

Article

Screening of biocatalysts for synthesis of the Wieland-Miescher Ketone

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Abstract: Lipases, a versatile class of biocatalysts, have been shown to function in non-aqueous media/organic solvents and to possess promiscuous catalytic activity for a wide range of organic transformations. In this study, we explore the biocatalytic properties of a library of commercially available lipases by screening them for catalysis of a one-pot synthesis of Wieland-Miescher Ketone, an important intermediate in the synthesis of biologically active compounds such as steroids and terpenoids, from methyl vinyl ketone and 2-methyl-1,3-cyclohexanedione. As a direct outgrowth of this screen, we have created an optimized procedure for Wieland-Miescher Ketone (WMK) synthesis using crude lipase preparations, characterizing both reaction yield and enantiomeric excess. We have also identified principal components of the crude lipase mixture through proteomics and present evidence for a non-lipolytic origin of the observed catalysis. Finally, using the optimized conditions developed in this study, we propose a general absorbance-based screening methodology for assessing biocatalytic potential of crude enzyme preparations for synthesis of WMK.

Keywords: lipase; biocatalysis; Wieland-Miescher ketone; biocatalyst screening; amylase; peptidase; enantiomeric excess; Robinson Annulation

1. Introduction

Metal-based catalysts have revolutionized synthetic chemistry, enabling challenging chemical reactions with exquisite stereochemical control.[1,2] Disadvantages of their frequent use, however, include high cost and potentially adverse environmental effects. As a result, chemists have explored biologically derived catalysts as less expensive and less toxic alternatives. [1–4] In particular, the pharmaceutical industry has adapted biocatalysts to ‘green’ large scale synthetic pathways.[3] Numerous enzymes, for example, have been evolved to synthesize pharmaceutical compounds, or their intermediates, with high yields and enantiomeric excess, improving overall process efficiency and environmental compatibility. [4]

Lipases, a versatile class of biocatalysts, have undergone extensive study over the past two decades.[5] They have been reported to function in non-aqueous media/organic solvents, possess the catalytic ability for a wide range of organic reactions, and provide recyclability via immobilization.[1,5–7] In their native (biological) setting, lipases catalyze the hydrolysis of esters from triacylglycerides. However, when placed under minimal or no-water conditions, they can catalyze various types of organic transformations such as Michael additions and Aldol additions.[6,8] Catalysis of these non-native transformations by enzymes has been previously deemed “promiscuous”.[9,10] The catalytic promiscuity of lipases has been the foundation on which various studies have been conducted to demonstrate a broad spectrum of lipase catalyzed organic transformations including the Robinson annulation, the Knoevenagel condensation, and the Morita-Baylis-Hillman reaction.[7,11,12]

Despite intensive investigation of the lipase biocatalysis, unanswered questions remain regarding fundamental aspects of their propensity for promiscuous catalysis.[13] Some studies have identified crude lipase preparations with biocatalytic properties presuming, but not demonstrating, that lipase, in particular the lipase active site responsible for the *in vivo* hydrolysis of fatty acids (Fig. 1A-B), is also responsible for promiscuous biocatalytic behavior observed in organic or aqueous/organic solvent mixtures (Fig. 1C).[12,14] While in many instances the lipase activity present in crude lipase preparations is directly responsible for promiscuous catalysis, this is not always the case.[15,16] In the absence of a more thorough characterization of these catalytic moieties, it is challenging to improve on the initial findings through the now-standard means of rational design and/or directed evolution.

Our approach to the characterization of biocatalysts involves screening crude preparations for biocatalytic activity with mass spectrometry followed by reaction optimization (Fig. 1D). In addition, we utilized mass spectrometry for identification of the primary components of biocatalytic lipase preparations. This opens the possibility of improving reproducibility and further development of biocatalytic methods. In this work, we describe methods to screen lipase preparations for biocatalytic activity towards a synthetically useful target, the Wieland-Miescher ketone. We describe how initial results from a mass spectrometry-based screen can be improved through a process of reaction optimization. We then take lead catalysts from the screen and further characterize their proteinaceous components, identifying non-lipase proteins that are potentially responsible for the observed catalytic activity. The methodology and fundamental characterization of biocatalysts described in this report have general utility in the screening and identification of biocatalysts in numerous organic transformations.

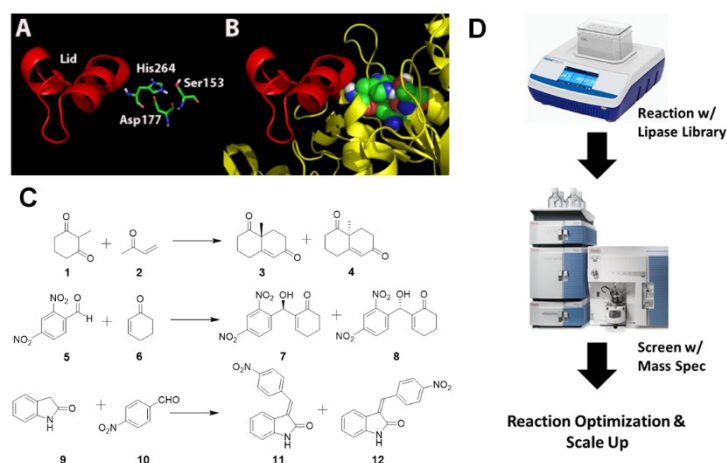


Figure 1. Porcine Pancreatic Lipase (PPL) is a model lipase for biocatalysis of C-C bond forming reactions. The active site (A; colored spheres) of PPL is presumed to catalyse the C-C bond forming reactions via carbonyl binding and activation in an oxy-anion hole. The lid domain (A; red ribbon) is presumed to undergo interfacial activation (assuming an open conformation in more hydrophobic solvents). B. Both lid domain and active shown in context of 3D protein ribbon structure (PDB ID 1ETH). C. Examples of carbon-carbon bond forming reactions catalysed by lipases (Robinson annulation, Morita-Baylis-Hillman, Knoevenagel condensation). D. Overview of screening methodology. Biocatalytic reactions were scaled down but otherwise conducted according to literature procedures. Mass spectrometry methodologies (LC/MS and MALDI) were used to screen the resulting biocatalytic reactions for product formation. Commercially available compounds were purchased for use as standards in mass spec calibration. Screening was then followed by reaction optimization and measurement of yield and stereoselectivity.

2. Results

We began by assembling a 14-member library of commercially available lipases preparations, selecting lipases previously reported to catalyze C-C bond forming reactions (Porcine pancreatic

lipase, *Candida antarctica* lipase B) [11,12,14,17], in addition to lipase preparations with unknown biocatalytic properties in promiscuous biocatalysis (Wheat germ lipase). Biocatalytic potential of the lipase library was then assessed for synthesis of the Wieland-Miescher ketone from methyl vinyl ketone and 2-methyl-1,3-cyclohexanedione using recently reported literature conditions. [12] In order to rapidly assess biocatalytic potential of the library, an LC/MS method was developed (Materials and Methods) that detected only the main fragmentation pattern Wieland-Miescher ketone product (Figure 2).

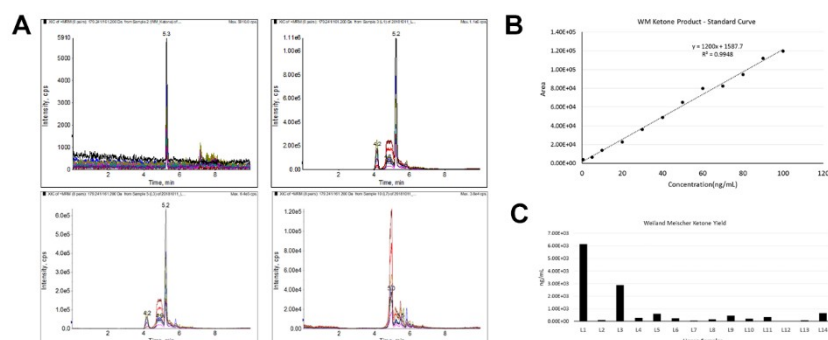


Figure 2. Development of an LC/MS method for biocatalyst screening. A. LC/MS data for standard and lipases L1, L3, and L7. B. Standard curve for Robinson Annulation product (3 & 4) generated with LC/MS method. C. Quantification of data for lipase library using standard curve from B. Reaction conditions: The following reagents/reactants were mixed in a centrifuge tube for synthesis of Wieland Miescher Ketone: methanol (1 mL), methyl vinyl ketone (81 μ L), 2-methyl-1,3-cyclohexanedione (23 mg), Deionized water (100 μ L), lipase (5 mg) After reactant assembly, centrifuge tubes were placed in a shaker (30 $^{\circ}$ C, 250 rpm) for 48 hours. Target mass (m/z) = 178.10.

Reaction yields were determined for each lipase using a standard curve (Fig. 2C and Table 1). While we were able to identify lipases (Entries 1, 3, and 5 from Table 1) with biocatalytic activity, the yields were less than 1%, in contrast to a literature report using these conditions.[12] As a result, we undertook optimization studies using our top hit from the screen (L1 - PPL) to attempt better reaction yields.

Table 1. Screening of Lipase Library for Synthesis of Wieland Miescher Ketone.

Entry	Lipase Code	Lipase Name	Yield(ng/ μ L) ^a
1	L1	Porcine Pancreatic Lipase (PPL)	6140
2	L2	Rhizopus Olyzae	92
3	L3	Wheat Germ Lipase	2880
4	L4	Candida Rugosa	269
5	L5	Aspergillus Niger	591
6	L6	Aspergillus Oryzae	242
7	L7	Pseudomonas Cepacia	59
8	L8	Candida Sp.	148
9	L9	Rhizopus Niveus	457
10	L10	Mucor Miehei	211
11	L11	Mucor Javanicus	331
12	L12	Candida Antarctica "B"	15
13	L13	Candida Antarctica "A"	77
14	L14	Pseudomonas Fluorescens	648

^a Yields determined by LC/MS.

In order to optimize this reaction, we turned to a previously described one-pot synthesis of WMK using L-proline in DMSO. [18] Therefore, in these initial optimization studies, we switched to DMSO as the primary solvent and included L-proline as a positive control. In the optimization

experiments, reaction yields were assessed using HPLC (Fig. 3C). We found that the PPL-catalyzed reaction in DMSO produced the triketone product 13 (Fig. 3A), whereas the L-Proline reaction formed the expected WMK products 3 & 4 (Fig. 3A). We hypothesized that L-Proline plays dual roles as a biocatalyst: one in which the amino acid promotes the necessary transition state for conversion of 13 to 14, and another in which the α -amino sidechain functions as a base to convert 14 to a mixture of products 3 & 4. Therefore, we reasoned that addition of a basic co-catalyst such as imidazole might better mimic the bi-functional L-Pro catalysis. We found that a combination of PPL and imidazole was able to produce a modest yield of 3 & 4 (Figure 3 and Table 2). By contrast, in reactions in which only PPL was present, or in which only imidazole was present, no WMK product was detected via HPLC (Fig. 3B and Table 2). Interestingly, the combination of BSA and imidazole also resulted in an appreciable yield of product (Table 2).

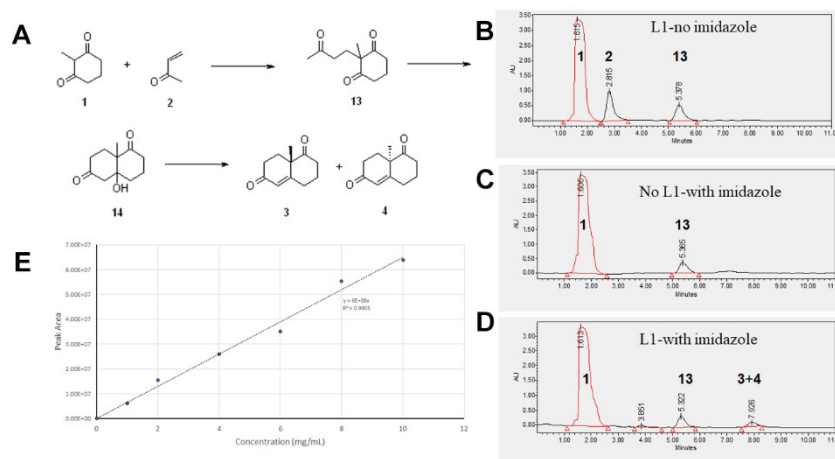


Figure 3. Investigation of reaction products in the presence and absence of imidazole and lipase preparation. A. Starting materials 2-methyl-1,3-cyclohexanedione (1) and methyl vinyl ketone (2) produce triketone intermediate (13), which proceeds through bicyclic species (14) to WMK products (3 & 4). B. HPLC chromatogram of reaction sample with L1 but no imidazole. C. HPLC chromatogram of reaction sample with imidazole but no L1. D. HPLC chromatogram of reaction sample with both L1 and imidazole. E. Standard curve generated using HPLC data obtained from commercially available Wieland-Miescher ketone. Reaction conditions taken from [18].

Table 2. Addition of Imidazole as Cocatalyst for Reaction Optimization.

Entry	Biocatalyst	Cocatalyst(Imidazole)	Yield(%) ^a
1	-	-	n.r.
2	-	+	n.r.
3	PPL (5 mg)	-	2.6
4	PPL (5 mg)	+	14.4
5	BSA (5 mg)	-	n.r.
6	BSA (5 mg)	+	13.4
7	L-Pro (6.4 mg)	-	68.8

^a Yields were determined by HPLC; n.r. = no reaction/product not detected by HPLC.

2.1. Reaction Optimization: Catalyst loading, solvent effects, enantiomeric excess.

Having identified the components necessary for forming modest amounts of WMK from protein and co-catalyst, we sought to further optimize the reaction by focusing on catalyst loading and solvent system. Setting up a series of reactions in DMSO, we varied catalyst loading from 5 mg to 35 mg/mL, finding that reaction yield increased up to 30 mg of catalyst per 1 mL reaction volume, but decreased with amounts in excess of 30 mg (Table 3; Fig. 4A). We then investigated reaction yields in solvents other than DMSO. Reactions in methanol produced similar yields to DMSO, yet reactions with increasingly longer chain alcohols (ethanol, propanol, butanol, pentanol) steadily decreased

reaction yields (Table 3). Enantiomeric excess of the Wieland-Miescher ketone products produced with the lipase/imidazole system in DMSO were substantially lower than those produced with L-Proline (Figure 4), whereas performing the PPL reaction in MeOH slightly improved enantiomeric excess. Reactions performed in longer chain alcohols also resulted in progressively lower ee values (Table 3, Lines 6 – 9).

Table 3. Determination of Optical Purity for Reactions with PPL and Imidazole .

Entry	Biocatalyst	Solvent	Yield(%) ^a	Ee(%) ^b
1	L-Proline	DMSO	68.8	70.1
2	PPL-5 mg	DMSO	14.4	9.8
3	PPL-35 mg	DMSO	51.9	17.7
4	PPL-5 mg	Methanol	7.3	24.1
5	PPL-30 mg	Methanol	55.1	21.7
6	PPL-30 mg	Ethanol	48.3	10.3
7	PPL-30 mg	Propanol	35.3	8.3
8	PPL-30 mg	Butanol	28.7	6.1
9	PPL-30 mg	Pentanol	29.6	3.6

^a Yields were determined by HPLC.

^b Determined by chiral HPLC; S-configuration.

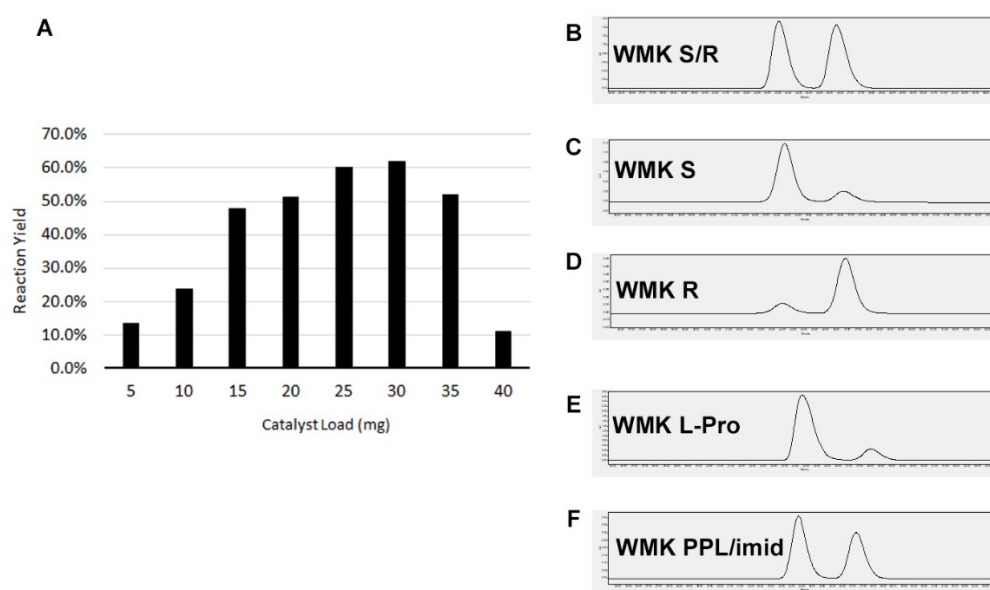


Figure 4. Effect of catalyst loading on reaction yield and determination of enantiomeric excess in L-Proline and PPL/imidazole catalyzed WMK reactions. A. PPL/imidazole reactions in DMSO with varying amounts of PPL catalyst. Chiral chromatograms are shown for B. racemic WMK standard; C. S-WMK enriched standard; D. R-WMK enriched standard; E. L-Proline catalyzed WMK product; and F. PPL/imidazole catalyzed WMK product.

2.2. Reaction Optimization: Exploration of co-catalysts

We next investigated the effects of co-catalysts other than imidazole on reaction yield. Dimethylaminopyridine (DMAP), which is more nucleophilic but less basic than imidazole, gave a substantially lower yield (Table 4, Entry 4). Alternately, ionic base co-catalysts sodium hydroxide and sodium bicarbonate gave less significantly reduced yields of WMK

Table 4. Screening Optimal PPL Loading with Different Cocatalyst and Solvents.

Entry	Cocatalyst	Solvent	Yield(%) ^a
1	Sodium Hydroxide	Methanol	43.0
2	Sodium Bicarbonate	Methanol	46.5
3	Imidazole	Methanol	55.1
4	DMAP	Methanol	24.5

^a Yields were determined by HPLC.

2.3. Determination of primary components of biocatalytic preparations.

Previous reports have demonstrated the presence of multiple biocatalytic entities present in commonly used lipase preparations.[19] Having optimized our reaction conditions with crude lipase preparations, we sought to more precisely define their composition. We separated the crude PPL mixture on a size exclusion column, isolating two primary components (Figure 5A). Bands were excised and subjected to proteolytic digest and subsequent characterization by mass spectrometry (see Materials and Methods). The higher molecular weight species was identified as porcine alpha-amylase, whereas the lower molecular weight species was identified as a porcine carboxypeptidase (Both carboxypeptidases A1 and B were present in the analysis). Surprisingly, neither of the principle components of this commonly used lipase preparation were porcine pancreatic lipase. This is significant, since numerous reports use this preparation of Porcine Pancreatic Lipase (Type II), and others, without further purification.[12,14,20,21] A second lipase from our library with biocatalytic activity for WMK production (L3; wheat germ lipase) was also characterized by this method (Fig. 5B; Fig. S1). Again, none of the principle components of this preparation were identified as wheat germ lipase.

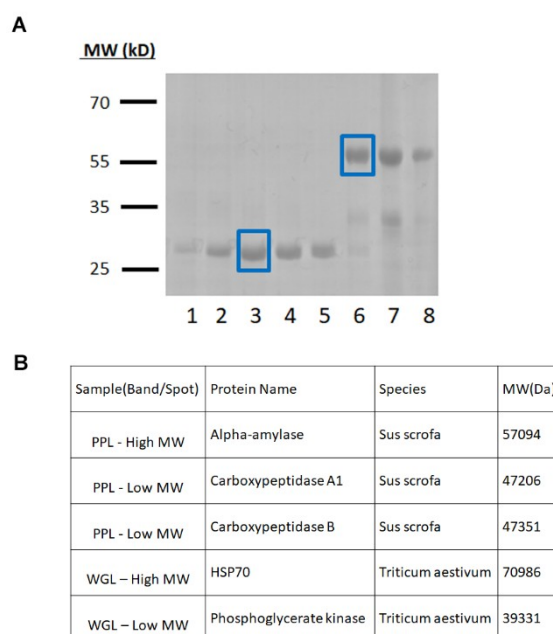


Figure 5. Identification of components of lipase L1 (Porcine Pancreatic Lipase) and L3 (Wheat Germ Lipase) preparations. A. PPL lipase preparation was purified using size exclusion chromatography. The fractions were also analysed using SDS-PAGE and two distinct bands were found to be present in the crude preparation of PPL and were subsequently characterized by mass spectrometry fingerprinting analysis. B. Peptides identified in the fingerprint analysis indicated that the band with lower molecular weight (PPL – Low MW; blue rectangle, Lane 3 in panel A) contained two proteins Carboxypeptidase A1 and Carboxypeptidase B) and the band with higher molecular weight (PPL – High MW; blue rectangle, Lane 6 in panel A) contained the protein Alpha-amylase. Electrophoretic mobility of the Low MW band corresponds to the mature (truncated) forms of Carboxypeptidase A1 and B. An identical analysis was conducted for the Wheat Germ Lipase (WGL) protein preparation, resulting in the identification of HSP70 and Phosphoglycerate kinase (only two of many *T. aestivum*

proteins present in this preparation. See Fig. S1.). The namesake “Porcine Pancreatic Lipase” or “Wheat Germ Lipase” were not identified as principal components in either analysis.

2.4. Reassessment of biocatalytic preparations with optimized reaction conditions.

Having identified optimized solvent, catalyst loading, and basic cofactor, we then rescreened our lipase library for biocatalytic activity towards WMK. In addition, as an outcome of this screen, we discovered that absorbance, in addition to mass spec, could be an effective means of screening for biocatalytic potential towards the WMK synthesis. In addition to select lipases from our initial screen, we also included commercially available proteins that were identified by proteomics as principal components of the crude PPL preparation (carboxypeptidase A (PA), carboxypeptidase B (PB), alpha amylase (P alphaS)) in addition to immobilized forms of *Candida antarctica* lipase B (L12-acr, L12-imo). By correlating absorbance (350 nm) of the crude reaction mixture with reaction yield (Fig. 6A), we demonstrate a correlation between biocatalytic potential and yield of WMK. Furthermore, plotting absorbance (350 nm) versus percent yield reveals that applying a colorimetric cutoff of $OD_{350} = 1.0$ would eliminate all of the reactions with yields below 10% (Fig. 6B). This indicates that colorimetric analysis could be a convenient basis for a large scale screen of biocatalysts for WMK production.

A more in-depth analysis of reaction yields from this screen reveals that the biocatalysts identified in the initial LC/MS screen (L1, L3, L5, L9, L14) also emerged as top biocatalysts in the colorimetric screen (Fig. 6 and Table 5). In addition, alpha amylase alone, which was identified as a primary component of the PPL preparation, produced a significant yield of WMK. The yield did not match that of the crude PPL preparation, indicating that multiple proteinaceous components, including porcine pancreatic lipase, carboxypeptidases, and amylases may be involved in the biocatalytic reaction. However, based on the ability of non-catalytic proteins (BSA) and individual amino acids (L-Pro) to promote product formation, we hypothesized that the endogenous catalytic activities of the component proteins were likely not contributing to product formation. To investigate this possibility, we then conducted our biocatalytic reactions in the presence of amylase and lipase inhibitors (Acarbose and orlistat, respectively; Table 6). In the presence of excess inhibitor, the yield of the PPL reaction was essentially unchanged. Similarly, the yield of the reaction with the amylase component was only modestly reduced in the presence of excess inhibitor. We also confirmed that intrinsic lipase activity of the PPL preparation was fully inhibited in the presence of orlistat by monitoring hydrolytic activity with p-nitrophenylpalmitate with and without inhibitor (Fig. S2). Based upon these results, we conclude that lipase activity is not responsible for the promiscuous catalysis observed with crude preparations of PPL. Furthermore, while the reaction can be catalyzed by multiple proteinaceous entities regardless of whether they possess intrinsic enzymatic activity, not all proteins are capable of catalyzing product formation.

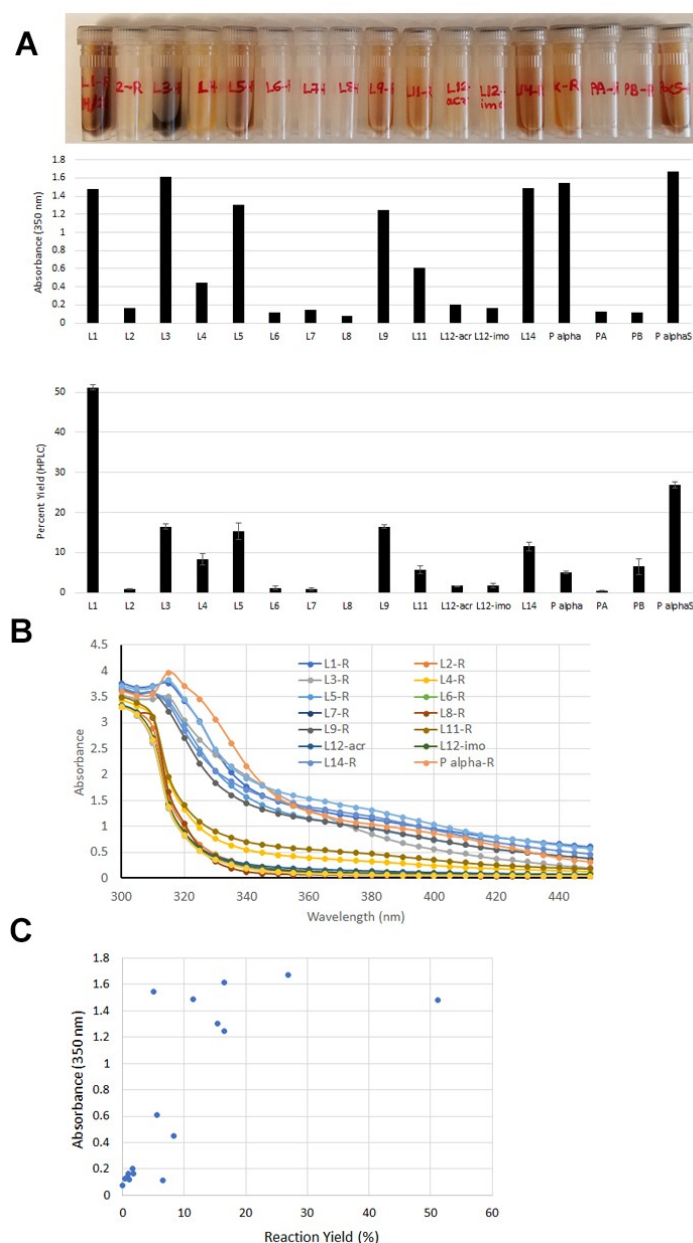


Figure 6. Rescreening of the lipase library under optimized conditions. A. Images of reactions after 89 hours (top panel); Absorbance of crude reaction mixtures at 350 nm (middle panel); Reaction yields from rescreening of each the lipase library (bottom panel). Error bars determined from three replicate HPLC injections. B. Wavelength scan of crude reaction absorbance. C. Plot of absorbance (350 nm) versus reaction yield.

Table 5. Rescreening of Lipase Library using Optimized Reaction Conditions.

Entry	Biocatalyst	Yield(%) ^a	ee(%) ^b
1	L1	51.9	21.7
2	L3	17.1	1.2
3	L5	13.5	19.9
4	L9	16.0	14.4
5	L14	10.9	(-)1.7
6	α -Amylase(solid)	26.4	19.3

^a Yields were determined by HPLC; n.r.= no reaction.

^b Determined by chiral HPLC (insert column information here); S-configuration.

Table 6. Addition of Catalyst Inhibitors

Entry	Biocatalyst	Inhibitor(Acarbose)	Inhibitor(Orlistat)	Yield(%) ^a
1	PPL	-	-	50.7
2	PPL	+	-	52.5
3	PPL	-	+	54.9
4	α -Amylase (solid)	-	-	31.0
5	α -Amylase (solid)	+	-	27.4
6	α -Amylase (solid)	-	+	27.6

^a Yields were determined by HPLC.

3. Discussion

In this study, we have screened lipase preparations for catalysis of a one-pot Wieland-Miescher ketone synthesis. Our initial screen detected minimal product formation in the presence of lipases alone. As a result, we turned to literature conditions used in the L-Pro catalyzed WMK synthesis for reaction optimization purposes. We found that inclusion of a basic co-catalyst was essential for promoting higher yields of WMK in the presence of a protein catalyst, in contrast to the L-Pro catalyzed reaction, which requires no co-catalyst. In addition, we used proteomics to characterize the primary components of a crude lipase mixture that has been previously used in numerous studies of biocatalysis. We anticipate that these results will inform future studies using this catalytic preparation. After further optimizing catalyst loading and solvent conditions, we then re-screened a lipase library, correlating absorbance of the crude reaction mixture with production of WMK product. This approach could easily be expanded to screen larger libraries of potential biocatalysts. Finally, we demonstrated that the innate enzymatic activities of the proteins present are not required for product formation. While the identification of the essential catalytic moieties remains incomplete (i.e., its shape complementarity?, surface exposed sidechains?), it is clear that future optimization work would be best focused on considerations other than intrinsic mechanisms of these promiscuous biocatalysts. For example, the poorer enantioselectivity of protein biocatalysts versus the individual amino acid L-Pro is intriguing, suggesting that the presence of a well defined active-site cavity contributes little to stabilization of the transition state. Yet, the advantages of a protein biocatalyst over an individual amino acid catalyst (more engineerable, more adaptable to immobilization, etc.) support future engineering efforts of these nascent catalytic entities. Furthermore, as not all lipase preparations tested supported promiscuous biocatalytic activity, there remain protein-specific features that remain to be discovered.

Promiscuity in biocatalysis remains an intriguing topic, often challenging chemists to think outside the comfort zone of well-established catalytic mechanisms. In this work, we have demonstrated that the assumptions regarding the role of intrinsic catalytic properties of promiscuous biocatalysts should be carefully reconsidered. In many studies, catalytic promiscuity has undoubtedly been misattributed to various properties of one enzyme or another, when, in actuality, the role of surface exposed amino acids, protein shape, or even influences on the dielectric environment may be more important. In order for promiscuous biocatalysts to fulfill their potential as inexpensive and reusable “green” reagents, it is critical that their modes of action be more precisely characterized. For example, in this study, we have identified alpha amylase as a key catalytic component of a crude lipase preparation from porcine pancreas. As a result, future studies can now be conducted on the optimization of this protein for promiscuous biocatalysis through rational design, directed evolution, or exploration of immobilization methodologies. Furthermore, this study establishes that colorimetric screening can be useful for identifying catalysts of WMK formation. In addition to this being a potentially effective platform for identifying new biocatalysts, we anticipate that this screen can be extended to other variations on the Robinson Annulation, enabling screening for substrate scope.

4. Materials and Methods

Chemicals, substrates, and solvents were purchased from commercial sources (Millipore-Sigma; TCI; VWR; and Fisher Scientific). Lipases, amylases, and carboxypeptidases were purchased from Sigma-Aldrich. Bovine serum albumin (BSA) was purchased from Fisher Scientific.

Lipase library components. Lipases were purchased from Sigma-Aldrich.

Lipase Code	Lipase	Catalog Number
L1	Porcine Pancreatic Lipase(PPL)	L3126
L2	Rhizopus Olyzae	62305
L3	Wheat Germ Lipase	L3001
L4	Candida Rugosa	L1754
L5	Aspergillus Niger	62301
L6	Aspergillus Oryzae	L4277
L7	Pseudomonas Cepacia	62309
L8	Candida Sp.	L3170
L9	Rhizopus Niveus	62310
L10	Mucor Miehei	62298
L11	Mucor Javanicus	L8906
L12	Candida Antarctica "B"	62288
L13	Candida Antarctica "A"	62287
L14	Pseudomonas Fluorescens	534730
L12-acr	Candida Antarctica "B"	L4777
L12-imo	Candida Antarctica "B"	52583
PA	Carboxypeptidase A	C9268
PB	Carboxypeptidase B	C9584
P alpha	Alpha amylase (suspension)	10102814001
P alpha S	Alpha amylase (powder)	A3176

LC/MS mass spectrometry method. Liquid chromatography (LC) was performed on a HPLC system (3200 Exion LC100; SCIEX Corporation). 5 μ L of samples were injected into the reversed-phase Gemini NX-C18 column (50 \times 2.0 mm, 3 μ m; Phenomenex, Torrance, CA, USA) by using auto sampler. The mobile phase composition for A was Water/Acetonitrile mixture(95/5, v/v) with 0.1% formic acid, and for B was Methanol. The gradient method was as follows: 100% A for 1 min, decrease to 15% over 4 min, hold at 15% for 1 min, increase back to 100% over 1 min, and hold at 100% for 3 min. The flow rate was 0.3 mL min⁻¹ and total run time was 10 min for each sample injection. The column temperature was kept at 35 °C. The LC elute was introduced into the Applied Biosystem Triple Quad API 3200, a triple-quadrupole tandem mass spectrometer equipped with a turbo spray ionization source, for quantification of compounds in positive ionization mode. Detection of the target molecule was performed in a multiple reaction monitoring (MRM) mode, and transition of *m/z* 179.24 to 161.20 was used for detection of WMK.

Biocatalysis general conditions. The following reagents/reactants were mixed in a microcentrifuge tube for synthesis of Wieland Miescher Ketone: Solvent (900 μ L), methyl vinyl ketone (19.8 μ L), 2-methyl-1,3-cyclohexanedione (20 mg), Deionized water (100 μ L), lipase (0 - 40 mg) or L-proline (6.4 mg), with or without imidazole (2.7 mg, 0.04 mmol). For reaction involving inhibitors, the biocatalyst was mixed with the inhibitor (5 mg) for 10 minutes at room temperature before adding the substrates and cocatalyst. After reaction assembly, capped microcentrifuge tubes were placed in a temperature-controlled shaker (New Brunswick Excella e24; 35 °C, 250 rpm) for 89 hours.

Cofactor and cosolvent optimization. The following reagents/reactants were mixed in a microcentrifuge tube for synthesis of Wieland Miescher Ketone: Alcohol solvent (900 μ L), methyl vinyl ketone (19.8 μ L), 2-methyl-1,3-cyclohexanedione (20 mg), Deionized water (100 μ L), lipase (L1, 30 mg), cocatalyst (0.04 mmol). After reaction assembly, capped microcentrifuge tubes were placed in a temperature-controlled shaker (New Brunswick Excella e24; 35 °C, 250 rpm) for 89 hours.

Lipase purification and proteomics. Solutions of porcine pancreatic lipase and wheat germ lipase were fractionated on a HiPrep 26/60 Sephacryl S-200 HR column connected to an Akta FPLC (GE Healthcare Life Sciences). Protein fractions were characterized via SDS-PAGE gels stained with SimplyBlue SafeStain (ThermoFisher). The bands of greatest intensity were excised and digested with trypsin overnight using a standard in-gel digestion protocol. The resultant tryptic peptides were desalted using a C18 Zip-Tip and applied to a MALDI target plate with α -cyano-4-hydroxycinnamic acid as the matrix. The sample was analyzed on an AB Sciex 5800 MALDI TOF/TOF mass spectrometer. MS/MS spectra were searched against Uniprot databases using Mascot version 2.3 (Matrix Science) within ProteinPilot software version 3.0 (AB Sciex). A significance score threshold was calculated in Mascot, with ion scores above the threshold considered positive IDs ($p < 0.05$).

Hi-Res Mass Spec analysis of triketone and WMK. Samples were analyzed with a Q Exactive HF-X (ThermoFisher, Bremen, Germany) mass spectrometer. Samples were introduced via a heated electrospray source (HESI) at a flow rate of 10 μ L/min. One hundred time domain transients were averaged in the mass spectrum. HESI source conditions were set as: nebulizer temperature 100 deg C, sheath gas (nitrogen) 15 arb, auxillary gas (nitrogen) 5 arb, sweep gas (nitrogen) 0 arb, capillary temperature 250 degrees C, RF voltage 100 V, spray voltage 3.5 KV. The mass range was set to 150-2000 m/z . All measurements were recorded at a resolution setting of 120,000. Solutions were analyzed at 0.1 mg/mL or less based on responsiveness to the ESI mechanism. Xcalibur (ThermoFisher, Bremen, Germany) was used to analyze the data. Molecular formula assignments were determined with Molecular Formula Calculator (v 1.2.3). All observed species were singly charged, as verified by unit m/z separation between mass spectral peaks corresponding to the ^{12}C and $^{13}\text{C}^{12}\text{C}_{-1}$ isotope for each elemental composition.

Characterization of triketone (2-methyl-2-(3-oxobutyl)-1,3-cyclohexanedione): Hi-Res mass spec: $m/z = 197.11761$ (mass error = 2.0 ppm) Assigned Chemical Formula: $\text{C}_{11}\text{H}_{17}\text{O}_3$ $[\text{M}+\text{H}]^+$; $^1\text{H-NMR}$ (400 MHz, CDCl_3): 2.60-2.77 (m, 4H), 2.35 (t, 2H), 2.11 (s, 3H), 2.04-2.08 (m, 2H), 1.89-1.95 (m, 2H), 1.25 (s, 3H).

Characterization of WMK ((S,R)-(+/-)-8a-methyl-3,4,8a-tetrahydro-1,6(2H,7H)-naphthalenedione): Hi-Res mass spec: $m/z = 179.10583$ (mass error = -4.6 ppm) Assigned Chemical Formula: $\text{C}_{11}\text{H}_{15}\text{O}_2$ $[\text{M}+\text{H}]^+$; $^1\text{H-NMR}$ (400 MHz): 5.85 (s, 1H), 2.67-2.76 (m, 2H), 2.38-2.52 (m, 4H), 2.09-2.19 (m, 3H), 1.65-1.78 (m, 1H), 1.45 (s, 3H).

HPLC analysis of reaction yield. Liquid chromatography was performed on a HPLC system (Waters Corporation) consisting of a binary pump (1525 Binary HPLC Pump), absorbance detector (2487 Dual Wavelength Absorbance Detector), and auto sampler (717-plus Autosampler). 5 μ L of sample were injected into the reversed-phase Xterra MS C18 column (4.6 \times 100 mm, 5 μ m; Waters, Milford, MA, USA) using the auto sampler. The samples were analyzed using an isocratic method: 80% Solvent A (water) and 20% Solvent B (methanol). The flow rate was 1.0 mL min^{-1} and the total run time was 13 minutes for each sample injection. The detector wavelength was set to 210 nm and the retention time for WMK was 7.9 minutes.

HPLC analysis of enantiomeric excess. Liquid chromatography for enantiomeric excess was performed on a HPLC system (Waters Corporation) consisting of binary pump (1525 Binary HPLC Pump), absorbance detector (2487 Dual Wavelength Absorbance Detector), and auto sampler (717-plus Autosampler). 5 μ L of sample were injected into the Chiralcel OD-H column (0.46 cm \times 25 cm; Daicel, Minato, Japan) using the auto sampler. Isopropanol/Heptane mixture (20/80, v/v) containing 0.1% formic acid was used as a mobile phase at a flow rate of 0.1 mL min^{-1} . Total run time was 111 minutes for each sample injection. The detector wavelength was set to 250 nm and the retention times for (S)-WMK and (R)-WMK were 43.5 and 46.3 minutes respectively.

5. Conclusions

In this work, we have characterized the biocatalytic potential of commercially available lipase preparations for catalysis of Wieland-Miescher Ketone synthesis. Optimization experiments demonstrated the importance of a basic co-catalyst for efficient production of Wieland-Miescher Ketone. Enantioselectivities were lower than a previously reported L-Pro catalyzed WMK synthesis.

We were also able to identify non-lipase proteins present in a commercially available lipase preparation with biocatalytic activity and establish a colorimetric screen for identification of biocatalysts. Taken together, these results promise for the engineering and characterization of optimized protein biocatalysts for the WMK synthesis.

Supplementary Materials: The following are available online at www.mdpi.com/xxx/s1, Figure S1: Survey of proteins identified from crude lipolytic preparation derived from Wheat Germ, Figure S2: Effect of inhibitors on lipolytic activity of a crude preparation from Porcine Pancreas.

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