

ESTHER, the database of the α/β -hydrolase fold superfamily of proteins: tools to explore diversity of functions

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ABSTRACT

The ESTHER database, which is freely available via a web server (<http://bioweb.ensam.inra.fr/esther>) and is widely used, is dedicated to proteins with an α/β -hydrolase fold, and it currently contains >30000 manually curated proteins. Herein, we report those substantial changes towards improvement that we have made to improve ESTHER during the past 8 years since our 2004 update. In particular, we generated 87 new families and increased the coverage of the UniProt Knowledgebase (UniProtKB). We also renewed the ESTHER website and added new visualization tools, such as the Overall Table and the Family Tree. We also address two topics of particular interest to the ESTHER users. First, we explain how the different enzyme classifications (bacterial lipases, peptidases, carboxylesterases) used by different communities of users are combined in ESTHER. Second, we discuss how variations of core architecture or in predicted active site residues result in a more precise clustering of families, and whether this strategy provides trustable hints to identify enzyme-like proteins with no catalytic activity.

INTRODUCTION

The α/β -hydrolase fold family of enzymes described in 1992 (1) is rapidly becoming one of the largest groups of structurally related proteins with diverse catalytic and non-catalytic functions. The fold is composed of a β -sheet of eight strands with the second strand antiparallel to the others and ordered as 12435678. The prototype of enzymes in the fold has a catalytic triad composed of a nucleophilic residue located at the top of a γ -turn between

the fifth β -strand and the following α -helix (the nucleophile elbow), an acidic (Glu or Asp) residue and a His. Since the initial description of this fold, few reviews have dealt with the increasing number of functions performed by members of the family (2–5). The ESTHER database, dedicated to proteins with an α/β -hydrolase fold, was created in 1995 (6), and since then, it has been upgraded regularly (7–11), for example, with incorporation of new tools serving different fields of research, such as analysis of mutations in humans associated with diseases (12) or mutations in pests developing resistance to insecticides (13).

Catalytic members (enzymes) in this superfamily include hydrolases (acetylcholinesterase, carboxylesterase, diene lactone hydrolase, lipase, cutinase, thioesterase, serine carboxypeptidase, proline iminopeptidase, proline oligopeptidase, epoxide hydrolase) along with enzymes that require activation of HCN, H₂O₂ or O₂ instead of H₂O for the reaction mechanism (haloalkane dehalogenase, haloperoxidase, hydroxynitrile lyase) (14). Known non-catalytic members include the neurologins, glutactin, neurotactin, the C-terminal domain of thyroglobulin, yolk proteins, the CCG1-interacting-factor-B and dipeptidylaminopeptidase VI. Some proteins are suspected to display more than one function and could be true moonlighting proteins (15). As the vast majority of members have not yet been characterized experimentally, many more new functions could emerge in the least known subfamilies (e.g. the domains of unknown function, or DUF) in the future. The relative ease of production of enzymes from bacteria or fungi encouraged the use of protein engineering to modify their substrate specificity, thermostability and enantioselectivity for biocatalysis applications. These proteins are used in various processes ranging from the selective isolation of enantiomeric precursors for biomedical applications to industrial products, such as additives to detergents or decontaminants. This adaptability is a longstanding story for families such as the lipases used in washing agents for decades, but this

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may prove useful for many other subfamilies as recently exemplified by efficient transformation of butyrylcholinesterase into a cocaine esterase (16). Furthermore, the natural substrate promiscuity displayed by members of the superfamily can now be exploited (17–20). However, to be efficient, these engineering techniques need information about active sites and the range of sequence and structure variability found in nature. In combining classifications and tools to compare available data on α/β -hydrolases, ESTHER may provide the bases for most of these requirements. Herein, we describe the main entry point of the database, that is, the Overall Table, and show how Active Site search and Family Tree building programs are implemented to classify families.

OVERALL TABLE AND FAMILY TREE

In the Pfam database, which groups most α/β -hydrolases, the AB_Hydrolase clan (CL0028) is subdivided into 66 families (21). The ESTHER database contains far more families. The number increased from 69 in 2004 to 89 in 2008 to now reach 148 families, some of them being grouped in 47 superfamilies. This large number of families results from the following two features: firstly, because we subdivided some large families according to sequence similarities and biological information available from seed sequences; secondly, because we included some new families in the time intervals between major Pfam releases. Pfam, InterPro (22) or Uniprot (23) databases are not always fully convergent, and we often use the subdivision of the database that uses the most detailed subdivision. Some of the divisions are specific to ESTHER. In the new Overall Table, totally redesigned compared with the original version, the families are organized into three levels of grouping, so that there are blocks and families of rank number 1 and rank number 2. These subdivisions already give an idea of relationships between families. The Overall Table contains not only links to other general database resources but also buttons with pop-up windows showing taxonomic coverage, mutations, structures and substrates. Other buttons lead to specialized databases or classifications as described later in the text. To make it easier to compare protein functions in different families, we designed a method to built a Family Tree based on Hidden Markov Model (HMM) comparison (11). The Tree can be visualized, and it is fully interactive.

CORRESPONDENCE AND OVERLAP WITH OTHER ENZYME CLASSIFICATIONS OR DATABASES

The families described in ESTHER have corresponding entries in major general classification databases (PIR, InterPro, Pfam) and Proteopedia (24) and in Wikipedia when annotated entries are sufficiently curated. A few other specialized databases or published classifications also overlap with the classification of ESTHER. All of them have their specific purpose and serve different communities of biologists. In ESTHER, the Overall Table now provides links to these databases and a page summarizes all the correspondences.

Bacterial lipases

The first bacterial lipases to be recognized as members of the superfamily were enzymes homologous to the mammalian hormone-sensitive lipases (25,26). Later on, bacterial lipases and esterases were ordered in eight families (27). Since then, this classification has been widely used and extended (28–38), although it has not been integrated in any web server or database. We introduced this classification in ESTHER (Table 1). Among the bacterial lipases, six of the eight families originally described are related to α/β -hydrolases, but not the other two, families II and VIII, which belong to the SGNH-hydrolase and β -lactamase folds, respectively. It is noteworthy that in many and even small families, bacterial enzymes have eukaryotic homologues. The classification is extended, for example, family V has been subdivided in three subfamilies according to ESTHER families.

Carbohydrate esterases

The CAZy database (Carbohydrate-Active Enzymes Database: <http://www.cazy.org>) describes the families of structurally related enzymes that degrade, modify or create glycosidic bonds (39). Among those, 6 families of 16 correspond to proteins belonging to the α/β -hydrolase fold (Table 2). The others belong to diverse folds, (β/α)₇ barrel,

Table 1. Correspondence of ESTHER families with the Arpigny and Jaeger classification

ESTHER families	Arpigny and Jaeger families
Bacterial_lipase	Family_I
Bacterial_lip_FamI.1	Family_I.1
Bacterial_lip_FamI.2	Family_I.2
Bacterial_lip_FamI.3	Family_I.3
Lipase_2	Family_I.4
Bacterial_lip_FamI.5	Family_I.5
Bacterial_lip_FamI.6	Family_I.6
Lipase_2	Family_I.7
Not α/β -hydrolase SGNH	Family_II
Polyesterase-lipase-cutinase	Family_III
Hormone-sensitive_lipase_like	Family_IV (25,26)
Hormone-sensitive_lipase_like_1	Family_IV
ABHD6-Lip	Family_V.1
Carboxymethylbutenolide_lactonase	Family_V.2
UCP031982	Family_V.3 (30)
LYsophospholipase_carboxylesterase	Family_VI
Carb_B_Bacteria	Family_VII
(not α/β -hydrolase β -lactamase)	Family_VIII
PHAZ7_phb_depolymerase	Family_IX (31)
Bacterial_EstLip_FamX	Family_X.1 (32)
Fungal-Bact_LIP	Family_X.2 (33)
Lipase_3	Family_XI (34)
Bact_LipEH166_FamXII	Family_XII (35)
CarbLipBact	Family_XIII (32)
PC-sterol_acyltransferase	Family_XIV (36)
Duf_3089	Family_XV (37,38)
Bacterial_Est97	Family_XVI (39)

The original A & J classification included only eight families (27). Two families contain enzymes corresponding to a different fold. In 2003, the first family was subdivided in seven subfamilies (28). Families IX to XIV were added recently (31–39). Here, we added family XV and subdivided family V.

Table 2. Correspondence of ESTHER families with the CAZy classification of carbohydrate hydrolases

Families in ESTHER	Families in CAZy
Antigen85c	CE-1
Acetylxylan_esterase	CE-5
Cutinase	CE-5
Acetyl-esterase_deacetylase	CE-7
Carb_B_Bacteria	CE-10
Pectinacetyl-esterase-Notum	CE-13
Glucuronoyl-esterase	CE-15
Folds of families which are not α/β -hydrolases	
(β/α) ₇ barrel	CE-4
(β)-helix	CE-8
(β/α) ₈ barrel	CE-9
2-layer-sandwich	CE-11
PIG-L	CE-14
SGNH	CE-2, CE-3, CE-6, CE-12, CE-16

(β)-helix, (β/α)₈ barrel, 2-layer-sandwich, PIG-L, SGNH-hydrolase. At least in one family (CE10), the vast majority (if not all) of the members are esterases acting on non-carbohydrate substrates. The information on this family is no longer updated in CAZy, but in ESTHER, it represents Block C, which contains some of the most populated families.

Serine peptidases

The MEROPS database (<http://merops.sanger.ac.uk/>) is an information resource for proteolytic enzymes acting on peptide bonds. Each clan is identified with two letters, of which the first represents the catalytic type of the families included in the clan (40) and the second is an arbitrary second capital letter. Among the 12 clans of serine peptidases, only one clan (SC) contains α/β -hydrolases. Yet, this clan is one of the most populated clans, with six families (S9 prolyl oligopeptidase, S10 carboxypeptidase Y, S15 Xaa-Pro dipeptidyl-peptidase, S28 lysosomal Pro-Xaa carboxypeptidase, S33 prolyl aminopeptidase, S37 PS-10 peptidase) that are all α/β -hydrolase families (Table 3).

Thioesterases

ThYme (Thioester-Active Enzyme Database: <http://www.enzyme.cbirc.iastate.edu>) is dedicated to enzymes involved in fatty acid and polyketide synthesis along with enzymes acting on thioester-containing substrates (41). Among the nine enzyme groups, the Thioesterase group contains 8 of 25 families that belong to the α/β -hydrolase fold and, as such, have counterpart entries in ESTHER (Table 5). The correspondence between the ESTHER and ThYme families was presented earlier (42). We extended the Overall Table and introduced the corresponding links into ESTHER.

Lipases

LED (Lipase Engineering Database: <http://www.led.uni-stuttgart.de/>) is an internet database, which integrates

Table 3. Correspondence of peptidase families in ESTHER with clans and families in MEROPS

Families in ESTHER	Families in MEROPS
Peptidase_S9	S9 prolyl oligopeptidase
Prolyl_oligopeptidase_S9	S9A
S9N_PPCE_Peptidase_S9	S9A
S9N_PREPL_Peptidase_S9	S9A
ACPH_Peptidase_S9	S9C
DPP4N_Peptidase_S9	S9B
Glutamyl_Peptidase_S9	S9D
PMH_Peptidase_S9	S9_upw
Carboxypeptidase_S10	S10 carboxypeptidase Y
Peptidase_S15	S15 Xaa-Pro dipeptidyl-peptidase
Cocaine_esterase	S15
Lactobacillus_peptidase	S15
Prolylcarboxypeptidase	S28 lysosomal Pro-Xaa carboxypeptidase
Proline_iminopeptidase	S33 prolyl aminopeptidase
Peptidase_S37	S37 PS-10 peptidase

information on sequence and structure of lipases and related proteins sharing the same α/β -hydrolase fold to facilitate protein engineering (43). Although this database overlaps with many ESTHER families, no correspondences were available between these classifications. Table 4 now fulfills this need, and the corresponding links have been introduced into ESTHER.

VARIATION IN CORE ARCHITECTURE

An important variation of the canonical core architecture was revealed by the crystal structure of *Penicillium finiculosum* polyhydroxybutyrate depolymerase, where the sequences dictating β -strands 1–4 in the central β -sheet are circularly permuted from the N- to the C-terminal of the protein (44). In the patatin family, which is closely related to the α/β -hydrolase fold family (45), a recent crystal structure of the acyltransferase domain of the iterative polyketide synthase (DynE8) from *Micromonospora chersina* shows an even more drastic split of the β -sheet core (46). Indeed, the first four β -strands are separated from the rest of the β -sheet and serve as a linker between the ketoacyl synthase and acyltransferase domains of the protein. Such a split of the central β -sheet has not been described in α/β -hydrolases, but it might occur for members known only by their genomic sequence where large insertions are found between predicted β -strands.

VARIATION IN PREDICTED ACTIVE SITE RESIDUES

The conservation degree of amino acid residues in individual sequences against the aligned sequences of the family is now visualized in ESTHER with a colour code; the darker the shade, the higher the conservation. The putative active site residues are also highlighted when this information can be derived from the best-characterized homologues within the family.

Table 4. Correspondence of ESTHER families with the classification in LED

Families in ESTHER	Families in LED
Block C and H	Class GGGX
Carboxylesterase	abH01 - Carboxylesterase
Fungal_carboxylesterase_lipase	abH02 - Yarrowia lipolytica lipase like - 03
Fungal_carboxylesterase_lipase	abH03 - Candida rugosa lipase like
Hormone-sensitive_lipase_like	abH04 - Moraxella lipase 2 like
Hormone-sensitive_lipase_like	abH05 - Hormone sensitive lipases
Hormone-sensitive_lipase_like	abH06 - Brefeldin A esterase like
	Class Y
DPPIV_Peptidase S9	abH27 - Dipeptidyl peptidase IV like
Prolylendopeptidase	abH28 - Prolyl endopeptidases
Lactobacillus_peptidase	abH29 - Dipeptidyl-peptidases
Cocaine_esterases	abH30 - Cocaine esterases
Fungal-Bact_LIP	abH38 - Candida antarctica lipase A like
	Class GX
Lipase_3	abH23 - Filamentous fungi lipases
Thioesterase	abH07 - Moraxella lipase 3 like
Lipase_2	abH18 - Bacillus lipases
Bacterial_lipase	abH15 - Burkholderia lipases
Lipase_3	abH37 - Candida antarctica lipase like
Bacterial_Lipase	abH24 - Pseudomonas lipases
Acidic_lipase	abH14 - Gastric lipases
Pancreatic_lipase	abH20 - Lipoprotein lipases
Cutinase	abH36 - Cutinases
Polyesterase-lipase-cutinase	abH25 - Moraxella lipase 1 like
Epoxyde_hydrolase Proline imminopeptidase	abH09 - Microsomal Hydrolases
Epoxyde hydrolase Haloalkane dehalogenase	abH08 - Cytosolic Hydrolases
Non-heme peroxydase Carbon-carbon_bond_hydrolase	
CIB-CCG1-interacting-factor-B	
Thioesterase_acyl-transferase	abH35 - Acyl-transferases
Lysophospholipase_carboxylesterase	abH22 - Lysophospholipase
Lipase_2	abH16 - Streptomyces lipases
CarbLipBact	abH11 - Carboxylesterases
LYsophospholipase_carboxylesterase	abH21 - Bacterial esterases
Not α/β -hydrolase	abH32 - Xylanase esterases
Antigen85c	abH33 - Antigen 85
Carboxypeptidase	abH34 - Lysosomal protective protein like
Acetyl-esterase_deacetylase	abH26 - Deacetylases
Bacterial_esterase	abH13 - Bacterial esterases
Hydroxynitrile_lyase	abH12 - Hydroxynitrile lyases
Palmitoyl-protein_thioesterase	abH19 - Thioesterases
Dienelactone_hydrolase	abH31 - Dienelactone Hydrolases
PGAP1	abH17 - Chloroflexus aurantiacus lipase-like
AlphaBeta_hydrolase	abH10 - Uncultured crenarchaeote

The most common active site nucleophilic residue in α/β -hydrolase enzymes is a Ser located within a GX SXG pentapeptide (where X denotes any residue), whereas the other two active site residues are an acidic residue (Glu or Asp) and a His as a general acid–base catalyst. In dienelactone hydrolases, the nucleophilic residue is a Cys, and in haloalkane dehalogenases, it is an Asp. In glycoside hydrolases, there are nearly always two catalytic residues, both either Asp or Glu. Recently, in *E. coli* RutD a His has been found to substitute the canonical Ser, to form a GHALG pentapeptide in the nucleophile elbow (47). Even more surprising, in tomato methyl ketone synthase 1 (MKS1), a 3-ketoacid decarboxylase that clearly clusters in a subclass of the hydroxynitrile lyase family, the nucleophilic residue is Ala87, central to the GHALG pentapeptide. This substitution seems to have occurred only recently in the *Lycopodium* lineage, as many close homologues from plants retain a Ser in their active site (48). Yet, the Ala87Ser mutant in MKS1

displays greater decarboxylation activity than the wild-type enzyme, arguing for a limited evolutionary history within a functionally constrained and structurally ‘highly conserved’ active site region. In all the active sites analysed, the only conserved residue is the His. However, this residue generally lies within a sequence that is not particularly well conserved between families, whereas it is often conserved in non-catalytic proteins. Hence, it cannot serve as a marker for inferring that a protein is an enzymatically active α/β -hydrolase.

However, even though extracting information from structural data for inferring the catalytic residues in all members of families remains arduous, we attempted to do so with the aim of improving the subdivision of families. For example, we used the occurrence of an Asp or a Glu as the putative active-site acidic residue to split the large family called Antigen85c in ESTHER and PF00756 Esterase in Pfam. As a result, this family is now separated in five subfamilies, three have an Asp

Table 5. Correspondence of ESTHER families with the thioesterase families in ThYme

Families in ESTHER	Families in ThYme
Acyl-CoA_Thioesterase	TE-2 (1) Acyl-CoA thioesterase (Acot) 1-6 (2) Bile acid-CoA amino acid N-acyltransferase (BAT) thioesterase
Thioesterase	TE-16 (1) TE domain of fatty acid synthase (FAS) or thioesterase I (2) TE domain of polyketide synthase (PKS) or non ribosomal peptide synthase (NRP), or type I thioesterases (TE I)
Thioesterase	TE-17 domain of polyketide synthase (PKS)
Thioesterase	TE-18 (1) S-acyl fatty acid synthetases/thioester hydrolases (Thioesterase II) (2) Type II thioesterase (TE II)
Thioesterase_acyl-transferase	TE-19 luxD
Palmitoyl-protein_thioesterase	TE-20 Palmitoyl-protein thioesterase (ppt1, ppt2)
Lysophospholipase_carboxylesterase	TE-21 (1) Acyl-protein thioesterase (apt1, apt2) (2) Phospholipase (3) Carboxylesterase
A85-EsteraseD-FGH	TE-22 (1) S-formylglutathione hydrolase (2) Esterase A (acetyl esterase) (3) Esterase D (carboxylesterase)

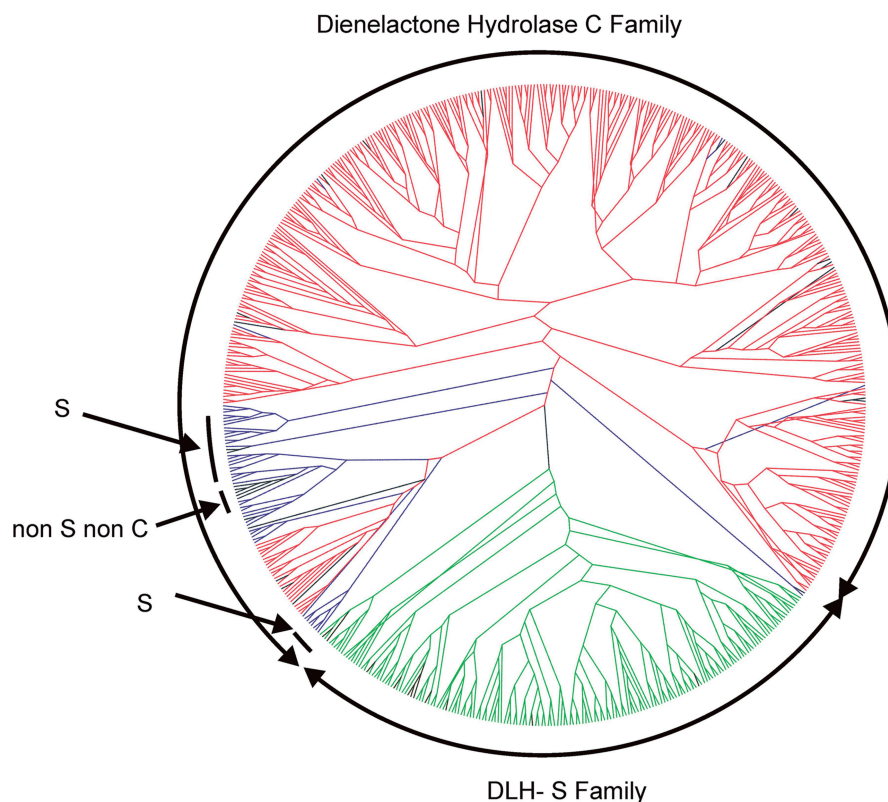


Figure 1. Sequences of putative dienelactone hydrolases where aligned with Clustal Omega (49). The Family Tree was built with FastTree 2 (50), and branches were coloured according to the nature of the nucleophilic residue in the active site. Two families were separated, in blue and red the Dienelactone_hydrolase family and in green the DLH-S family. Blue branches are sequences that have a Cys suitably positioned to be the nucleophilic active site residue, whereas green and red branches have a Ser. Black branches are sequences that have neither Cys nor Ser.

(A85-EsteraseD-FGH, A85-Feruloyl-Esterase, A85-Est-Putative) and two have a Glu (A85-IroE-IroD-Fes-Yiel, A85-Mycolyl-transferase). Another example is diene-lactone hydrolase, whose active-site nucleophilic residue is a Cys; yet, a recent crystal structure of a related *Anabaena variabilis* enzyme shows a Ser in the homologous position (Protein Data Bank code, 2O2G). This sequence is annotated as a diene-lactone hydrolase, and many sequences issued from whole genome sequencing are annotated accordingly. We analysed together sequences from the original diene-lactone hydrolase family and sequences homologous to the newly annotated sequence and built a Family Tree from the aligned sequences (Figure 1). This procedure led us to clearly separate a family, which we called DLH-S (with S for Ser). Although members of this family are annotated diene-lactone hydrolases, too little data are available for inferring a clear function for this new family. The Tree also shows some sequences truly homologous to well characterized diene-lactone hydrolases that have a Ser in the active site, or groups of sequences that have neither Ser nor Cys in homologous position of the active-site nucleophilic residue. Overall, this kind of analysis can help clarify the strategy and results from genome annotation and sequence analysis. We plan to refine, extend and standardize this approach to other ESTHER families of in the near future.

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