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Herein, we report on an artificial nickel chlorinase (ANCCase) resulting from anchoring a biotinylated nickel-based cofactor within streptavidin (Sav). The resulting ANCCase was efficient for the chlorination of diverse C(sp³)–H bonds. Guided by the X-ray analysis of the ANCCase, the activity of the artificial chlorinase could be significantly improved. This approach opens interesting perspectives for late-stage functionalization of organic intermediates as it complements biocatalytic chlorination strategies.

Halogen-containing motifs are ubiquitously present in natural products due to the bioavailability of halides in the environment.¹ The incorporation of a halogen atom can significantly modulate the properties of a molecule, potentially influencing its bioactivity, metabolism, and pharmacokinetic profile.² Accordingly, halogenated compounds play an important role in both the pharmaceuticals³ and agrochemicals sectors,⁴ Scheme 1a. Moreover, halogenated compounds can serve as versatile substrates in a myriad of coupling reactions, thus facilitating the synthesis of valuable advanced synthetic intermediates.⁵ Noteworthy, chlorinated natural products dominate the halogenated compound landscape with over 1000 examples.⁶ Therefore, developing efficient chlorination methodologies has attracted increasing attention in recent decades.

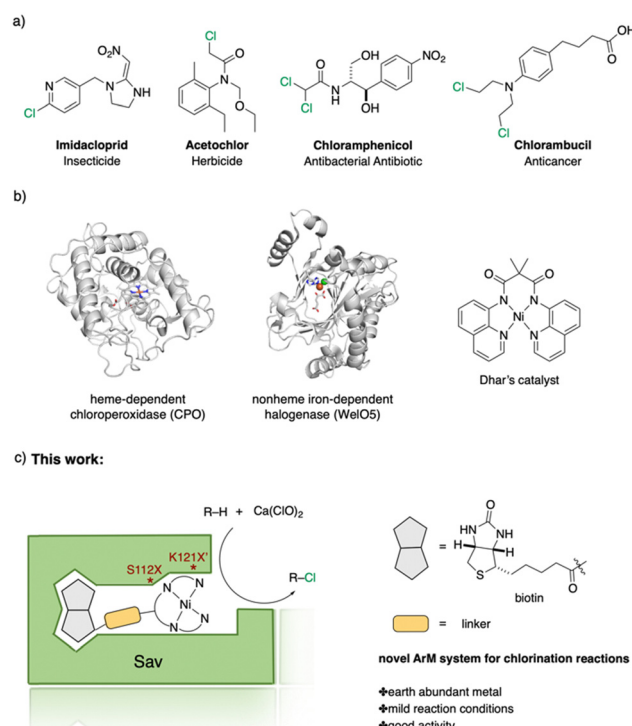
Given the ubiquity of chlorinated natural products in the environment, the initial research endeavors focused on elucidating the biosynthetic pathway underpinning these molecules. To date, four distinct classes of halogenases have been identified:⁷ heme-/vanadium-dependent haloperoxidases, flavin-dependent halogenases, nonheme iron-dependent halogenases and *S*-adenosyl-*L*-methionine (SAM)-dependent halogenases. Among them, the α -KG-dependent nonheme iron halogenase, heme-dependent

An artificial nickel chlorinase based on the biotin–streptavidin technology†

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haloperoxidases, and non-heme halogenases SyrB2 offer versatile means to chlorinate inert C(sp³)–H bonds, Scheme 1b.

Recently, several groups reported their achievements in utilizing repurposed halogenases.⁸ Benefiting from the development of directed evolution technologies, the substrate scope of chlorination has been significantly expanded. Drawing inspiration from natural chlorinases, chemists have developed several metal complexes that catalyze chlorination reactions. To



Scheme 1 Selected examples of chlorine-containing compounds and recently-reported Ni-catalyst for chlorination reactions. (a) Selected examples of chlorine-containing agrochemicals and pharmaceuticals. (b) Selected examples of enzymes and homogeneous catalyst for chlorination. (c) Artificial Nickel Chlorinase (ANCCase) catalyzed chlorination of alkanes.

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cite a few, Mn-,⁹ Fe-¹⁰ and Ni-based¹¹ complexes have been reported to be efficient catalysts. Importantly, however, competition between the formation of chlorinated products and oxygenated side-products persists. Recently, the Dhar group reported on Ni-based complexes relying on tetradentate amido-quinoline as a ligand, which exhibited promising chlorination activity, Scheme 1b.

Artificial metalloenzymes (ArMs) result from incorporating a metallocofactor within a host protein. Over the years, ArMs have been shown to combine the advantages of both homogeneous and enzymatic catalysts and endow the protein with new-to-nature reactivity. Since the foundational study by Wilson and Whitesides in the late 1970s,¹² many research groups have contributed to the development of ArMs displaying diverse catalytic properties.¹³ Building upon the biotin-Sav technology, our team and others¹⁴ have developed various ArMs. The pronounced affinity of biotin for Sav ensures the tight binding of a biotinylated metallocofactor to the Sav scaffold. Depending on the biotinylated metallocofactor, the assembled ArMs exhibit a diverse range of activities. Herein, we report on an artificial nickel chlorinase (ANCCase) for the C(sp³)-H chlorination of alkanes, taking advantage of the biotin-Sav technology, Scheme 1c.

Inspired by the chlorination homogeneous catalyst recently reported by Dhar and coworkers, we set out to synthesize a related biotinylated Ni-complex.

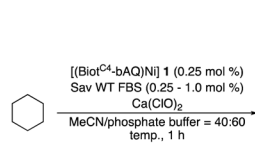
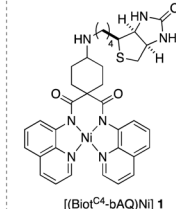
To maintain the C₂-symmetry of the metallocofactor, we set out to attach the biotin anchor to the bridging carbon atom flanked by the two amido-quinoline moieties. Additionally, we converted the original dimethyl group to a cyclohexyl entity to minimize the influence on the dihedral angle between the two amido-quinoline planes. Accordingly, the cofactor [(Biot^{C4}-baQ)Ni] **1** was synthesized, see, Scheme S1 (ESI[†]).

With the cofactor **1** at hand, we set out to confirm its incorporation within Sav. For this purpose, we assessed the quantitative anchoring of the [(Biot^{C4}-baQ)Ni] **1** into Sav WT by a HABA displacement titration using CD spectroscopy as readout.¹⁵ The determined dissociation constant was $K_d = 0.29 \pm 0.05 \mu\text{M}$ (see ESI[†]). Next, we evaluated its activity for the chlorination of cyclohexane. In the absence of Sav, [(Biot^{C4}-baQ)Ni] **1** yielded 21 turnovers (TON), Table 1, entry 1.

Upon incorporation into Sav WT, [(Biot^{C4}-baQ)Ni] **1**-Sav WT hereafter, the resulting ArM afforded significantly higher TON (TON = 54), underscoring the positive influence of the second coordination sphere provided by Sav on chlorinase activity. We then optimized the reaction conditions. Sav is a homotetrameric protein with four free biotin-binding sites (four FBS) consisting of two pairs of close-lying bis-biotin binding sites. In select cases, the occupancy of the adjacent monomers has been shown to affect the catalytic performance of the ArM.¹⁶

Based on experimental evaluations, we determined the optimal ratio to be [(Biot^{C4}-baQ)Ni] **1**/Sav FBS = 1:2, Table 1, entries 2–4. Additionally, both temperature and pH substantially influenced TON, Table 1, entries 5–9. The highest TON was obtained at 25 °C and pH = 9. Importantly, an alkaline reaction environment proved essential to minimize the

Table 1 Optimization of reaction conditions for ANCCase-catalyzed chlorination of cyclohexane^a

| Entry | [Ni]/Sav FBS | Temperature (°C) | pH | Ca(ClO) ₂ (equiv.) | TON |
|-------|--------------|------------------|----|-------------------------------|---------|
| 1 | Without Sav | 25 | 9 | 10 | 21 ± 2 |
| 2 | 1:2 | 25 | 9 | 10 | 54 ± 11 |
| 3 | 1:1 | 25 | 9 | 10 | 52 ± 8 |
| 4 | 1:4 | 25 | 9 | 10 | 43 ± 7 |
| 5 | 1:2 | 15 | 9 | 10 | 38 ± 1 |
| 6 | 1:2 | 35 | 9 | 10 | 31 ± 7 |
| 7 | 1:2 | 25 | 6 | 10 | 15 ± 3 |
| 8 | 1:2 | 25 | 8 | 10 | 39 ± 5 |
| 9 | 1:2 | 25 | 10 | 10 | 34 ± 2 |
| 10 | 1:2 | 25 | 9 | 5 | 32 ± 4 |
| 11 | 1:2 | 25 | 9 | 20 | 40 ± 7 |

^a Reaction conditions: [cyclohexane] = 20 mM, [(Biot^{C4}-baQ)Ni] **1** = 50 μM, MeCN/phosphate buffer (100 mM) = 200/300 μL, 1 h. Sav WT FBS = streptavidin wild-type free binding sites. Varying amounts of dichlorination products were detected in all cases.

demetallation of the cofactor. The concentration of the hypochlorite markedly influenced TON, Table 1, entry 10 and 11. Striking the right balance in Ca(ClO)₂ concentration was instrumental in ensuring reaction efficiency. Importantly, across most experiments, only trace amounts of oxygenated side-products were detected, along with dichlorination products. Following the chlorination reaction, we observed the precipitation of

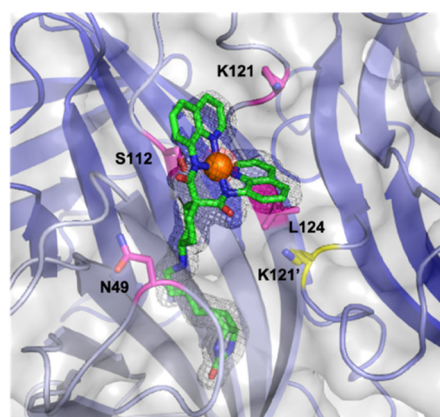


Fig. 1 Close-up view of the structure of the [(Biot^{C4}-baQ)Ni] **1**-Sav WT (PDB: 8QQ3). The cofactor [(Biot^{C4}-baQ)Ni] **1** is displayed as sticks (atoms are color-coded; nitrogen = blue, oxygen = red, carbon = green or cyan, chloride = green and sulfur = yellow) with the nickel as an orange sphere. The protein is represented as a cartoon and transparent surface model. The monomers are color-coded in different shades of blue. The residues S112 and K121 are displayed as magenta sticks (atoms are color-coded; nitrogen = blue, oxygen = red and carbon = magenta and yellow for the adjacent monomer). The 2F_o-F_c difference map is displayed as a dark grey mesh (1σ) and the anomalous electron density is displayed as red mesh (5σ). The occupancy of the nickel was set to 69%.



[(Biot^{C4}-bAQ)Ni] 1-Sav K121Y. Notably, Sav S112F-K121Y emerged as the most active, affording TON = 92. The double mutant ANCase [(Biot^{C4}-bAQ)Ni] 1-Sav S112P-K121Y: TON = 87.

We then attempted to determine Michaelis parameters for [(Biot^{C4}-bAQ)Ni] 1-Sav WT and [(Biot^{C4}-bAQ)Ni] 1-Sav S112Y. However, all attempts did not reveal a saturation kinetic profile. This suggests that the radical-chlorination mechanism does not proceed *via* the formation of a Michaelis-Menten complex, followed by the rate-determining step. This is consistent with the mechanism suggested by Dhar and coworkers for the related homogeneous catalyst.^{11e}

Finally, the substrate scope was explored using a focused library of Sav mutants combined with cofactor [(Biot^{C4}-bAQ)Ni] 1, Table 2. The ANCases were efficient for the chlorination of diverse C(sp³)-H bonds. Key findings from this screening included: (i) for toluene, apart from the expected chlorination products, additional derivatives arising from chlorination of aromatic C(sp²)-H bonds were identified. (ii) in the case of ethylbenzene, the dominant product was acetophenone, and a relatively low yield of chlorinated product, *i.e.* 2-(chloroethyl)-benzene, was observed. No difference in TON and product distribution was observed for the chlorination of ethylbenzene using [(Biot^{C4}-bAQ)Ni] 1-Sav S112F-K121Y upon carrying out the reaction under anaerobic conditions. We hypothesize that the reason for the high yield of oxygenated product is the low oxidation potential of the secondary benzyl radical.¹⁸ Additionally, no enantioselectivity was achieved in this reaction.

In conclusion, we have engineered an ANCase by incorporating the [(Biot^{C4}-bAQ)Ni] 1 within a Sav scaffold, thereby catalyzing the chlorination of C(sp³)-H bonds. Genetic optimization led to the identification of a double mutant ANCase [(Biot^{C4}-bAQ)Ni] 1-Sav S112F-K121Y which was most active for the chlorination of various C(sp³)-H bonds. Current efforts focus on achieving regioselective chlorination, which would provide valuable building blocks for organic synthesis.

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Conflicts of interest

There are no conflicts of interest to declare.

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