolution	29.96-1.85	19.44-2.46	19.77-1.95
no. of reflections	20926	10125	20623
$R_{\rm work}/R_{\rm free}$	0.201/0.231	0.194/0.240	0.188/0.220
no. of atoms	2306	2195	2182
nonsolvent	2065	2103	2068
solvent	241	92	114
B-factors			
protein	29.2	38.3	23.4
water	43.4	39.0	30.2
rmsd			
bond lengths (Å)	0.014	0.005	0.013
bond angles (deg)	1.24	0.97	1.58

 $<sup>^{</sup>a}\!\mathrm{Values}$  in parentheses are for the highest resolution shell.

 Table 3

 Data Collection and Refinement Statistics for FIK-F36A Structures

	FIK-F36A	FIK-F36A products
data collection		
X-ray source	ALS 8.3.1	ALS 8.3.1
space group	C2	C2
cell dimensions		
<i>a, b, c</i> (Å)	141.9, 89.3, 71.4	141.0, 88.6, 70.9
$\alpha, \beta, \gamma(\deg)$	90.0, 118.0, 90.0	90.0, 118.0, 90.0
wavelength	1.116	1.116
resolution <sup>a</sup>	19.75–2.30 (2.48–2.30)	72.48–1.90 (1.95–1.90)
$R_{\rm merge}$ (%) <sup>a</sup>	12.6 (40.3) <sup>b</sup>	8.9 (66.3)
I/σI <sup>a</sup>	6.0 (2.5)	13.37 (2.22)
completeness (%) <sup>a</sup>	99.7 (99.4)	95.9 (90.9)
redundancy <sup>a</sup>	4.1 (4.2)	4.3 (4.1)
refinement		
resolution	19.75-2.30	72.48-2.00
no. of reflections	34432	48041
$R_{\rm work}/R_{\rm free}$	0.220/0.247	0.218/0.251
no. of atoms	6660	6351
nonsolvent	6092	6081
solvent	563	270
B-factors		
protein	33.7	32.6
water	38.2	40.1
rmsd		
bond lengths (Å)	0.018	0.006
bond angles (deg)	1.64	1.03

<sup>&</sup>lt;sup>a</sup>Values in parentheses are for the highest resolution shell.

bue to high noncrystallographic symmetry,  $R_{\text{pim}}$  is reported.