Synthetic pathways to tetrahydrocannabinol (THC): an overview

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The therapeutic effects of molecules produced by the plant species Cannabis sativa have since their discovery captured the interest of scientists and society, and have spurred the development of a multidisciplinary scientific field with contributions from biologists, medical specialists and chemists. Decades after the first isolation of some of the most bioactive tetrahydrocannabinols, current research is mostly dedicated to exploiting the chemical versatility of this relevant compound class with regard to its therapeutic potential. This review will primarily focus on synthetic pathways utilised for the synthesis of tetrahydrocannabinols and derivatives thereof, including chiral pool-based and asymmetric chemo- and biocatalytic approaches.

1. Introduction

The plant species Cannabis sativa has captured for centuries the attention of society because of its therapeutic and psychoactive properties. Dating back to 440 B.C., the first literary mention by the philosopher Herodotus was about the use of hemp by the Scythians. The plant was processed into clothing fibres and the hemp-seed was thrown on red-hot stones producing vapours for delight and joy. In the following centuries, cannabis remained a useful plant for society and has often been described in literature. In the 17th century, cannabis-based therapies were prescribed to treat disorders such as depression, joint pain and epilepsy. Centuries later, in 1964 Mechoulam et al. described the isolation of tetrahydrocannabinols and a partial synthesis of the molecular scaffold. This seminal paper sparked more intensive research into cannabinoids and led to the scientific field that we know today. The major phytocannabinoids isolated from Cannabis sativa include cannabidiol (CBD, 5), Δ8- and Δ9-tetrahydrocannabinol (THC) - an overview.

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The biosynthesis of phytocannabinoids and structurally related molecules in *Cannabis sativa* has been described extensively over the past decades. In most cannabinoids a structural similarity can be identified comprising a resorcinol (A-ring) and terpinoid moiety (C-ring, Fig. 1).13 Terpenoids are formed starting from the (non)-mevalonate pathways which produce dimethylallyl pyrophosphate (DMAPP) and isopentenyl pyrophosphate (IPP).14 Both components are coupled by geranyl pyrophosphate synthase to form geranyl pyrophosphate (GPP, 1) which is a precursor for a broad range of terpenoids (Scheme 1). Coupling of olivetolic acid (2) with GPP by geranyl transferase yields cannabigerolic acid (CBGA, 3), which is the biosynthetic starting point for most cannabinoids.1,12 Via a subsequent stereoselective ring-closure induced either by CBDA or THCA synthase, cannabidiolic acid (CBDA, 4) and Δ^9^-tetrahydrocannabinolic acid (Δ^9^-THCA, 6) are formed, respectively. In addition, numerous other cannabinoid subvariants are formed from CBGA (3) including cannabidiol (CBD, 5), Δ^9^-tetrahydrocannabinol (Δ^9^-THC, 7) and cannabinol (CBN, 8). Until now various synthetic cannabinoids have been synthesised and high affinity CB1 and CB2 agonists and antagonists have been identified, which hold considerable promise for pharmaceutical applications.4,5 Although a large collection of synthetically prepared cannabinoids has been reported, the synthetic approaches employed have not yet been systematically reviewed for tetrahydrocannabinol.15 Here we will provide an overview of the most prominent synthetic pathways to the THC scaffold. This is of interest from both a medicinal chemistry viewpoint, as it will shed light on possible new cannabinoid derivatives to be synthesised, but also from a synthetic organic chemistry angle, as the synthesis of the terpenoid C-ring of THCs is especially challenging requiring creative synthetic solutions (Fig. 1).

2. Terpinoid chiral pool approaches

The biosynthesis of THC relies on stereoselective enzymatic conversions using linear, achiral polyketides and alkenyl phosphates. From a synthetic point of view, utilising chiral pool feedstocks is preferred over asymmetric approaches since it avoids typically more complex asymmetric transformations to control the stereochemistry.

2.1 Cannabinoid synthesis using (−)-verbenol

The first isolation and structural NMR characterisation of THC and (−)-CBD was reported by Gaoni and Mechoulam in 1964.3 This pioneering research is considered as one of the most influential publications since it sparked the scientific field of

**Scheme 1** Biosynthesis of several major cannabinoids from *Cannabis sativa*.
cannabinoid chemistry and biology. Previously, Fahrenholtz and co-workers attempted to obtain enantiomerically pure THC, but were unsuccessful in resolving a THC racemate.\(^{16,17}\) Three years after the initial publication by Mechoulam and Gaoni, the first fully stereoselective synthesis of \((-\text{-trans-}\Delta^8\text{-THC})\) was described by Mechoulam, Braun and Gaoni.\(^{18}\)

Friedel–Crafts alkylation of olivetol (11) with \((-\text{-verbenol})\) (10), catalysed by \(p\)-TSA or boron trifluoride to produce the intermediate allylic cation, provided olivetylverbenyl (12), which after repeated treatment with boron trifluoride provided \((-\text{-trans-}\Delta^8\text{-THC})\) (9) (Scheme 2). A direct approach using Lewis acid catalysis, combining both steps into one, led to similar overall yields of up to 35\% of \((-\text{-trans-}\Delta^8\text{-THC})\) (9). Subsequent isomerisation through chlorination and base-induced elimination provided \((-\text{-trans-}\Delta^9\text{-THC})\) (7).\(^{19}\)

Interestingly, five years later the authors gave a mechanistic rationale describing the allylic carbocation intermediate.\(^{20}\) By using deuterated \((-\text{-verbenol})\) (10) the corresponding labelled \((-\text{-trans-}\Delta^8\text{-THC})\) (9) product was prepared, effectively demonstrating the stereochemistry of the prepared C–C bond. The authors stated that due to the challenging isolation of pure cannabinoids from plants, they considered stereoselective synthesis a preferred approach to obtain single, well-defined cannabinoids in enantiopure form.

Recently, Dethe and co-workers also demonstrated the use of \((-\text{-verbenol})\) (10) in the synthesis of phytocannabinoids...
isomeric cannabinoids 14 alkyla

tion of carbazole ring opened using LiAlH4. In addition, the authors succeeded in isolating the corresponding epoxide isomers, which were both subsequently oxidised with m-CPBA to give the regioisomeric cannabinoids 14 and 15 selectively using different amounts of catalyst. The natural products 16 and 17 were obtained upon m-CPBA oxidation of olefin 15 to give the corresponding epoxide isomers, which were both subsequently ring opened using LiAlH4. In addition, the authors successfully extended this approach to (α)-verbenol leading to ent-murrayamine-O and -P (not shown).

2.2 p-Mentha-2,8-dien-1-ol for both THC isomers

Shortly after the publication of Mechoulam, the use of enantiopure chiral pool terpenoids became common practice in cannabinoid synthesis. Elaborating on an earlier procedure, Petzilka and co-workers reported a stereoselective synthesis of (−)-murrayamine-O (16) and (−)-murrayamine-P (17, Scheme 3).21 Previous reports showed that BF3-mediated activation of verbenol leads to the thermodynamic Δ8-isomer.18 In this synthesis, however, the authors describe that with small amounts of BF3·OEt2 and short reaction times the kinetic Δ9-isomer was obtained. This was demonstrated by Friedel–Crafts alkylation of carbazole 13 with verbenol (10) to give the regioisomeric cannabino
doids 14 and 15 selectively using different amounts of catalyst. The natural products 16 and 17 were obtained upon m-CPBA oxidation of olefin 15 to give the corresponding epoxide isomers, which were both subsequently ring opened using LiAlH4. In addition, the authors successfully extended this approach to (α)-verbenol leading to ent-murrayamine-O and -P (not shown).

Besides bis-adducts and iso-THC derivatives, there was no Δ8-THC encountered under these conditions (Scheme 4B). Despite the challenging purification and rather low yield, this pioneering work is still used in the production of (−)-trans-Δ9-THC derivatives because of the high efficiency and mild reaction conditions of the one-pot process.

The stereoselective preparation of Δ8/9-THC scaffolds using p-mentha-2,8-dien-1-ol can also be achieved by using a direct C-ring conjugation and acetate 19. Previous Lewis or Brønsted acid approaches often lacked regiochemical selectivity, were highly sensitive to reaction conditions and afforded many side products.25 In contrast, most C-ring arylation strategies rely on metal-catalysed activation of resorcinol structures and allow for a broad range of aryl substrates.26–29

Rickards and Rönneberg introduced the organometallic synthetic approach in the synthesis of Δ2-THC brominated analogues (Scheme 5).30 It commenced with the preparation of homocuprate olivetol by Petzilka in the synthesis of Δ2-THC (Scheme 4A).19

The successful isolation and direct synthesis of (−)-trans-Δ9-THC (7) was reported shortly thereafter by Razdan and co-workers.24 Instead of the isomerisation of (−)-trans-Δ9-THC (9) to its Δ2-isomer using gaseous hydrochloric acid as shown by Mechoulam, Razdan described a direct synthesis starting from the same starting materials but now using 1% of boron trifluoride as catalyst and magnesium sulfate as drying agent.
Deprotection of the methyl ethers appeared challenging and was unsuccessful using boron tribromide. Alternatively, the authors proposed to mask the alkene bonds via addition of hydrogen bromide, forming the highly unstable product 22. The brominated isopropenyl moiety unexpectedly underwent mono-demethylation and concomitant formation of the B-ring system at ambient temperature to afford a methyl ether protected THC scaffold. Fortunately, the final demethylation could be completed using boron tribromide, to give halogenated product 23. (−)-trans-Δ⁹-THC (7) was successfully obtained by following Mechoulam’s intermolecular dehalogenation with potassium tert-butoxide (Scheme 2). The authors also investigated a one-pot procedure to convert 21 into 23 without intermediate purification. Compound 21 was subjected to a stoichiometric excess of HBr at −15 °C, then mono-demethylated, cyclised at ambient temperature, and finally deprotected using boron tribromide to afford crude 23. The crude product was subjected to potassium tert-butoxide to form (−)-trans-Δ⁹-THC in 75% overall yield from (−)-trans-CBD dimethyl ether 21.

2.3 p-Menth-2-ene-1,8-diol for 6-hydroxy-CBD and Δ⁹-THC

In continuation of previous research, Razdan and co-workers investigated the use of other chiral monoterpenoids in the preparation of (−)-trans-Δ⁹-THC (7). A structurally comparable starting material is p-menth-2-ene-1,8-diol (25), which is activated in a similar fashion using BnBr or Lewis acid catalysis. Conversions of up to 51% were observed and isolated yields of 28% of (−)-trans-Δ²-THC (7) were obtained using anhydrous ZnCl₂ instead of the earlier reported boron trifluoride etherate, (Scheme 6). Notably, with this zinc-based Lewis acid the reactions were successfully conducted on multi-gram scale. Despite the one-step procedures of both p-mentha-2,8-dien-1-ol (18) and p-menth-2-ene-1,8-diol (24), the reaction produced a large variety of side products which limits large scale application.

Stoss and Merrath attempted to optimise the synthesis of (−)-trans-Δ⁹-THC (7) by crystallising the (−)-trans-6-hydroxy-CBD (25) intermediate. This crucial step allowed a cleaner cyclisation reaction with ZnBr₂ to form (−)-trans-Δ⁹-THC (7). Despite the two-step sequence, the product was obtained in higher isolated yield, because the final product purification was more efficient. Recently, Cabaj and co-workers published a new synthetic route for the large scale preparation of p-menth-2-ene-1,8-diol (24). This discovery may contribute to industrial application of 24 in the synthetic preparation of cannabinoids.³³

2.4 Syntheses using (+)-trans-2- and (+)-trans-3-carene epoxides

Razdan and co-workers additionally investigated other methods to readily prepare enantiopure THC, and were inspired by the seminal work of Ohloff and Giersch.³⁴ By utilising (+)-trans-2-carene epoxide (26), structurally similar to the strained bicyclic (−)-verbenol (10), another one-step synthesis to (−)-trans-Δ⁹-THC (7) was discovered (Table 1).³⁵ Both boron trifluoride and p-TSA were effective in the activation of carene epoxide, but both catalysts resulted in mixtures of the (−)-trans-Δ²-THC (7) and (−)-cis-Δ²-THC (23) stereoisomers. The observed stereochromical outcome was explained via a carbocation intermediate or concerted pathway by Brønsted or Lewis acid catalysis, respectively. Later, Jamieson and co-workers investigated the mechanism of (+)-trans-2-carene epoxide (21) and compared it with p-mentha-2,8-dien-1-ol (18).³⁶ Interestingly, reactions of 2-carene epoxides (26) with olivetol (11) were lower yielding and produced more byproducts than p-mentha-2,8-dien-1-ol. Reactions using (+)-trans-2-carene epoxides did not afford cannabidiols, and it was hypothesised that the initial Friedel–Crafts alkylation did not occur. The authors postulated a phenolic nucleophilic substitution reaction of olivetol (11) on the cyclopropane moiety of carene, directly followed by an acidic cyclisation reaction.³⁷

Additionally, the isomeric (+)-trans-3-carene epoxide (27) was reacted with p-TSA and the authors observed a p-mentha-dienol-like pathway.³⁵ The proposed mechanism proceeds through acid-catalysed epoxide ring-opening forming a tertiary cation, followed by elimination to an allylic carbocation.³⁸ The (−)-trans-3-carene (27) only gave (−)-trans-Δ⁸-THC (9) in low yield, while forming a large variety of side products.³⁹

2.5 cis-Chrysanthanol in the synthesis of (−)-trans-Δ⁹-THC

Razdan and co-workers also sought to start from a different enantiopure verbenol-based terpenoid, which would allow a

![Scheme 6 Preparation of Δ⁹-THC using p-menth-2-ene-1,8-diol (24).](image-url)
Phenolic deprotection, p-TSA-mediated cyclisation and methyl ether deprotection finally afforded enone 33. This was transformed into the corresponding enol triflate to access the oxidised C-ring of ∆^8-THCs. Due to the sp^2-hybridised alkene in enone 33, the final stereochemistry was installed through a Birch reduction with poor selectivity. Ultimately, triflated (-)-trans and (-)-cis-∆^8-THC derivatives were obtained in a 3:1 ratio (not shown).^{26,43} Despite the low stereoselectivity in the apoverbenone route, the resulting building block thus provides straightforward access to oxidised C-ring cannabinoids. Alternatively, the synthesis of enantiomerically enriched oxidised (-)-trans-∆^8-THC (9) using (+)-apoverbenone (31) was published by Tius and Kannangara (Scheme 8B).^{27} After successfully preparing olivetolic lithiocuprate from ethoxycarbonyl-protected olivetol (37), (+)-apoverbenone (31) was reacted with the lithiocuprate to form a ketone intermediate. This was then transformed into an enol triflate and deprotected to obtain enol ether 34. Treatment with boron trifluoride etherate led to synthon 35, which is a precursor for C-ring-modified (-)-trans-∆^8-THC derivatives.

2.7 CBD synthesis using (R)-2,9-dibromocamphor

Vaillancourt and Albizati developed a one-step pathway to cannabidiols with a higher efficiency than earlier methods.^{28} The monoterpenoid (R)-2,9-dibromocamphor (39) was postulated as a suitable starting point, as it would facilitate stereoselective arylation and allow for high regioselectivity. Methoxy-protected olivetol 36 was transformed into homocuprate 38 and reacted with dibromide 39 to form adduct 40 (Scheme 9). Keto-enol isomerisation, ring-opening and subsequent deprotection were conducted in one-pot using sodium naphthalenide and diethyl chlorophosphate in a high yield of 89%. The resulting enolphosphate 41 was demethylated and reduced with an excess of lithium in methyamine to form (-)-trans-CBD (5) and its monomethylated CBD counterpart in 35 and 43% yield, respectively.

2.8 Carveol for the synthesis of hexahydrocannabinols

Hexahydrocannabinols, such as (+)-machaeriol and (+)-machaeridol were prepared using aryl cuprates and chiral
C-ring oxidised cannabinoid 50 after acidic hydrolysis. Subsequent keto reduction, radical Barton–McCombie deoxygenation and acidic cyclisation reaction afforded (−)-trans-Δ⁸-THC (9).

2.9 α-Iodocyclohexenones

Kobayashi and co-workers revisited the preparation of (−)-trans-Δ⁸-THC and (−)-trans-CBD using aryl cuprates. The authors aimed to use α-iodocyclohexenones and a variety of resorcinols and unveil the arylation of these cyclohexenones (Scheme 12).47 They observed that 1,4-addition of bulky diaryl cuprates to enones was only successful in the presence of BF₃·OEt₂. The stereochemistry of α-iodocyclohexenone 52 facilitated the exclusive formation of trans-configured alylated products. After preparing α-iodoketone 52 from 51, the product was transformed into a dehalogenated enol Grignard, which opened synthetic strategies towards various 9′ derivatisations (not shown). Thereafter, the reactive enol Grignard intermediate was transformed into the corresponding enol phosphate 53. A sequence of methylation with MeMgBr in the presence of Ni(acac)₃, methyl ether deprotection using sodium ethanethiolate and cyclisation with ZnBr₂ afforded (−)-trans-Δ⁸-THC (7) in 28% yield. Notably, the 1,4-arylation to enones was reported and optimised a year later.48 This invaluable publication broadened the aryl substrate scope, led to new C-ring oxidised substrates, and the synthesis of α-iodocyclohexenones using (+)-β-pinene.

The successful stereoselective route to tetrahydrocannabinol using α-cyclohexenone was extended to (−)-trans-CBD (5), C-ring oxidised derivatives, and even alkenyl derivatives on the B-ring (Scheme 12).49 The route started from cyclohexenediol monoacetate 54, which appeared more versatile and allowed for more facile preparation of different α-cyclohexenones. The alkylation of monoacetate 54 was investigated using various reagents, additives, and catalysts. Isopropenylmagnesium bromide, zinc chloride, TMEDA and NiCl₂(tpp)₂ were shown to be most reactive, resulting in yields of up to 95% with minimal regioisomeric sideproducts. Subsequent Jones oxidation and iodination afforded 55. The consecutive coupling of diaryl...
cuprate 38, transformation to enol Grignard and final coupling to diethyl chlorophosphate afforded 56, the precursor of \((\pm)-\text{trans-CBD}\) (5).

The same group also published direct 1,4-arylation of diaryl cuprates onto non-iodinated cyclohexenones, promoted solely by boron trifluoride.\(^{50}\) This direct approach is considered advantageous for organic synthesis, but synthetically challenging because of the strongly stabilised boron enolate. To circumvent the low reactivity, MeLi was used to create the lithio enolate intermediates which are more reactive towards electrophiles. To demonstrate the principle, cyclohexenone 51 was activated with boron trifluoride, arylated with 38, transformed into the enolate using MeLi, and finally phosphorylated using diethyl chlorophosphate to afford 53.

### 3. Concerted approaches

The earliest approaches to the tetrahydrocannabinol scaffold were based on chiral pool syntheses and hence afforded enantiopure products, albeit often with formation of side products and in moderate to low yields. Alternatively, more elaborate synthetic pathways using asymmetric catalysts were employed, leading to high levels of enantioselectivity and additional access to unnatural THC enantiomers. Synthesis of unsaturated six-membered rings can be achieved using Diels–Alder \([4 + 2]\) cycloadditions, which have also been used to prepare cannabinoids.\(^{51}\) The earliest report of a Diels–Alder cyclisation in the preparation of CBD was by Korte, Dlugosch and Claussen in 1966.\(^{52}\) They conducted an intermolecular reaction with methyl vinyl ketone and dimethyl ether-protected 3-methyl-1,3-butadiene-olivetol. Without the use of any chiral auxiliary, CBD analogues were obtained.\(^{53,54}\)

The first asymmetric synthesis of the unnatural isomer \((\pm)-\text{trans-}\Delta^2\text{THC}\) (59) using a Diels–Alder reaction was reported by Evans, Shaughnessy and Barnes (Scheme 13).\(^{55}\) Key-step was the synthesis of the acetylated cycloadduct 57 on 50 mmol scale. Employing the enantiopure Cu(II)-bisoxazoline catalyst C1 led to an \(\text{endo}\)-transition state, resulting in a \(\text{trans}\)-configuration of crystalline cycloadduct 57.\(^{56-58}\) Transesterification into the benzyl ester, nuclophilic methylation and ester hydrolysis provided \(p\)-menth-1-ene-3,8-diol (58). Activation of the diol using \(p\)-TSA in the presence of olivetol (11) resulted in the cannabinol analogue which was cyclised using anhydrous ZnBr\(_2\) to obtain \((\pm)-\text{trans-}\Delta^2\text{THC}\) (59) in modest yield. Two years later, the key-step Diels–Alder cyclisation catalysed by C1 was further investigated in a mechanistic study with a significant larger substrate scope.\(^{59}\)

Subsequent seminal work of Avery and co-workers represented the first entry into a direct synthesis of the B- and C-rings of hexahydrocannabinol derivatives through a Diels–Alder cyclisation reaction (Scheme 14).\(^{60}\) Though this approach utilises \((S)\)-citronellal, a chiral pool terpinoid, a concerted double cyclisation approach was required to provide the tricyclic cannabinoid scaffold. Olefin 61 was obtained as a 1:1 mixture of diastereomers via the reaction of lithiated methoxymethyl ether (MOM) protected triol 60 with \((S)\)-citronellal. The following hetero-Diels–Alder cyclisation reaction surprisingly proceeded using 4% aqueous HCl in methanol at ambient temperature in moderate yield, while normally these reactions are conducted at elevated temperatures. The stereo-selectivity of the cyclisation reaction was explained by invoking intermediate 62, and afforded tricyclic scaffold 63. The subsequent phenolic triflation provided hexahydrocannabinol 64, suitable for further functionalisation. To ensure successful palladium-catalysed cross-coupling, Avery and co-workers first protected the phenolic position with a MOM group. Finally, a double cyclisation approach was required to provide the tricyclic scaffold 65. The following hetero-Diels–Alder cyclisation reaction surprisingly proceeded using 4% aqueous HCl in methanol at ambient temperature in moderate yield, while normally these reactions are conducted at elevated temperatures. The stereo-selectivity of the cyclisation reaction was explained by invoking intermediate 62, and afforded tricyclic scaffold 63. The subsequent phenolic triflation provided hexahydrocannabinol 64, suitable for further functionalisation. To ensure successful palladium-catalysed cross-coupling, Avery and co-workers first protected the phenolic position with a MOM group. Finally, a double cyclisation approach was required to provide the tricyclic scaffold 65.
hexahydrocannabinols (+)-machaeriol A and B (65 and 66, respectively) were obtained via Suzuki–Miyaura cross-coupling reactions and subsequent hydrolysis. Notably, Avery and co-workers did not comment on the stereospecificity, and specific rotations were not described. Nevertheless this pioneering work demonstrated the possibility to perform late-stage derivatisation of a cannabinoid scaffold.

The group of Sherburn demonstrated the synthesis of Δ⁹-THC with a late stage trans-selective Diels–Alder cyclisation. To be able to optimise this reaction towards the desired trans-product it was first investigated in silico. DFT calculations showed that the stereoselectivity is heavily influenced by conjugative effects between the diene and the aromatic ring substituents. The method was tested using aluminium tris(2,6-diphenylphenoxide) (ATPH) as Lewis acid catalyst on olefins 66 and 67. In contrast to earlier DFT calculations, ATPH dramatically inversed stereoselectivity to afford 68 and 69 in moderate trans-diastereoselectivity (Scheme 15). 69 was finally treated with methylmagnesium chloride and cyclised using ZnBr₂ to obtain (−)-trans-Δ⁹-THC.⁷²

Minuti and co-workers applied late stage hyperbaric Diels–Alder reactions giving rise to a variety of Δ⁸-THC derivatives.⁷⁴ The authors performed this reaction with six benzylidene acetone dienophiles (70) and two diene coupling partners (71a, 71b) promoted by mildly Lewis acidic conditions. This resulted in excellent yields, full trans-stereoselectivity, however, with additional formation of a regioisomeric product. The hyperbaric Diels–Alder reaction was applied to 1-olivetolic acid ((+)- and (−)-trans-configured bicyclic products 72 (Scheme 16, top route). The bicyclic intermediate was subsequently treated with MeMgBr, selectively deprotected with NaSMe, cyclised using ZnBr₂ and finally deprotected with additional NaSMe to afford rac-Δ⁸-THC. The authors resolved intermediate (±)-72 via [(S)-1-amino-2-(methoxymethyl)pyrrolidine (SAMP)] resolution, separated the diastereoisomers and followed with a subsequent Bronsted acid-mediated hydrazine hydrolysis. After obtaining both (R,S)-(−)-72 and (S,S)(+)-72 the intermediate products were transformed into the corresponding diastereomerically pure (−)- and (+)-trans-Δ⁸-THCs (9 and 76, respectively).

The influence of solvents in the Diels–Alder reaction and the preparation of Δ⁹-THC and derivatives was also demonstrated later. The previously used aryl alkene substituent 70 was transformed into dienophile 73 using a Wittig olefination, and reacted with alkenes⁶⁵ and alkynes⁶⁶ coupling partners in a hyperbaric Diels–Alder cyclisation reaction (Scheme 16, bottom route). The optimal results with methyl vinyl ketone (74a) or methyl acrylate (74b) were obtained in ethanol at 9 kbar providing olefin 75 in high cis-selectivity. This result is in strong contrast with the previous trans-selective Diels–Alder reaction (70 to 72). Therefore, the authors investigated a small variety of dienophiles, which were even less stereoselective. To demonstrate the applicability of the system, both (±)-cis- and (±)-trans-Δ⁸-THC were prepared via the Diels–Alder approach using the endo- and exo-cycloadduct of 75, respectively. Additionally, the single (−)- and (−)-enantiomers of trans-Δ⁸-THC were obtained via the SAMP-hydrazine route (59 and 7, respectively). Despite the laborious and low yielding approach, all stereoisomers of Δ⁸- and Δ⁹-THCs were obtained, thereby widening the scope to prepare synthetic cannabinoid derivatives.

4. Other approaches

A different approach to create the BC-ring system of cannabinoids proceeds via pericyclic reactions. The first intramolecular cyclisation reaction was shown by Childers and Pinnick. Using a Claisen rearrangement at room temperature, the authors obtained mixtures of (±)-trans-Δ⁵-THCs. Another approach was reported by Kirschleger and co-workers who published a one-step Bronsted acid-mediated double cyclisation giving rise to (±)-cis/trans-Δ⁵-THCs.⁶⁷ Alternatively, the synthesis of (±)-cis-11-nor-Δ⁹-THC-9-carboxylic acid was reported by Tius and Gu, using a similar key-step cationic cyclisation reaction.⁶⁸ Despite the low stereoselectivity of these concise synthesis routes, they offer fast access to new pharmacologically relevant cannabinoids and aid in the development of new potential drugs.⁶⁹
One of the most recent stereoselective cyclisations was the preparation of (−)-perrottetinene (86), a phenethyl-substituted (−)-cis–Δ^9-THC analogue. Kim and co-workers synthesised the cannabinoid using a key-step Ireland–Claisen rearrangement (Scheme 17A). Dihydropinosylvin (77) was regioselectively iodinated, and phenol protected to a furfuryl aryl iodide (78). The following Stille cross-coupling with tributylstannyl (S)-methyl vinyl carbinol (79) led to the isolation of 80, which was subsequently coupled with 5-methylhex-5-enoic acid (81) to obtain olefin 82. The following Ireland–Claisen rearrangement was conducted using LDA and TBSCI resulting in a silyl enol ether intermediate. The addition of HMPA enhances formation of the (Z)-configured silyl enol ether, which then rearranges through the chair-like transition state 83 to form the (S,R)-configured product 84 with high diastereoselectivity (>20:1). Precursor 85 was formed after esterification, treatment with MeMgBr and Grubbs olefin metathesis. (−)-Perrottetinene (86) was obtained after p-TSA-mediated cyclisation.

The Ireland–Claisen rearrangement by Kim and the molybdenum-catalysed allylic alkylation reaction by Trost and Dogra, inspired Leahy and co-workers to design a chemoenzymatic synthesis of (−)-trans–Δ^9-THC (7, Scheme 17B). The synthesis followed the methylation and formylation of olivetol 11 described by Trost and Dogra, but was then followed by an acetone aldol condensation reaction to afford the enone-substituted product 87. The enone was reduced using the Corey–Bakshi–Shibata oxazaborolidine, or via a sequence of NaBH₄ reduction, vinyl butyrate acylation in presence of Savinase 12T and basic hydrolysis to obtain 88. Ester 89 was produced by coupling 88 with 5-methylhex-5-enoic acid (83) in a Steglich DCC coupling. Critically, as opposed to Kim’s approach, the absence of HMPA resulted in formation of the (E)-configured silyl enol ether leading to the formation of chair-like intermediate 90. Ireland–Claisen rearrangement then afforded the (R,R)-configured product 91. Leahy demonstrated the versatility of 91 by transforming it into CBD or (−)-trans–Δ^9-THC. CBD was prepared using a sequence of Grubbs olefin metathesis, Wittig olefination with Ph₃P=CH₂ and final deprotection and methylation using MeMgl (not shown). (−)-trans–Δ^9-THC (7) was formed using an esterification, ring-closing metathesis, deprotection using MeMgl and a final Lewis acid-based cyclisation with ZnBr₂.

Previously, the SN2 arylation of α-iodocyclohexanone was demonstrated by Kobayashi, in the preparation of (−)-trans–Δ^9-THC and -CBD. Alternatively, Zhou and co-workers later found

Scheme 17 Ireland–Claisen rearrangement by Kim and co-workers to afford (−)-perrottetinene (86), and the inverted Ireland–Claisen rearrangement by Leahy and co-workers to afford (−)-trans–Δ^9-THC (7).
that α-iodocyclohexanones are effective candidates in Suzuki-Miyaura cross-coupling reactions (Scheme 18).73 The key step in the publication is a high-pressure asymmetric ketone reduction with ruthenium catalyst (S,R,R)-C2. Following one of their earlier publications,74 the authors set out to investigate the reduction of rac-2,6-dimethylphenylcyclohexanone, which appeared synthetically challenging due to the steric hindrance caused by the two ortho-methoxy groups. Fortunately, by changing one methoxy into a fluoride atom, the reaction proceeded more efficiently. The chemistry was applied in the synthesis of both (−)-trans-Δ⁹- and Δ⁸-THC isomers. Initially, the Suzuki-Miyaura cross-coupling of α-iodocyclohexanone 94 with boronic acid 93 and hydrogenation on Pd/C provided racemic 95. The crucial high-pressure asymmetric ketone reduction with (S,R,R)-C2 and concomitant racemisation were applied to substrate 95 to afford (−)-cis-configured alcohol 96. To obtain the target trans-configured cannabinoid scaffolds, a sequence of saponification, Wittig olefination and basic olefination was employed. From product 96 still 10 steps were required to produce both the (−)-trans-Δ⁹- and Δ⁸-THC isomers, in a total yield of 43 and 51%, respectively.

The group of Lupton has investigated an NHC-catalysed (4 + 2) annihilation reaction to a biologically relevant molecules, but there is still a need for more efficient access to new cannabinoid derivatives, as many examples are based on multistep organic transformations. A recent effective approach to obtain synthetic cannabinoids was published by Westphal and co-workers, who demonstrated the late stage derivatisation of a functionalised (−)-trans-Δ⁹-THC scaffold.77 Currently, this late-stage approach is attractive to efficiently prepare synthetic cannabinoid derivatives and is a hot topic in the scientific field.78–80 Alternatively, the groups of Carreira and Studer have disclosed stereodivergent syntheses of cannabinoïds, affording a variety of cannabinoids by varying catalyst ligands.81–84 This latter approach may provide fast access into more stereodiverse cannabinoid scaffold 99, demonstrated by Ametovski and Lupton.

5. Outlook and conclusion

In conclusion, we have outlined several chiral pool and asymmetric catalytic approaches to THC and derivatives. Chiral pool approaches generally provide a fast entry into cannabinoids, but the final product mixtures are challenging to purify, having a limited substrate scope and may lack regioselectivity. Alternatively, asymmetric catalysis excels in substrate scope, high regio- and stereoselectivity, but requires more elaborate synthesis routes.

The published approaches are generally well applicable and already provided several potential drug candidates. However, more research is required to afford straightforward and efficient access to new cannabinoid derivatives, as many examples are based on multistep organic transformations. A recent effective approach to obtain synthetic cannabinoids was published by Westphal and co-workers, who demonstrated the late stage derivatisation of a functionalised (−)-trans-Δ⁹-THC scaffold.77 Currently, this late-stage approach is attractive to efficiently prepare synthetic cannabinoid derivatives and is a hot topic in the scientific field.78–80 Alternatively, the groups of Carreira and Studer have disclosed stereodivergent syntheses of cannabinoïds, affording a variety of cannabinoids by varying catalyst ligands.81–84 This latter approach may provide fast access into more stereodiverse cannabinoid derivatives.

Without question, organic synthesis is most suitable for the preparation of biologically relevant molecules, but there is still a need for more efficient preparation techniques. At this moment, only one continuous flow process has been applied in the synthesis of ortho-substituted cannabinoids.84 Generally, the current preparation of cannabinoids and pharmaceuticals such as Nabilone occurs via batch chemistry,27,68,85 but may well be changed to flow processes because of the excellent scalability, safety and efficiency.

Besides the synthetic effort to obtain facile access to new cannabinoid derivatives, another goal is to elucidate the bio-
logical mechanisms of the endocannabinoid system. Despite numerous publications in literature, it is still unclear how the CB1, CB2 (and possibly CB3) receptors interact in various diseases.4–9 The receptors have shown to play a key role in various health conditions such as nausea during chemotherapy, (neuropathic) pain, multiple sclerosis and Tourette syndrome.

The preparation of new and specialised cannabinoids might help to further elucidate the biological mode of action of these receptors and enable treatment of these and other diseases. Clearly, the multidisciplinary scientific field of cannabinoids, with the continuous development of new synthetic methods, remains to hold considerable promise for the development of novel pharmaceuticals.

Conflicts of interest
There are no conflicts to declare.

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