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## Bifluoride-catalysed sulfur(VI) fluoride exchange reaction for the synthesis of polysulfates and polysulfonates

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### Abstract

Polysulfates and polysulfonates possess exceptional mechanical properties making them potentially valuable engineering polymers. However, they have been little explored due to a lack of reliable synthetic access. Here we report bifluoride salts ( $Q^+[FHF]^-$ , where  $Q^+$  represents a wide range of cations) as powerful catalysts for the sulfur(VI) fluoride exchange (SuFEx) reaction between aryl silyl ethers and aryl fluorosulfates (or alkyl sulfonyl fluorides). The bifluoride salts are significantly more active in catalysing the SuFEx reaction compared to organosuperbases, therefore enabling much lower catalyst-loading (down to 0.05 mol%). Using this chemistry, we are able to prepare polysulfates and polysulfonates with high molecular weight, narrow polydispersity and excellent functional group tolerance. The process is practical with regard to the reduced cost of catalyst, polymer purification and by-product recycling. We have also observed that the process is not sensitive to scale-up, which is essential for its future translation from laboratory research to industrial applications.

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#### Author contributions

K.B.S. and P.W. led the project. J.D. and B.G. designed the experiments. B.G., J.D. and L.Z. carried out the experiments. F.Z., L.M.K., J.L. and Y.L. collected and analysed the TGA and DSC data of all polymers and provided helpful suggestion on the project. B.G. wrote the manuscript. P.W., Q.Z., J.D., Y.L. and K.B.S. edited the manuscript.

#### Competing financial interests

A patent application covering this work has been filed by TSRI (US patent application no. 62/182755, International patent application no. PCT/US2016/038701).

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Click chemistry features the formation of carbon–heteroatom bonds to generate molecules with desired functionality using highly efficient, selective, modular and benign reaction processes<sup>1</sup>. From a synthetic perspective, polymer chemistry depends on many of the same basic principles as click chemistry —modularity and efficiency are inherent in practical polymer syntheses<sup>2</sup>, which explains the growing impact of click chemistry on materials science over the past decade<sup>3</sup>, especially in the field of polymer preparation and modification<sup>4,5</sup>.

Recently, we introduced sulfur(VI) fluoride exchange (SuFEx) as another embodiment of click chemistry<sup>6</sup>. The SuFEx click chemistry allows the activation of S<sup>VI</sup>-F bond with a 'Si<sup>+</sup>' species in the presence of organosuperbase catalysts, such as 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) and 2-*tert*-butylimino-2-diethylamino-1,3-dimethylperhydro-1,3,2-diazaphosphorine (BEMP, a phosphazene base—full name cited from Reaxys). With these catalysts, the reaction between aryl fluorosulfates (or aryl sulfonyl fluorides) and aryl silyl ethers exclusively affords diaryl sulfates (or diaryl sulfonates). The SuFEx reaction has been successfully applied to the synthesis of aromatic polysulfates and polysulfonates by our group<sup>7,8</sup>, and to post-polymerization modifications by others<sup>9–11</sup>.

Polysulfates and polysulfonates with their sulfur(VI) linkages (–SO<sub>2</sub>–) possess excellent mechanical properties and high chemical stability, as needed for engineering polymers<sup>12–15</sup>. For instance, the bisphenol A (BPA) polysulfate has slightly higher tensile modulus, similar yield stress and significantly lower oxygen permeability compared to its polycarbonate counterpart<sup>7</sup>. Nevertheless, they have been rarely used for industrial applications due to a lack of reliable and scalable synthetic access. Our SuFEx process represents the first efficient protocol for the preparation of these polymers. Considering the availability of bisphenols (BPA has an annual production of millions of tonnes, most of which are used for polycarbonates and epoxy resins industry) and sulfonyl fluoride gas (SO<sub>2</sub>F<sub>2</sub>, widely used as fumigant) as commercial feed-stocks, it is attractive to further explore this process for potential industrial applications.

However, several disadvantages of the base-catalysed SuFEx process have hampered its scale-up. First, the catalyst loading is quite high. For a typical polymerization reaction, no less than 10 mol% of DBU has to be used. One limitation of high catalyst-loading is that a tedious purification procedure is required to remove catalyst residue prior to polymer processing. Second, BEMP requires only 1.0 mol% of catalyst-loading, but is quite expensive and lacks bulk accessibility. Third, the strong basic nature of the catalysts restricts the substrate scope of the process. For instance, alkyl sulfonyl fluoride monomers are not well compatible with this process due to the facile deprotonation of their  $\alpha$ -H when subjected to strong bases<sup>16,17</sup>.

Addressing these issues, we report a new set of catalysts for SuFEx-based polysulfate and polysulfonate synthesis, namely the bifluoride salts (Q<sup>+</sup>[FHF]<sup>–</sup>, where Q<sup>+</sup> refers to a wide range of organic and inorganic cations)<sup>18,19</sup>. The bifluoride species has been little appreciated by the synthetic community. Its presence is nearly certain wherever a fluoride ion is exposed to any potential proton sources<sup>20</sup>. The 4-electron-3-centered anion ([FHF]<sup>–</sup>) has one proton trapped between two fluoride anions via the strongest hydrogen bond ever

recorded<sup>21</sup>. Such a formation gives the bifluoride ion modest acidity ( $\text{pH} \approx 3.0$  for the saturated aqueous solution of potassium bifluoride at room temperature)<sup>22</sup>, moderate nucleophilicity<sup>23</sup> and excellent stability<sup>21</sup>. While the strongly basic fluoride sources, such as cesium fluoride, only exhibited moderate activity in our SuFEx reaction<sup>7</sup>, we hypothesized that the acidic bifluoride species could facilitate SuFEx reactivity.

## Results and discussion

We first investigated potassium bifluoride ( $\text{K}^+[\text{FHF}]^-$ ) as catalyst for the polycondensation of **A-1** and **B-1** in *N*-methyl-2-pyrrolidone (NMP) at 130 °C (Fig. 1a). The polymerization reaction catalysed by DBU (10 mol%) and BEMP (1 mol%) served as controls (Fig. 1b; catalyst loadings are given in molar ratio to the total amount of both monomers). The number-average molecular weight ( $M_n^{ps}$ ) (ps, polystyrene) and polydispersity (PDI) were determined by gel permeation chromatography (GPC), which was calibrated with commercial polystyrenes. Our results indicated that 2.0 mol% of potassium bifluoride was indeed able to promote the polysulfate formation. But it was much slower compared to the reaction catalysed with DBU or BEMP (17 h versus 1 h), affording the polysulfate **P-1** with a low molecular weight ( $M_n^{ps} = 28$  kDa). However, the molecular weight of **P-1** could be dramatically improved to as high as 100 kDa by combining 2.0 mol% of potassium bifluoride with 1.0 mol% of tetrabutylammonium chloride or 18-crown-6 ether as catalyst. In all entries, the PDI remained narrow (1.4~1.6, Fig. 1c).

Subsequently, we investigated the impact of cations on the activities of the bifluoride catalysts ( $\text{Q}^+[\text{FHF}]^-$ , Fig. 1d). Various onium bifluoride salts were prepared through ion-exchange of their chloride/bromide precursors ( $\text{Q}^+\text{X}^-$ , X = Cl, Br) with silver bifluoride ( $\text{Ag}^+[\text{FHF}]^-$ ), and were used as 0.1 M stock solution in acetonitrile (Supplementary Section 2-3-1). We found that a wide variety of onium bifluorides were highly active at promoting the reaction, requiring only 0.1 mol% loading in most cases (the minimum amount required to produce **P-1** with  $M_n^{ps} > 20$  kDa). Upon the addition of the bifluoride catalyst, the reaction was triggered instantly with the release of *tert*-butyldimethylsilyl fluoride (TBSF, b.p. 89 °C)<sup>24</sup>, and yet not much heat was released. Within one hour **P-1** was obtained with  $M_n^{ps}$  ranging from 30 kDa to 100 kDa (PDI = 1.4~1.7). We found that tris(dimethylamino)sulfonium bifluoride (**Q-5**)<sup>25</sup> was the most effective catalyst—at a 0.05 mol% catalyst loading it was able to generate **P-1** with a molecular weight up to 100 kDa. Hexamethyl guanidinium bifluoride (**Q-6**) also exhibited remarkable activity at 0.08 mol% catalyst loading (PDI = 2.0). However, a confined, five-membered cyclic version of the cation (**Q-7**) had decreased activity (0.25 mol% loading).

Inspired by the extraordinary activity of bifluoride salts in catalysing the SuFEx polymerization reaction, we investigated whether poly(hydrogen fluoride) salts ( $\text{Q}^+[\text{H}_n\text{F}_{n+1}]^-$ ,  $n > 1$ , Fig. 1e) could promote the transformation as well<sup>18</sup>. The dihydrogen trifluoride salt **Q-11** was prepared via ion-exchange of its chloride precursor with excess  $\text{K}^+[\text{FHF}]^-$  (ref. 26). Thereafter, equilibrations of **Q-11** with hydrofluoric acid (48 wt.% in  $\text{H}_2\text{O}$ ) in acetonitrile gave catalysts **Q-12–Q-14** *in situ*, which were used directly without the removal of water (Supplementary Section 2–3). We found **Q-11–Q-13** were excellent catalysts for **P-1** formation, although slightly higher catalyst-loadings were required (0.5 mol% in most

cases). Intriguingly, we confirmed the formation of **P-1** with a moderate molecular weight ( $M_n^{PS} = 20$  kDa) even when the HF:F<sup>-</sup> ratio was 9:1 (**Q-14**), suggesting that the polymerization reaction has an excellent tolerance for the presence of a few per cent of water and HF in the system. In contrast, we only obtained a mixture of oligomers and partly hydrolysed monomers when triethylamine trihydrofluoride, pyridinium poly(hydrogen fluoride)<sup>27</sup>, or hydrofluoric acid were employed as catalysts (see Supplementary Section 2–4 for more information on catalyst evaluation).

To evaluate the fidelity of the bifluoride catalytic system, we examined monomers with diverse functional groups (Table 1). The polysulfates **P-2**, **P-5** and **P-6** had previously been prepared under the DBU-catalysed polycondensation reaction<sup>7</sup>. When the bifluoride catalyst **Q-5** was used, we obtained these polysulfates with comparable molecular weights and polydispersities. For instance, the  $M_n^{PS}/PDI$  of **P-2** was 46 kDa/1.5 with the DBU catalyst versus 84 kDa/1.6 with the **Q-5** catalyst. Under the same reaction conditions, monomers **A-3/B-3**, **A-4/B-4** and **A-8/B-8** were converted to polysulfates with molecular weights ranging from 36 to 52 kDa. Even the sterically hindered monomer **B-9** underwent cross-condensation with **A-1** giving polymer **P-9** with high molecular weight and good polydispersity ( $M_n^{PS} = 58$  kDa, PDI = 1.7), alternatively **A-9** reacted with **B-1** giving **P-9** with a slightly lower molecular weight ( $M_n^{PS} = 23$  kDa, PDI = 1.4). The orthogonal nature of the SuFEx click reaction to many other standard chemical reactions enables us to incorporate latent functional groups on the emergent polymer backbones for post-polymerization modification. For example, propargyl groups in **P-12** can serve as handles for the CuAAC-mediated functionalization<sup>5</sup>. Alternatively, the methoxy groups in **P-11a** were converted into fluorosulfates (**P-11b**) to undergo further SuFEx modification<sup>11</sup>.

We have also examined **Q-5** (0.1 mol%) catalysed polycondensation of **AB-1**, in which the two functional groups, –OSO<sub>2</sub>F and –OTBS, were residents in a single molecule (**AB-1** is bench-stable for at least one year at room temperature). Under the same reaction conditions, polysulfate **P-1** was obtained with a  $M_n^{PS}$  of 65 kDa (PDI = 1.5), which is comparable to polymers obtained from the **A-1/B-1** polycondensation reaction (Fig. 2a). Intriguingly, we were also able to realize the direct polycondensation of **A-1** with SO<sub>2</sub>F<sub>2</sub> gas catalysed by **Q-5**, affording **P-1** with moderate molecular weight ( $M_n^{PS} = 22$  kDa, PDI = 1.4, Fig. 2b). This strategy is generally applicable to other monomers like **A-7**, with which polysulfate **P-7** was prepared in near quantitative conversion ( $M_n^{PS} = 28$  kDa, PDI = 1.5, Fig. 2c). In these cases, aryl fluorosulfate intermediates (–C<sub>6</sub>H<sub>4</sub>OSO<sub>2</sub>F) were generated *in situ* via the reaction between aryl silyl ether (–C<sub>6</sub>H<sub>4</sub>OTBS) and sulfuryl fluoride gas catalysed by **Q-5**, which underwent further condensation with another molecule of aryl silyl ether to extend the backbone and finally resulted in the polysulfate.

When DBU or BEMP was employed as the catalyst, the reactions of aryl silyl ether with alkyl sulfonyl fluoride afforded a mixture of oligomers and decomposition adducts of both monomers and oligomers, which might be due to the deprotonation of the α-H of the sulfonyl fluoride group<sup>17</sup>. Hydrogen fluoride catalysts, however, provide an acidic reaction environment that is compatible with those alkyl sulfonyl fluorides<sup>22</sup>. To demonstrate this, we combined the aliphatic bis(sulfonyl fluorides) **B-13**, **B-14** and **B-15** respectively with **A-1** in

NMP in 1:1 ratio and in the presence of 5 mol% catalyst **Q-5** at 130 °C. The resulting polysulfonates **P-13**, **P-14** and **P-15** were afforded in quantitative yields and moderate molecular weights (Table 2). On the other hand, the polycondensation of the aromatic bis(sulfonyl fluoride) **B-17** required lower catalyst-loading (0.5 mol%) and gave polysulfonate with a higher molecular weight. The bifluoride-catalysed SuFEx polycondensation reaction is also applicable to the synthesis of the fluoropolymer **P-16** from the perfluorobutane-1,4-disulfonyl fluoride **B-16** and the silyl ether **A-1** ( $M_n^{PS}$  = 23 kDa, PDI = 1.3). Fluoropolymers often possess excellent physical and chemical properties, such as low dielectric constants and coefficients of friction<sup>28</sup>.

For laboratory use of the bifluoride-catalysed SuFEx polymerization or ligation, there are several options for catalyst preparation summarized in Supplementary Section 2–3. For industrial interests, however, bulk supplies of high-quality catalysts at low cost is required. Fortunately, a reliable protocol for the large-scale preparation of **Q-5** from commercially available SF<sub>4</sub> has already been reported (Fig. 3a)<sup>25,29</sup>. In addition, there is a practical access to other bifluoride salts which involves the ion-exchange of onium halides (Q<sup>+</sup>X<sup>-</sup>, X = Cl, Br, I) with anhydrous hydrogen fluoride (Fig. 3b). Anhydrous HF and many onium halides are bulk chemicals, therefore practical industrial scale syntheses of bifluoride catalysts is expected. Demonstrating this, we carried out a 5-gram scale preparation of 1-ethyl-3-methylimidazolium bifluoride (EMI<sup>+</sup>[FHF]<sup>-</sup>) (modified procedure based on refs 30,31). (Caution! Anhydrous HF is dangerous for its strong corrosiveness and toxicity, and must be manipulated in specific apparatus under safety guidance. See Supplementary Section 2-3-6).

Having established the optimal conditions for the bifluoride-catalysed SuFEx polymerization reaction, as well as the protocols for catalysts preparation, we were further interested in enlarging the synthesis of polysulfate **P-1** to 100-gram scale in our laboratory. A bulk polymerization with 0.2 mol of **A-1** and 0.2 mol of **B-1** was carried out at ~120 °C (internal temperature) in NMP (50 ml), using only 0.05 mol% of catalyst **Q-5** (Fig. 3c). Polysulfate **P-1** was obtained as white fibrous materials in near quantitative yield (112.7 g, Fig. 3d). GPC analysis gave a  $M_n^{PS}$  of 110 kDa (PDI = 1.7, Supplementary Fig. 8), which was even higher than the **P-1** product obtained at the 2.0 mmol scale ( $M_n^{PS}$  = 92 kDa, PDI = 1.7, Fig. 1d). Differential scanning calorimetry and thermal gravimetric analysis results (Supplementary Section 2–8) were comparable with the data obtained from the BEMP-catalysed bulk polycondensation<sup>7</sup>, indicating these two batches of polymers had the same quality. We also examined the end-capping effect, which is a standard strategy for molecular weight-control (Supplementary Section 2–6)<sup>32</sup>. The only by-product of the SuFEx-polymerization process is the TBSF (Fig. 3e). Nevertheless, we found it can be used in place of *tert*-butyl-dimethylsilyl chloride for the production of the silyl ether monomer **A** and hence recycled (Supplementary Section 2–9).

## Conclusion

In summary, we have developed bifluoride salts as a new class of catalysts for the SuFEx click reaction, and have applied this chemistry to the preparation of polysulfates and polysulfonates. Compared to organosuperbase catalysts (for example, DBU and BEMP), the bifluoride ion [FHF]<sup>-</sup> is acidic. It allows the SuFEx polymerization reaction to be extended

to a broader substrates scope, such as aliphatic sulfonyl fluorides. Most importantly, the bifluoride catalyst is significantly more active, therefore the polymerization reaction requires much lower catalyst-loading, which is essential for cost-effective industrial scale production. Our large-scale experiment further confirms the high fidelity of the bifluoride catalytic process for scale-up. Moreover, we have also provided practical solutions for catalyst preparation and for by-product recycling (TBSF). This work sets a foundation for future translation of polysulfate synthesis from laboratory research to industrial applications.

## Methods

### Process for the bulk preparation of polysulfate P-1

Monomers **A-1** (91.36 g, 0.2 mol) and **B-1** (78.48 g, 0.2 mol) were combined in a dry 1 l three-neck round-bottom flask equipped with a Teflon-coated magnetic stir bar, a thermometer and a reflux condenser (Fig. 3c). The flask was purged with nitrogen gas, into which dry NMP (50 ml) was then added. Next, it was placed in a pre-heated oil bath (130 °C, oil temperature). With stirring, monomers gradually dissolved in NMP, and the internal temperature stabilized at 123 °C. Under vigorous stirring, catalyst **Q-5** (0.1 M in dry CH<sub>3</sub>CN, 2.0 ml, 0.0002 mol) was added. The reaction initiated instantaneously. The internal temperature rose to 135 °C, then quickly dropped to 100–110 °C due to the refluxing of TBSF. The reaction was allowed to run for 1 h, during which the reaction mixture turned highly viscous and the internal temperature stabilized at around 120 °C. The flask was subsequently cooled to room temperature. TBSF was then distilled out from the reaction mixture as colourless liquid at 80 °C (diaphragm pump; 49.1 g product, Fig. 3e). After the distillation, 250 ml DMF was added to the reaction flask to dissolve the polymer at 120 °C. The warm DMF solution was then slowly poured into 2 l methanol with vigorous mechanical stirring at room temperature. White fibres of **P-1** crashed out and aggregated like a bird's nest in methanol, which was collected and dried at 70 °C for 12 h to give 112.7 g dry polymer ( $M_n^{PS}$  = 110 kDa, PDI = 1.7, Fig. 3d). Full experimental details and characterization of compounds are given in Supplementary Information.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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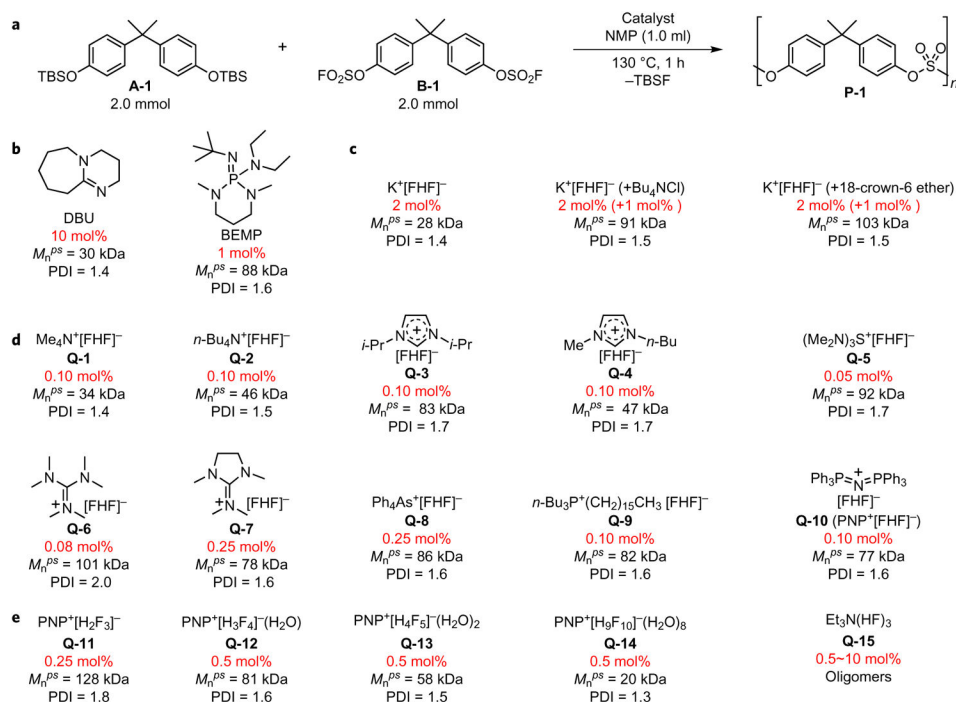


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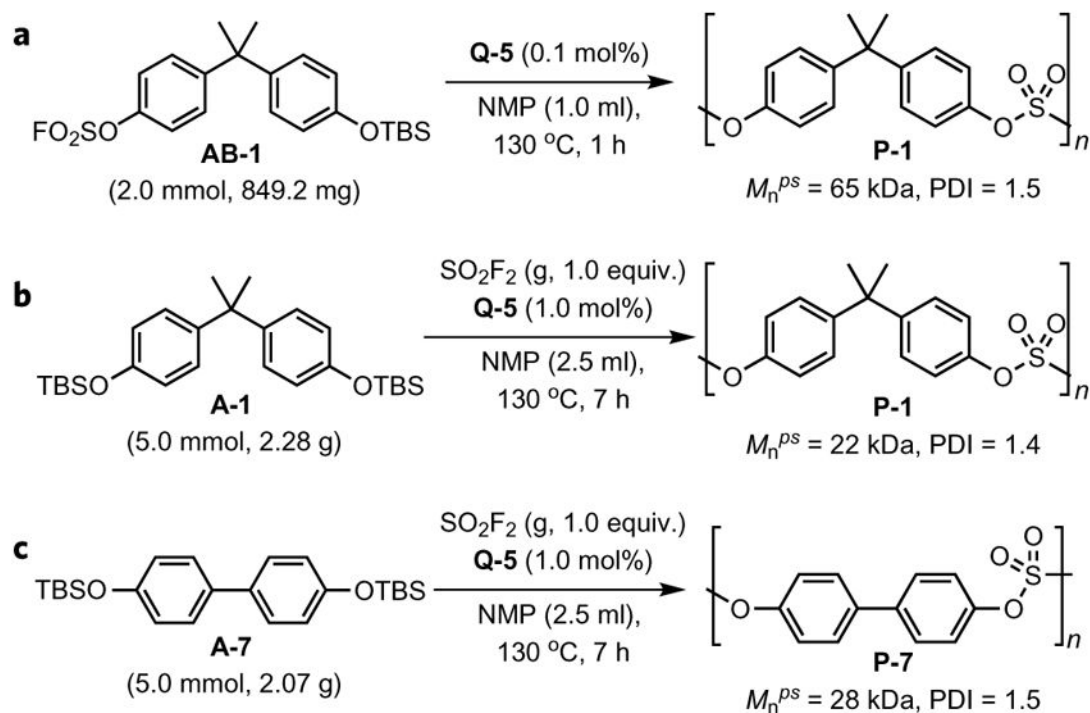


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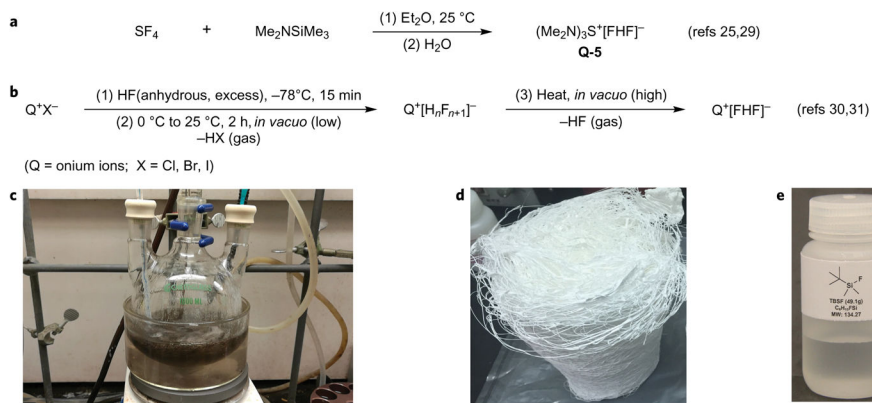
**Figure 1. Catalyst screening for the SuFEx-based synthesis of polysulfates**

**a**, The synthesis of **P-1** from monomers **A-1** and **B-1** is employed as a model reaction for catalyst screening. **b**, Control experiments with DBU or BEMP as catalysts. **c**, Evaluation of  $K^+[FHF]^-$  (with/without phase transfer reagents) as the catalyst with a reaction time of 17 h. **d**, Evaluation of onium bifluoride ( $Q^+[FHF]^-$ ) salts as catalysts. **e**, Evaluation of poly(hydrogen fluoride) salts as catalysts. NMP, *N*-methyl-2-pyrrolidone; TBS, *tert*-butyldimethylsilyl;  $M_n^{PS}$ , number-average molecular weight with polystyrene as standard; PDI, polydispersity index.



**Figure 2. Two alternative strategies for the synthesis of polysulfates via the bifluoride-catalysed SuFEx reaction**

**a**, The synthesis of **P-1** from the bifunctional monomer **AB-1**. **b**, The synthesis of **P-1** from the polycondensation of **A-1** with sulfuryl fluoride gas ( $\text{SO}_2\text{F}_2$ ). **c**, The synthesis of **P-7** from the polycondensation of **A-7** with sulfuryl fluoride gas ( $\text{SO}_2\text{F}_2$ ). NMP, *N*-methyl-2-pyrrolidone; TBS, *tert*-butyldimethylsilyl;  $M_n^{ps}$ , number-average molecular weight with polystyrene as standard; PDI, polydispersity index.

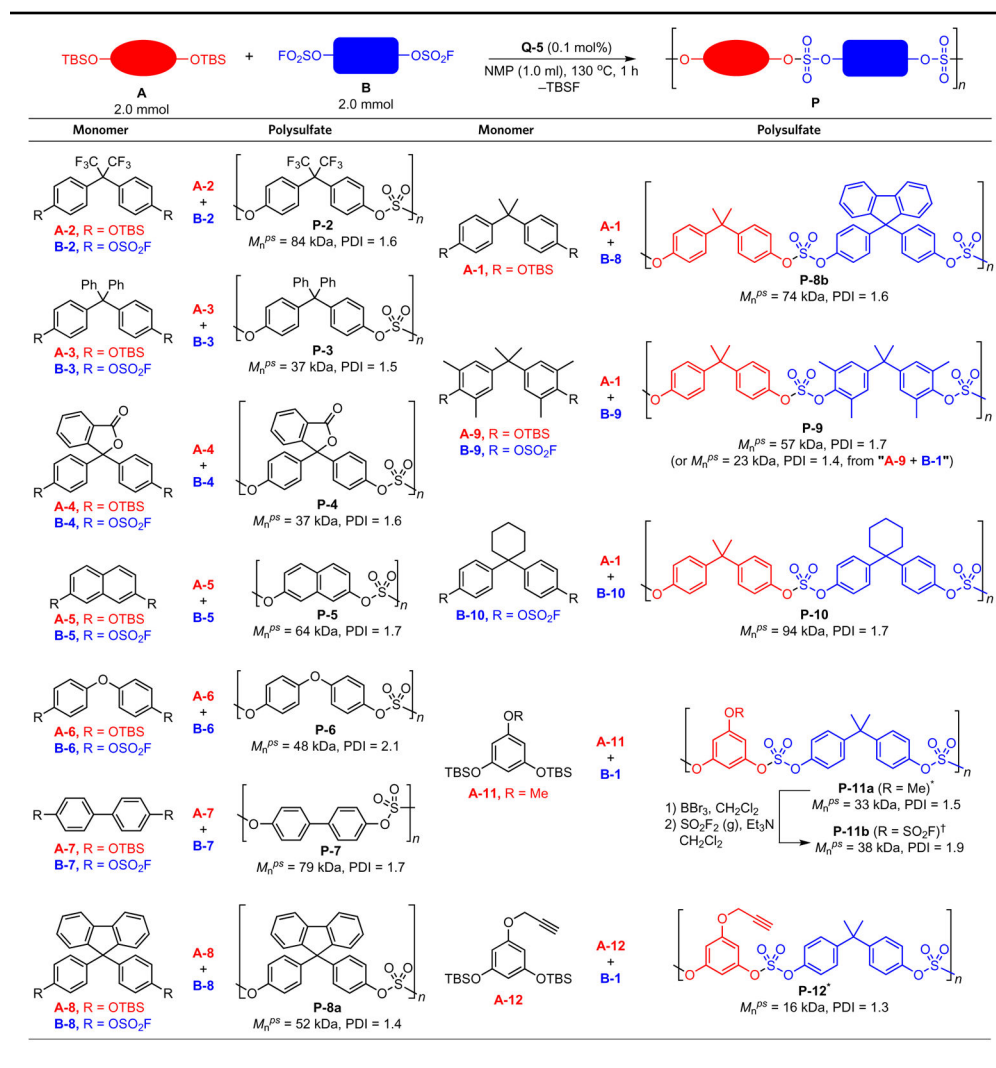


**Figure 3. Catalysts preparation and bulk synthesis of P-1**

**a**, Reported procedure for the preparation of catalyst **Q-5**. **b**, A schematic protocol for the preparation of bifluoride catalysts via the ion-exchange reaction of onium halides ( $\text{Q}^+\text{X}^-$ , X = Cl, Br, I) with anhydrous hydrogen fluoride (see Supplementary Section 2-3-6 for details). **c**, The 100-gram scale preparation of **P-1** from the polycondensation of **A-1** (0.2 mol) and **B-1** (0.2 mol) catalysed by 0.05 mol% of **Q-5** in NMP (50 ml). **d**, 112.7 g **P-1** was obtained from bulk polymerization ( $M_n^{PS} = 110$  kDa, PDI = 1.7). **e**, 49.1 g TBSF was recycled from the bulk polymerization process.

Table 1

Synthesis of polysulfates from diverse building blocks.



Typical reaction conditions: **A** (2.0 mmol) and **B** (2.0 mmol) in NMP (1.0 ml) at 130 °C for 1 h with **Q-5** (0.004 mmol) as the catalyst.

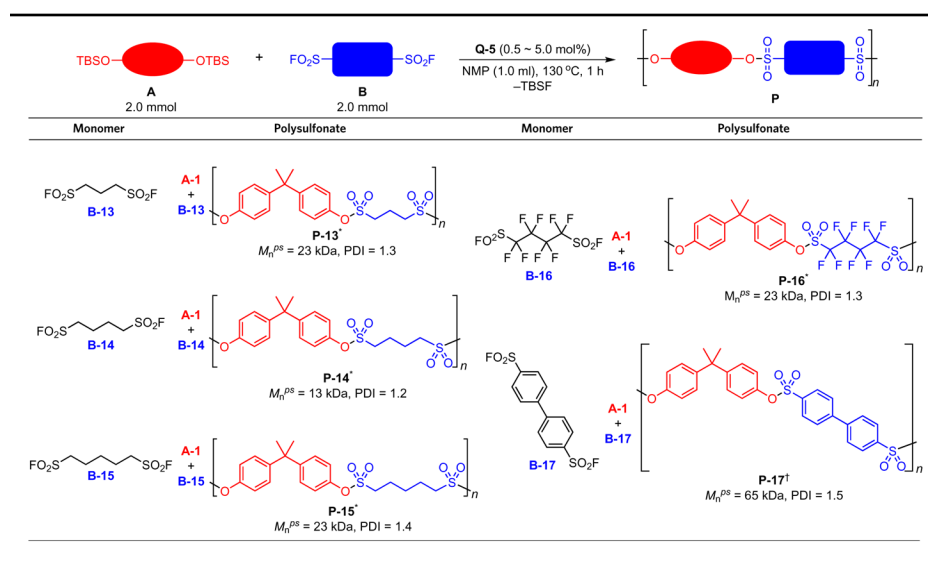
\* 5.0 mol% of **Q-5** was used.

† **P-11a** was treated with BBr<sub>3</sub> (10.0 equiv.) in dichloromethane first, then with SO<sub>2</sub>F<sub>2</sub> (balloon)/Et<sub>3</sub>N in dichloromethane.

NMP, *N*-methyl-2-pyrrolidone; TBS, *tert*-butyldimethylsilyl; *M<sub>n</sub><sup>PS</sup>*, number-average molecular weight with polystyrene as standard; PDI, polydispersity index.

Table 2

Synthesis of polysulfonates from aliphatic and aromatic bis(sulfonyl fluoride).



Typical reaction conditions: **A** (2.0 mmol) and **B** (2.0 mmol) were polymerized in NMP (1.0 ml) at 130 °C for 1 h with **Q-5** as catalyst.

\* 5.0 mol% of **Q-5** was used.

† 0.5 mol% of **Q-5** was used.

NMP, *N*-methyl-2-pyrrolidone; TBS, *tert*-butyldimethylsilyl;  $M_n^{PS}$ , number-average molecular weight with polystyrene as standard PDI, polydispersity index.