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Bio-click chemistry: a bridge between biocatalysis and click chemistry

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The fields of click chemistry and biocatalysis have rapidly grown over the last two decades. The development of robust and active biocatalysts and the widespread use of straightforward click reactions led to significant interactions between these two fields. Therefore the name bio-click chemistry seems to be an accurate definition of chemoenzymatic reactions cooperating with click transformations. Bio-click chemistry can be understood as the approach towards molecules of high-value using a green and sustainable approach by exploiting the potential of biocatalytic enzyme activity combined with the reliable nature of click reactions. This review summarizes the principal bio-click chemistry reactions reported over the last two decades, with a special emphasis on small molecules. Contributions to the field of bio-click chemistry are manifold, but the synthesis of chiral molecules with applications in medicinal chemistry and sustainable syntheses will be especially highlighted.

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1. Introduction

Chemoenzymatic syntheses are powerful methodologies for the creation of complex molecular structures and functionalities.¹ The development of processes which combine biocatalysis and chemical synthesis has rapidly increased over recent years. The two main pillars of innovation in the field are the generation of more active, selective, and stable biocatalysts,² and the development of new synthetic procedures in water, allowing for highly efficient, and sustainable one-pot procedures.³

The term click chemistry was introduced by Sharpless in 2001, comprising highly efficient and reliable reactions enabling the rapid construction of structural and functional diversity through the union of small building blocks. Commonly these reactions have very high inherent driving

forces, thus avoiding tedious chromatographic work-up.⁴ Evidently, the concept of click chemistry is largely inspired by nature's "synthetic toolbox", and it is not surprising that the merger of click chemistry and biocatalysis exhibits great potential for the synthesis of complex molecular structures.

The increasing demand for sustainable processes is fostering the development and establishment of new synthetic tools. Among these tools, click chemistry and biocatalysis are of utmost importance, since they are characterized by their high selectivity and orthogonality, mild reaction conditions avoiding protecting groups, efficiency, reliability, and user-friendliness.⁵ The growing interest in click chemistry and biocatalysis in recent years is reflected by the drastic increase in the number of publications on the topic (Fig. 1). It is not surprising that the combination of these two disciplines, which we propose to call



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Dr Kamal Dua, Lecturer, Discipline of Pharmacy, Graduate School of Health, University of Technology Sydney, has a research experience of more than 12 years working in the field of drug delivery targeting inflammatory diseases. Dr Dua is also a Node Leader of Drug Delivery Research in the Centre for Inflammation at Centenary Institute/UTS, where the targets identified from the research

projects are pursued developing novel formulations as the first step towards translation into clinics. Dr Dua research in two complementary areas; drug delivery and immunology, specifically addressing how these disciplines can advance one another helping community to live longer and healthier.



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and he is set to transfer the technologies developed to pharma and nutraceutical industry. In 2020, he received Dr B.C. Deb memorial award by Indian Science Congress Association and the Bharat Shiksha Ratan Award given by Global Society for Health and Educational Growth, India.



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bio-click chemistry, has given rise to a myriad of highly interesting developments and has led to a significant impact on the synthesis of structurally diverse molecules, such as chiral amino alcohols, triazoles, amides, cycloadducts, thioethers, and others. In addition, bio-click methodologies show a synergy oftentimes allowing for shorter, stereoselective and efficient synthetic routes.

The present review focuses on syntheses in which biocatalytic processes and click reactions were used in combination. Recent reviews have focused on specific applications of click chemistry and biocatalysis in the *in situ* generation of inhibitors,^{6,7} the immobilization and modification of enzymes,⁸ or the generation of semisynthetic enzymes.⁹ However, the discrete topic of bio-click chemistry has yet to be approached in depth. We seek to fill this gap and discuss the most important aspects of the development of bio-click methodologies, as well as their main advantages and limitations.

2. Bio-click chemistry

Organic chemistry has a sheer endless set of reactions allowing for the generation of molecular diversity and complexity. Chlorophyll,¹⁰ taxol,^{11,12} and human lysozyme¹³ are only a few examples of highly important complex molecules obtained through elegant chemical syntheses. Albeit highly sophisticated, organic synthesis is undergoing constant improvements¹⁴ especially through the development of more selective, efficient, and sustainable reactions. The interconnection of different disciplines, such as biocatalysis and click chemistry, is essential to achieve these goals (Fig. 2).

Click chemistry is generally outlined by a group of reactions with desirable characteristics for biocompatible reactions or bioconjugations: reliability, effectiveness, protecting group-free, high yielding, and easy to purify (often without any chromatography). The fact that click chemistry has become an integral part of the synthetic toolbox is demonstrated by its success in chemical synthesis, drug discovery, medicinal chemistry,¹⁵ molecular labelling,¹⁶ polymer synthesis,¹⁷ and others.

The cutting-edge example for click chemistry is (i) the copper-catalyzed 1,3-dipolar cycloaddition of terminal alkynes

and organic azides (CuAAC). Nevertheless, many other transformations are classified as click reactions as well (ii) the Diels–Alder reaction, (iii) additions to alkenes, (iv) nucleophilic opening of strained rings, (v) non-aldol carbonyl transformations, and (vi) addition reactions to alkynes.¹⁸ It has to be clarified, that although click reactions are regio- and stereo-specific, they are not necessarily enantioselective.⁴ This shortcoming can be overridden by the merger of click chemistry with enantioselective biocatalysts such as enzymes.

Biocatalysis comprises the use of purified enzymes, cell-free extracts or whole microorganisms as catalysts in a vast array of transformations, rendering them as excellent and sustainable methods for the construction of complex molecular structures, while fulfilling green chemistry principles.¹⁹ However, enzymes have inherent limitations often preventing their widespread use in synthesis: rather low operational stability, inhibition by substrates or products, limited tolerance of non-natural substrates and limited applicability in organic solvents. Innovative technologies, such as protein engineering and enzyme immobilization, have emerged to overcome these drawbacks and limitations.² Many enzymes obtained by protein engineering are more robust, stable and capable to transform non-natural substrates. Immobilized enzymes generally present outstanding operational stability, an easy removal from the reaction medium and the capability of repetitive recycling.

Enzymatic processes allowing for the synthesis of complex molecules under highly sustainable and green conditions are increasingly popular. Various drug molecules commonly synthesized through traditional catalytic processes are eventually produced by chemoenzymatic methods, thus improving process efficiencies and substantially reducing waste generation.^{20,21}

A state-of-the-art example is the biocatalytic synthesis of the HIV treatment candidate islatravir, as reported by Huffman and co-workers.²² In this process, five enzymes obtained by directed evolution (two immobilized enzymes) and four auxiliary enzymes were applied towards the generation of islatravir from simple achiral building blocks in a three-step cascade reaction (Fig. 3). The target molecule was obtained in 51% overall yield, with high atom economy, using water as solvent, avoiding intermediate purification and in less than half of the number of steps compared to the conventional chemical synthesis.²²

Although, Caruso *et al.* used the term bio-click chemistry in a study towards the functionalization of polymer drug vehicles,²³ a more exhaustive review of the interface between click chemistry and biocatalysis is necessary. Considering the potential for the development of highly efficient, sustainable, and selective processes, a profound impact of bio-click methods on the field can be envisaged.

It should be noted that the term “bio-click chemistry” differs markedly from click chemistry, as it interconnects the latter with biocatalytic processes. In many cases, this allows for the development of greener processes. However, it must be clarified that the examples covered in this review are based not only on sustainability or higher yields, but rather cover a broad spectrum of cases allowing us to show the current advantages and limitations of bio-click chemistry.



Fig. 1 The number of results to the search for click chemistry (blue) and biocatalysis (green) in Scopus per year between 2000–2020.



Bio-Click Chemistry



Fig. 2 (a) General click chemistry and biocatalysis reactions, (b) bio-click chemistry interface and (c) schematic representation of bio-click chemistry cooperation for the synthesis of small molecules.

3. Bio-click reactions

Click chemistry is a dynamic and growing field, continually leading to new transformations. Many of these developments are predominantly focused on bioconjugation and applications in materials science.²⁴ Examples for new transformations characterised as click reactions are the sulfur(vi) fluoride exchange (SuFex)²⁵ and the synthesis of azides from primary amines²⁶ respectively. Nevertheless, interactions of the latter with biocatalysis for the creation of new molecules are still missing. Significant cooperation between click chemistry and biocatalysis has been reported for CuAAC-, Diels-Alder-, epoxide-opening- and thiol-Michael reactions, and these studies will be discussed in further detail.

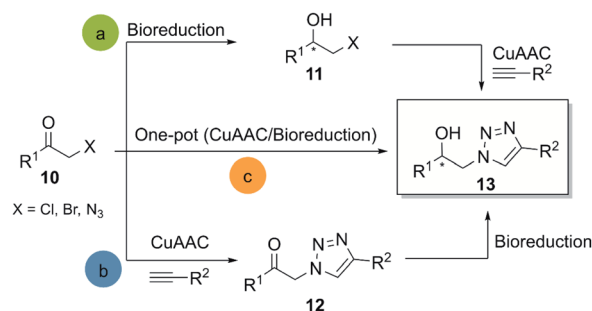
3.1 CuAAC and biocatalysis

CuAAC is probably the most representative click reaction. It allows for the synthesis of 1,2,3-triazoles from organic azides and alkynes in the presence of a copper(I)-catalyst (Scheme 1). The 1,2,3-triazole ring has shown great value as a pharmacophore, and several reviews focused on the potential of this invaluable motif in medicinal chemistry.^{15,27} Regarding the bio-click approach, the main focus in literature is the combination of CuAAC with oxidoreductases and hydrolases respectively.

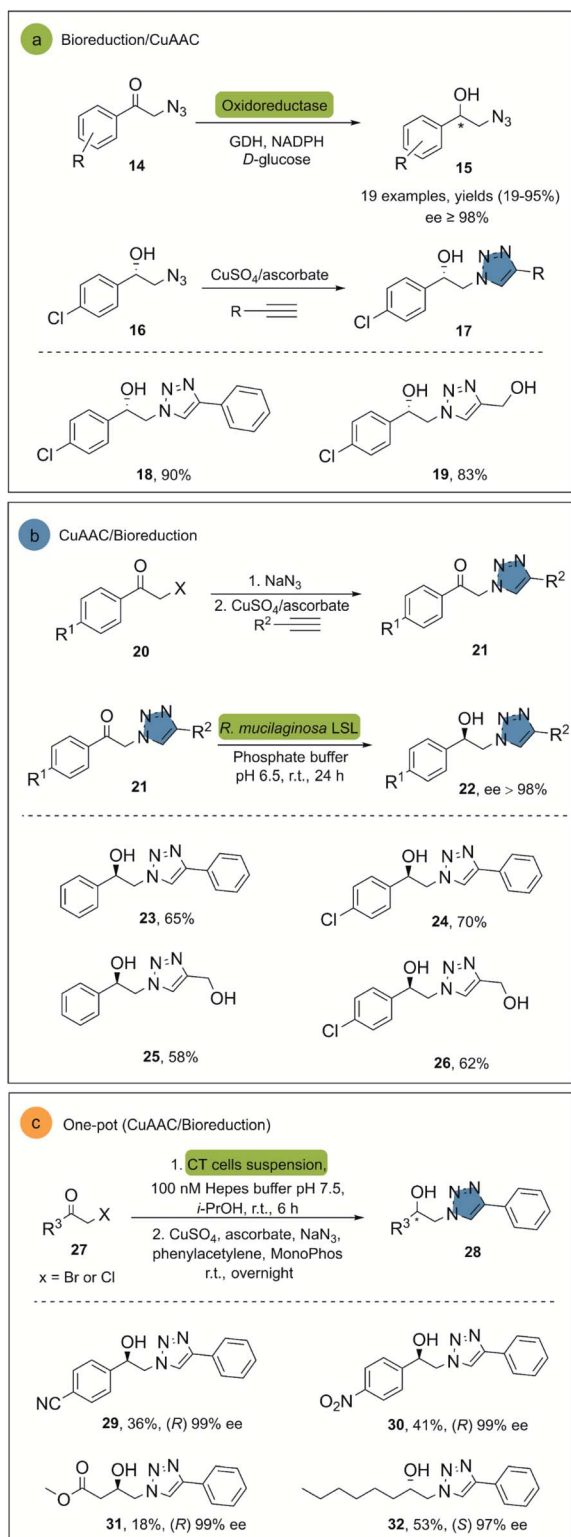
3.1.1 CuAAC and oxidoreductases. The most common application of click reactions combined with oxidoreductases is for the synthesis of highly enantiopure β -hydroxytriazoles **13**. These compounds present excellent pharmacological

properties, as they are able to act as potential HIV protease inhibitors²⁸ and β -adrenergic receptor blockers.²⁹ Enantiopure β -hydroxytriazoles can be generally obtained by three distinct approaches: (a) bioreduction of α -chloro/bromo-acetophenones or α -azidoacetophenones with subsequent CuAAC reaction, (b) the generation of β -ketotriazoles and subsequent bioreduction, (c) a simultaneous one-pot bioreduction and triazole formation (Scheme 1).

In 2008, Hua and co-workers reported the use of a recombinant carbonyl reductase from *Candida magnoliae* (CMCR) and an alcohol dehydrogenase from *Saccharomyces cerevisiae* (Ymr226c), with glucose dehydrogenase as cofactor-regeneration enzyme, for the enzymatic reduction of α -azidoacetophenone.²⁹ Both enzymes were able to catalyze the formation of the



Scheme 1 General methodologies for the chemo-biocatalytic synthesis of β -hydroxytriazoles.



Scheme 2 (a) Synthesis of β -blocker analogues containing 1,2,3-triazoles moieties, (b) stereoselective reduction of β -ketotriazoles with *R. mucilaginosa* LSL, (c) one-pot whole-cell catalysis and CuAAC for the conversion of α -haloketones to β -hydroxytriazoles.

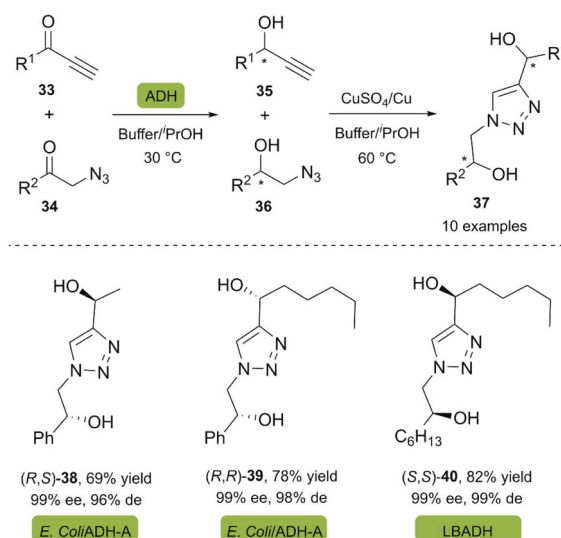
corresponding chiral alcohol in excellent optical purities, good yields and with wide substrate scope. After the successful

bioreduction of 2-azidoacetophenones **14**, the desired 1,2,3-triazoles **17** were obtained in high yields, using CuSO_4 and sodium ascorbate as catalytic system (Scheme 2a).

Another interesting example is the direct synthesis of enantioenriched β -hydroxytriazoles from the corresponding haloketones and β -ketotriazoles.³⁰ In this approach, a whole-cell procedure of wild type *Rhodotorula mucilaginosa* LSL successfully catalyzed the one-pot enantioselective bioreduction of the substrate in water at room temperature, although only four examples were reported under these conditions (Scheme 2b). Notably, the reactants necessary for the ketotriazole formation, such as sodium azide and copper sulfate, did not affect the *R. mucilaginosa* LSL system in its catalytic activity. This represents an important advantage, avoiding purification steps and the manipulation of organic azides, often exhibiting explosive properties.³¹

Janssen and co-workers described an alternative system for the synthesis of β -hydroxytriazoles using a one-pot methodology.³² The developed system consists of the *E. coli* strain MC1061, which overexpresses enzymes such as alcohol dehydrogenases (ADH) and halohydrin dehalogenases (HHDH) with different stereospecificities. The construct that overexpressed (*R*)-stereospecific (ADH) and (HHDH) was called CT-cells and produce (*R*)-enantiomers, while the BT-cells form the (*S*)-enantiomers in high enantiomeric purity, albeit only moderate yields. The catalyst for the click reaction was CuSO_4 with sodium ascorbate in the presence of MonoPhos as ligand, increasing the CuAAC reaction rate³³ (Scheme 2c).

Chiral diols are a group of compounds widely used as organocatalysts and intermediates for the preparation of chiral heterocycles.^{34,35} Gotor and co-workers reported the synthesis of dihydroxytriazoles **37** through a highly efficient chemoenzymatic procedure under mild reaction conditions.³⁶ In this two-stage one-pot procedure, the recombinant enzyme ADH-A from *Rhodococcus ruber* overexpressed in *E. coli* (Prelog-



Scheme 3 Synthesis of dihydroxytriazoles from chemoenzymatic approach.



enzyme) and a commercially available LBADH from *Lactobacillus brevis* (anti-Prelog-enzyme) were used for the bioreduction of several alkynones **33** and α -azido ketones **34** in a phosphate buffer with 2-propanol. A copper wire was used as a reusable catalyst for the click reaction, supplemented by a catalytic amount of copper sulfate (Scheme 3).

A powerful methodology for asymmetric ketone reduction was reported by Omori and co-workers in 2013, using whole cells from plant tissue from carrots (*Daucus carota*). The main advantage of this approach is its simplicity, economy of the process and the easy availability. The reaction features the one-pot bioreduction of azidoacetophenones (*p,m*-substituted) **41** with the subsequent CuAAC reaction using the Sharpless–Fokin catalyst (Scheme 4).³⁷ The products were obtained in moderate yields but excellent enantiomeric excess (>99%). However, a disadvantage of this process is the rate of the bioreduction, requiring 5–7 days for completion. This impedes the implementation of a single-stage one-pot procedure, since the click reaction occurs very fast, and non-reduced triazoles are insoluble in water.

Dioxygenases are versatile biocatalysts that have been used in the chemoenzymatic synthesis of chiral hydroxyazides. In 2011, the Stefani group reported the enzymatic dihydroxylation of arenes **45** catalyzed by whole-cell *P. putida* F39/D en route to chiral azides **47**. The subsequent click reaction between **47** and several alkynes was achieved with the Sharpless–Fokin catalyst in *t*-BuOH : H₂O or toluene : H₂O,³⁸ and the corresponding triazolyconduritols **48** were obtained in good to excellent yields (Scheme 5).

3.1.2 CuAAC and hydrolases. Over the last decade, chemoenzymatic methods based on the combination of hydrolases and the click CuAAC reaction have been successfully implemented for peptide-triazole synthesis **55** and the kinetic resolution (KR) of racemic secondary alcohols **56** (Scheme 6).

Enzymatic aminolysis is an excellent environmentally-friendly alternative to coupling reactions generally used in amide synthesis, and its importance is expected to increase over the coming years.³⁹ However, the enzymatic formation of the peptide bond usually requires a high amount of biocatalyst, which is a major drawback. The sustainability of these



Scheme 5 Chemo-enzymatic synthesis of the bromoazidocondurititol derivative, and synthesis of triazolyconduritols.



Scheme 6 Representative structures obtained by chemoenzymatic hydrolases/CuAAC reactions in (a) peptidomimetic synthesis (b) kinetic resolution of racemates.

processes was greatly improved with the implementation of immobilized enzymes, which are usually very stable and easily removed from the reaction medium, playing a crucial role in green amide syntheses.

Due to their structural and electronic similarity, 1,2,3-triazoles can act as peptidomimetic amides. Specifically, the 1,4-disubstituted 1,2,3-triazoles are bioisosteres of *trans*-amide compounds.⁴⁰ Due to the importance of peptides as pharmacologically active compounds, the triazol-ring represents an excellent alternative for the synthesis of peptides with increased metabolic stability, as well as biologically-active natural product analogs difficult to obtain through conventional synthesis.

In 2013, the Müller group reported the first enzymatic aminolysis catalyzed with lipases and a subsequent CuAAC reaction.⁴¹ For the aminolysis reaction, different commercially available immobilized lipases were studied. However, only Novozym® 435 and Immobead® 150 successfully catalyzed the reaction of methyl esters **57** with propargyl amine **58**. Novozym® 435 (*Candida antarctica* lipase B [CALB] immobilized in Lewatit VP OC 1600) proved to be superior and achieved 68% conversion in 24 h. The optimized reaction conditions can be applied to a broad range of substrates and generally achieve good yields. The biocatalytic synthesis of propargylamides was then coupled to the CuAAC reaction using CuO₂/benzoic acid as



Scheme 4 A chemoenzymatic methodology for the synthesis of chiral hydroxytriazole.





Scheme 7 Synthesis of propargyl amides with CAL-B (Novozym® 435) and consecutive three-component synthesis of the amido methylsubstituted 1,2,3-triazoles.

catalyst, and H₂O : MeOH (1 : 1) as solvent system. The amide-triazole products **66–67** were obtained with yields between 51–85% (Scheme 7).

Based on the aforementioned studies, the Müller group reported the consecutive seven-component synthesis of triamides with a triazole moiety in 2019.⁴² This approach incorporated the biocatalytic aminolysis-CuAAC reaction in a new 5-stage multi-component one-pot synthesis. The reactions featured (i) the synthesis of a diamide with a methyl ester group, from a 4-component Ugi-reaction, (ii) the propargylamide synthesis catalyzed by Novozym® 435, (iii) a CuAAC reaction and (iv) a Suzuki cross-coupling to the final product. This sequential one-pot procedure allowed to synthesize compound **70** in 36% overall yield. The described method allows for the generation of



Scheme 8 Consecutive seven-component sequence: Ugi reaction, CALB-catalyzed aminolysis, CuAAC, Suzuki coupling for the synthesis of product **70**.

compound libraries, in an easy way without the need for intermediate purification (Scheme 8).

A second application of the hydrolases combined with CuAAC are the kinetic resolutions of racemic mixtures. In this context, Büyükdali *et al.* described the one-pot synthesis of chiral benzothiophenyltriazoles and benzofuranyltriazoles in 2015.⁴³ This bio-click approach produces the enantiomerically enriched homopropargylic alcohols (+) **72** using the commercially available TLIM (immobilized *Thermomyces lanuginosus* lipase). TLIM proved superior to other lipases, such as Novozym® 435 concerning both, enantioselectivity and reaction time. Furthermore, the reaction occurs in vinyl acetate, which acts as an acyl donor and solvent (Scheme 9). Unfortunately, the amount of biocatalyst necessary was 1 : 1% w/w with respect to the substrate, and recycling experiments of the catalyst were not reported. The concatenation of the TLIM-mediated enzymatic resolution and a click reaction was achieved through a subsequent one-pot multicomponent CuAAC reaction using CuSO₄·5H₂O/Na ascorbate, NaN₃, L-proline, and Na₂CO₃, furnishing the corresponding triazole derivatives (+) **79** with good yields (Scheme 9). The *in situ* generation of the organic azide eliminates the need of an intermediate purification process.

In 2018, Moisă *et al.* reported a new bio-click approach for the separation of racemic secondary alcohols **82** by simple extraction.⁴⁴ This methodology uses the enantioselectivity of the lipases of Novozym® 435 and *Pseudomonas fluorescens* (L-AK) for the catalysis of an O-acylation of heteroarylethanol with different esters **83a–d** as acylating agents. The corresponding



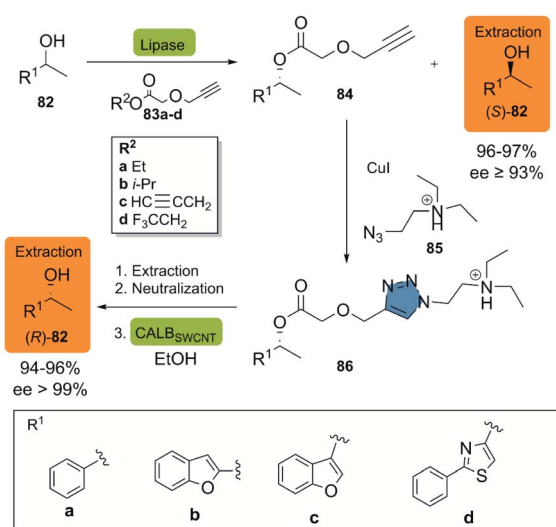
Scheme 9 Enzymatic resolution of racemic homopropargylic alcohols with TLIM and multicomponent synthesis of 1,4-disubstituted 1,2,3-triazole derivatives.

(*R*)-product **84** is then subjected to a CuAAC reaction with azido-functionalized tertiary amine **85** in the presence of CuI, furnishing the triazole quantitatively. The ionizable triazole ester was efficiently extracted with an aqueous acetic acid solution, while the (*S*)-**82** alcohol was recovered from the organic phase. The alcoholysis of the triazole derivative in the presence of ethanol was catalyzed by CALB_{SWCNT} (CALB covalently immobilized on single-walled carbon nanotubes) and allowed for the obtention of the (*R*)-**82** alcohol (Scheme 10). Up-scaling of the process to a multi-gram scale revealed, that there was neither a change in the selectivity of the enzymes nor in the efficiency, as reflected in virtually unchanged isolated yields of the products (>99%).

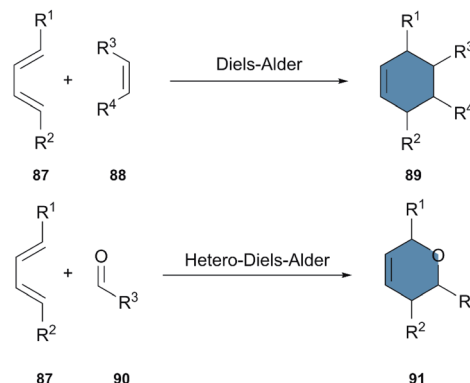
3.2 Diels–Alder reactions and biocatalysis

The Diels–Alder (DA) reaction is an outstanding tool for the synthesis of cyclohexenes and other ring structures. Involving the cycloaddition of a diene and a dienophile, it has been extensively used for the synthesis of complex molecules and is one of the most powerful transformations in organic chemistry (Scheme 11). Moreover, the DA reaction is stereospecific, shows a high atom economy and reliability, and can be carried out in environmentally-friendly solvents.^{45,46} A hetero-Diels–Alder-reaction (HAD) is equally well studied, and is a reliable alternative for the synthesis of ring structures including heteroatoms.

In contrast to the CuAAC reaction, the DA reaction can be carried out without a catalyst. However, the use of Lewis acids allows to decrease the activation energy of the DA reaction, increasing its rate and lowering the necessary reaction temperature.^{47–49} A further increase of the reaction rate can be achieved by the incorporation of EWG-groups in the dienophile and EDG-groups in the diene, favoring the electronic distribution necessary for the reaction to occur.

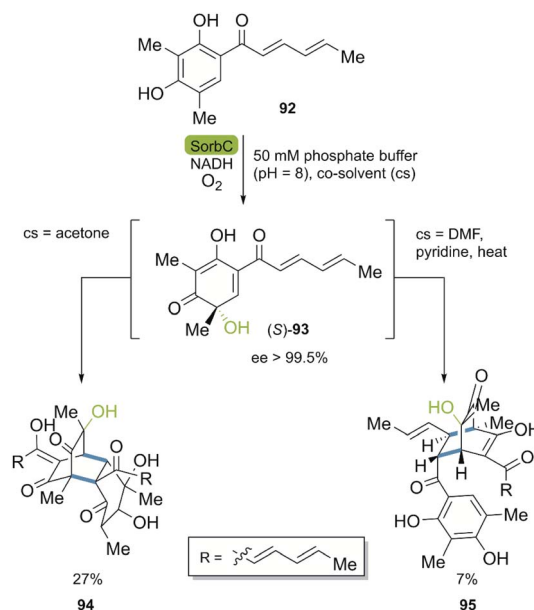


Scheme 10 Production of the two enantiomeric forms of (hetero)arylethanols by an enzymatic KR/click reaction-based separation process. This figure has been Adapted from ref. 44 with permission from the Royal Society of Chemistry.

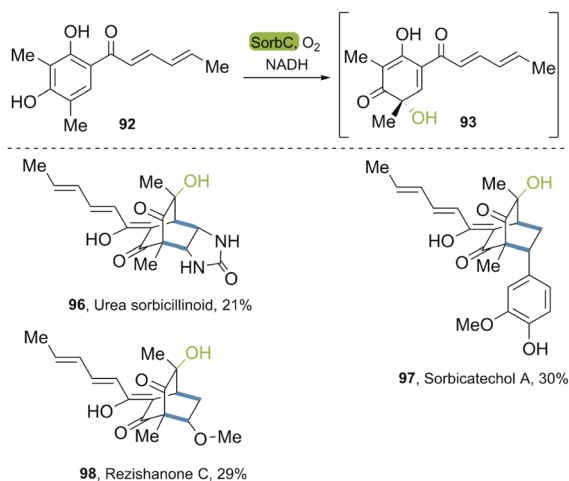


Scheme 11 Diels–Alder reaction and hetero-Diels–Alder reaction.

3.2.1 DA/HDA and oxidoreductases. The combination of DA reactions with oxidoreductases has been successfully applied in the stereoselective synthesis of complex molecules. The bio-click approach has been used for the synthesis of sorbicillin derivatives, an important family of compounds from fungi, exhibiting promising biological activities.⁵⁰ In 2017, the Gulder group reported the total synthesis of bisorbicillinoids by an enzymatic oxidative dearomatization of sorbicillin **92** and a subsequent DA reaction (Scheme 12).⁵¹ In the first part of the synthesis, the enzyme SorbC catalyzed the enantioselective formation of sorbicillinol (*S*)-**93** (ee > 99.5%), followed by the rapid dimerization to the product through a DA reaction. The solvent polarity proved to be crucial for the stability of sorbicillinol (*S*)-**93**, when CH₂Cl₂ was used to quench the reaction, the bisorbicillinol **94** was obtained in a 27% yield in 40 minutes, without any other dimeric analogs. Different co-solvents could be used for the control of the selective formation of different



Scheme 12 Key step of the stereoselective enzymatic total synthesis of bisorbicillinol **94** and sorbiquinol **95**.



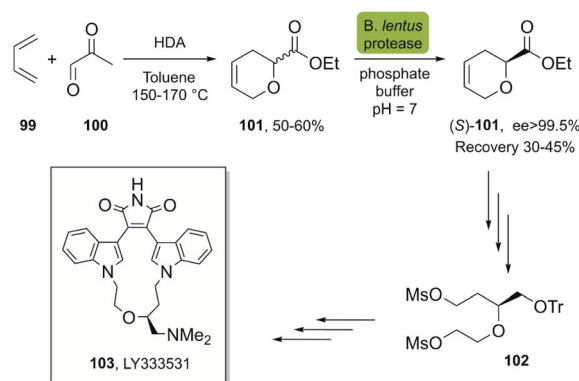
Scheme 13 Chemoenzymatic synthesis of the sorbicillinoids via enantioselective oxidative dearomatization of sorbicillin with SorbC and DA reaction.

byproducts. As an example, the use of DMF combined with the work-up comprising extraction with CH_2Cl_2 , a fast evaporation of CH_2Cl_2 after 4 h, and the subsequent heating in the presence of pyridine furnished sorbiquinol **95** in 7% yield. This chemoenzymatic approach proved to be an excellent alternative in comparison to the chemical syntheses reported to date. For example, the Deng group reported an elegant enantioselective total synthesis of bisorbicillinol **94** with a global yield of 27%, however applying 10 steps.⁵²

In 2018, the groups of Gulder and Narayan independently reported a chemoenzymatic approach towards the synthesis of new sorbicillin derivatives.^{53,54} In these investigations, the use of the SorbC enzyme once again furnished sorbicillinol **93** by oxidative dearomatization as a key step. The use of CH_2Cl_2 enabled the extraction of **93**; and its subsequent reaction with different dienophiles allowed the production of sorbicillin derivatives in a simple chemoenzymatic process. The scope of products was very versatile, giving access to urea sorbicillinoid **96**, sorbatechol A **97**, and rezishanone C **98** (Scheme 13).

The Narayan-group synthesized the urea sorbicillinoid **96** via a DA reaction between sorbicillin and bisacylated urea and following by the addition of LiOH. Furthermore, they described different monooxygenases, such as Trop B and AzaH, for the catalysis of the oxidative dearomatization of resorcinols, allowing for the synthesis of valuable *ortho*-quinol products,⁵⁴ a process that was adapted to gram scale.

3.2.2 DA/HDA and hydrolases. In 2002 Caille *et al.*⁵⁵ reported the chemoenzymatic synthesis of ethyl (*S*)-3,6-dihydro-2H-pyran-2-carboxylate **101**, which is a useful intermediate in the synthesis of macrocyclic bisindolylmaleimide (LY333531) **103**, a potent inhibitor of protein kinase C β (PKC β).⁵⁶ The bio-click approach developed by Caille *et al.* took advantage of inexpensive starting materials, namely butadiene **99** and ethyl glyoxylate **100**, producing 2-carboethoxy 3,6-dihydro-2H-pyran **101** in a HAD-reaction. The subsequent resolution of the racemic ester by selective enzymatic hydrolysis using *B. lentus*

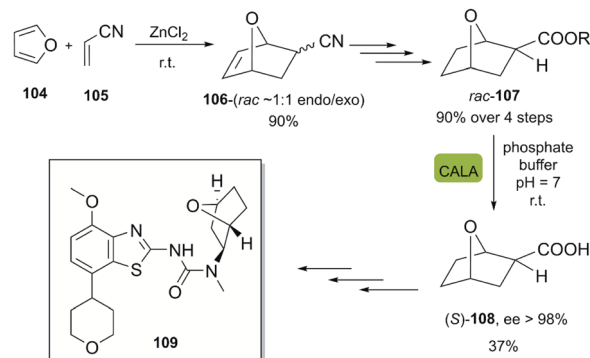


Scheme 14 Chemoenzymatic synthesis of (*S*)-**101**, a key intermediate in the synthesis of LY333531, a protein kinase C inhibitor (PKC β).

protease was carried out on a gram-scale, and after 8 h the ee of the (*S*)-configured ester was >99.5% (Scheme 14).

In 2010, Wirz *et al.* reported the scalable enantioselective synthesis of a benzothiazole derivative **109**, which was evaluated as an A2 receptor antagonist for the treatment of major depression.⁵⁷ The developed chemoenzymatic process consists of the DA-reaction of furan **104** with acrylonitrile **105**, activated by ZnCl_2 , furnishing bicyclic product **106**. The following steps gave rise to the racemic product **107**, which was hydrolyzed by CALA lipase in a stereoselective fashion, furnishing the (*S*)-enantiomer **108** with excellent enantioselectivity (Scheme 15).

Abacavir is a carbocyclic nucleoside with antiviral activity, acting as a reverse transcriptase inhibitor (see Scheme 16). It is used in combinations with other nucleoside analogues, such as lamivudine, for the treatment of HIV-infection.⁵⁸ Crimmins *et al.* described an asymmetric synthesis of abacavir in 1996, making use of a first-generation Grubbs catalyst.⁵⁹ An alternative synthetic route (Scheme 16a), used the Vince lactam **111**, which can be obtained by the DA reaction of cyclopentadiene **110** and tosyl cyanide.⁶⁰ The subsequent enzymatic kinetic resolution of the racemic Vince lactam with lactamase furnished the corresponding amino acid **113** (with (–) lactamase) or the (–) Vince lactam **112** (using (+) lactamase), which was then hydrolyzed into the desired amino acid **113**.⁶¹ This key



Scheme 15 Bio-click approach for the synthesis of (*S*)-**108**. Precursor in the synthesis of A2 receptor antagonist **109**.





Scheme 16 Chemoenzymatic synthesis of abacavir. (a) Using a lactamase by kinetic resolution of *rac*-Vince lactam and (b) using a CALB lipase in the kinetic resolution of aminocyclopentenol derivative.

intermediate can then be converted into the desired product abacavir.⁶²

The Vince lactam is not only a useful intermediate in the synthesis of abacavir, but also in the synthesis of peramivir and carbovir. In fact, numerous groups have reported the application of different lactamases for the resolution of the Vince lactam, en route to these powerful antiviral agents.⁶³ A work carried out by Tardibono *et al.*⁶⁴ presents a bio-click methodology towards homoabacavir **116b** (Scheme 16b). In this approach, the hetero Diels Alder-reaction between cyclopentadiene and the transient nitroso-compound of *N*-Boc-hydroxylamine is used for the formation of intermediate **114**. The subsequent reduction of the N–O bond furnished the racemic aminocyclopentenol derivative **115**, which was then successfully subjected to a kinetic resolution with CALB lipase, using vinyl acetate with posterior basic hydrolysis, furnishing **115**(–) (80% ee) as a key intermediate for the further syntheses of homoabacavir **116b** and homocarbovir.⁶⁴

3.3 Epoxide-opening and biocatalysis

Epoxides are valuable and versatile building blocks and synthetic intermediates common in nature and synthesis.^{65,66} The outstanding reactivity of the three-membered ring is mainly caused by the special geometry and angle strain, endowing the ring opening reaction with a favourable thermodynamic driving force (24 to 28 kcal mol^{–1}).^{67,68} The S_N2 ring opening of epoxides shows a variety of desirable characteristics, such as reliability, stereospecificity and regioselectivity.⁴ The congener aziridines present similar characteristics, however with the advantage of the presence of a nitrogen atom, allowing subsequent chemical transformations in a somewhat easier way.^{69,70} Nevertheless, fewer efficient methods for the direct synthesis of aziridines are available,⁷¹ also reflected in the scarce number of respective bio-click approaches incorporating these substrates. The recent development of methods for the enzyme-catalyzed olefin aziridination⁷² could change these shortcomings and foster the development of a greater number of bio-click approaches incorporating aziridines.

The ring-opening of epoxides has been extensively applied for the synthesis of complex molecules.⁷³ In fact, the synthesis of numerous Active Pharmaceutical Ingredients (APIs) uses ring-opening of epoxides as a key step.^{74,75} The bio-click approach towards the ring-opening reaction of epoxides is a field of increasing interest.

Generally, the bio-click approach incorporating epoxides is applied in two ways: (a) the selective enzymatic oxidation of alkenes, furnishing epoxides in green conditions for the subsequent ring opening; (b) the oxidation of alkenes, a subsequent enzymatic resolution of the racemic epoxides and the ring-opening of the desired candidate (Scheme 17).

3.3.1 Epoxide-opening and oxidoreductases. The oxidoreductases are useful for the synthesis of enantiopure epoxides.⁷⁶ Also, the enantioselective biooxidation of alkenes has been used as a key step in the bio-inspired synthesis of natural products.^{77,78} The directed evolution and rational design of enzymes has allowed the optimization of selectivity, catalytic efficiency and stability of oxidoreductases towards the levels necessary for industrial applications.^{79,80} Regarding the bio-click approach, the direct or indirect synthesis of epoxides using oxidoreductases is predominant. However, the biooxidation, chemical epoxidation and ring opening approach has been used in the total synthesis of molecules with multiple contiguous stereocenters.



Scheme 17 General bio-click approach in epoxide-opening reactions.



cis-1,2-Dihydrocatechols **46** are obtained by the enzymatic dihydroxylation of arenes **45** and represent an excellent platform for the synthesis of alkaloids, sugars, cyclitols, prostaglandins, terpenes, polymers and others (Scheme 18a).⁸¹ As an example, the enantioselective synthesis of *cis*-1,2-dihydrocatechols is carried out in a whole-cell fermentation, which provides an efficient cofactor regeneration system.⁸² *E. coli* JM109 (pDTG601) which contains the genes from *Pseudomonas putida* F1 for the overexpression of the toluene dioxygenase system (TDO),⁸³ is the most common biocatalyst applied for the enantioselective synthesis of *cis*-1,2-dihydrocatechol derivatives. Careful optimization of the reaction conditions in a biphasic system can achieve yields as high as 35 g L⁻¹.⁸⁴ This is quite remarkable, considering that no efficient chemical method is known for the production of this class of compounds at scale.^{79,81}

The *cis*-1,2-dihydrocatechols are highly versatile and a great platform for enantioselective synthesis⁸⁵ (Scheme 18b). Monoepoxidation, for example, is a transformation that can be easily applied to *cis*-1,2-dihydrocatechols **121** and combined with

enantio- and regioselective **122** ring-openings, which can be efficiently applied in total synthesis.⁸⁶

For the ring opening of the epoxides on dihydrocatechols **122** the general preference is towards the cleavage in the 5a position (see Scheme 18b), mainly associated to two factors: (a) the greater stabilization of the transient carbocation in the allylic position; (b) less steric hindrance at position 5a as compared to 6a. In this way, the cleavage of epoxides is generally achieved with high regio- and enantioselectivity, providing the corresponding products of the *trans*-1,2 addition efficiently.⁸⁶

Representative examples for the opening of monoepoxides **122** (obtained from biocatalytic enzymatic *cis*-dihydroxylation of benzene derivatives **45**) with different nucleophiles are ubiquitous (see Scheme 18, vitamin C,⁸⁷ codeine,⁸⁸ narseronine,⁸⁹ C2 epimer of aminocyclitol⁹⁰) and crucial in the synthesis of natural products.

The indirect epoxidation of α -haloketones using ketoreductases (KREDs) has become a common strategy to obtain chiral epoxides, also reflected in the stereoselective synthesis of different APIs obtained on a kg-scale. The development of new



Scheme 18 (a) Biocatalytic dihydroxylation of arenes by *E. coli* JM109 (pDTG601), (b) chemical reactivity of *cis*-1,2-dihydrocatechols. Representative molecules obtained from bio-click methodologies (TDO-ring opening). (c) Vitamin C 1998 by Longmore *et al.*,⁸⁷ (d) codeine 2009 by Leisch,⁸⁸ (e) narseronine 2011 by Schwartz *et al.*,⁸⁹ (f) C2 epimer of aminocyclitol 2019 by Carrau.⁹⁰



Scheme 19 KRED mediated epoxidation of α -bromo ketones and ring opening: (a) synthesis of BMS-960, (b) synthesis of MK-7145.

variants of KREDs through protein engineering has widely enhanced scope, robustness, and selectivity of these biocatalysts. The fact that they are an excellent green alternative to conventional catalysts is further underscored by the easy workup, generally avoiding chromatographic product purifications.

In 2017, Hou *et al.* reported a chemo-enzymatic synthesis of BMS960, a powerful S1P_1 -receptor agonist.⁹¹ Its synthesis involved the enzymatic reduction of α -bromoketone **134** (100 g in 5 h at 40 °C) to the corresponding alcohol **135** using the commercially available KRED-NADH-110 (substrate enzyme ratio 200 : 1). The chiral alcohol was then extracted with MTBE and reacted with sodium *tert*-butoxide towards the epoxide with 93% overall yield and 100% ee (*S*-enantiomer). Subsequently, a regio- and stereospecific ring opening of the epoxide **136** with (*S*)-ethyl piperidine-3-carboxylate in the presence of a catalytic amount of DMAP at 50 °C, provided the product **137** in up to 77% yield and with an ee of 99.6% after recrystallization. Four successive chemical transformations furnished BMS-960 in 23–33% overall yield (Scheme 19a).

In 2020, Ruck *et al.* reported a kg-scale synthesis of MK-7145, a clinical candidate for the treatment of hypertension and associated heart failures.⁹² The synthesis involved the enzymatic reduction of a bromo-ketone **139** to the corresponding bromohydrin as a key step. The commercial KRED-MIF-20-induced reduction and a subsequent ring closure furnished the corresponding enantiopure epoxide in 82% yield and excellent enantiopurity. Subsequently the target molecule MK-7145 **141** was obtained through a bis-epoxide opening with piperazine at 140 °C, furnishing the product in 75% yield (Scheme 19b).

Similar to the abovementioned biocatalytic *cis*-dihydroxylation of arenes **142**, the Hollmann group recently described the direct biocatalytic epoxidation of naphthalene derivatives.⁹³ The methodology described is based on the recombinantly

evolved peroxygenase variant PaDa-I from *Agrocybe aegerita* (rAaeUPO), a self-sufficient enzyme. Careful control of the reaction time of the nucleophilic addition allows for the generation of the ring-opening products **144** using nucleophiles such as sodium azide (Scheme 20).

In this way, the synthesis of eleven derivatives in yields ranging from 6 to 73% was reported. These arene oxides are excellent building blocks for the synthesis of *trans*-disubstituted cyclohexadienes and can be further functionalized towards triazoles, amino alcohols, and other valuable entities. Epichlorohydrin **145** is a versatile molecule with multiple applications,⁹⁴ and a useful starting material for the synthesis of different APIs.

The synthesis of rivaroxaban, an anticoagulant widely used in the prophylaxis of cardiovascular diseases,⁹⁵ has been described by numerous groups using (*R*)-epichlorohydrin **145** as a building block.⁹⁶ In a bio-click approach, rivaroxaban was synthesized *via* (*R*)-**145**, which can be obtained either by an enantioselective biooxidation of 3-chloropropene **146** using chloroperoxidase CPO from *Caldariomyces fumago*,⁹⁷ or by resolution of racemic **145** using epoxyhydrolase EH ArEH⁹⁸ from *Agrobacterium radiobacter* (Scheme 21). Further functionalisation of the desired epoxide with phthalimide under basic conditions using a phase transfer agent and a subsequent regio- and enantiospecific ring opening of epoxide **148** with arylamine



Scheme 20 Bio-click synthesis of *trans*-disubstituted cyclohexadiene derivatives using peroxygenase PaDa-I, and chemical ring-opening with different nucleophiles.



Scheme 21 Bio-click approach for the synthesis of rivaroxaban.

149, furnished the product 150 smoothly en route to rivaroxaban.⁹⁹

3.3.2 Epoxide-opening and hydrolases. In 2012, Porcar *et al.*¹⁰⁰ reported the stereoselective chemoenzymatic synthesis of imidazoles under continuous flow conditions for the generation of new chiral ionic liquids.¹⁰¹ The first step is the epoxide opening of cyclohexene oxide 152 with imidazole 153 in a continuous flow reactor with microwave (MW) irradiation (Scheme 22). The resulting cyclohexanol derivative (\pm)-*trans*-154 is then subjected to the second step, an acylation with vinyl acetate catalyzed by the commercially available immobilized lipases Novozym® 435 or PSL-CI (*Pseudomonas cepacia* lipase), allowing an efficient kinetic resolution. The applied continuous-flow methodology demonstrated high efficiency in a batch process, and the lipases provided the corresponding acylated kinetically-resolved product 155 with excellent enantioselectivities (>99%) and virtually complete conversion.

In 2015, Villar *et al.* reported the chemoenzymatic synthesis of optically pure, orthogonally protected *trans*-3-amino-4-hydroxypiperidines, potential molecules for the synthesis of chiral bioactive compounds (Scheme 23).¹⁰² The developed bio-click methodology developed, starts with the regioselective epoxide-opening with diallylamine, providing access to the racemic *trans*-cyclohexanol 157. The subsequent enzymatic kinetic resolution *via* transesterification with vinyl acetate was performed with Novozym® 435, which gave the desired product 158 with the highest conversion (47%) and enantioselectivity (ee > 99%) among the enzymes tested.

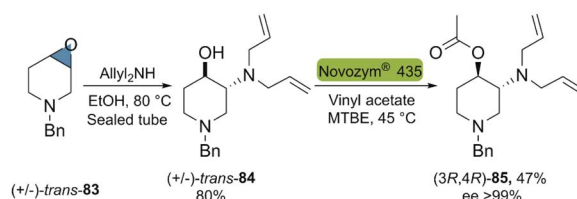
Propranolol is a versatile beta-adrenergic receptor antagonist used in the treatment of several cardiovascular such as hypertension, cardiac arrhythmias, *etc.*¹⁰³ Quite remarkably, (*S*)-propranolol is 100 times more potent β -adrenergic receptor blocker than its enantiomer.¹⁰⁴ In 2015, Dong *et al.*¹⁰⁵ reported

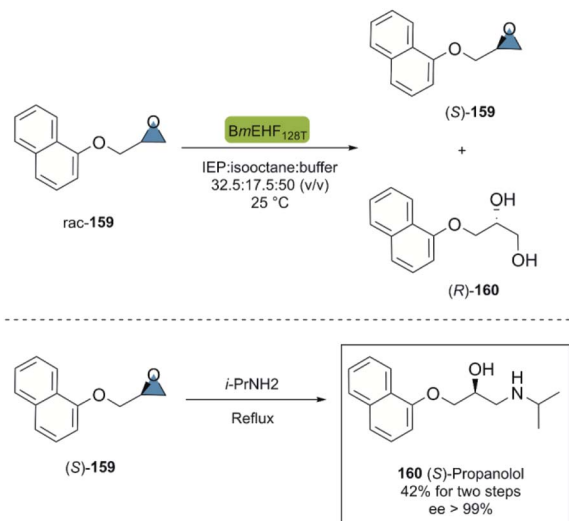
Scheme 22 Synthesis of *trans*-2-(1*H*-imidazol-1-yl)cyclohexanol (b) enantioselective acylation of (\pm)-*trans*-2-(1*H*-imidazol-1-yl)cyclohexanol 154 with vinyl acetate catalyzed by Novozym® 435 or PSL-CI. This figure was used with permission.¹⁰⁰

the chemoenzymatic two-step synthesis of (*S*)-propranolol (Scheme 24). The enzymatic kinetic resolution of racemic epoxide 159 with the hydrolase BmEH128T (from *Bacillus megaterium*), smoothly furnished the (*S*)-configured product 159 smoothly after centrifugation of the undesired enantiomer. The subsequent regioselective ring-opening of (*S*)-159 with isopropylamine under reflux conditions gave (*S*)-propanolol 160 in an overall yield of 42% and excellent enantioselectivity after recrystallization (ee > 99%).

3.4 Thiol-Michael reactions and biocatalysis

Thiol-Michael click reactions, either *via* free-radical, or catalyzed Michael-additions, are valuable transformations widely

Scheme 23 Chemoenzymatic synthesis of optically pure orthogonally protected *trans*-3-amino-4-hydroxypiperidines.



Scheme 24 Chemoenzymatic synthesis of (S)-propanolol.

recognized in polymer science and bioconjugation chemistry, since they allow for the efficient formation of C–S bonds from thiols and alkenes (Scheme 25).^{106–108} They are usually highly efficient, work under mild reaction conditions, are regioselective and atom-economic, and only generate few or no by-products which are generally easily separated.^{109,110} Bio-click reactions approaches featuring thiol-Michael chemistry are scarce to date, despite the advantages of chemoenzymatic reactions in this context.

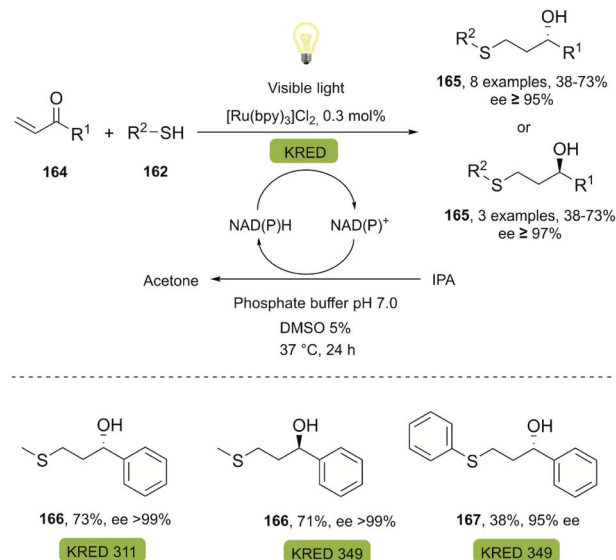
3.4.1 Thiol-Michael reactions and oxidoreductases. The spatial and temporal control of light as an energy source for thiol-Michael click reactions proved valuable in tandem bioclick reactions, as reported in 2018 by Lauder *et al.* for the synthesis of volatile chiral 1,3-mercaptoalkanol **166–167**.¹¹¹ The reported one-pot methodology comprises a thiol-Michael reaction photocatalyzed by visible light combined with [Ru(bpy)₃]Cl₂ (0.3 mol%), fully compatible with the conditions of the subsequent bioreduction using KREDs (see Scheme 26). The enzymes used furnished the corresponding (S)- or (R)-enantiomers in yields (38–73%) and high enantiomeric excess.

4. Summary and outlook

This review summarizes developments in a field of research that combines biocatalytic transformations and click reactions. The proposal of the term bio-click chemistry comes quite naturally if one considers the varied and efficient syntheses of complex molecules explored. A clear classification of the different chemoenzymatic transformations will undoubtedly promote the



Scheme 25 Thiol-Michael click chemistry.



Scheme 26 One-pot photo-biocatalytic (thiol-Michael/KRED reduction).

development of more sustainable and efficient processes in this area (see Table 1).

Bio-click chemistry reactions are demonstrated to be remarkably complementary, as is the case of click chemical reactions, which are regiospecific by nature but not necessarily enantioselective. The joint use of enzymes/microorganisms



Fig. 3 Biocatalytic synthesis of islatravir. (This figure has been adapted from ref. 22. With permission from Science Journal, copyright 2020).

Click reaction	Enzyme or microorganism	Structure	Function	Reference
CuAAC	Carbonyl reductase from <i>Candida magnoliae</i> (CMCR)		β -Adrenergic receptor blocker analogue	29
	(+)- γ -Lactamase, (-)- γ -lactamase from <i>Bradyrhizobium japonicum</i> USDA 6 <i>Candida antarctica</i> B lipase (CALB)		Carbocyclic nucleoside with antiviral activity	58, 60, 61 and 112 64
Diels-Alder or hetero-Diels-Alder		Abacavir		
Oxidoreductase SorbC			Antiviral activity against H1N1	53, 54 and 113
		Sorbicatechol A		
Epoxide-opening	<i>E. coli</i> JM109 (pDTG601)		Painkiller	88
		Codeine		
Ketoreductase, KRED-NADH-110			S1P1 receptor agonist	91
		BMS-960		
Thiol-Michael	Ketoreductase, KRED-311		Volatile sulfur compound	111
		1,3-mercaptoalkanol		

It is essential to emphasize, that click reactions, biocatalysis or their combination do not generate greener processes *per se*, and it is always necessary to carefully evaluate the exact methodology, the implementation in a reaction sequence and the expected overall efficiency and sustainability for future improvements.

Conflicts of interest

There are no conflicts to declare.

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