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Guillaume Arcile, Pascal Retailleau, Jamal Ouazzani, Jean-François Betzer. Total Synthesis of the fungal metabolite Trienylfuranol A via Nucleophilic Diastereodivergent Additions to Oxocarbenium Ions. *European Journal of Organic Chemistry*, Wiley-VCH Verlag, 2021, 2021 (13), pp.2050-2054. 10.1002/ejoc.202100265 . hal-03215484

HAL Id: hal-03215484

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Submitted on 3 May 2021

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Total Synthesis of the fungal metabolite Trienylfuranol A via Nucleophilic Diastereodivergent Additions to Oxocarbenium Ions

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Abstract: Herein, we describe the first total synthesis of trienylfuranol A, a fungal triene-substituted tetrahydrofuran metabolite. The stereoselectivity of the chiral center bearing the trienyl side chain was diastereodivergently controlled by the addition of nucleophilic species on substituted γ -butyrolactone. Remarkably, the C-nucleophilic species or hydride addition onto oxocarbenium intermediate leads to the opposite diastereoselectivity reported for Kishi or Woerpel models. The use of Hantzsch ester (HEH) as organic hydride donor has enabled us to access to the desired stereochemistry. The total synthesis of trienylfuranol A was achieved in 8 steps from acetaldehyde and pyruvic acid.

We previously isolated the sporochartine family of natural compounds from the fungus *Hypoxylon monticulosum* CLL 205,^[1] now referred as *Hypomontagnella spongiphila*.^[2] According to the structure of the sporothriolide 1^[3] and the trienylfuranol A 2,^[4] we suggested that sporochartines A-D 3a-d might derive from a Diels-Alder type reaction between 1 and 2 (Figure 1). The cultivation of *Hypoxylon monticulosum* CLL 205 offers significant quantity of 1 and 3a-d, while 2 is only detected in very low amount. In the context of our ongoing research program on biosynthetic pathway of sporochartine 3a-d natural products and in order to elucidate the mechanism of reaction between 1 and 2, we engaged the total synthesis of trienylfuranol A 2; sporothriolide 1 is easily isolated in gram-scale from the cultivation broth.

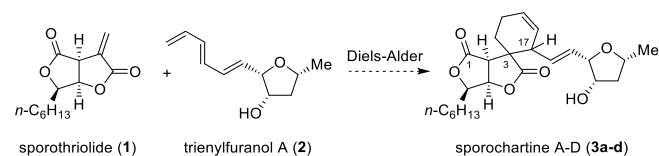
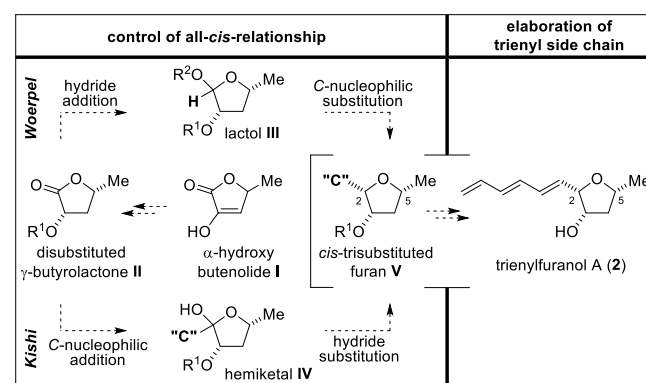


Figure 1. Plausible biosynthetic pathway of sporochartines through Diels-Alder cycloaddition between sporothriolide and trienylfuranol A.

In order to provide an efficient synthetic pathway of trienylfuranol A 2, and in respect to the stability of this natural metabolite which is highly susceptible to polymerization or degradation upon synthesis or purification,^[4] we considered to perform the synthesis involving two major milestones. First, we planned to synthesize a trisubstituted and functionalized furan V, bearing the three stereogenic centers in all-*cis*-relationship, and secondly the introduction of the terminal trienic side chain (Scheme 1). The α -

hydroxy- γ -butenolide I can be smoothly hydrogenated over palladium to produce the disubstituted- γ -butyrolactone II as a single diastereoisomer. This lactone II should be converted into *cis*-trisubstituted furan V. This will be achieved through the introduction of a functionalized side chain followed by a reduction at the same site, operations that can be performed in any order. The last substitution stage, which involves the formation of oxocarbenium intermediate, set up the C-2 stereogenic center. Even if stereoselective addition of nucleophilic species on oxocarbenium ions in furanoside series is well documented, it remains not fully predictable.^[5] The stereochemical outcome of addition reactions on furanosyl oxocarbenium is dictated by stereoelectronic factors and highly dependent on the substrate structures.^[6] For polysubstituted substrates, the conformational stability and reactivity of oxocarbenium ion is controlled by the interplay between the ring substituents and their relative effects.

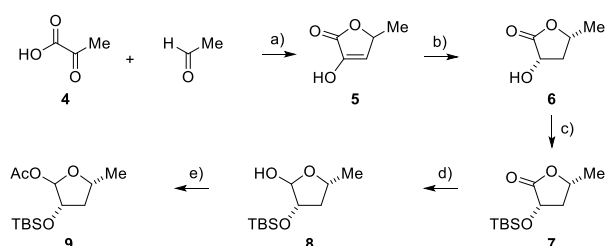


Scheme 1. Retrosynthetic pathway of trienylfuranol A (2).

Indeed, in contrast with six-membered ring cation series, a predictive stereochemical model of addition reactions on five-membered ring oxocarbenium ions is not available.^[7] The stereoselectivity remains elusive, difficult to predict and occurred often with counterintuitive selectivity. As we are not able to anticipate the cumulative stereochemical effects of the 1,2- and 1,4-stereoiduction of the disubstituted γ -butyrolactone II, we have considered two approaches to introduce the functionalized side chain, precursor of the trienyl moiety. The first one is based on a well-established C-nucleophilic substitution on lactol III,

which is obtained by hydride reduction of lactone **II**. This general approach has been studied by several groups with different nucleophiles and various Lewis acid. Woerpel and co-workers have proposed stereochemical models to account stereodirecting substituent effects on five- and six-membered ring oxocarbenium ions and to explain the stereoselective formation of C-furanosides^[8] and C-pyranosides^[9] series, respectively. The second approach, developed by Kishi and co-workers for the formation of C-glycopyranoside, consists of the same in a reverse sequence.^[10] First, C-nucleophile species was added on lactone type compound **II** to lead to hemiketal intermediate **IV**, which undergoes subsequent deoxygenation by treatment with hydrosilane and boron trifluoride etherate. This formation of C-glycoside was extended to furanoside series^[11] and commonly applied to the formation of C-nucleosides.^[12]

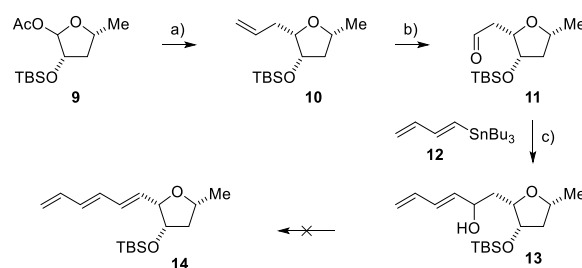
We first prepared the disubstituted γ -butyrolactone **7** and its reduced O-acetyl lactol **9** counterpart (Scheme 2). The synthesis of these compounds is based on previous reports^[13] where pyruvic acid **4** was condensed with acetaldehyde through a gram-scale acid-catalyzed aldol/crotonisation/lactonisation sequence to provide the α -hydroxy- γ -butenolide **5**.^[14] This unsaturated lactone **5** was reduced under hydrogenation over 10% Pd/C in EtOAc to set up the hydroxyl function and methyl substituents in *cis* relative configuration *via* a 1,3-induced asymmetric hydrogenation.^[15] The disclosed secondary alcohol of **6** was protected as its TBS ether in 93% yield and the protected lactone **7** was reduced with DIBAL-H to yield the lactol **8** and acetylated to provide the more stable O-acetyl lactol **9** in good yields.



Scheme 2. Preparation of disubstituted γ -butyrolactone and their reduced lactol counterparts. Reagents and conditions: a) H_2SO_4 conc., 0 °C to RT, 30 min., 31%. b) H_2 , Pd/C 10%, EtOAc, RT, 24 h, 70%. c) TBSOTf, 2,6-di-*tert*-butylpyridine, CH_2Cl_2 , 0 °C to RT, 1.5 h, 93%. d) DIBAL-H, CH_2Cl_2 , -78 °C to -15 °C, 2.5 h, 41%. e) Ac_2O , DMAP cat., CH_2Cl_2 , 0 °C, 1.5 h, 85%.

We started the synthesis of the targeted *cis*-trisubstituted furan **V** from the O-acetyl substrate **9** by a homologation reaction with allyltrimethylsilane as C-nucleophile in presence of Lewis acid.^[16] We have used SnBr_4 as Lewis acid which is widely employed by Woerpel and co-workers for five-membered-ring oxocarbenium ions^[8a] and used recently by Ghosh and co-workers for the synthesis of amphirionin-4^[17] and monocerin,^[18] while other Lewis acids such as $\text{BF}_3\cdot\text{OEt}_2$, TiCl_4 or TMSOTf are described as less efficient. In our hands, with these conditions, the nucleophilic substitution of O-acetyl moiety of compound **9** with allyltrimethylsilane provided to the desired allylated compound **10** in good yield with an excellent α -selective substitution as previously observed by Ghosh and co-workers (Scheme 3).^[17] The oxidative cleavage of the corresponding alkene function by ozonolysis leads to the aldehyde **11** in 75% yield. Aldehyde **11** was reacted with (*E*)-1-lithio-1,3-butadiene, generated from the corresponding tributylstannane derivative **12**^[19] by transmetalation,^[19a] to form the (*E*)-butadienylated adduct **13** in

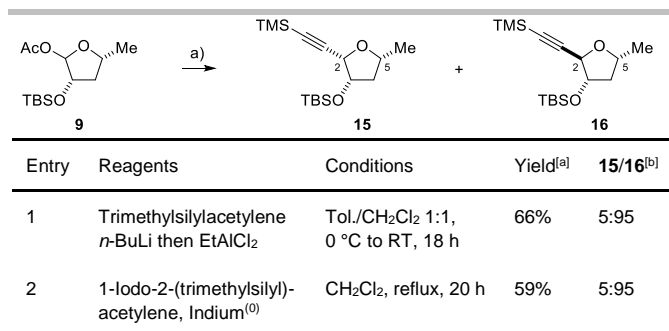
an optimized yield of 62% yield and as a mixture of epimers in a 1:1 ratio. However, the dehydration of 1,3-butadienyl carbinol **13** into the targeted compound **14** with the trienic side chain was more difficult than expected. All attempts of direct dehydration of **13** or reactions involving the introduction of better leaving groups failed. We have only observed the consumption of starting material and extensive formation of decomposition products. This would suggest that elimination reaction conditions are not compatible with the unusual and very sensitive terminal trienyl moiety.^[4] We consequently designed an alternative mild synthetic pathway to preserve the trienyl moiety.



Scheme 3. Nucleophilic substitution with allyltrimethylsilane. Reagents and conditions: a) Allyltrimethylsilane, SnBr_4 , CH_2Cl_2 , RT, 30 min., 80%. b) O_3 , CH_2Cl_2 , -78 °C, 5 min. then PPh_3 , -78 °C to RT 16 h, 75%. c) (*E*)-Buta-1,3-dien-1-yltributylstannane **12**, *n*-BuLi, THF, -78 °C to RT, 16 h, 62%.

We turned out to an approach which implies the use of the same (*E*)-buta-1,3-dien-1-yltributylstannane **12** for the construction of the trienic unit, in milder conditions by Pd-catalyzed cross-coupling reactions. To achieve this objective, we plan to prepare a furan **V** bearing a functionalized vinyl unit. This partner for the targeted cross-coupling reaction could be obtained from an alkyne derivative. Thus, in place of allyltrimethylsilane for the nucleophilic substitution on O-acetyl compound **9**, we have considered alkynyl species as C-nucleophile (Table 1). To realize this nucleophilic substitution, we intend to use organoalane reagents, successfully used for the introduction of the alkynyl groups at the anomeric position for nucleoside synthesis.^[20] In our case, the most suitable organoalane reagent is ethyl[trimethylsilyl(ethynyl)]aluminum chloride, prepared *in situ* by transmetalation of lithio-ethynyltrimethylsilane with ethylaluminum dichloride. An excess of this reagent was used to assist the departure of acetyl moiety on the one hand, and to perform the addition on the generated oxocarbenium on the other.^[21] The reaction of O-acetyl compound **9** with the *in situ* prepared organoalane reagent furnished cleanly the alkynylated product in 66% yield. However, to our great surprise we have observed a highly β -selective reaction, in 95:5 C-2 epimeric mixture in favor of the β -isomer **16**. The relative stereochemistry of this major β -isomer **16** and minor α -isomer **15** were independently assigned on the basis of the 2D NOESY spectra. For the β -isomer **16**, key NOE cross-peaks was observed between H-2 and CH_3 -5 while the desired minor α -isomer **15** display a correlation between H-2 and H-5 (see supporting information). Other conditions using indium-mediated alkynylation under Barbier conditions^[22] lead to the same outcome in terms of stereoselectivity.

Table 1. Nucleophilic substitution of **9** with alkynyl species.

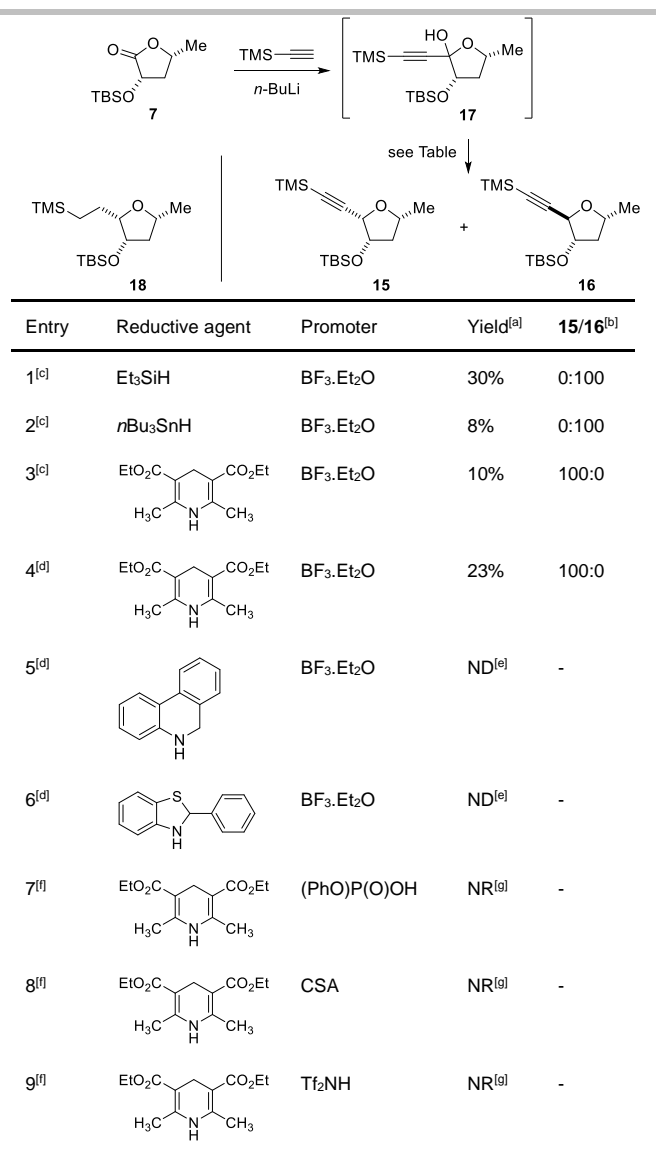


[a] Yield of isolated product. [b] Determined by ¹H NMR analysis of the crude reaction mixtures.

As the undesired β-isomer **16** was obtained during the previous Woerpel approach, we turned out to the complementary Kishi sequence which consists to introduce firstly the alkynyl moiety and to reduce the resulting hemiketal **IV** in the last step.^[10] The lactone **7** was treated with the *in situ* prepared lithium(trimethylsilyl)acetylide at -45 °C to -25 °C for two hours to provide the hemiketal **17**, isolated as a crude product in 95% yield (Table 2). All the trials to purify hemiketal **17** induced a significant fall of material and gave lower quality NMR spectra. The following reductive-deoxygenation of the crude hemiketal **17** was initially performed by the classical triethylsilane as reducing agent with BF₃.Et₂O as promoter. Unfortunately, once again the undesired β-isomer **16** was produced (Table 2, entry 1). Other hydrosilanes as hydride sources or the use of tributyltin hydride also led to the same β-isomer **16** in lower yields (entry 2). With these disappointing results, we turned to the use of organic derivatives as reducing agent. First we have performed the reaction with Hantzsch ester (HEH),^[23] a simplest NAD(P)H mimics widely used as reductant in the hydrogenation of unsaturated bonds and successfully used to trap *in situ* oxonium species.^[24] To our delight, even if the yield is low, the reaction with hemiketal **17** and HEH as a reducing agent for oxocarbenium reduction provided the desired α-isomer **15** (entry 3). This diastereodivergence for triethylsilane and HEH was previously observed by MacMillan and co-workers during the synthesis of C-nucleoside compounds in ribose series.^[25] Taking into account the low stability of oxocarbenium intermediate and working at lower temperature, the desired α-isomer **15** was isolated in 23% yield for two steps from lactone **7** (entry 4). The use of other synthetic NAD(P)H analogue like 9,10-dihydrophenanthridine^[26] (entry 5) or benzothiazoline^[27] (entry 6) lead to a complex mixture of several compounds, where the desired reduced compounds **15** or **16** were not observed. Switching from Lewis to Brønsted acid to generate the oxocarbenium ion intermediate turned out to be detrimental for reactivity. The use of diphenylphosphate^[28] or more acidic promoters such as camphorsulfonic acid or triflimide did not lead to any reaction (entries 7-9).^[29]

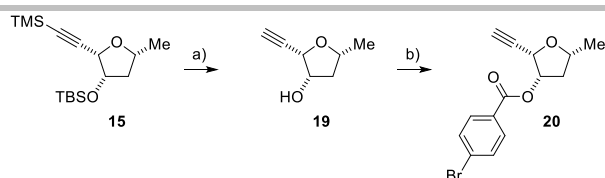
On large scale synthesis, small quantity of over-reduced compound **18** was observed and can be cleanly isolated. The hydrogenation of alkyne function to alkane derivatives with HEH as hydrogen donors was reported under palladium catalysis^[30] but, to the best of our knowledge, this result represents the first example of alkyne reduction with HEH under metal-free conditions.

Table 2. Introduction of alkynyl moiety using Kishi's protocol.



Reagents and conditions for the first step: Trimethylsilylacetylene, *n*-BuLi, -45 °C to -25 °C, 2 h., 95% (crude product). [a] Overall yield of isolated products for the two steps. [b] Determined by ¹H NMR analysis of the crude reaction mixtures. [c] Reductive agent (4 equiv), promoter (4 equiv), CH₂Cl₂, -40 °C, 2 h. [d] Reductive agent (2.5 equiv), promoter (2.5 equiv), CH₂Cl₂, -60 °C to -40 °C, 2 h. [e] ND=product **15** or **16** not detected. [f] Reductive agent (2 equiv), promoter (0.2 equiv), 4 Å molecular sieves, Toluene, RT, 16 h and 50 °C, 6 h. [g] NR=no reaction.

As the nature and configuration of functional groups on furan ring determines the stability and reactivity of oxocarbenium ions intermediate, but also the conformation of the formed product, we decided to assign unambiguously the relative stereochemistry by X-ray crystallography. The silyl protecting groups of **15** were removed by TBAF treatment to give the corresponding deprotected compound **19** (Scheme 4). The free secondary alcohol was derivatized into *p*-bromobenzoyl ester **20** and recrystallization of this solid compound from slow evaporation of its solution in methanol yielded single crystals suitable for X-ray crystallography. The crystallographic analysis showing an all-*cis* orientation of the substituents on the furan ring and confirming the relative stereochemistry assignment deduced from NOE correlations (Figure 2).^[31]



Scheme 4. Derivatization of α -isomer **15** as its *para*-bromobenzoate **20**. Reagents and conditions: a) TBAF, THF, RT, 2 h., 28%. b) *para*-bromobenzoic acid, DCC, DMAP, CH₂Cl₂, 0 °C to RT, 18 h, 45%.

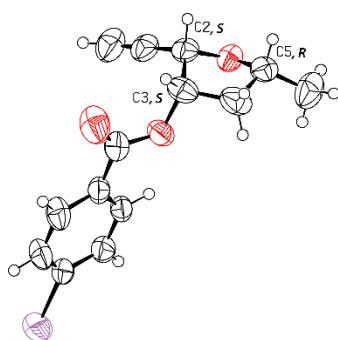
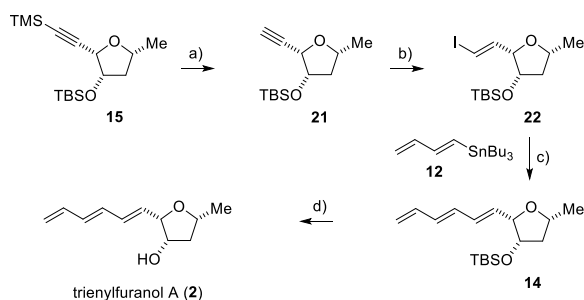


Figure 2. X-ray crystal structure of **20**.

After carrying out the synthesis of desired functionalized furan **15**, the last stage to complete the synthesis of trienylfuranol A involve the introduction of the trienyl side chain. For this purpose, the TMS group of **15** was selectively removed by treatment with methanol under basic conditions to afford the terminal alkyne **21**, which was converted into (*E*)-vinyl iodide **22** by a hydrozirconation-iodination sequence with Schwartz reagent^[32] prepared in situ from Cp₂ZrCl₂ and DIBAL-H^[33] and then followed by iodine treatment (Scheme 5). A Stille cross-coupling reaction^[34] between vinyl iodide **22** and the previously prepared dienylstannane **12**^[19] proceed under mild conditions at room temperature to afford the expected terminal triene **14** in satisfactory 71% yield. Subsequent desilylation of TBS protecting group with TBAF produced the targeted trienylfuranol A (**2**).



Scheme 5. Completion of the total synthesis of trienylfuranol A (**2**). Reagents and conditions: a) K₂CO₃, MeOH, RT, 3 h., 63%. b) Cp₂ZrCl₂, DIBAL-H, 0 °C to RT, 50 min., then I₂, -78 °C to RT 2 h., 63%. c) (*E*)-Buta-1,3-dien-1-yltributylstannane **12**, PdCl₂(CH₃CN)₂, RT, 3 h., 71%. d) TBAF, THF, 0 °C to RT, 3 h., 43%.

In order to identify the natural isomer of trienylfuranol A **2** in this racemic mixture, the two enantiomers were separated by Supercritical Fluid Chromatography (SFC) on a semi-preparative chiral column, and their $[\alpha]_D^{20}$ measured (see supporting

information). Among the two chromatographic peaks, the natural isomer has a retention time of 5.6 min and a specific optical rotation $[\alpha]_D^{20} +1.7$ (*c*=0.10, CH₃CN) [reported +1.9 (*c*=1.10, CH₃CN)]. The unnatural antipode, with a retention time of 12.3 min, gives an opposite optical rotation $[\alpha]_D^{20} = -2.0$ (*c*=0.10, CH₃CN).

In conclusion, the first total synthesis of trienylfuranol A was achieved in 8 steps, from acetaldehyde and pyruvic acid. The synthetic route was adapted to provide a concise and efficient approach for the synthesis of this natural metabolite taking into account of the very sensitive terminal trienyl moiety. Moreover, we have developed a diastereodivergent approach for the stereogenic center bearing this trienic side chain. Starting from γ -lactone as substrate, the addition reactions of alkyne nucleophilic species and hydride reagent on the carbonyl moiety allow access, with high selectivity, both *cis* and *trans* diastereoisomers, which are unambiguously assigned by X-ray studies. The ordering of the reaction sequence but also and above an appropriate choice of the reducing agent allow to ensure the stereocontrol. A separation by Supercritical Fluid Chromatography on chiral stationary phase allow the obtaining and characterization of the enantiomerically natural pure (+)-antipode. While we recently contributed to the elucidation of the biosynthesis pathway of the sporothriolide **1**,^[35] the biosynthesis of the trienylfuranol A **2** and sporochartines **3** remains to be elucidated, in particular the involvement of a Diels-Alder type reaction. We will benefit from the availability of the synthesized trienylfuranol A and sporothriolide to engage chemo-enzymatic reactions and establish unambiguously the sporochartines formation pathway. This work is ongoing in our laboratory.

Acknowledgements

SFC separation of compound **2** enantiomers was performed by the HPLC platform of ICSN, CNRS, Gif-sur-Yvette, France. The authors are grateful to Nathalie Hue and Vincent Steinmetz.

Keywords: Natural products • Total synthesis • Diastereoselectivity • Hydrides • Reduction

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- [31] Crystallographic data for the compound **20** have been deposited at the CCDC, Cambridge, UK, with the deposit number 2052218. The supplementary crystallographic data can be obtained free of charge from The Cambridge Crystallographic Data Centre via URL www.ccdc.cam.ac.uk/data_request/cif.
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