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Iodine-enabled organoelectrocatalysis: enantioselective cross dehydrogenative coupling of sulfides and aldehydes†

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A method for the direct asymmetric α-sulfenylation of aldehydes with sulfides was developed. By merging electrochemistry and asymmetric organocatalysis, we obtained α-sulfenylated aldehydes with good to excellent enantioselectivities. Mechanistic investigations indicated a pivotal role of iodine as a catalytic mediator, not only facilitating redox transformations but also possibly contributing to the formation of sulfenyl iodide (RSI) intermediates derived from electrochemically generated disulfides. Our metal-free protocol offers a sustainable and efficient route to access a wide array of α-sulfenylated aldehydes. Remarkably, these transformations take place at room temperature, obviating the need for additional stoichiometric oxidants, thus exemplifying an environmentally friendly and practical synthetic strategy.
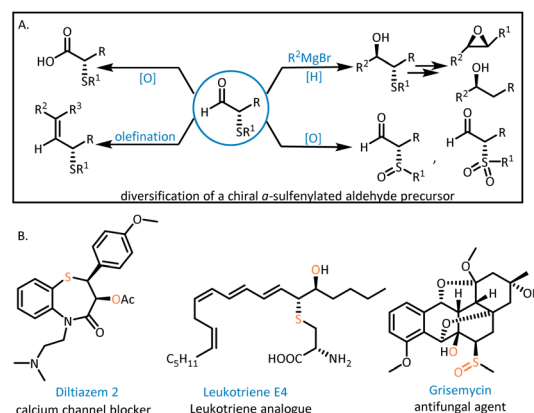
Introduction

Enantiomerically enriched α-sulfenylated aldehydes are key intermediates for preparing important chiral building blocks,[1] such as β-hydroxysulfides existing widely in natural products and pharmaceuticals,[2] chiral secondary alcohols and 1,2-disubstituted epoxides[3] (Fig. 1A). Furthermore, thioethers can be converted to sulfoxides and sulfones, which are prevalent in active pharmaceutical ingredients (APIs) and natural products[4] (Fig. 1B). Consequently, many efficient methods have been devised to construct the α-sulfenylated aldehyde motif.

In the most common approach, enamines react with sulfur-based electrophiles to give chiral α-sulfenylated aldehydes.[5] Most sulfenylation reactions feature reagents with a weak N–S bond as a "S⁺" equivalent. As a consequence, large nitrogen-based leaving groups are formed as by-products thus reducing the reaction's atom economy[6] (Scheme 1a). A complementary approach has been reported by the Jørgensen group utilizing

an oxidative umpolung strategy *via* α-substituted O-bound quinol ethers (Scheme 1b).[7] This method is very effective for α-branched aldehydes but also requires DDQ or other stoichiometric oxidants. The direct dehydrogenative C–H/S–H cross-coupling is the most attractive and environmentally benign approach to construct chiral α-sulfenylated aldehydes[8] (Scheme 1c). Unfortunately, a rapid and effective enantioselective method using unmodified thiols is less developed.

Recently, merging organocatalysis and electrocatalysis has helped to realize dehydrogenative cross-couplings in asymmetric synthesis.[9] With the tunability of the redox potential, traditional oxidants or reductants can be avoided. For instance, Jørgensen demonstrated the possibility of a direct intermolecular α-arylation of aldehydes using electron-rich aromatic compounds wherein a transformation is impossible by Friedel–Crafts reactions.[10] Luo used an asymmetric enamine addition to an anode-generated benzyne intermediate to construct α-arylated and α-cyclohexenylated cyclic β-ketocarbonyls.[11] Mei reported an unusual asymmetric Shono-type oxidation with acyclic amines by means of anodic

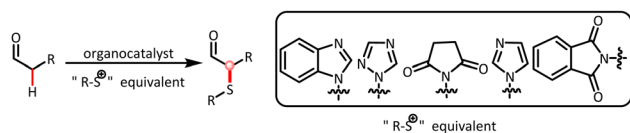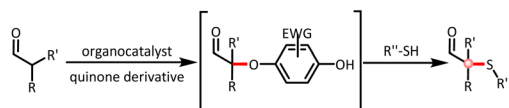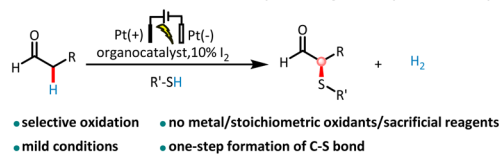*Freie Universität Berlin Institute of Chemistry and Biochemistry, Takustr. 3, 14195 Berlin, Germany. E-mail: mathias.christmann@fu-berlin.de*

Fig. 1 Diversification of chiral α-sulfenylated aldehydes (A) and derivatives of chiral α-sulfenylated aldehydes in natural products and pharmaceuticals (B).

Scheme 1 Methods for the synthesis of enantiomerically enriched α-sulfenylated aldehydes.

oxidation and organocatalysis.[12] In related studies, Meggers,[13] Lin,[14] and Guo[15] have demonstrated the effectiveness of employing transition-metal catalysts towards this goal.

Despite these significant advancements, the direct synthesis of chiral α-sulfenylated aldehydes using thiols remains challenging. First, nucleophilic thiols are not reactive toward nucleophilic enol or enamine intermediates. To address this limitation, it is required to identify a suitable mediator effecting a polarity reversal at sulfur. Second, thiols immediately form disulfides at the anode according to a literature report,[16] and the amino catalyst can possibly react with disulfides.[17] In addition, thiol radicals could be generated in the reaction system. The amino catalyst and enamine intermediate can form radical intermediates in an oxidative environment that may lead to undesired side reactions.[18] Finally, the α-sulfenylated product could be further oxidized to the sulfoxides and sulfones.

Inspired by iodine-catalyzed electrooxidative cross-coupling reactions[19] and our continuing efforts in asymmetric organic synthesis,[20] we envisioned merging organo- and electrocatalysis. With the anode oxidation, we propose that iodine as a mediator assists the electrophilic sulfenylation by converting sulfides/disulfides into an electrophilic sulfur species 7 (Fig. 5). The hydrogen iodide byproduct of the reaction is electrochemically converted into hydrogen with concomitant regeneration of iodine. Furthermore, the redox mediator iodine can decrease the oxidation potential of the reaction, avoiding the overoxidation of the sulfenylated product. Herein, we report our study on the direct enantioselective α-sulfenylation of aldehydes between thiols and aldehydes *via* organoelectrocatalysis with catalytic amount of iodine (Scheme 1c). Compared with traditional multi-step methods,[3,5] the outstanding features of our protocol for the preparation of α-sulfenylated aldehydes include mild conditions, no additional stoichiometric oxidants, and high atom economy, which all together are in line with the 12 principles of green chemistry.[21]

Results and discussion

Reaction optimization

We commenced our studies by using hydrocinnamaldehyde (**1a**) and 4-methyl thiophenol (**2a**) as substrates, I_2 and secondary amine (**3a**) as catalysts, n-Bu_4NPF_6 (2.0 equiv.) as the electrolyte and 300 : 1 (v/v) $MeCN/H_2O$ as the solvent in an undivided cell equipped with two platinum plate electrodes to screen the optimized reaction conditions (Table 1). Several commonly used chiral amine catalysts were tested first. To our delight, the desired product **4a** was obtained in 20% yield with 80% ee value using (2S,5R)-2-(*tert*-butyl)-3,5-dimethyl-imidazolidin-4-one (**3a**) as the catalyst (entry 1). In contrast, L-proline **3b** provided the product as a racemate (entry 2). The primary amine **3e** did not afford any desired product (entry 5). Increasing the constant current to 5 mA afforded the product with high yield and enantioselectivity (entry 6). Further increase of the current lowered the yield and the enantiomeric excess (entry 7). As solvent, 300 : 1 (v/v) $MeCN/H_2O$ was found to give the optimal yields and highest ee values. H_2O was an essential factor for good ee value but further increasing the amount of H_2O had a deleterious effect on the yield (entries 8–10). This phenomenon may be attributed to the fact that an appropriate amount of H_2O facilitates the hydrolysis of

Table 1 Optimization of the reaction conditions[a]

Entry	3	Electric current	Solvent	Yield[b] (%)	ee[c,d] (%)
1	3a	2 mA	300 : 1 MeCN/H₂O	20	81
2	3b	2 mA	300 : 1 MeCN/H₂O	40	0
3	3c	2 mA	300 : 1 MeCN/H₂O	31	0
4	3d	2 mA	300 : 1 MeCN/H₂O	34	44
5	3e	2 mA	300 : 1 MeCN/H₂O	0	—
6	3a	5 mA	300 : 1 MeCN/H₂O	84(80)[c]	85
7	3a	6 mA	300 : 1 MeCN/H₂O	52	74
8	3a	5 mA	MeCN	86	37
9	3a	5 mA	400 : 1 MeCN/H₂O	84	75
10	3a	5 mA	200 : 1 MeCN/H₂O	40	89
11	3a	5 mA	Toluene	0	0
12	3a	5 mA	DMF	Trace	—
13[e]	3a	5 mA	MeCN	n.d.	—
14	—	5 mA	MeCN	n.d.	—
15	3a	—	MeCN	n.d.	—

[a] General conditions: **1a** (0.90 mmol), **2a** (0.30 mmol), **3** (30 mol%), I_2 (10 mol%), n-Bu_4NPF_6 (0.6 mmol), MeCN (3 mL) and H_2O (10 μL) in an undivided cell with two platinum electrodes (each 1.0×1.0 cm²) electrolysis at room temperature for 6 h. [b] The yield of the aldehyde product was determined by ^1H NMR analysis with CH_2Br_2 as an internal standard. [c] The aldehyde product was reduced by $NaBH_4$ to the corresponding alcohol and then isolated yield (two steps and one pot) and ee value was calculated. [d] Enantioselectivities were determined by chiral HPLC analysis. n.d. = not detected. [e] Without iodine.

iminium intermediates while excess H_2O leads to catalyst deactivation.[22]

After extensive efforts in screening reaction parameters, an optimized system employing Pt as the electrodes with 5 mA constant current at room temperature and using **3a** (30 mol%) as organocatalyst, I_2 (10 mol%) as mediator, three equivalents of aldehyde **1a** in a 0.1 M mixture of MeCN and H_2O (300 : 1 (v/v) MeCN/H_2O) containing n-Bu$_4$NPF$_6$ (2.0 equiv., 0.2 M) as electrolyte for 6 h, was able to give **5a** with 80% isolated yield and 85% ee. Control experiments showed that the reaction did not proceed in the absence of I_2, organocatalysts or electric current (entries 13–15). These results imply that these conditions are crucial for a successful transformation.

Substrate scope

With the optimized conditions in hand, we investigated the reaction scope with respect to the thiol substrates by using the hydrocinnamaldehyde as the aldehyde first (Table 2). Thiophenols bearing electron-neutral substituents, such as 4-methyl, 3,4-dimethyl and hydrogen (**2a**, **2e** and **2f**) showed good yields (65–80%) with high to excellent enantioselectivity (86–91% ee). A slight decrease in yield was observed when the steric bulk in *para* position of the thiophenol increased (**2b** and **2c**) with little influence on the enantioselectivity. Notably,

4-hydroxyl thiophenol failed to afford the corresponding product under the standard reaction conditions, but 4-methoxy thiophenol gave the product **5d** with 78% yield and 91% ee. Furthermore, thiophenols bearing electro-withdrawing substituents (**2g**–**2j**) proceeded smoothly to afford the α-sulfenylated products in moderate to good yields (48–84%) and enantioselectivities (61–87% ee). Aliphatic thiols could be employed as substrates (**2k** and **2l**) to furnish the products under this mild condition albeit with lower yields. These substrates are not commonly employed in electrochemistry because of their instability.[23] Interestingly, the corresponding products can be obtained in excellent enantioselectivity (>99% and 95% ee). Moreover, the substrate scope can be extended to the hetaryl thiol. 2-Thiophenothiol (**2m**) could react smoothly to furnish the desired product in 76% yield with 89% ee.

Next, we explored the asymmetric α-sulfenylation reaction with different aldehydes (Table 2). Different substitutions on the phenyl ring of phenylpropanals are explored firstly. Electron-donating group (**1n**, –OMe) and halogen (**1o** and **1p**) lead to a slight decrease on the yield. (35–45%) and ee value (31–76%). The reaction was also found to be amenable to aldehyde with different chain lengths, which gave the desired products in good yields and enantioselectivities. For example, two unbranched aldehydes afforded the corresponding α-sulfenylated products **5q** and **5r** in 67% and 53% yields with 71% and 84% ee, respectively. Moreover, cyclohexyl- (**1s**–**t**) and phenyl-substituted aldehydes (**1u**) can also react smoothly, affording products **5s**–**5u** in 64–85% yields and 71–85% ee. The introduction of alkene, alkyne groups were also well tolerated under the standard conditions, in 65% and 60% yields with moderate enantioselectivity (53% and 29% ee, respectively). Nitrogen heterocycles are reactive substrates often used in electrochemistry. They are prone to Shono-type oxidations followed by nucleophilic attack. Gratifyingly, we observed that the *N*-Boc-protected piperidine aldehyde could be used as a substrate to afford **4x** in moderate yield and 71% ee. Finally, a derivative of **5s** (Fig. 2) was synthesized and its absolute configuration was determined by a single-crystal X-ray diffraction study of **6a**. The absolute configurations of other products were determined by depicted in analogy.

To test the scalability of this protocol, a synthesis of **5a** on a 3.0 mmol-scale was carried out, smoothly affording the desired product **5a** in 65% yield with 80% ee. Furthermore, considering the practicability, we conducted the reaction on gram-scale (9.0 mmol) using graphite electrodes instead of platinum electrodes, which afforded 2.16 g of **5a** (93% yield and 57% ee) (see the ESI† for detailed information). We further carried out

Table 2 Evaluation of substrate scope[a]

[a] Reactions were performed on a 0.30 mmol scale under the standard conditions. Unless noted otherwise, the α-sulfenylated aldehyde products were reduced to the corresponding alcohol by NaBH$_4$ and shown are their total isolated yields for the two steps. The ee values were determined by chiral HPLC analysis. [b] The yields and ee values were determined by the aldehyde form.
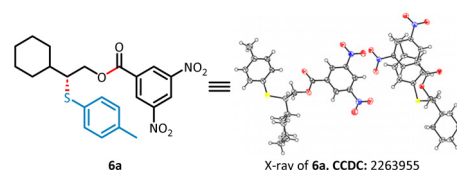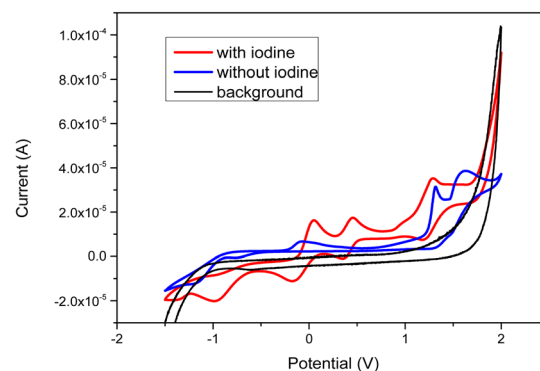

Fig. 2 Absolute configuration of **6a**.

simple nucleophilic addition to directly obtain β-hydroxysulfide **6b** with 69% yield and 94 : 6 dr value (relative configuration *tentatively* was assigned in analogy to reference).[24] Olefination with triethylphosphonoacetate afforded (*E*)-ester **6c** with minor racemization (65%, 83% ee) (Fig. 3).
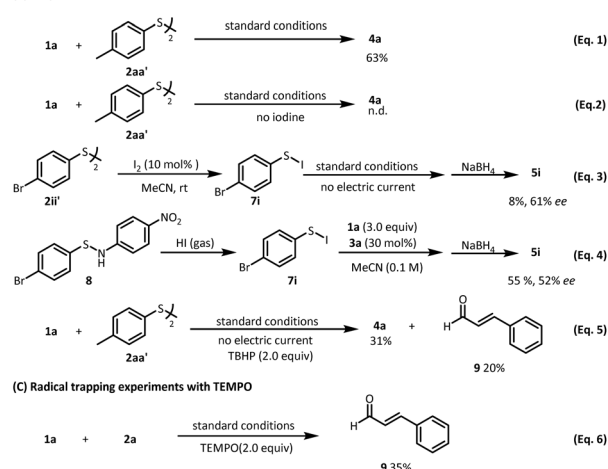
Mechanistic studies

Next, a series of control experiments were conducted to support the proposed reaction pathway depicted in Fig. 5. First, a cyclic voltammetry experiment shows a significant decrease in the highest oxidation potential of the reaction mixture with the addition of I_2 to the reaction mixture (Fig. 4A, from 1.44 V to 1.09 V *vs.* Fc$^+$/Fc), which implies that I_2 can decrease the reaction oxidation potential *via* accelerating the electron transfer at the electrode/electrolyte interface. Furthermore, an oxidation peak of product **4a** in acetonitrile was observed at 1.31 V *vs.* Fc$^+$/Fc, where other substrates are preferred to be oxidized under the current of 5 mA (see the ESI, Fig. S2 and S4†), explaining why the sulfenylated product will not suffer overoxidation to the sulfoxide or sulfone directly under the standard conditions. In addition, according to our monitoring of the reaction process with ^{1}H NMR, the anodic oxidation will quickly oxidize the thiol in 1.5 h to the corresponding disulfide (see ESI Fig. S6† for the details), which is accordance with literature reports.[16] Hence, we replaced the thiophenol with disulfide and subjected it to the standard reaction conditions. As expected, the sulfenylated product **4a** was formed though with a slightly decreased yield of 63% (eqn (1)). Moreover, to rule out the attack of enamine intermediate to disulfide straightly, the reaction was carried out without iodine in the presence of disulfide under standard conditions. As expected, no desired product formed (eqn (2)). In addition, according to our hypothesis, the sulfenylated iodide 7 is a key intermediate. The proposed reactive intermediate was generated *in situ* by reaction of disulfide of **2ii′** and I_2. After they were stirred for 10 minutes, aldehyde and catalyst **3a** were then added to the above mixture. The reaction was stirred overnight in the absence of electric current. After the reduction workup with NaBH$_4$, we could separate the corresponding alcohol product in 8% yield while the expected yield is 10% with 0.1 equiv. I_2 participating the reaction (eqn (3)). To further validate the presence of the key intermediate in our reaction, we syn-

Fig. 4 (A) Cyclic voltammograms (CV): using Pt wire as anode and cathode, ferrocene as the internal reference electrode, 0.1 M *n*-Bu₄NPF₆ with scanning rate 100 mV s^{-1}. All the potential values were adjusted relative to Fc/Fc$^+$. Yields of **4a** were determined by ^{1}H NMR analysis with CH₂Br₂ as an internal standard. The isolated yields of **5i** and **9** were given and the enantioselectivities were determined by chiral HPLC analysis. (B) Subjected intermediates to standard reaction conditions. (C) Radical trapping experiments with TEMPO (2,2,6,6-tetramethyl-1-piperidinyloxy).

thesized intermediate **7i** using a direct approach (see ESI† for the details). Subsequently, we exposed the intermediate to a mixture of aldehyde and catalyst, and the corresponding alcohol product could be finally separated with 55% yield and 52% ee (eqn (4)). In addition, we replaced the electrochemical oxidation by using TBHP (*tert*-butyl hydroperoxide) as the oxidant with other conditions unchanged. The sulfenylated product was formed in 31% yield and α,β-unsaturated product **9** was also isolated with 20% yield, which was probably formed by the elimination of the corresponding sulfoxide product (eqn (5)). Furthermore, when TEMPO was used as the radical trapping reagent using thiophenol as the substrate under the standard conditions, the product was completely oxidized to then furnish the α,β-unsaturated product **9** with 35% yield finally (eqn (6)).

Based on these above experiments and previous literature,[25] we proposed a novel reaction pathway (Fig. 5). Initially, con-
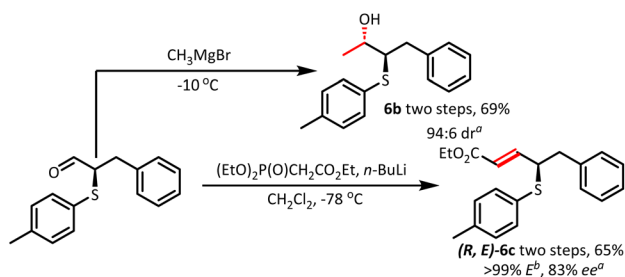
Fig. 3 Derivation of α-sulfenylated aldehydes. ^{a} Determined by chiral HPLC analysis. ^{b} The ratio of *E/Z* was determined by ^{1}H NMR spectroscopy.
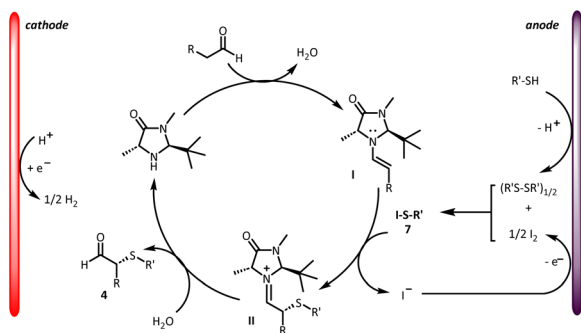
Fig. 5 Proposed mechanism for the iodine-mediated α-sulfenylation of aldehydes.

densation of aldehyde with amine catalyst **3a** produces enamine intermediate **I**. Concomitantly, the thiol is converted to the corresponding disulfide *via* anodic oxidation. Reaction of I$_2$ with the disulfide is then proposed to generate sulfenyl iodide (R'S-I) **7** as the reactive electrophile. The iminium intermediate **II** is then produced by the nucleophilic attack of **I** to **7**. Iminium hydrolysis affords the α-sulfenylated adduct **4** with regeneration of organocatalyst **3a**. The iodide anion which will be oxidized to iodine under the anodic oxidation conditions.

Conclusions

In conclusion, we have developed a direct and highly enantio-selective method to construct the α-sulfenylated aldehydes from aldehydes and thiophenols by merging anodic oxidation and organocatalysis. This mild and metal-free protocol operates simply in one step and does not require additional stoichiometric oxidants. Mechanistic investigations indicate that the presence of catalytic amounts of iodine enables the formation of the key RSI intermediate *in situ* and take effect as a redox mediator to avoid the sulfenylated product to be overoxidized. This atom-economic method is currently being extended in our laboratory to other asymmetric α-functionalization reactions.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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