

Tetrahedron report number 777

Synthetic applications of fluorous solid-phase extraction (F-SPE)

Wei Zhang^{a,*} and Dennis P. Curran^{b,*}

^aFluorous Technologies, Inc., University of Pittsburgh Applied Research Center, 970 William Pitt Way, Pittsburgh, PA 15238, USA

^bDepartment of Chemistry, University of Pittsburgh, Pittsburgh, PA 15260, USA

Received 12 June 2006

Available online 1 November 2006

Contents

1. Introduction	11838
1.1. Synthesis is reaction and separation	11838
1.2. Light fluorous chemistry	11838
2. Concept of F-SPE	11839
2.1. Classification of F-SPE	11839
2.2. Standard F-SPE	11839
2.3. Reverse F-SPE	11840
3. F-SPE methods	11840
3.1. Pressure and gravity F-SPE	11840
3.2. Common F-SPE systems	11841
4. Tabular summary of F-SPE	11842
5. Practical aspects of F-SPE	11860
5.1. A typical F-SPE procedure	11860
5.2. F-SPE demonstration with dyes	11860
5.3. Common issues related to F-SPE	11860
5.3.1. Loading solvents	11860
5.3.2. Elution solvents	11861
5.3.3. Fluorous silica gel reuse	11861
5.3.4. Gravity F-SPE with large fluorous silica gel	11861
6. Conclusions	11861
Note added in proof	11862
Acknowledgements	11862
References and notes	11862
Biographical sketch	11865

Keywords: Fluorous; Light fluorous; Fluorous silica gel; Fluorous solid-phase extraction (F-SPE); Reverse F-SPE; Plate-to-plate SPE; Parallel synthesis; High-throughput synthesis; Solution-phase synthesis; Separation; Reagents; Reactants; Catalysts; Chiral auxiliaries; Scavengers; Protecting groups; Displaceable tags; FluoroFlash[®]; VacMaster[®]; FlashMaster[™]; Ex-Block; RapidTrace[®]; Whatman[®].

Abbreviations: BINAP, 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl; BINOL, 1,1'-bi-2,2'-naphthol; Boc, *tert*-butyloxycarbonyl; BTF, benzotrifluoride; Cbz, benzyloxycarbonyl; CDMT, 2-chloro-4,6-dimethoxy-1,3,5-triazine; *m*-CPBA, *meta* chloroperbenzoic acid; DCT, 2,4-dichloro-1,3,5-triazine; dba, dibenzylideneacetone; DEAD, diethyl azodicarboxylate; DIPEA, diisopropylethylamine; DCC, dicyclohexylcarbodiimide; DBU, 1,8-diazabicyclo[5.4.0]undec-7-ene; dppp, 1,2-bis(diphenylphosphino)propane; DMF, *N,N*-dimethylformamide; DMAP, 4-dimethylaminopyridine; DMSO, dimethylsulfoxide; dppf, 1,1'-bis(diphenylphosphino)ferrocene; FC-72, perfluorohexanes; F-SPE, fluorous solid-phase extraction; FTI, Fluorous Technologies, Inc.; HFE-7100, perfluorobutyl methyl ether; HOAT, 1-hydroxy-7-azabenzotriazole; HOBT, 1-hydroxybenzotriazole; HPLC, high-performance liquid chromatography; LLE, liquid-liquid separation; MOM, methoxymethyl; μ w, microwave; NIS, *N*-iodosuccinimide; Oxone, potassium peroxydisulfate; Rf, perfluoroalkyl group; Rfh, perfluoroalkyl group with CH₂ spacer; TEA, triethylamine; TFA, trifluoroacetic acid; THF, tetrahydrofuran; THP, 2-tetrahydropyranyl; TLC, thin-layer chromatography; TMS, trimethylsilyl; TPP, triphenylphosphine.

* Corresponding authors. Tel.: +1 4126248240; fax: +1 4126249861 (D.P.C.); e-mail addresses: w.zhang@fluorous.com; curran@pitt.edu

1. Introduction

1.1. Synthesis is reaction and separation

The enterprise of organic synthesis is a collective endeavor involving reaction and separation followed by identification and analysis.¹ The ‘reaction and separation stage’ produces a new organic molecule while the ‘identification and analysis stage’ proves what the molecule is and how pure it is. The yield of every organic reaction depends on both the efficiency of the underlying reaction and the ability to recover the target product in pure form from the reaction mixture. Reaction methods have evolved and improved dramatically over the past decades and continue to do so. On the other hand, while they continued to be refined, the core separation methods that synthetic chemists routinely use have not changed for a decade or more.

The concept of the ‘ideal reaction’ serves as an inspiring, if unattainable, goal for basic research in synthesis.² In the trenches (the labs), reactions are usually a means to an end (chemical and drug discovery), and chemists tend to spend more time searching, hoping and wishing for the ‘ideal separation’.^{1a,b} It is rare that a new, generally applicable separation method comes along.

The purpose of this report is to provide an overview of the increasingly popular new separation technique of fluorosolid-phase extraction (F-SPE). Though it is still less than a decade old,³ the technique has matured rapidly and is now ready for prime time, as illustrated by the expanding uses over the last 2–3 years.⁴ After a brief introduction of two varieties of F-SPE—standard and reverse—we provide comprehensive tabular collections of published uses in small molecule synthesis that are intended to illustrate both the scope of the method and the diverse array of reagents and materials that are now available. We close by providing interested readers with practical information on how to conduct an F-SPE.

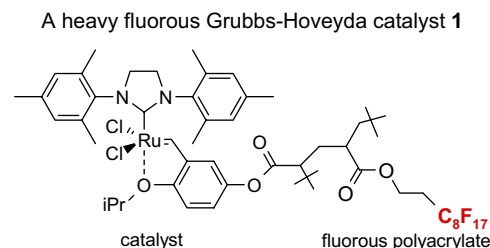
1.2. Light fluoros chemistry

The bifurcation of fluoros chemistry into ‘heavy’ and ‘light’ branches in 1999 was a direct result of the introduction of F-SPE. The earliest work in the fluoros field focused on introducing large (therefore ‘heavy’) fluoros tags onto organic and organometallic reaction components (catalysts, reagents, reactants, etc.).^{5,6} These tags then rendered the resulting tagged reaction components soluble in fluorocarbon and other highly fluorinated solvents, and enabled powerful techniques like fluoros biphasic and triphasic reactions, biphasic and triphasic liquid–liquid extractions (LLEs),⁷ thermomorphic reactions, and more.⁸ The exciting branch of heavy fluoros chemistry, whose techniques are especially suitable for large-scale processes, continues to forge ahead at a rapid pace today.

Heavy fluoros tags are often called ponytails because they usually sprout several fluoroalkyl chains bearing 39 or more (often many more) fluorines. From the outset, workers in the field were bent on giving these ponytails a haircut. The resulting light fluoros molecules (typically with 9–17 fluorines) are cheaper and more readily available, and are much

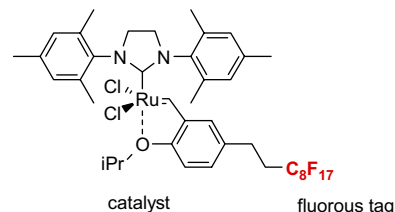
more soluble in common organic solvents. But therein lies the rub—they are also much less soluble in fluoros solvents so the LLE breaks down. The enabling advance for light fluoros molecules was the replacement of the LLE with an SPE.^{3,9,10}

Figure 1 compares and contrasts a pair of related heavy and light fluoros alkene metathesis¹¹ catalysts. The heavy fluoros catalyst **1** is a copolymer of a catalyst component and the fluoros acrylate.¹² (However, many molecular heavy fluoros catalysts are also known.) It is freely soluble in FC-72 but is insoluble in organic solvents like CH₂Cl₂ and EtOAc. Following a reaction with a substrate in C₆H₅CF₃–CH₂Cl₂ the catalyst is extracted away from the products with FC-72. Multiple cycles of recovery and reuse were conducted. In principle, it makes little difference that heavy catalysts like **1** have low or even no solubility in the organic



ratio catalyst/fluoros acrylate ~1/10
1 mmol 5.3 g, 61% fluorine
soluble in C₆F₁₄, C₆H₅CF₃;
not soluble in CH₂Cl₂, EtOAc

A light fluoros Grubbs-Hoveyda catalyst **2**



1 mmol = 1.0 g, 31% fluorine
soluble in BTF, CH₂Cl₂, EtOAc,
many other organic solvents;
not soluble in C₆F₁₄

Light catalyst **2** in action

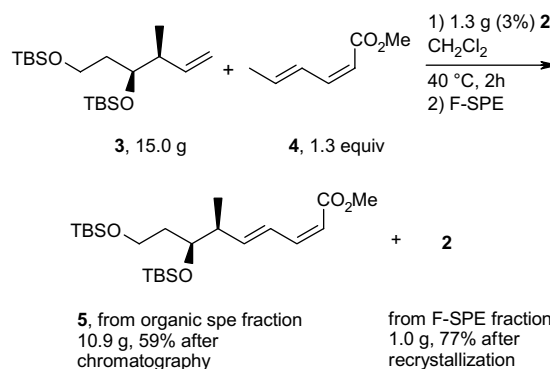


Figure 1. Comparing and contrasting light and heavy fluoros Grubbs-Hoveyda.

reaction solvent, so long as the reaction works. In practice, however, it makes a big difference, especially in small-scale discovery chemistry. Target reactions may fail either because they are too slow to occur at all or because other side reactions occur more rapidly. The rate of a target reaction often depends directly on the concentration of a reactant in the reaction solution—in other words, on its solubility. So having soluble reaction components is a huge advantage, especially when reactions are not understood in great detail, as is always the case with new reactions.

Light fluororous catalyst **2** has very different physical properties from heavy cousin **1**.¹³ A green crystalline solid, it is freely soluble in most common organic reaction solvents but has low solubility in fluororous solvents like FC-72. These properties are advantageous for running reactions since one can simply use the standard conditions for non-fluororous reagents without modification.

For example, cross metathesis of **3** and **4** has been conducted at University of Pittsburgh on large scale for a preparative total synthesis of analogs of the anti-cancer agent dictyostatin.¹⁴ In a typical run, 15 g of alkene **3** and diene **4** (1.3 equiv) were combined with 1.3 g of catalyst **2** (3%) in 50 mL of dichloromethane. The homogeneous mixture was warmed to 40 °C for 2 h and cooled prior to evaporation of the reaction solvent. The mixture was then subjected to F-SPE over 50 g of fluororous silica gel to provide an organic fraction of cross- and self-metathesis products. Purification of this fraction provided the cross-coupled product **5** in 60% yield. The fluororous fraction was primarily the recovered catalyst **2** and could be used as such; however, we prefer to recrystallize this product to ensure high catalyst quality for the next use in either the same or a different reaction. Recrystallization of the crude fluororous product from this reaction provided 1.0 g (77%) of recovered catalyst, which was of comparable appearance and purity to the original sample.

Comparing and contrasting this type of light fluororous reaction with traditional solution-phase methods and also solid-phase methods highlights the advantages of the fluororous approach.¹⁵ In reaction, identification, and analysis phases, the light fluororous approach resembles traditional solution-phase methods rather than solid-phase methods because the fluororous reaction components are molecules, not materials. Light fluororous molecules are often soluble in a broad range of standard organic reaction solvents and exhibit reactivity comparable to their non-fluororous parents.¹⁶ In other words, their reaction features are readily predicted. Fluororous molecules can be routinely analyzed by all standard spectroscopic methods and separated by both fluororous and non-fluororous techniques.

The advantage over traditional solution-phase chemistry comes at the separation stage, because the separation of fluororous compounds by F-SPE is a reliable and generic procedure that resembles more a filtration than a chromatography. The separation depends primarily on the presence or absence of a fluororous tag, not polarity or other molecular features that control traditional chromatography. In many respects, light fluororous methods capture the best features of traditional solution-phase chemistry, yet still provide a facilitated separation.

2. Concept of F-SPE

2.1. Classification of F-SPE

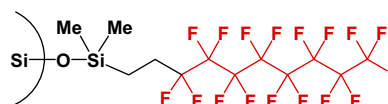
F-SPE separations can be grouped into two classes: standard and reverse. The standard or original solid-phase extraction is much more common and involves the partitioning between a fluororous solid phase and a fluorophobic liquid phase.⁴ This technique has been used in many settings (manual SPE, automated SPE, plate-to-plate SPE, automated flash chromatography, HPLC) and has proven generality. In contrast, the nascent technique of reverse fluororous solid-phase extraction¹⁷ uses a fluorophobic solid phase (standard silica gel) and a fluororous liquid phase. While there are currently only a few examples, the reverse F-SPE technique has considerable potential. Both techniques have been described in detail^{4,17} and we provide here a brief summary. Section 5 of the report on practical aspects is for those planning to use the standard F-SPE technique in the lab.

SPE¹⁸ and chromatography are related because both involve partitioning between solid and liquid phases, and the transition zone between the two techniques is grey. In chemical analysis, an SPE is usually used to help concentrate a very dilute sample; however, in synthesis it is used to partition a concentrated sample rapidly into two fractions. In general, the synthetic SPE resembles a filtration more than a chromatography, and has higher loading levels and lower solvent volumes. After elution of a first fraction with a first solvent, a second solvent of stronger eluting power is added to elute a second fraction. That second fraction is simply too well adsorbed on the solid phase to be eluted by the first solvent in a practical time frame. Multiple fraction collection and analysis are not required—there is one wash for unretained molecules and one wash for retained molecules. In chromatography, elution of successive fractions is typically a ‘time-dependent’ process—sooner or later, all of the fractions elute. In SPE, elution of successive fractions is a ‘solvent-dependent’ process.

Among different packing materials for SPE, ion-exchange resins have good retention selectivity, give ‘mass-controlled’ separation, and are little affected by the volume of the loading solvent. In contrast, polarity-based normal- and reverse-phase silica gels are much less selective and are more sensitive to the polarity and volume of the loading solvent.¹⁸ Fluororous silica gel has strong and selective fluorine–fluorine interaction with fluororous molecules.

2.2. Standard F-SPE

Standard F-SPE was introduced in 1997³ and involves the use of a fluororous solid phase and a fluorophilic (but not fluororous) solvent, as illustrated in Figure 2. The fluororous solid phase is typically silica gel with a fluorocarbon bonded phase ($-\text{SiMe}_2(\text{CH}_2)_2\text{C}_8\text{F}_{17}$), and this is commercially available from Fluororous Technologies, Inc. under the trade name of FluoroFlash[®].¹⁹



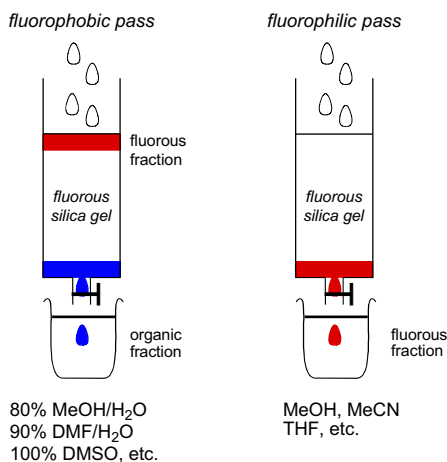


Figure 2. A cartoon of a 'standard' F-SPE. The organic fraction is blue and the fluororous fraction is red.

Briefly, a crude reaction mixture containing both fluororous and non-fluorous reaction components is charged onto fluororous silica gel and then the silica is eluted with a fluorophobic solvent like 70–80% MeOH–H₂O, 50–60% CH₃CN–H₂O, 80–90% DMF–H₂O, or 100% DMSO. In this 'fluorophobic pass', non-fluorous (organic) compounds typically move at or near the solvent front and elute immediately, while fluororous compounds are retained on the silica gel. In the ensuing 'fluorophilic pass', elution with one of many organic solvents (water-free MeOH or CH₃CN, THF, among others) then provides a fluororous fraction containing those compounds bearing the fluororous tag.

The procedure is simple, general and reliable, and has now been used many times in diverse settings. Importantly, it does not seem to be very sensitive to the polarity of either the fluororous component or the organic component. Thus, the standard F-SPE is a very attractive separation technique in library settings since products with very different characteristics will all exhibit substantially the same behavior.

2.3. Reverse F-SPE

Reverse F-SPE is a new technique with few examples to date,¹⁷ but with considerable potential. The polarities of the solid phase and liquid phase are reversed, and standard silica gel is used as the polar solid phase while blends of fluororous²⁰ and organic solvents are used as the fluorophilic liquid phase. The concept is illustrated in Figure 3. A sample containing fluororous and non-fluorous components is charged to regular silica gel with standard solvents, and then the silica is eluted with a fluorophilic solvent to remove a fluororous fraction (fluorophilic pass). In our first paper,¹⁷ we used FC-72 (perfluorohexanes) and ether, among other combinations, but we now more often use HFE-7100 (perfluorobutyl methyl ether) blended with ethyl acetate, ether or another organic cosolvent. Following that, a fluorophobic pass can be conducted with any standard organic solvent.

Perfluoroalkyl alkyl ethers like HFE-7100 are preferred over fluorocarbons because the fluorocarbons have very poor

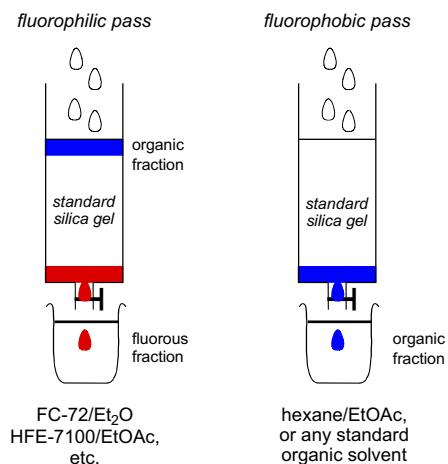


Figure 3. A cartoon of a 'reverse' F-SPE. The organic fraction is blue and the fluororous fraction is red.

eluting power (even for most light fluororous compounds) and they have limited miscibility in organic solvents. Because of this, the range of fluorophilic blends with such solvents is limited.

The reverse F-SPE is attractive for removing fluororous reagents, catalysts, and other byproducts from standard organic target products because after the fluorophilic pass is complete, the organic product absorbed on the head of the silica column can simply be purified by standard flash chromatography. It is also attractive because solvent elution conditions and prospects for success can be readily assessed by using standard silica gel TLC plates.

However, because standard silica gel is used, the behavior of reverse fluororous methods can be significantly affected by the polarity of both the fluororous and the non-fluorous components. While we still have limited experience, we currently feel that reverse F-SPE exhibits the most power when used to separate relatively nonpolar fluororous components from relatively polar organic components. Of course, such kinds of separations might also be conducted by standard chromatography with traditional solvents like hexane–EtOAc. However, the replacement of nonpolar solvent (hexane) of this combination with a fluororous solvent will often provide a better separation because this will significantly retard the elution of the organic fraction without retarding nearly as much (and perhaps even promoting) the elution of the fluororous component.

3. F-SPE methods

3.1. Pressure and gravity F-SPE

Depending on the type of fluororous silica gel that is used, F-SPEs can be driven with light pressure (positive or negative) or by gravity. Commercially available FluoroFlash[®] SPE silica gel²¹ has 40–60 μm particle sizes. Cartridges packed with this size silica gel require positive pressure on the top or negative pressure under the bottom to drive the elution process. When particle size is increased to around 120 μm, gravity SPE is possible.²²



Figure 4. 2×12 Vacuum SPE manifold.

3.2. Common F-SPE systems

A basic unit for conducting 1–24 SPEs shown in Figure 4 is commercially available from Supelco. Other companies such as Fisher and Waters sell comparable units. The unit is a 2×12 manifold and employs negative pressure, which is convenient for SPE cartridges with 2–10 g silica gel. Fractions are collected in a 10–15 mL test tube. For F-SPE with big cartridges (6–10 g silica gel), more than one tube is needed to collect both non-fluorous and fluorous fractions.

Since the F-SPE process is highly reproducible and functional group independent, it can be easily automated or used in a plate-to-plate format. For parallel synthesis, plate-to-plate F-SPE can significantly increase the throughput. Samples are loaded onto a plate whose cartridges are packed with fluorous silica gel and fractions are collected in a matched receiving plate. The silica gel plate may be cartridge- or well-formatted. Figure 5 shows a 24-cartridge plate and a 24-well plate of VacMaster® from Biotage.²³ Similar systems are also available from United Chemical Technologies, Supelco, and Waters. If 40 μm fluorous silica gel is used, then the receiving plate is connected to a vacuum pump. The 24-channel plate has the following technical



Figure 5. 24-Cartridge (left) and 24-well (right) SPE plates.

features: (1) each cartridge has 6 mL, and each well has 10 mL volume, which can be charged with 3–4 g of fluorous silica gel leaving ~3–5 mL top space for elution solvent; (2) each receiving well has 10 mL volume for collecting fraction; (3) a six-channel pipette is used for parallel sample loading and solvent loading; and (4) the Whatman® receiving plate^{24a} has a standard footprint, which can be directly concentrated in a Genevac vacuum centrifuge. The 24-well plate is good for parallel purification of 10–100 mg quantity of products. This system has been demonstrated in the purification of small libraries produced involving amine scavenging reactions with fluorous isatoic anhydride, amide coupling reactions with F-CDMT, and amide coupling reactions with a fluorous Mukaiyama condensation reagent.^{25a}

The 96-well F-SPE plate is more suitable for parallel synthesis of larger number but smaller quantity of samples. Figure 6 shows a pair of 96-well Ex-Block plates poised for plate-to-plate F-SPE.^{24b} Each well in the top block has 3 mL volume and is charged with 1 g of fluorous silica gel. The bottom receiving well also has 3 mL volume. The 96-well plate F-SPE system has been demonstrated in gravity F-SPE with 120 μm size silica gel for fluorous scavenging reactions and amide coupling reactions.²² There are several similar 96-well plates commercially available. However, most of them only have 2 mL well volume.

The F-SPE process can also be automated. The RapidTrace® SPE workstation has been widely used in biology labs for sample preparation,²⁶ but it is less popular in synthetic labs (Fig. 7). The workstation can have up to 10 modules arranged in parallel and attached to a computer to control cartridge conditioning, sample loading, cartridge elution, and fraction collection. The automated sample loading can handle solutions and slurries containing small amounts of solid. Pump-controlled solvent delivery gives accurate solvent volume and flow rate. Each module conducts 10 SPEs sequentially. A maximum of 10×10=100 SPE separations can be finished in 1–2 h unattended. Each SPE cartridge has 3 mL volume, which can be charged with 1.5 g of fluorous silica gel for the separation of 10–100 mg samples. Relative to the plate-to-plate SPE, the RapidTrace® unit has higher upfront instrument cost but significantly saves manpower and provides consistent results.^{25b}

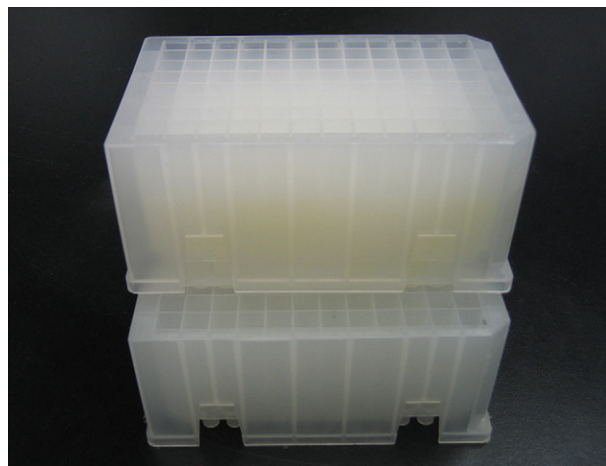


Figure 6. Ex-Block for 96-well plate-to-plate F-SPE.



Figure 7. RapidTrace[®] SPE workstation (left, single unit; right, 10 parallel units).

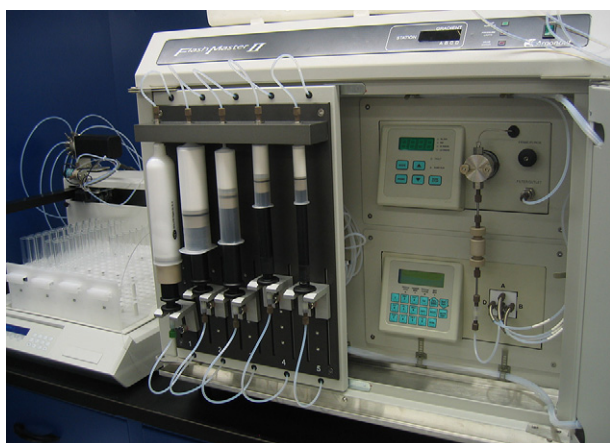


Figure 8. Biotage FlashMaster[™] II for variable-scale F-SPE.

Commercial systems from Isco and Biotage are available for large-scale F-SPE or flash chromatography. Among them, the Biotage FlashMaster[™] II can handle up to 200 psi back pressure, which is more suitable for fluoros separations using MeOH–H₂O and MeCN–H₂O as the elution solvents. This system has 10 channels; cartridge sizes from 5 to 100 g can be easily fit in (Fig. 8). The FlashMaster[™] system also has many features of standard HPLC including gradient solvent mixing, flow control, and UV-triggered fraction collection.

4. Tabular summary of F-SPE

This tabular section is intended to provide a comprehensive collection of the published uses of F-SPE for separation in small molecule synthesis from its inception in 1997 up to early 2006. Small molecule synthesis is not the only use of F-SPE, but applications in oligonucleotide synthesis,²⁷ peptide synthesis,²⁸ proteomics²⁹ and other areas³⁰ are not covered here. Nor do we cover other uses for fluoros silica

gel including HPLC demixing in fluoros mixture synthesis³¹ and catalyst/reagent support applications.³²

The tables are organized so that readers can easily scan them for relevant fluoros reaction components (left column) and allied transformations (center column). References are provided in the right column. In almost all cases, papers report multiple examples of the use of F-SPE, but we often extract only a single representative example. In the case of chiral auxiliaries and protecting groups, for example, the F-SPE may be used in multiple steps, including tagging, reaction of tagged substrate, and detagging. Here, we typically focus on the key reactions. In the case of library synthesis, we summarize the steps and show a generic example of the library core with R groups to give readers a sense of the scope and substitution pattern of the library.

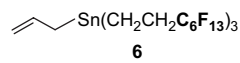
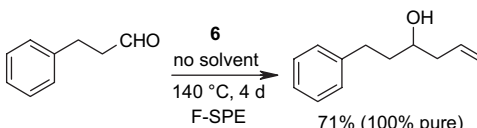
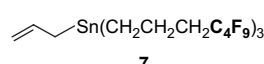
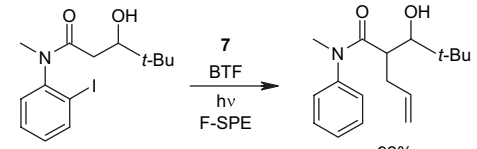
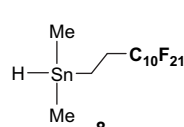
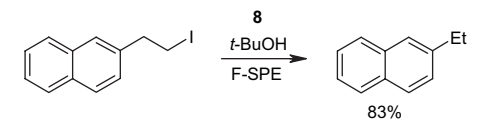
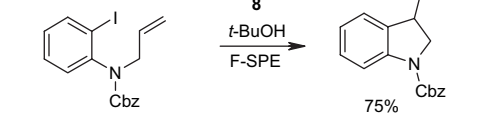
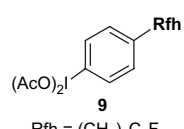
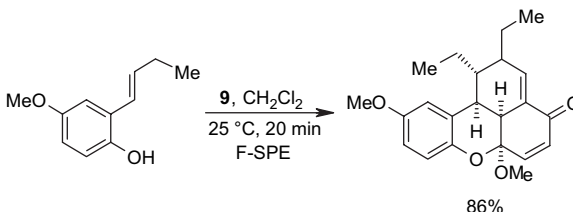
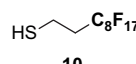
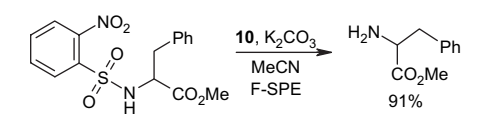
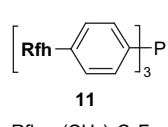
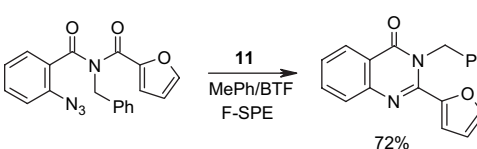
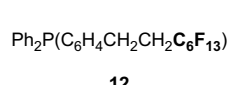
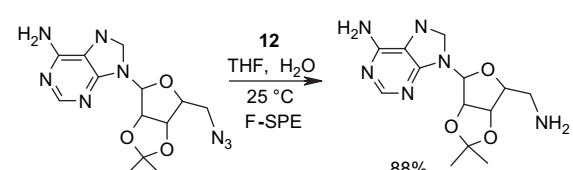
The tables are organized according to fluoros reaction component under the following headings:

- Reagents,³³ Table 1: One or more fluoros reagents are used in at least stoichiometric quantities, providing fluoros byproducts. The precursor and the target product are organic.
- Reactants, Table 2: A fluoros reactant is incorporated into the product, but it is not a chiral auxiliary or a protecting group.
- Catalysts,³⁴ Table 3: The fluoros reaction component is a catalyst or precatalyst. The precursor and the target product are organic.
- Chiral auxiliary, Table 4: The substrate and product bear a chiral auxiliary with a fluoros tag.
- Scavengers,³⁵ Table 5: A fluoros reagent is used to consume (scavenge) some undesired reaction component (usually an unreacted starting reagent) and the scavenged product is separated from the organic target product.
- Protecting groups,³⁶ Table 6: The substrate and/or product bear a fluoros version of a common protecting group.

- Displaceable tags,³⁶ Table 7: The substrate and derived intermediates bear a tag that is displaced with other functionalities, usually in a diversity oriented synthesis setting.

A number of the more popular fluororous reaction components shown in the tables are commercially available from Fluorous Technologies, Inc.,¹⁹ Aldrich, Fluka, and Wako (in Japan).

Table 1. Reactions with fluororous reagents

Reagent	Transformation	Ref.
 <p>6 $\text{Sn}(\text{CH}_2\text{CH}_2\text{C}_6\text{F}_{13})_3$</p>	 <p>6 no solvent 140°C, 4 d F-SPE 71% (100% pure)</p>	3
 <p>7 $\text{Sn}(\text{CH}_2\text{CH}_2\text{CH}_2\text{C}_4\text{F}_9)_3$</p>	 <p>7 BTF $h\nu$ F-SPE 92%</p>	37
 <p>8 $\text{C}_{10}\text{F}_{21}$</p>	 <p>8 $t\text{-BuOH}$ F-SPE 83%</p>  <p>8 $t\text{-BuOH}$ F-SPE 75%</p>	38
 <p>9 $\text{Rfh} = (\text{CH}_2)_2\text{C}_6\text{F}_{17}$</p>	 <p>9, CH_2Cl_2 25°C, 20 min F-SPE 86%</p>	39
 <p>10 C_8F_{17}</p>	 <p>10, K_2CO_3 MeCN F-SPE 91%</p>	40
 <p>11 $\text{Rfh} = (\text{CH}_2)_3\text{C}_8\text{F}_{17}$</p>	 <p>11 MePh/BTF F-SPE 72%</p>	41
 <p>12 $\text{Ph}_2\text{P}(\text{C}_6\text{H}_4\text{CH}_2\text{CH}_2\text{C}_6\text{F}_{13})$</p>	<p style="text-align: center;">Staudinger reactions</p>  <p>12 THF, H_2O 25°C F-SPE 88%</p>	42

(continued)

Table 1. (continued)

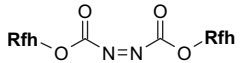
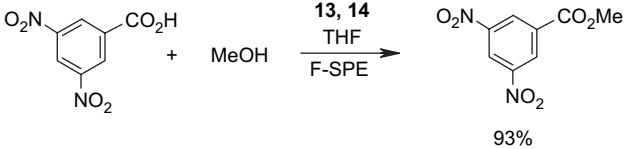
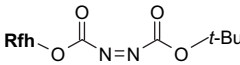
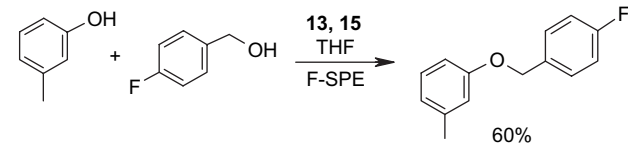
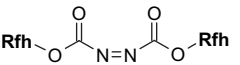
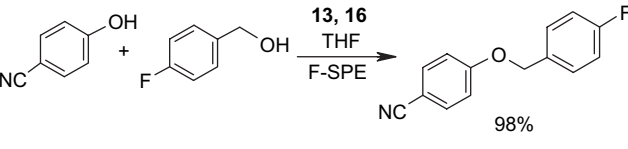
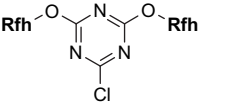
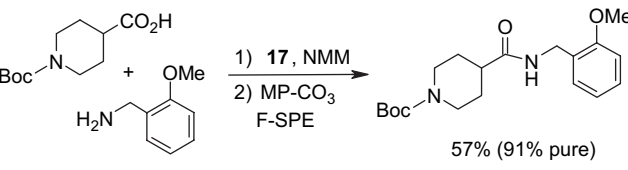
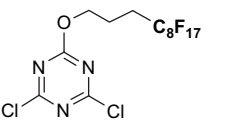
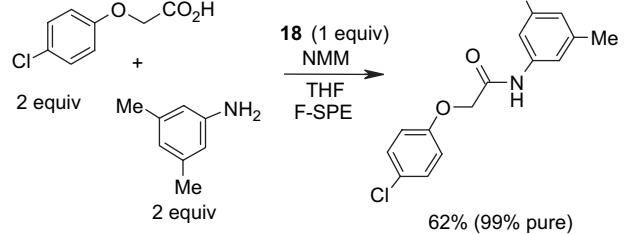
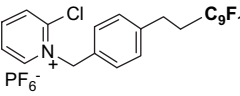
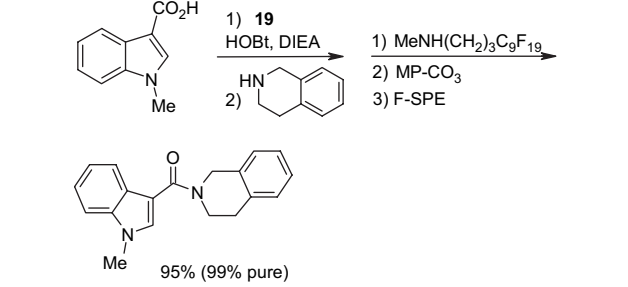
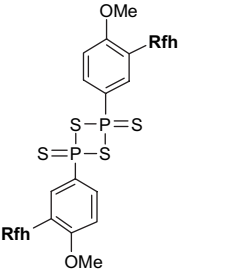
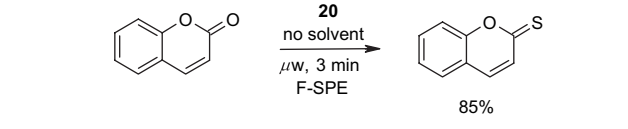
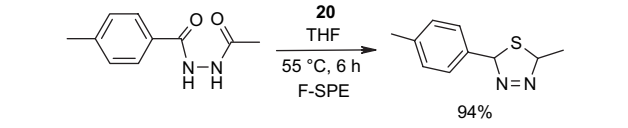
Reagent	Transformation	Ref.
$\text{Ph}_2\text{P}(\text{C}_6\text{H}_4\text{CH}_2\text{CH}_2\text{C}_8\text{F}_{17})$ 13, F-TPP  14, F-DEAD-1 $\text{Rfh} = (\text{CH}_2)_2\text{C}_6\text{F}_{13}$	Mitsunobu reactions 	
 15, F-DEAD-2 $\text{Rfh} = (\text{CH}_2)_3\text{C}_6\text{F}_{13}$		43–45
 16, F-DEAD-3 $\text{Rfh} = (\text{CH}_2)_3\text{C}_6\text{F}_{13}$		
 17, F-CDMT $\text{Rfh} = (\text{CH}_2)_3\text{C}_6\text{F}_{13}$		25
 18, F-DCT		46
 19, F-Mukaiyama condensation reagent		47
 20, F-Lawesson's reagent $\text{Rfh} = (\text{CH}_2)_5\text{C}_8\text{F}_{17}$ $(\text{CH}_2)_4\text{C}_8\text{F}_{17}$ $(\text{CH}_2)_4\text{C}_6\text{F}_{13}$	 	48,49

Table 2. Reactions with fluororous reactants

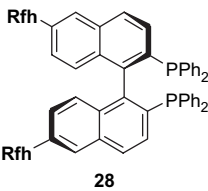
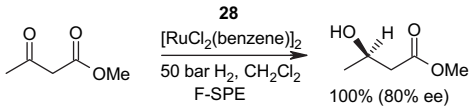
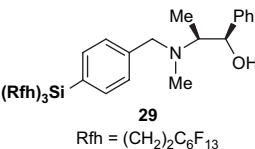
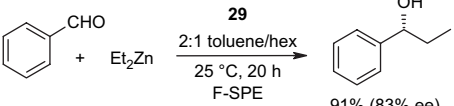
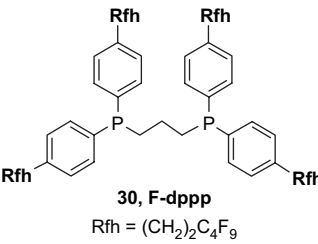
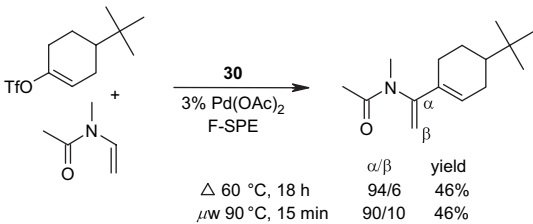
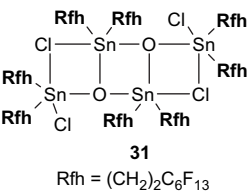
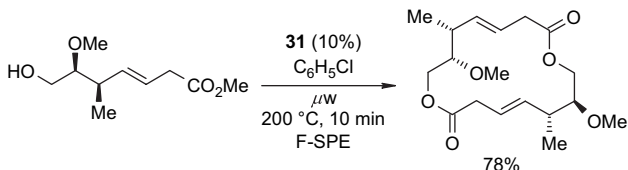
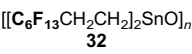

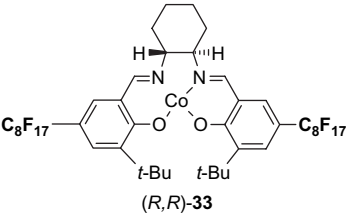
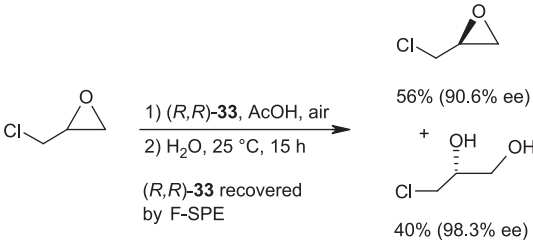
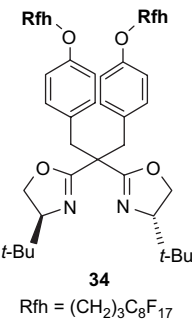
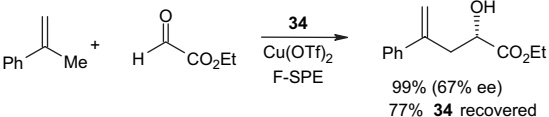
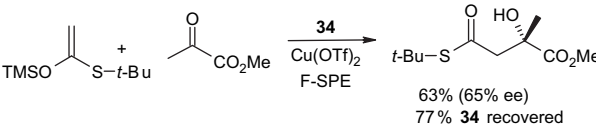
Reactants	Transformation	Ref.
<p>21</p>	<p>21</p> <p>THF-Et₂O [(cod)CuCl] F-SPE</p> <p>97% (3:1)</p>	50
<p>22</p>	<p>22</p> <p>AIBN (10%) hexane 80 °C, 2 h F-SPE</p> <p>95%</p>	51
<p>23</p>	<p>23</p> <p>Cu(OAc)₂ ascorbic acid <i>i</i>-Pr₂NH, MeOH F-SPE</p> <p>95%</p>	52

Table 3. Reactions with fluororous catalysts

Catalyst	Transformation	Ref.
<p>24</p> <p>Rfh = (CH₂)₂C₆F₁₃</p>	<p>24</p> <p>BTF, 60 °C F-SPE</p> <p>100% (100% pure)</p>	9
<p>25</p> <p>Rfh = (CH₂)₂C₆F₁₃</p>	<p>25, Sml₂</p> <p>THF -45 °C, 2 h F-SPE</p> <p>82% (81% ee) 98% 25 recovered</p>	53
<p>26, (R)-F-BINOL</p> <p>Rfh = (CH₂)₂C₆F₁₃</p>	<p>26</p> <p>Ti(O-<i>i</i>Pr)₄ BTF 0 °C, 1 h F-SPE</p> <p>95% (85% ee) 100% 26 recovered</p>	54
<p>27</p>	<p>27: Ti(O-<i>i</i>Pr)₄ (2:1)</p> <p>hexane 0 °C, 61 h F-SPE</p> <p>88% (75% ee)</p>	55

(continued)

Table 3. (continued)

Catalyst	Transformation	Ref.
 <p>28 Rfh = (CH₂)₂C₆F₁₃</p>	 <p>28 [RuCl₂(benzene)]₂ 50 bar H₂, CH₂Cl₂ F-SPE</p>	<p>100% (80% ee)</p> <p>56</p>
 <p>29 Rfh = (CH₂)₂C₆F₁₃</p>	 <p>29 2:1 toluene/hex 25 °C, 20 h F-SPE</p>	<p>91% (83% ee) 100% 29 recovered</p> <p>57</p>
 <p>30, F-dppp Rfh = (CH₂)₂C₄F₉</p>	 <p>30 3% Pd(OAc)₂ F-SPE Δ 60 °C, 18 h μW 90 °C, 15 min</p>	<p>α/β yield 94/6 46% 90/10 46%</p> <p>58</p>
 <p>31 Rfh = (CH₂)₂C₆F₁₃</p>	 <p>31 (10%) C₆H₅Cl 200 °C, 10 min μW F-SPE</p>	<p>78%</p> <p>59</p>
 <p>32</p>	 <p>32 (2%) Et₃N-TsCl CH₂Cl₂ 25 °C, 1 h F-SPE</p>	<p>70%</p> <p>60</p>
 <p>(R,R)-33</p>	 <p>1) (R,R)-33, AcOH, air 2) H₂O, 25 °C, 15 h (R,R)-33 recovered by F-SPE</p>	<p>56% (90.6% ee) 40% (98.3% ee)</p> <p>61</p>
 <p>34 Rfh = (CH₂)₃C₈F₁₇</p>	 <p>34 Cu(OTf)₂ F-SPE</p>	<p>99% (67% ee) 77% 34 recovered</p> <p>62</p>
	 <p>34 Cu(OTf)₂ F-SPE</p>	<p>63% (65% ee) 77% 34 recovered</p>

(continued)

Table 3. (continued)

Catalyst	Transformation	Ref.
$[\text{Ni}(\text{F}_3\text{CC}(\text{O})\text{CHC}(\text{O})\text{C}_6\text{F}_{13})_2]$ 35		63
 36 Rfh = $(\text{CH}_2)_3\text{C}_6\text{F}_{13}$		13
 2 Rfh = $(\text{CH}_2)_3\text{C}_8\text{F}_{17}$		13
 37		64
 38		65

Table 4. Reactions with fluororous chiral auxiliaries

Chiral auxiliary	Transformation	Ref.
 39		66

(continued)

Table 4. (continued)

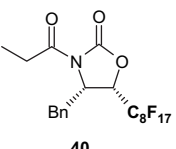
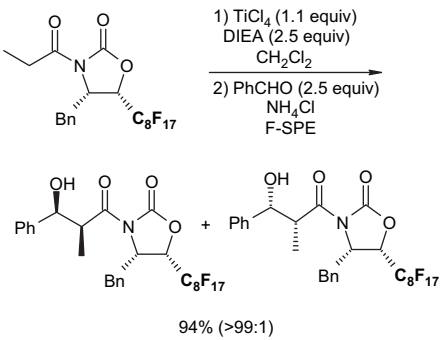
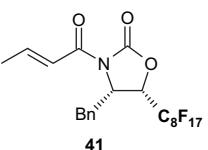
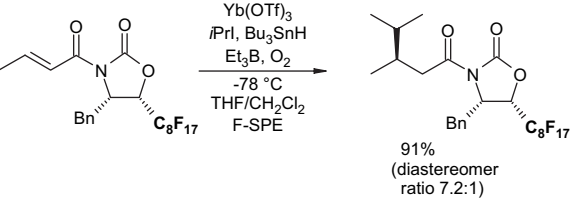
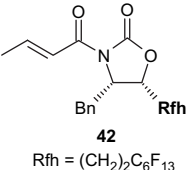
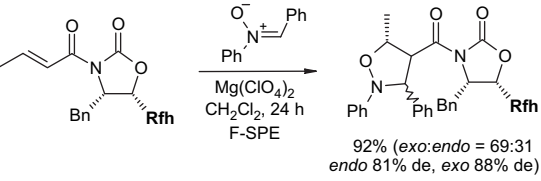
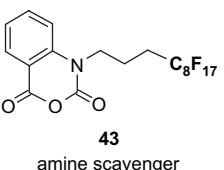
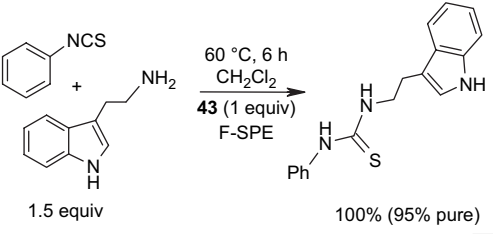
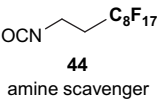
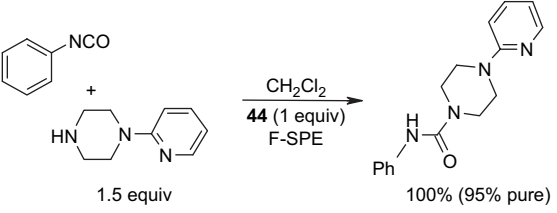
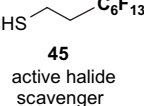
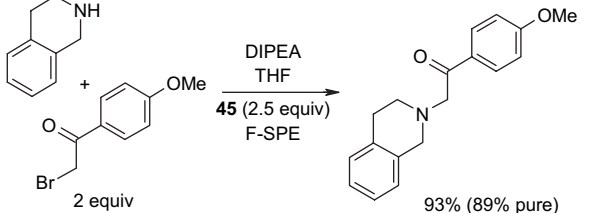
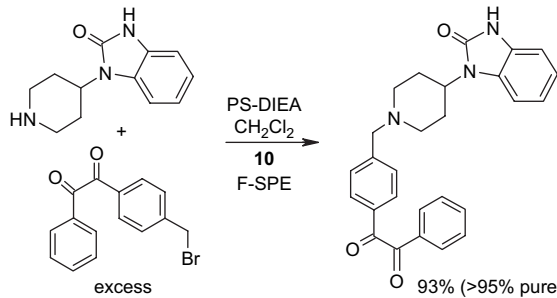
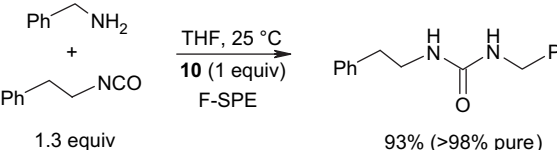
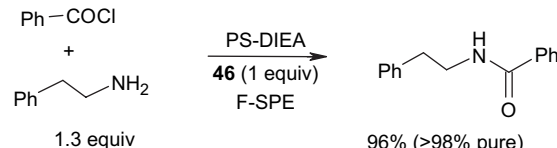
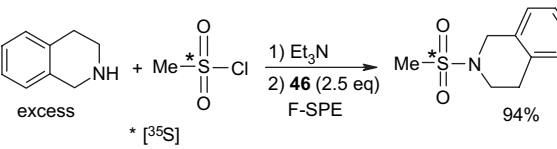
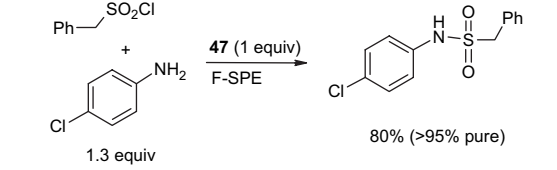
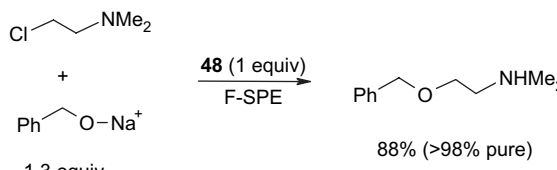
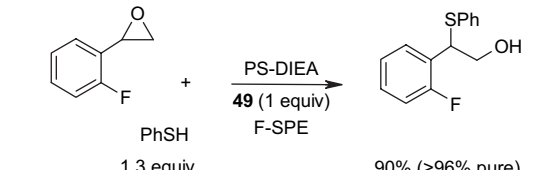
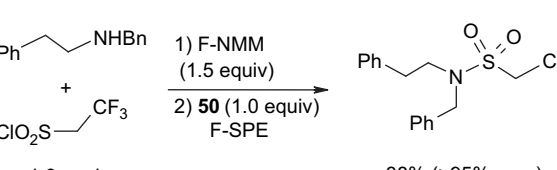
Chiral auxiliary	Transformation	Ref.
 <p>40</p>	 <p>1) TiCl_4 (1.1 equiv) DIEA (2.5 equiv) CH_2Cl_2 2) PhCHO (2.5 equiv) NH_4Cl F-SPE</p> <p>94% (>99:1)</p>	67
 <p>41</p>	 <p>$\text{Yb}(\text{OTf})_3$ <i>i</i>PrI, Bu_3SnH Et_3B, O_2 -78 °C THF/CH_2Cl_2 F-SPE</p> <p>91% (diastereomer ratio 7.2:1)</p>	68
 <p>42 Rfh = $(\text{CH}_2)_2\text{C}_6\text{F}_{13}$</p>	 <p>$\text{O}^- \text{N}^+ \text{Ph}$ Ph' $\text{Mg}(\text{ClO}_4)_2$ CH_2Cl_2, 24 h F-SPE</p> <p>92% (<i>exo:endo</i> = 69:31 <i>endo</i> 81% de, <i>exo</i> 88% de)</p>	69

Table 5. Reactions with fluorous scavengers

Scavenger	Transformation	Ref.
 <p>43 amine scavenger</p>	 <p>60 °C, 6 h CH_2Cl_2 43 (1 equiv) F-SPE</p> <p>100% (95% pure)</p>	70
 <p>44 amine scavenger</p>	 <p>CH_2Cl_2 44 (1 equiv) F-SPE</p> <p>100% (95% pure)</p>	71
 <p>45 active halide scavenger</p>	 <p>DIEPA THF 45 (2.5 equiv) F-SPE</p> <p>93% (89% pure)</p>	72

(continued)

Table 5. (continued)

Scavenger	Transformation	Ref.
<p>HS-CH₂-CH₂-C₈F₁₇ 10 active halide scavenger</p>	 <p>PS-DIEA CH₂Cl₂ 10 F-SPE excess 93% (>95% pure)</p>	73,74
<p>HS-CH₂-CH₂-C₈F₁₇ 10 isocyanate scavenger</p>	 <p>THF, 25 °C 10 (1 equiv) F-SPE 1.3 equiv 93% (>98% pure)</p>	75
<p>Cl-C(=O)-C₇F₁₅ 46 amine scavenger</p>	 <p>PS-DIEA 46 (1 equiv) F-SPE 1.3 equiv 96% (>98% pure)</p>	74
<p>Cl-C(=O)-C₇F₁₅ 46 amine scavenger</p>	 <p>excess Me-S(=O)₂-Cl 1) Et₃N 2) 46 (2.5 eq) F-SPE * [³⁵S] 94%</p>	75
<p>HO₃S-C₈F₁₇ 47 amine scavenger</p>	 <p>47 (1 equiv) F-SPE 1.3 equiv 80% (>95% pure)</p>	74
<p>ClO₂S-C₆F₁₃ 48 base scavenger</p>	 <p>48 (1 equiv) F-SPE 1.3 equiv 88% (>98% pure)</p>	74
<p>C₆F₁₃ 49 thiol scavenger</p>	 <p>PS-DIEA 49 (1 equiv) F-SPE PhSH 1.3 equiv 90% (>96% pure)</p>	74
<p>H₂N-CH₂-CH₂-N(C₆F₁₃)₃-CH₂-CH₂-NH₂ 50, F-Tris sulfonyl chloride scavenger</p>	 <p>1) F-NMM (1.5 equiv) 2) 50 (1.0 equiv) F-SPE 1.3 equiv 88% (>95% pure)</p>	76

(continued)

Table 5. (continued)

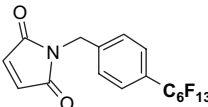
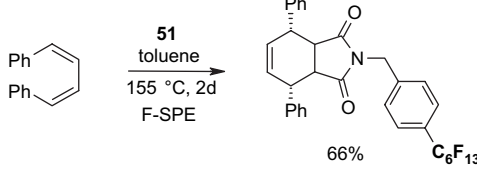
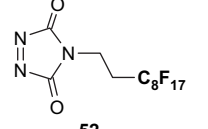
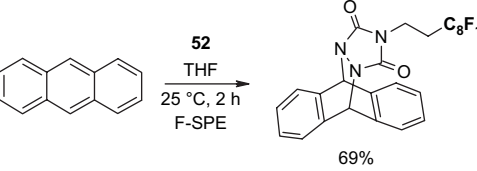
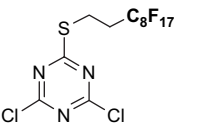
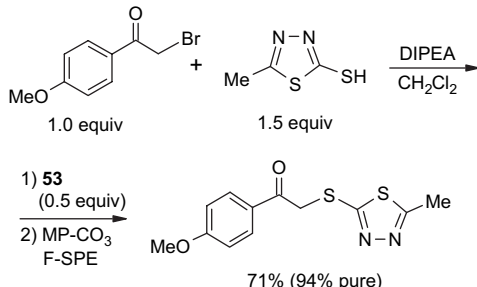
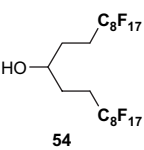
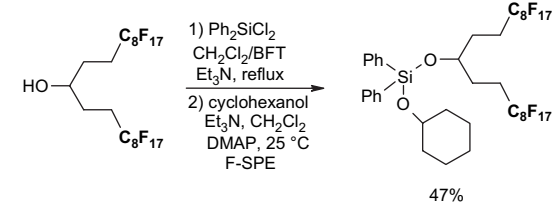
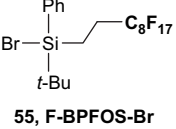
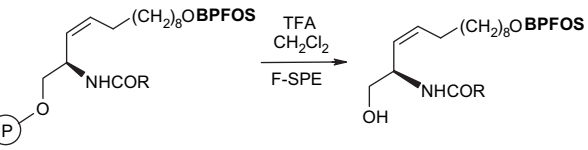
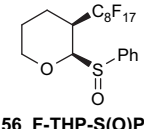
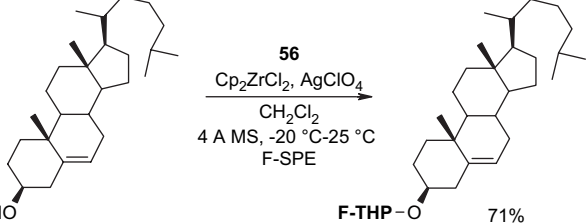
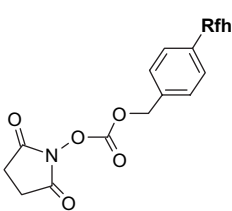
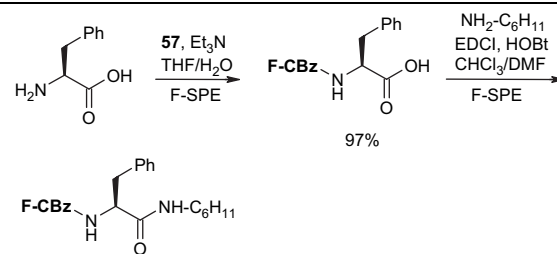
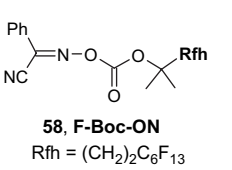
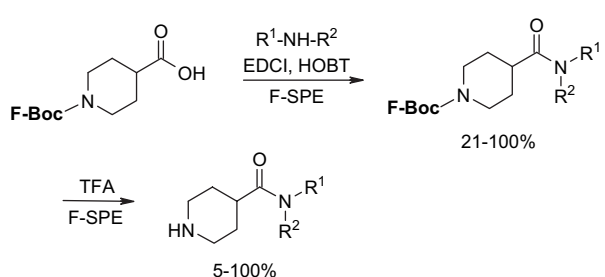
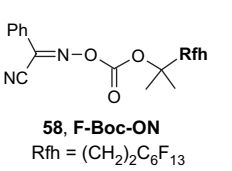
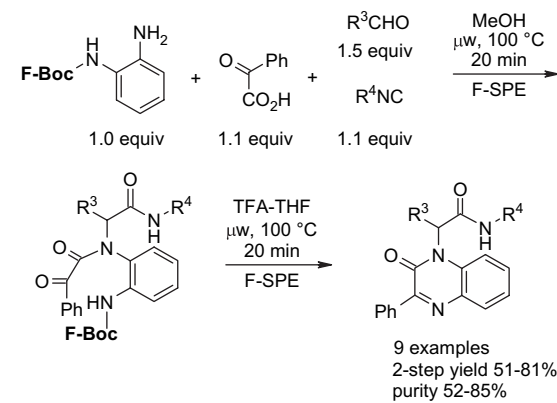
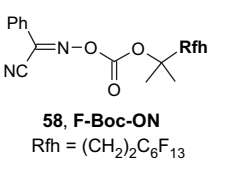
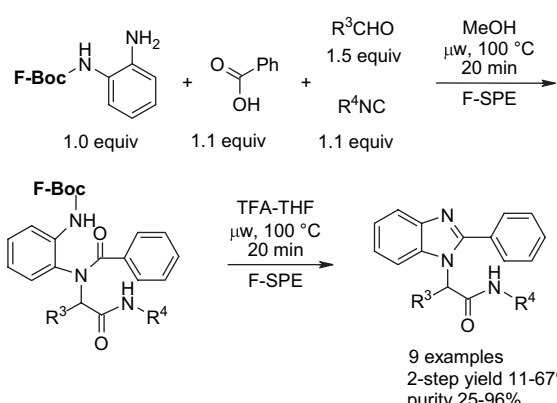
Scavenger	Transformation	Ref.
 51 dienophile scavenger		77
 52 dienophile scavenger		77
 53, F-DCT thiol scavenger		78

Table 6. Reactions with fluorous protecting groups

Protecting group	Transformation	Ref.
 54		79
 55, F-BPFOS-Br		80
 56, F-THP-S(O)Ph		81

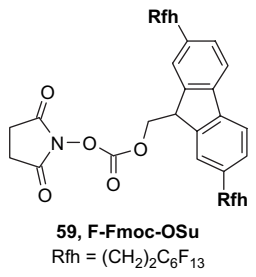
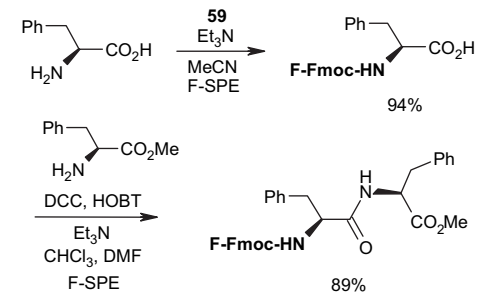
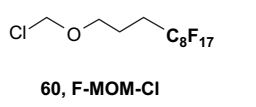
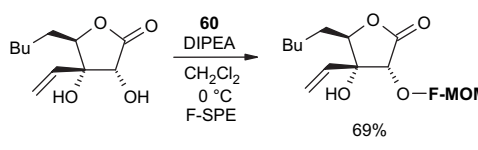
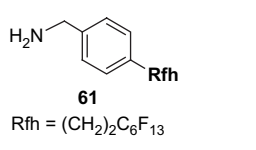
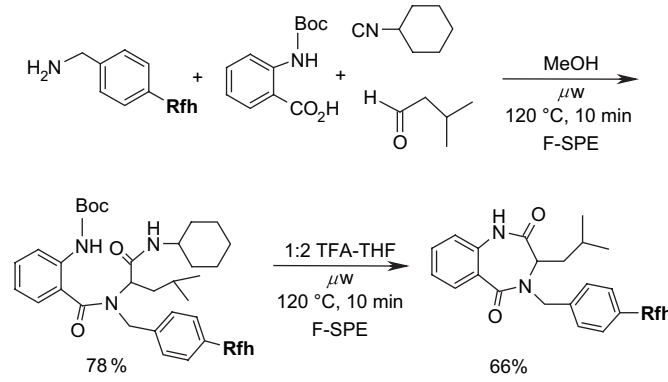
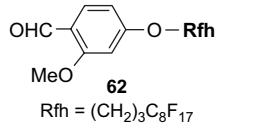
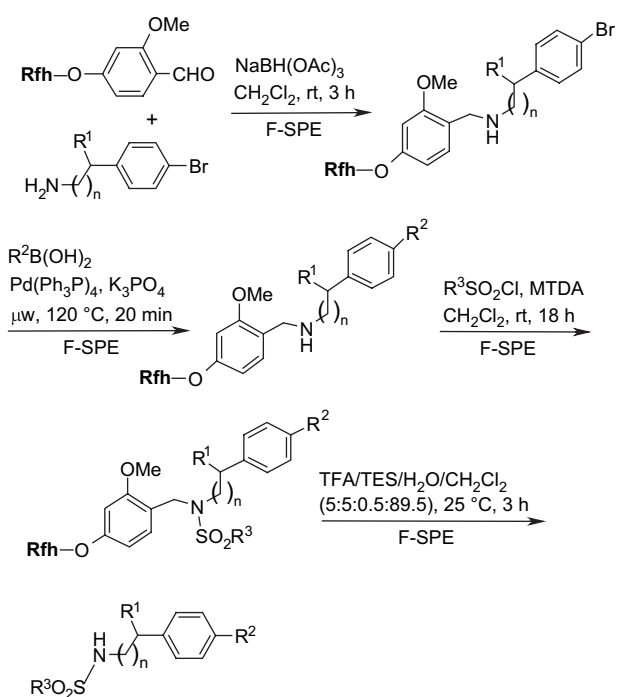
(continued)

Table 6. (continued)

Protecting group	Transformation	Ref.
 <p>57, F-CBz-OSu Rfh = (CH₂)₂C₆F₁₇</p>	 <p>97% 62%</p>	82
 <p>58, F-Boc-ON Rfh = (CH₂)₂C₆F₁₃</p>	 <p>21-100% 5-100%</p>	83
 <p>58, F-Boc-ON Rfh = (CH₂)₂C₆F₁₃</p>	 <p>1.0 equiv 1.1 equiv 1.5 equiv MeOH, 100 °C, 20 min, F-SPE TFA-THF, 100 °C, 20 min, F-SPE 9 examples 2-step yield 51-81% purity 52-85%</p>	84
 <p>58, F-Boc-ON Rfh = (CH₂)₂C₆F₁₃</p>	 <p>1.0 equiv 1.1 equiv 1.5 equiv MeOH, 100 °C, 20 min, F-SPE TFA-THF, 100 °C, 20 min, F-SPE 9 examples 2-step yield 11-67% purity 25-96%</p>	84

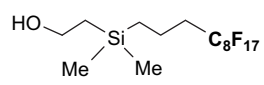
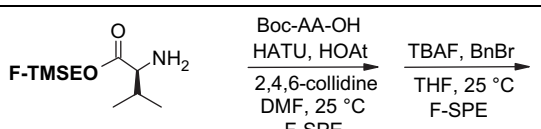
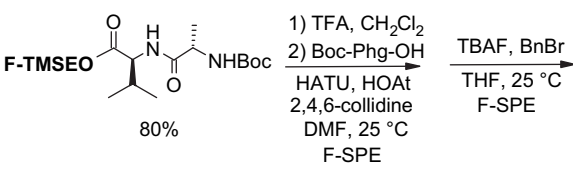
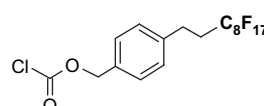
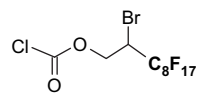
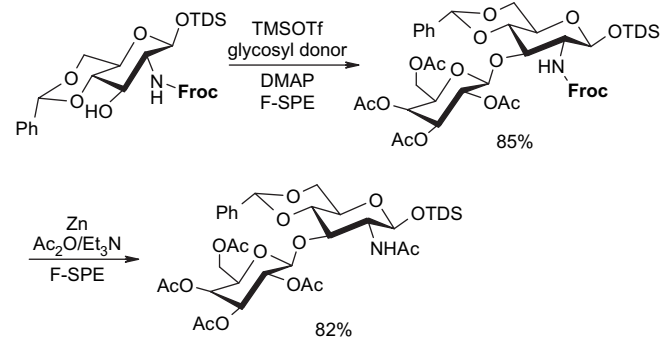
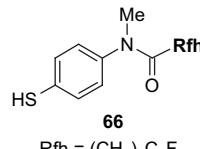
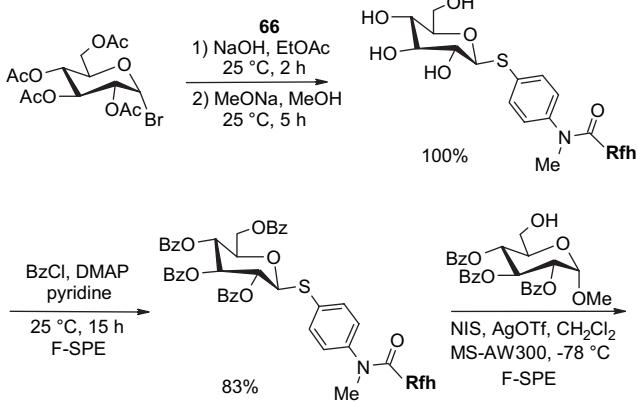
(continued)

Table 6. (continued)

Protecting group	Transformation	Ref.
 <p>59, F-Fmoc-OSu Rfh = (CH₂)₂C₆F₁₃</p>		85
 <p>60, F-MOM-Cl Rfh = (CH₂)₂C₆F₁₃</p>		86
 <p>61 Rfh = (CH₂)₂C₆F₁₃</p>		87
 <p>62 Rfh = (CH₂)₃C₈F₁₇</p>		88

(continued)

Table 6. (continued)

Protecting group	Transformation	Ref.	
 <p>63, F-TMS-OH</p>	 <p>Boc-AA-OH HATU, HOAt 2,4,6-collidine DMF, 25 °C F-SPE</p> <p>TBAF, BnBr THF, 25 °C F-SPE</p>		
	 <p>1) TFA, CH₂Cl₂ 2) Boc-Phg-OH HATU, HOAt 2,4,6-collidine DMF, 25 °C F-SPE</p> <p>TBAF, BnBr THF, 25 °C F-SPE</p> <p>80%</p> <p>63%</p>	89	
	 <p>64, F-Cbz-Cl</p>	<p>Peptides</p> <p>GCCSLPPCALNNPDYCamide 59% (91% pure)</p> <p>RQIKIWFQNRRMKWKKamide 7% (72% pure)</p>	28b
 <p>65, F-Froc-Cl</p>	 <p>TMSOTf glycosyl donor DMAP F-SPE</p> <p>Zn Ac₂O/Et₃N F-SPE</p> <p>85%</p> <p>82%</p>	90	
 <p>66 Rfh = (CH₂)₂C₈F₁₇</p>	 <p>1) NaOH, EtOAc 25 °C, 2 h 2) MeONa, MeOH 25 °C, 5 h</p> <p>BzCl, DMAP pyridine 25 °C, 15 h F-SPE</p> <p>NIS, AgOTf, CH₂Cl₂ MS-AW300, -78 °C F-SPE</p> <p>100%</p> <p>83%</p> <p>76% 75% 66 recovered</p>	91	

(continued)

Table 6. (continued)

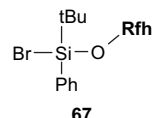
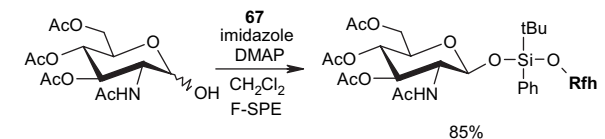
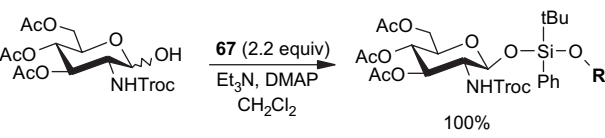
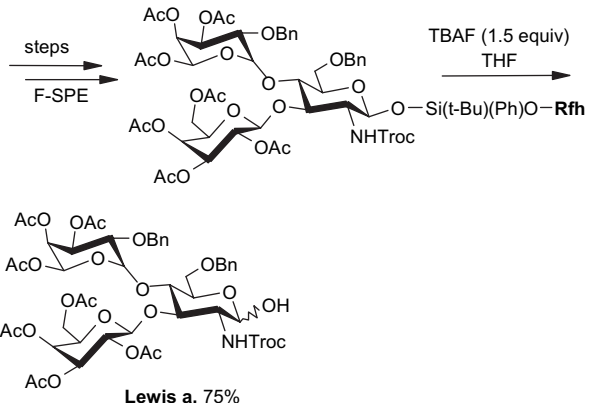
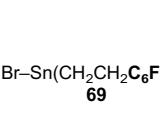
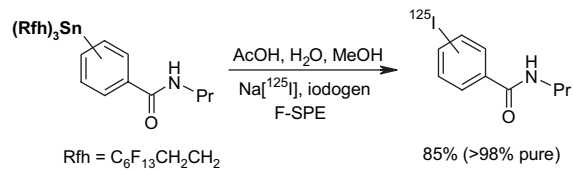
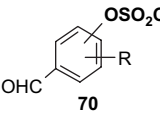
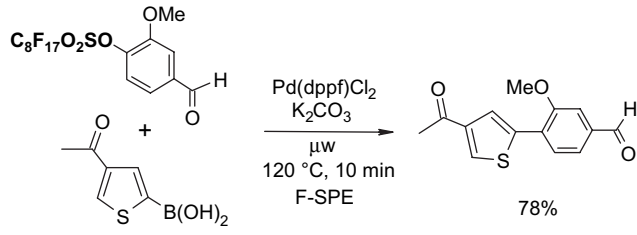
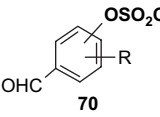
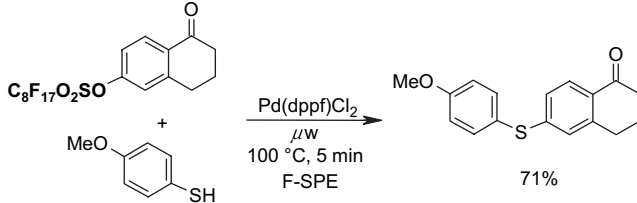
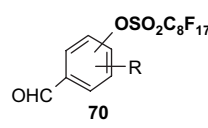
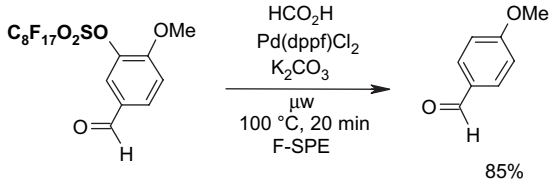
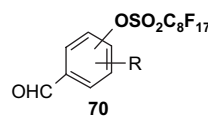
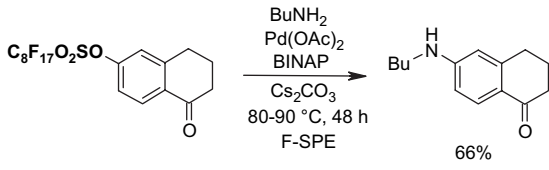
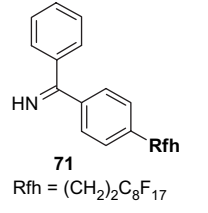
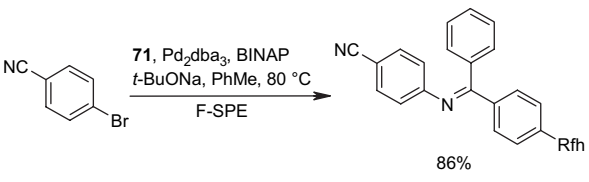
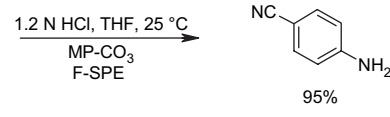
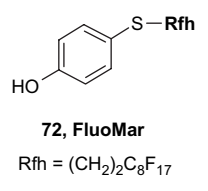
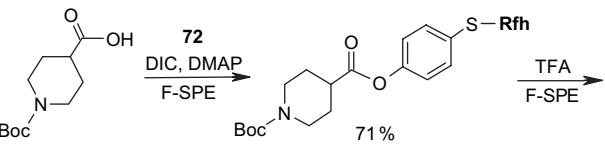

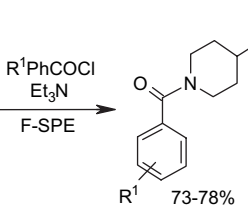
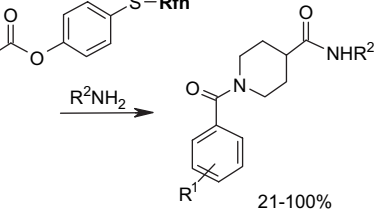
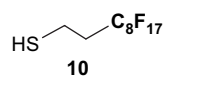
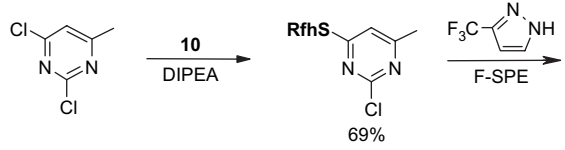
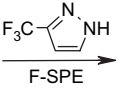
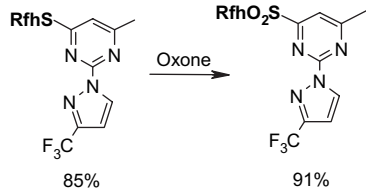
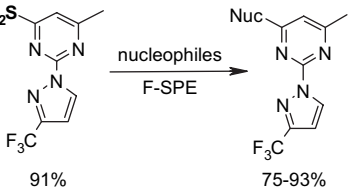
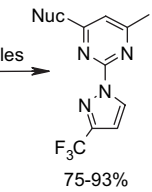
Protecting group	Transformation	Ref.
 <p>67 Rfh = (CH₂)₂C₈F₁₇</p>	 <p>67 imidazole DMAP CH₂Cl₂ F-SPE</p> <p>85%</p>	92
	 <p>67 (2.2 equiv) Et₃N, DMAP CH₂Cl₂</p> <p>100%</p>	
	 <p>steps F-SPE</p> <p>TBAF (1.5 equiv) THF</p> <p>Lewis a, 75%</p>	93

Table 7. Reactions with fluorous displaceable tags

Displaceable tags	Transformation	Ref.
 <p>69 Rfh = C₆F₁₃CH₂CH₂</p>	 <p>(Rfh)₃Sn AcOH, H₂O, MeOH Na[¹²⁵I], iodogen F-SPE</p> <p>85% (>98% pure)</p>	94
	Suzuki reactions	
 <p>70</p>	 <p>C₈F₁₇O₂SO Pd(dppf)Cl₂ K₂CO₃ μW 120 °C, 10 min F-SPE</p> <p>78%</p>	95
 <p>70</p>	 <p>C₈F₁₇O₂SO Pd(dppf)Cl₂ μW 100 °C, 5 min F-SPE</p> <p>71%</p>	96

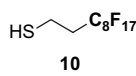
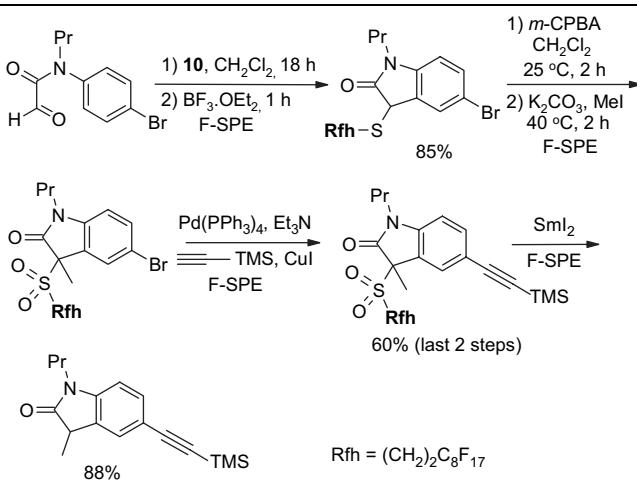
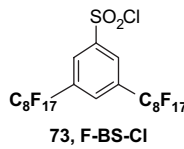
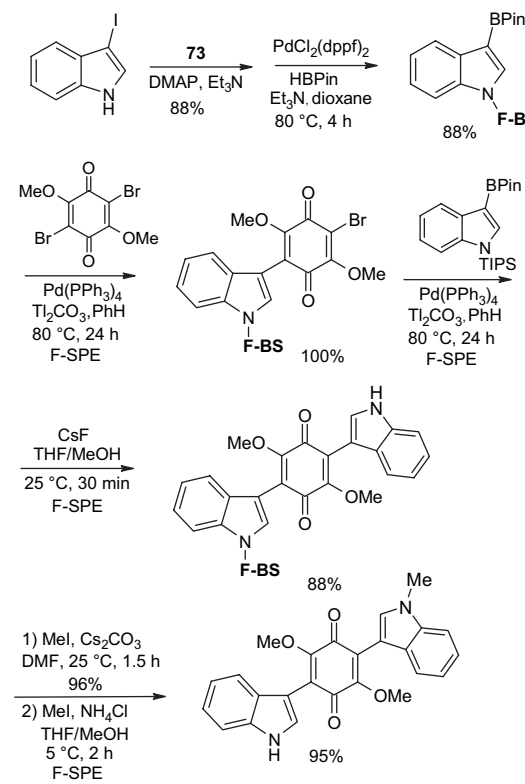
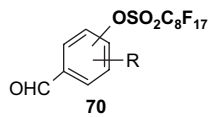
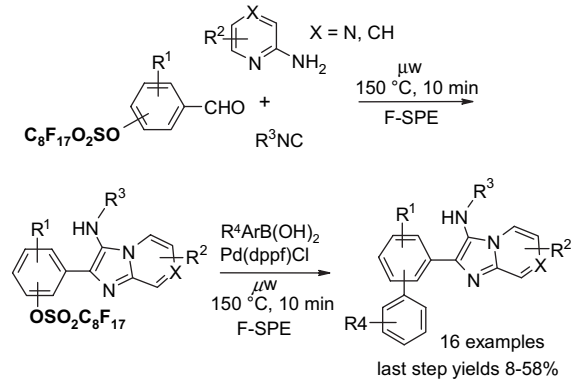
(continued)

Table 7. (continued)

Displaceable tags	Transformation	Ref.
 <p>70</p>	 <p>85%</p>	97
 <p>70</p>	 <p>66%</p>	98
 <p>71 Rfh = (CH₂)₂C₈F₁₇</p>	 <p>86%</p>  <p>95%</p>	99
 <p>72, FluMar Rfh = (CH₂)₂C₈F₁₇</p>	 <p>71%</p>   <p>73-78%</p>  <p>21-100%</p>	100
 <p>10</p>	 <p>69%</p>   <p>85%</p>  <p>91%</p>  <p>75-93%</p> <p>Rfh = (CH₂)₂C₈F₁₇</p>	101

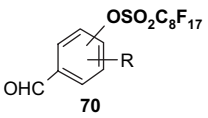
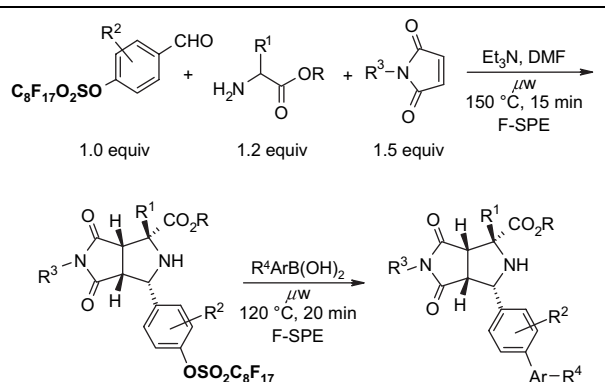
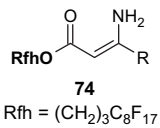
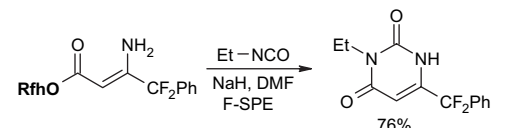
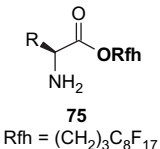
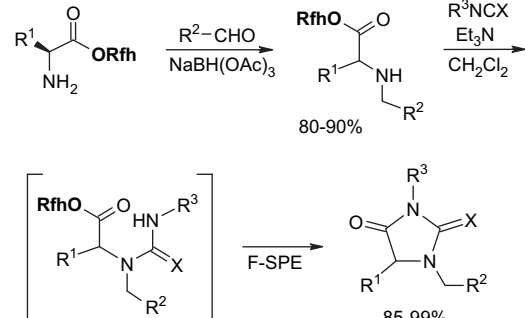
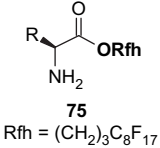
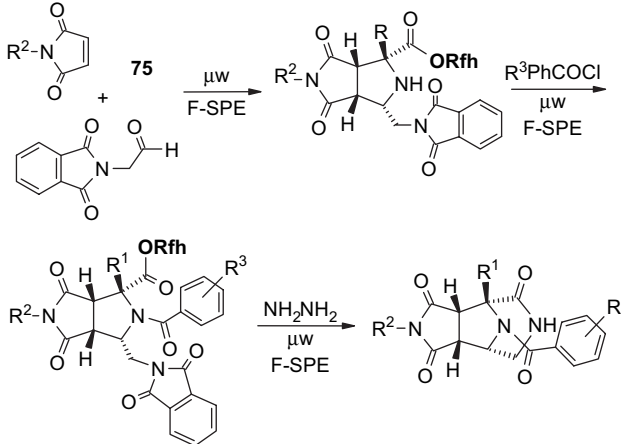
(continued)

Table 7. (continued)

Displaceable tags	Transformation	Ref.
 <p>10</p>		102
 <p>73, F-BS-Cl</p>		103
 <p>70</p>		104

(continued)

Table 7. (continued)

Displaceable tags	Transformation	Ref.
 <p>70 OSO₂C₈F₁₇</p>	 <p>1.0 equiv 1.2 equiv 1.5 equiv</p> <p>Et₃N, DMF 150 °C, 15 min F-SPE</p> <p>R⁴ArB(OH)₂ 120 °C, 20 min F-SPE</p>	105
 <p>74 Rfh = (CH₂)₃C₈F₁₇</p>	 <p>Et-NCO NaH, DMF F-SPE</p> <p>76%</p>	106
 <p>75 Rfh = (CH₂)₃C₈F₁₇</p>	 <p>R²-CHO NaBH(OAc)₃</p> <p>80-90%</p> <p>R³NCX Et₃N CH₂Cl₂</p> <p>F-SPE</p> <p>85-99%</p>	107
 <p>75 Rfh = (CH₂)₃C₈F₁₇</p>	 <p>R²-N-pyrrolidone + 75</p> <p>microwaves F-SPE</p> <p>R³PhCOCl microwaves F-SPE</p> <p>NH₂NH₂ microwaves F-SPE</p>	108

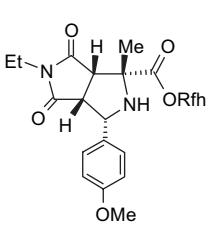
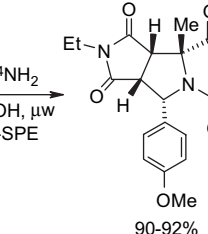
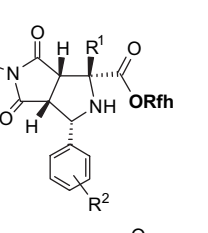
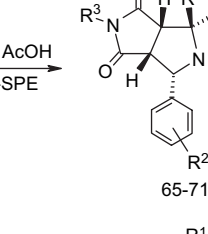
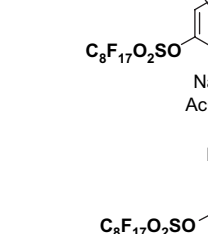
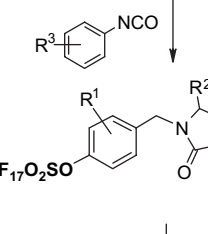
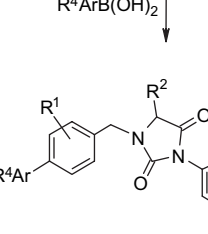
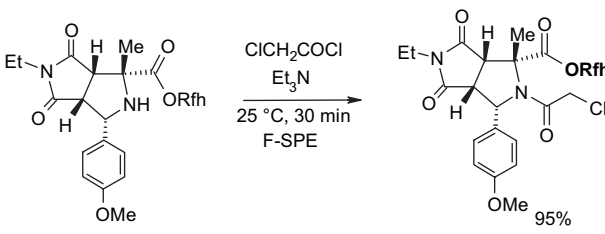
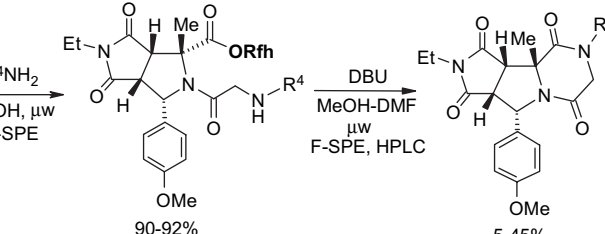
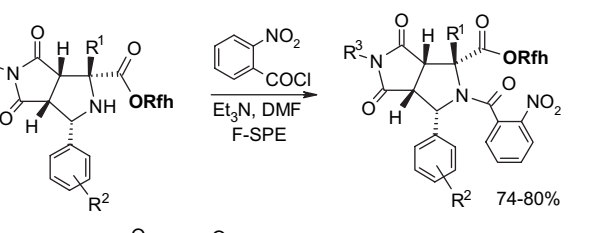
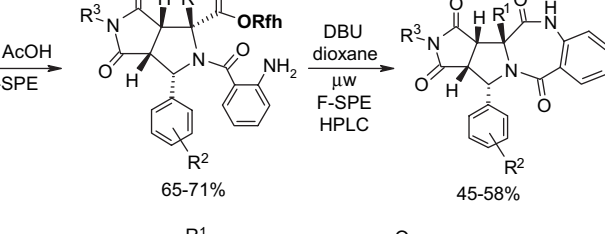
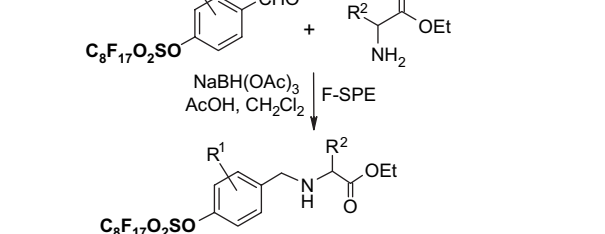
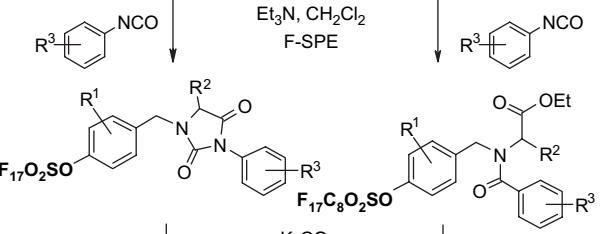
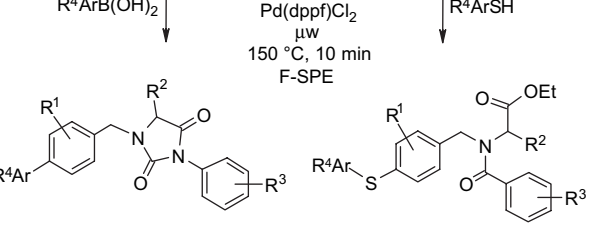
(continued)

Table 7. (continued)

Displaceable tags	Transformation	Ref.
<p>75 1.5 equiv</p>	<p>3 equiv F-SPE</p> <p>H₂, Pd/C 12 h F-SPE</p> <p>3 equiv F-SPE</p> <p>μw 150 °C, 15 min</p>	109
<p>75 Rfh = (CH₂)₃C₈F₁₇</p> <p>H₂N-CH₂-CO-ORfh</p> <p>Et₃N toluene</p> <p>150 °C, 10 min μw F-SPE HPLC</p> <p>4 equiv</p> <p>39%</p>		110
<p>75 Rfh = (CH₂)₃C₈F₁₇</p> <p>CHO</p> <p>1.0 equiv</p> <p>72</p> <p>1.2 equiv</p> <p>Et₃N, DMF</p> <p>130 °C, 20 min μw F-SPE</p> <p>1.5 equiv</p>		111
<p>75 Rfh = (CH₂)₃C₈F₁₇</p> <p>R⁴-PhNCX (5 equiv)</p> <p>DMAP</p> <p>μw F-SPE</p> <p>86-92%</p>		111
<p>75 Rfh = (CH₂)₃C₈F₁₇</p> <p>K₂CO₃ DMF</p> <p>F-SPE HPLC</p> <p>75-85%</p> <p>X = O, S</p>		111

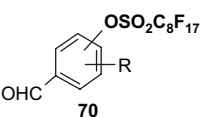
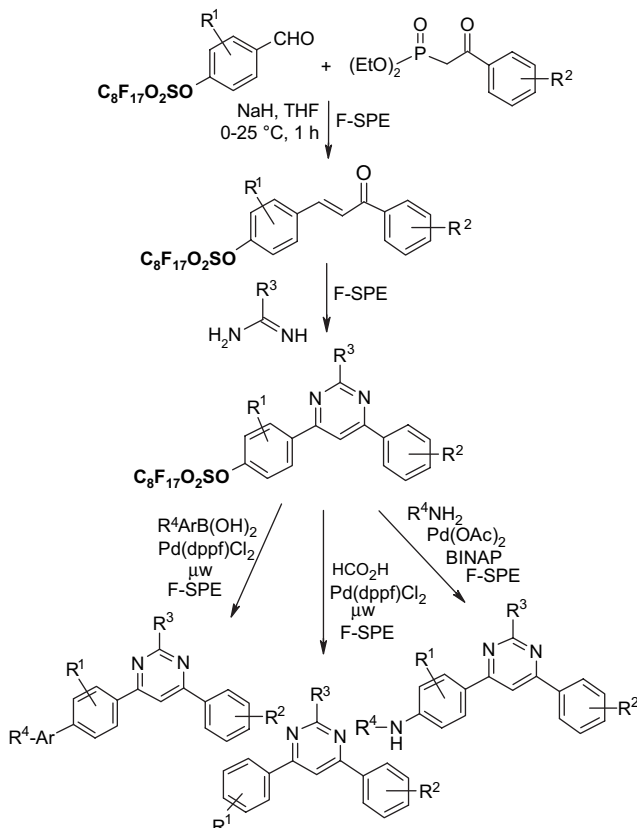
(continued)

Table 7. (continued)

Displaceable tags	Transformation	Ref.
      	<p>  </p> <p>  </p> <p>  </p> <p>  </p> <p>  </p> <p>  </p> <p>  </p>	<p>111</p> <p>111</p> <p>95,96</p>

(continued)

Table 7. (continued)

Displaceable tags	Transformation	Ref.
 70		95,97,98

5. Practical aspects of F-SPE²¹

5.1. A typical F-SPE procedure

F-SPEs in cartridge format are very easy to conduct and have the following general steps: cartridge washing (for new cartridges only), preconditioning, sample loading, fluorophobic elution, fluorophilic elution, and final washing for cartridge reuse (optional). A typical F-SPE procedure for the separation of a reaction mixture with a 2 g SPE cartridge is as follows:²¹

Step 1—Cartridge washing: Wash a new cartridge with 1 mL of DMF under a vacuum or positive pressure depending on your SPE manifold. This step can be omitted with recycled cartridges.

Step 2—Preconditioning: Pass through 6 mL of 80:20 MeOH–H₂O to condition the cartridge. Discard the preconditioning eluent.

Step 3—Sample loading: Dissolve sample (100–300 mg) in 0.4 mL of DMF and load onto the cartridge by using vacuum or positive pressure to ensure the sample is completely adsorbed onto the cartridge (see Table 8 for alternative loading solvents).

Step 4—Fluorophobic elution: Wash with 6–8 mL of 80:20 MeOH–H₂O to obtain the fraction containing the organic compounds.

Step 5—Fluorophilic elution: Wash with 8 mL of MeOH to obtain the fraction containing the fluororous compounds.

Step 6—Final washing (optional): To regenerate the SPE cartridge for reuse, wash with 6 mL of THF or acetone and air dry.

5.2. F-SPE demonstration with dyes

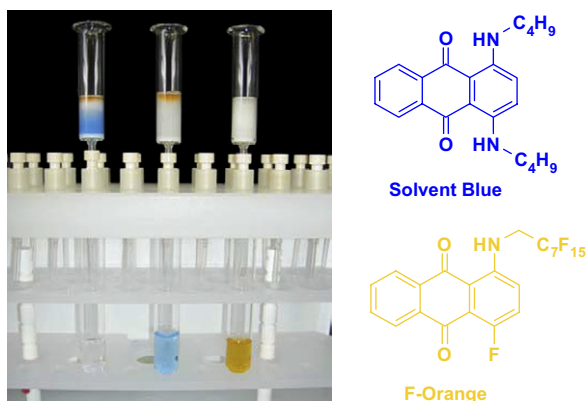
The following dye separation demonstrates how F-SPE works. The non-fluorous compound is Solvent Blue[®] dye; the fluororous compound is F-orange dye. These two dyes have similar polarities. Figure 9 shows fluororous cartridges containing the dye mixture in three different stages of elution. The left-hand test tube illustrates how the F-SPE cartridge appears after loading a mixture of the two dyes and elution with a small amount of 80:20 MeOH–H₂O. The center tube shows how the non-fluorous components (blue fraction) are washed from the cartridge by using more 80:20 MeOH–H₂O. The adsorbed fluororous dye is not eluted even with extensive flushing with 80:20 MeOH–H₂O and remains on the cartridge. Finally, the orange fluororous dye is easily eluted with 100% MeOH or THF, as shown by the third tube.

5.3. Common issues related to F-SPE

5.3.1. Loading solvents. Many different solvents can be used for sample loading; however, the more fluorophilic the solvent is the smaller its volume should be to prevent breakthrough. A list of solvents with increasing fluorophilicity is as follows: H₂O < DMSO < DMF < MeOH < MeCN < THF < HFC-7100 (C₄F₉OCH₃) < FC-72. For normal F-SPE,

Table 8. Suggested maximum loading solvent volumes for C₈F₁₇-tagged substrates

Solvent	Maximum loading volume		
	2 g cartridge (mL)	5 g cartridge (mL)	10 g cartridge (mL)
THF	0.2	0.5	1.0
CH ₂ Cl ₂	0.2	0.5	1.0
MeCN	0.2	0.5	1.0
MeOH	0.2	0.5	1.0
DMF	0.4	1.0	2.0
DMSO	0.6	1.5	3.0

**Figure 9.** F-SPE with blue (organic) and orange (fluorous) dyes. *Left tube:* beginning of fluorophobic wash (80:20 MeOH–H₂O); *center tube:* end of fluorophobic wash; *right tube:* end of fluorophilic wash (100% MeOH).

the mass loading (weight of crude sample compared to the weight of fluorous silica gel) is suggested to be around 5–10%.²¹ With the least fluorophilic DMSO, the solvent loading (volume of solvent compared to the volume of fluorous silica gel) can be as high as 30%, whereas with high fluorophilic THF, the solvent loading should be less than 10% to avoid fluorous sample breakthrough. Table 8 provides recommendations for some common loading solvents and cartridge sizes.

The usual symptom of breakthrough is that a small amount of the fluorous compound comes off early in the organic fraction, but the bulk of the fluorous compound is retained on the cartridge. This happens because the loading solvent elutes the fluorous compounds, but the elution stops as soon as the fluorophobic solvent elutes the loading solvent from the cartridge. Breakthrough problems can often be solved by using less volume of loading solvent. Other solutions are to use a more fluorophobic loading solvent, to use a larger cartridge, or to lower the sample mass loading.

A good loading solvent should have low fluorophilicity and good dissolving power for organic compounds. Direct loading of a reaction mixture onto a fluorous cartridge for SPE is sometimes possible. However, in common practice, the reaction mixture is usually filtered first to remove insoluble solid and catalysts. The concentrated crude mixture is then dissolved in an appropriate loading solvent and loaded onto a cartridge preconditioned with a fluorophobic solvent. In large-scale F-SPE, an aqueous workup of reaction mixture is recommended. This removes water-soluble materials and preserves the lifetime of cartridge for reuse.

5.3.2. Elution solvents. An F-SPE has two solvent passes, the first one uses the fluorophobic solvent and the second one uses the fluorophilic solvent. The fluorophobic solvent is usually a water-miscible organic solvent with certain amount of water to reduce fluorophilicity. Solvents such as 70:30 MeCN–H₂O, 80:20 MeOH–H₂O, and 90:10 DMF–H₂O are the common choices. Acetone–H₂O and THF–H₂O can also serve the purpose. If a component in the reaction mixture is water sensitive, then 100% DMSO can be used for fluorophobic wash. All the non-fluorous components are expected to elute with the fluorophobic solvent in 3–5 column volumes, while fluorous components are retained on the cartridge. If organic components have low solubility in elution solvent, this can occasionally generate a precipitate and block the cartridge during F-SPE. Reduced mass loading and slightly increased the percentage of organic solvent can minimize this problem. After the elution of non-fluorous components, a more fluorophilic solvent such as MeOH, acetone, MeCN, or THF is used to wash out the fluorous component retained on the cartridge in 3–5 column volumes.

5.3.3. Fluorous silica gel reuse. To control the cost spent on fluorous silica gel and reduce waste disposal, the cartridges can be washed thoroughly and conditioned for reuse. The fluorous stationary phase can be cleaned by washing with fluorophilic solvents (acetone, MeCN, and THF) or with a mixture of MeCN–H₂O containing 0.5% TFA. To extend cartridge lifetime, crude samples containing strongly acidic or basic compounds, or having insoluble solids or a large amount of salts are not recommended for directly load onto the cartridge without pretreatment. However, if the cartridge will be discarded after use, the precautions are not necessary.

5.3.4. Gravity F-SPE with large fluorous silica gel. Compared to normal F-SPE with 40 μm fluorous silica gel, the resolution of gravity F-SPE with 120 μm silica gel is reduced to some extent. Cartridges with the large size fluorous silica gel need to be carefully conditioned to remove air bubbles. In the case of plate-to-plate F-SPE, the plate loaded with a high boiling solvent such as DMF or DMSO is degassed in a vacuum chamber (20–30 mmHg) for 3–5 min to remove the air bubbles.

6. Conclusions

The results summarized in this report show that the new separation technique of fluorous solid-phase extraction (F-SPE) has successfully debuted and is now ready for prime time.

While there is still more to be learned, basic F-SPE techniques are well understood and have proven generality. The predictability and generic ‘fluorous/non-fluorous’ nature of F-SPE’s make them especially attractive in research settings with single compounds or compound libraries. The learning curve is not steep; indeed, you can be up and running in the lab with your first F-SPE in as little as 15–30 min. Fluorous silica gel is now commercially available in an assortment of sizes and formats, and an increasing number of fluorous reaction components (reagents, catalysts, tags, scavengers, protecting groups) are also sold commercially. Thus, we expect that the usefulness of F-SPE will continue to expand as it is applied to more and different problems, and we intend that this report will help to fuel that expansion.

Note added in proof

A special issue of fluorous chemistry containing several papers dealing with fluorous solid-phase extraction has just appeared: Zhang, W. *QSAR Comb. Sci.* **2006**, *25*, 679.

Acknowledgements

We thank the National Institutes of Health, National Institutes of General Medical Science, for generous grant funding (1R43GM062717-01, 2R44GM062717-02, 1R43GM066415-01, 1R43GM067326-01, and 2R43GM067326-01).

References and notes

- (a) Curran, D. P. *Chemtracts: Org. Chem.* **1996**, *9*, 75–87; (b) Curran, D. P. *Angew. Chem., Int. Ed.* **1998**, *37*, 1175–1196; (c) Yoshida, J.; Itami, K. *Chem. Rev.* **2002**, *102*, 3693–3716; (d) Tzschucke, C. C.; Markert, C.; Bannwarth, W.; Roller, S.; Hebel, A.; Haag, R. *Angew. Chem., Int. Ed.* **2002**, *41*, 3964–4000.
- Wender, P. A.; Handy, S. T.; Wright, D. L. *Chem. Ind. (London)* **1997**, 765–767.
- Curran, D. P.; Hadida, S.; He, M. *J. Org. Chem.* **1997**, *62*, 6714–6715.
- Previous reviews covering aspects of fluorous silica gel and F-SPE: (a) Curran, D. P. *Synlett* **2001**, 1488–1496; (b) Curran, D. P. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 101–127.
- (a) Horvath, I. T.; Rabai, J. *Science* **1994**, *266*, 72–75; (b) Horvath, I. T. *Aqueous-Phase Organometallic Catalysis*, 2nd ed.; Cornils, B., Hermann, W. A., Eds.; Wiley: Weinheim, 2004; pp 646–654.
- (a) Overview of the field: *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; (b) Scope and definition of terms: Horvath, I. T.; Curran, D. P.; Gladysz, J. A. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 1–4.
- Nakamura, H.; Linclau, B.; Curran, D. P. *J. Am. Chem. Soc.* **2001**, *123*, 10119–10120.
- Gladysz, J. A.; Correa de Costa, R. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 24–40.
- (a) Curran, D. P.; Luo, Z. Y. *J. Am. Chem. Soc.* **1999**, *121*, 9069–9072; (b) Zhang, Q.; Luo, Z.; Curran, D. P. *J. Org. Chem.* **2000**, *65*, 8866–8873.
- Reviews of light fluorous chemistry: (a) Zhang, W. *Tetrahedron* **2003**, *59*, 4475–4489; (b) Zhang, W. *Chem. Rev.* **2004**, *104*, 2531–2556; (c) Curran, D. P. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 128–156; (d) Curran, D. P. *Aldrichimica Acta* **2006**, *39*, 3–9.
- Grubbs, R. H. *Handbook of Metathesis*; Wiley-VCH: Chichester, UK, England, 2003.
- Yao, Q. W.; Zhang, Y. L. *J. Am. Chem. Soc.* **2004**, *126*, 74–75.
- Matsugi, M.; Curran, D. P. *J. Org. Chem.* **2005**, *70*, 1636–1642.
- (a) Fukui, Y.; Brueckner, A. M.; Shin, Y.; Balachandran, R.; Day, B. W.; Curran, D. P. *Org. Lett.* **2006**, *8*, 301–304; (b) Moura-Letts, G.; Curran, D. P. *Org. Lett.*, in press.
- These and many other separation techniques have been compared and contrasted in detail in the setting of Mitsunobu reactions: (a) Dandapani, S.; Curran, D. P. *Chem.—Eur. J.* **2004**, *10*, 3130–3138; (b) Dembinski, R. *Eur. J. Org. Chem.* **2004**, 2763–2772.
- (a) Gladysz, J. A. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 41–55; (b) Curran, D. P.; Wang, X. A.; Zhang, Q. S. *J. Org. Chem.* **2005**, *70*, 3716–3719; (c) Chen, C. H.-T.; Zhang, W. *Mol. Divers* **2005**, *9*, 353–359.
- Matsugi, M.; Curran, D. P. *Org. Lett.* **2004**, *6*, 2717–2720.
- (a) Majors, R. E. *LC-GC* **1998**, 8–18; (b) Handley, A. J.; McDowall, R. D. *Extraction Methods in Organic Analysis*; Handley, A. J., Ed.; Sheffield Academic: Sheffield, UK, 1999; pp 54–74; (c) For a special issue on SPE for sample preparations, see: Poole, C. F.; Wilson, I. D. *J. Chromatogr. A* **2000**, *885*, 1–471.
- (a) www.fluorous.com; (b) DPC founded FTI and owns an equity interest in the company.
- Reviews covering fluorous solvents: (a) Gladysz, J. A.; Emnet, C. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 11–23; (b) Ryu, I.; Matsubara, H.; Emnet, C.; Gladysz, J. A.; Takeuchi, S.; Nakamura, Y.; Curran, D. P. *Green Reaction Media in Organic Synthesis*; Mikami, K., Ed.; Blackwell: Oxford, 2005; pp 59–124.
- For details of F-SPE procedures, see FTI Application Notes: http://fluorous.com/download/FTI_AppNote_F-SPE.pdf.
- FTI unpublished results.
- VacMaster[®]-96 SPE manifold and FlashMaster[™] II are products of Biotage (www.biotage.com).
- (a) Whatman[®] 24-well (10 mL) receiving plates, product number 7701-51022 (www.whatman.com); (b) 96-Well Ex-Block is a product of Exelixis, (www.exelixis.com).
- (a) Zhang, W.; Lu, Y.; Nagashima, T. *J. Comb. Chem.* **2005**, *7*, 893–897; (b) Zhang, W.; Lu, Y. *J. Comb. Chem.*, ASAP article.
- RapidTrace[®] system is available from Caliper Life Sciences (www.caliperls.com).
- (a) Pearson, W. H.; Berry, D. A.; Stoy, P.; Jung, K. Y.; Sercel, A. D. *J. Org. Chem.* **2005**, *70*, 7114–7122; (b) Beller, C.; Bannwarth, W. *Helv. Chim. Acta* **2005**, *88*, 171–179.
- (a) Filippov, D. V.; van Zoelen, D. J.; Oldfield, S. P.; van der Marel, G. A.; Overkleef, H. S.; Drijfhout, J. W.; van Boom, J. H. *Tetrahedron Lett.* **2002**, *43*, 7809–7812; (b) de Visser, P. C.; van Helden, M.; Filippov, D. V.; van der Marel, G. A.; Drijfhout, J. W.; van Boom, J. H.; Noort, D.; Overkleef, H. S. *Tetrahedron Lett.* **2003**, *44*, 9013–9016.

29. Brittain, S. M.; Ficarro, S. B.; Broack, S.; Peters, E. C. *Nat. Biotechnol.* **2005**, *23*, 463–468.
30. Palmacci, E. R.; Hewitt, M. C.; Seeberger, P. H. *Angew. Chem., Int. Ed.* **2001**, *40*, 4433–4437.
31. (a) Luo, Z. Y.; Zhang, Q. S.; Oderaotoshi, Y.; Curran, D. P. *Science* **2001**, *291*, 1766–1769; (b) Short review: Zhang, W. *ARKIVOC* **2004**, (i), 101–109.
32. Horn, J.; Michalek, F.; Tzschucke, C. C.; Bannwarth, W. *Top. Curr. Chem.* **2004**, *242*, 43–75.
33. Dandapani, S. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 175–182.
34. (a) Barthel-Rosa, L. P.; Gladysz, J. A. *Coord. Chem. Rev.* **1999**, *190–192*, 587–605; (b) de Wolf, E.; van Koten, G.; Deelman, B.-J. *Chem. Soc. Rev.* **1999**, *28*, 37–41; (c) Cavazzinni, M.; Montannari, F.; Pozzi, G.; Quici, S. *J. Fluorine Chem.* **1999**, *94*, 183–187.
35. Lindsley, C. W.; Leister, W. H. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 236–246.
36. (a) Zhang, W. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 222–235; (b) Zhang, W. *Curr. Opin. Drug Discov. Devel.* **2004**, *7*, 784–797.
37. Curran, D. P.; Luo, Z.; Degenkolb, P. *Bioorg. Med. Chem. Lett.* **1998**, *8*, 2403–2408.
38. Curran, D. P.; Hadida, S.; Kim, S. Y.; Luo, Z. Y. *J. Am. Chem. Soc.* **1999**, *121*, 6607–6615.
39. Lindsley, C. W.; Zhang, Z. *Handbook of Fluorous Chemistry*; Gladysz, J. A., Curran, D. P., Horvath, I. T., Eds.; Wiley-VCH: Weinheim, 2004; pp 371–373.
40. Christensen, C.; Clausen, R. P.; Begtrup, M.; Kristensen, J. L. *Tetrahedron Lett.* **2004**, *45*, 7991–7993.
41. Barthelemy, B.; Schneider, S.; Bannwarth, W. *Tetrahedron Lett.* **2002**, *43*, 807–810.
42. Lindsley, C. W.; Zhao, Z.; Newton, R. C.; Leister, W. H.; Strauss, K. A. *Tetrahedron Lett.* **2002**, 4467–4470.
43. Dandapani, S.; Curran, D. P. *Tetrahedron* **2002**, *58*, 3855–3864.
44. Dandapani, S.; Curran, D. P. *J. Org. Chem.* **2004**, *69*, 8751–8757.
45. Desai, B.; Dallinger, D.; Kappe, C. O. *Tetrahedron* **2006**, *62*, 4651–4664.
46. Zhang, W.; Lu, Y. *QSAR Comb. Sci.* **2006**, *25*, 724–727.
47. Nagashima, T.; Lu, Y. M.; Petro, M. J.; Zhang, W. *Tetrahedron Lett.* **2005**, *46*, 6585–6588.
48. Dalicsek, Z.; Pollreisz, F.; Gomory, A.; Soos, T. *Org. Lett.* **2006**, *8*, 1093–1095.
49. Kaleta, Z.; Makowski, B. T.; Soos, T.; Dembinski, R. *Org. Lett.* **2006**, *8*, 1625–1628.
50. Kainz, S.; Luo, Z. Y.; Curran, D. P.; Leitner, W. *Synthesis* **1998**, 1425–1427.
51. Ryu, I.; Kreimerman, S.; Niguma, T.; Minakata, S.; Komatsu, M.; Luo, Z. Y.; Curran, D. P. *Tetrahedron Lett.* **2001**, *42*, 947–950.
52. Kaleta, Z.; Egyed, O.; Soos, T. *Org. Biomol. Chem.* **2005**, *3*, 2228–2230.
53. Nakamura, Y.; Takeuchi, S.; Ohgo, Y.; Curran, D. P. *Tetrahedron* **2000**, *56*, 351–356.
54. Nakamura, Y.; Takeuchi, S.; Okumura, K.; Ohgo, Y.; Curran, D. P. *Tetrahedron* **2002**, *58*, 3963–3969.
55. Fawcett, J.; Hope, E. G.; Stuart, A. M.; West, A. J. *Green Chem.* **2005**, *7*, 316–320.
56. Hope, E. G.; Stuart, A. M.; West, A. J. *Green Chem.* **2004**, *6*, 345–350.
57. Nakamura, Y.; Takeuchi, S.; Okumura, K.; Ohgo, Y. *Tetrahedron* **2001**, *57*, 5565–5571.
58. Vallin, K. S. A.; Zhang, Q. S.; Larhed, M.; Curran, D. P.; Hallberg, A. *J. Org. Chem.* **2003**, *68*, 6639–6645.
59. Beeler, A. B.; Acquilano, D. E.; Su, Q.; Yan, F.; Roth, B. L.; Panek, J. S.; Porco, J. A., Jr. *J. Comb. Chem.* **2005**, *7*, 673–681.
60. Bucher, B.; Curran, D. P. *Tetrahedron Lett.* **2000**, *41*, 9617–9621.
61. Cavazzini, M.; Quici, S.; Pozzi, G. *Tetrahedron* **2002**, *58*, 3943–3949.
62. Simonelli, B.; Orlandi, S.; Benaglia, M.; Pozzi, G. *Eur. J. Org. Chem.* **2004**, 2669–2673.
63. Croxtall, B.; Hope, E. G.; Stuart, A. M. *Chem. Commun.* **2003**, 2430–2431.
64. Curran, D. P.; Fischer, K.; Moura-Letts, G. A. *Synlett* **2004**, 1379–1382.
65. Dalicsek, Z.; Pollreisz, F.; Gomory, A.; Soos, T. *Org. Lett.* **2005**, *7*, 3243–3246.
66. Hein, J. E.; Geary, L. M.; Jaworski, A. A.; Hultin, P. G. *J. Org. Chem.* **2005**, *70*, 9940–9946.
67. Hein, J. E.; Hultin, P. G. *Synlett* **2003**, 635–638.
68. Hein, J. E.; Zimmerman, J.; Sibi, M. P.; Hultin, P. G. *Org. Lett.* **2005**, *7*, 2755–2758.
69. Hein, J. E.; Hultin, P. G. *Tetrahedron: Asymmetry* **2005**, *16*, 2341–2347.
70. Zhang, W.; Chen, C. H.-T.; Nagashima, T. *Tetrahedron Lett.* **2003**, *44*, 2065–2068.
71. Zhang, W.; Curran, D. P.; Chen, C. H.-T. *Tetrahedron* **2002**, *58*, 3871–3875.
72. Lindsley, C. W.; Zhao, Z.; Leister, W. H.; Robinson, R. G.; Barnett, S. F.; Defeo-Jones, D.; Jones, R. E.; Huber, H. E. *Bioorg. Med. Chem. Lett.* **2005**, *15*, 761–764.
73. Lindsley, C. W.; Bogusky, M. J.; Leister, W. H.; McClain, R. T.; Robinson, R. G.; Barnett, S. F.; Defeo-Jones, D.; Hartman, G. D. *Tetrahedron Lett.* **2005**, *46*, 2779–2782.
74. Lindsley, C. W.; Zhao, Z.; Leister, W. H. *Tetrahedron Lett.* **2002**, *43*, 4225–4228.
75. Zhang, A. S.; Elmore, C. S.; Egan, M. A.; Melillo, D. G.; Dean, D. C. *J. Labelled Compd. Radiopharm.* **2005**, *48*, 203–208.
76. Lindsley, C. W.; Zhao, Z.; Leister, W. H.; Strauss, K. A. *Tetrahedron Lett.* **2002**, *43*, 6319–6323.
77. Werner, S.; Curran, D. P. *Org. Lett.* **2003**, *5*, 3293–3296.
78. Lu, Y.; Zhang, W. *QSAR Comb. Sci.* **2006**, *25*, 728–731.
79. Rover, S.; Wipf, P. *Tetrahedron Lett.* **1999**, *40*, 5667–5670.
80. Wipf, P.; Reeves, J. T.; Rover, S. U.S. Patent 06,673,539, 2004.
81. Wipf, P.; Reeves, J. T. *Tetrahedron Lett.* **1999**, *40*, 4649–4652.
82. Curran, D. P.; Amatore, M.; Guthrie, D.; Campbell, M.; Go, E.; Luo, Z. Y. *J. Org. Chem.* **2003**, *68*, 4643–4647.
83. Luo, Z. Y.; Williams, J.; Read, R. W.; Curran, D. P. *J. Org. Chem.* **2001**, *66*, 4261–4266.
84. Zhang, W.; Tempest, P. *Tetrahedron Lett.* **2004**, *45*, 6757–6760.
85. Matsugi, M.; Yamanaka, K.; Inomata, I.; Takekoshi, N.; Hasegawa, M.; Curran, D. P. *QSAR Comb. Sci.* **2006**, *25*, 732–735.
86. Curran, D. P.; Ogoe, C. *QSAR Comb. Sci.* **2006**, *25*, 713–715.
87. Zhang, W. Unpublished results.
88. Villard, A.-L.; Warrington, B. H.; Ladlow, M. *J. Comb. Chem.* **2004**, *6*, 611–622.

89. Fustero, S.; Sancho, A. G.; Chiva, G.; Sanz-Cervera, J. F.; del Pozo, C.; Aceña, J. L. *J. Org. Chem.* **2006**, *71*, 3299–3302.
90. Manzoni, L.; Castelli, R. *Org. Lett.* **2006**, *8*, 955–957.
91. Jing, Y.; Huang, X. *Tetrahedron Lett.* **2004**, *45*, 4615–4618.
92. Manzoni, L. *Chem. Commun.* **2003**, 2930–2931.
93. Manzoni, L.; Castelli, R. *Org. Lett.* **2004**, *6*, 4195–4198.
94. Donovan, A.; Forbes, J.; Dorff, P.; Schaffer, P.; Babich, J.; Valliant, J. F. *J. Am. Chem. Soc.* **2006**, *128*, 3536–3537.
95. Zhang, W.; Chen, C. H.-T.; Lu, Y.; Nagashima, T. *Org. Lett.* **2004**, *6*, 1473–1476.
96. Zhang, W.; Lu, Y.; Chen, C. H.-T. *Mol. Divers.* **2003**, *7*, 199–202.
97. Zhang, W.; Nagashima, T.; Lu, Y.; Chen, C. H.-T. *Tetrahedron Lett.* **2004**, *45*, 4611–4613.
98. Zhang, W.; Nagashima, T. *J. Fluorine Chem.* **2006**, *127*, 588–591.
99. Cioffi, C. L.; Berlin, M. L.; Herr, R. J. *Synlett* **2004**, 841–845.
100. Chen, C. H.-T.; Zhang, W. *Org. Lett.* **2003**, *5*, 1015–1017.
101. Zhang, W. *Org. Lett.* **2003**, *5*, 1011–1014.
102. McAllister, L. A.; McCormick, R. A.; Brand, S.; Procter, D. J. *Angew. Chem., Int. Ed.* **2005**, *44*, 452–455.
103. Kasahara, T.; Kondo, Y. *Chem. Commun.* **2006**, 891–893.
104. Lu, Y.; Zhang, W. *QSAR Comb. Sci.* **2004**, *23*, 827–835.
105. Zhang, W.; Chen, C. H. T. *Tetrahedron Lett.* **2005**, *46*, 1807–1810.
106. Fustero, S.; Catalán, S.; Flores, S.; Jiménez, D.; del Pozo, C.; Aceña, J. L.; Sanz-Cervera, J. F.; Mérida, S. *QSAR Comb. Sci.* **2006**, *25*, 753–760.
107. (a) Zhang, W.; Lu, Y. *Org. Lett.* **2003**, *5*, 2555–2558; (b) Lu, Y.; Zhang, W. *Mol. Divers.* **2005**, *9*, 91–98.
108. Zhang, W.; Nagashima, T., unpublished results.
109. Nagashima, T.; Zhang, W. *J. Comb. Chem.* **2004**, *6*, 942–949.
110. Zhang, W.; Lu, Y.; Geib, S. *Org. Lett.* **2005**, *7*, 2269–2272.
111. (a) Zhang, W.; Lu, Y.; Chen, C. H.-T.; Curran, D. P.; Geib, S. *Eur. J. Org. Chem.* **2006**, 2055–2059; (b) Zhang, W.; Lu, Y.; Chen, C.-H.; Zeng, L.; Kassel, D. B. *J. Comb. Chem.* **2006**, *8*, 687–695.

Biographical sketch

Wei Zhang received his B.S. from Nanjing University and his Ph.D. in 1993 from the University of Pittsburgh. After a two-year appointment at the University of Pittsburgh as a Research Assistant Professor, he joined DuPont Agricultural Chemicals in 1995. In early 2001, he joined the newly founded Fluorous Technologies, Inc. and later became the Director of Discovery Chemistry. His professional interests include free radical chemistry, fluororous chemistry, microwave-assisted synthesis, and development of new high-throughput technologies for synthesis and purification of compound libraries. He has over 80 peer-reviewed publications in these research areas.



Dennis P. Curran received his B.S. in 1975 from Boston College and his Ph.D. in 1979 from the University of Rochester. After a two year post-doctoral stay at the University of Wisconsin, he joined the Chemistry Department at the University of Pittsburgh in 1981 and is now a Distinguished Service Professor and Bayer Professor of Chemistry. He is the founder of Fluorous Technologies, Inc., and has received the Morley Medal (2006), the Pittsburgh Magazine Innovators Award (2003), the American Chemical Society Award for Creativity in Organic Synthesis (2000), the ACS Cope Scholar Award (1988), and the Janssen Prize for Creativity in Organic Synthesis (1998). He is currently an ISI Highly Cited Researcher (www.isihighlycited.com), and is well known for his research in synthetic radical chemistry and fluororous chemistry.