A NEW TACTIC FOR TOCOPHEROL SYNTHESIS USING INTRAMOLECULAR BENZYNE TRAPPING BY AN ALCOHOL

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Abstract – A formal total synthesis of (S)-α-tocopherol, the major component of natural Vitamin E has been achieved using intramolecular benzyne trapping as a key step to form the chroman ring. The synthesis also features an efficient new method for benzotriazole N-amination using an oxaziridine; chiral, non-racemic intermediates are generated using asymmetric dihydroxylation.

This paper is dedicated to Professor Lutz Tietze on his 75th birthday, with all best wishes.

INTRODUCTION

α-Tocopherol 1 is the major component of naturally occurring Vitamin E, the remaining components of which generally consist of less methylated analogues and examples containing alkenes in their side chains.\(^1\)

![Figure 1. α-Tocopherol, the major component of the natural Vitamin E mixture](image-url)

The acetate derived from tocopherol is manufactured on a multi-ton scale by coupling trimethylhydroquinone and isophytol followed by acetylation and is used primarily as an anti-oxidant in a very wide variety of foods, pharmaceuticals, cosmetics, indeed almost any product which can suffer from the deleterious effects of oxidation upon prolonged storage.\(^2\) More recent investigations...
have revealed that tocopherol is distinguished by its ability to inhibit oxytosis, 5-lipoxygenase and phospholipase A₂ along with various kinases as well as being capable of activating protein phosphatase 2A and of gene regulation.³ It has been strongly suggested that vitamin E should be studied much more seriously, in view of these discoveries.³ The anti-oxidant activity usually involves generation of a phenoxide radical followed by quinone formation and concomitant opening of the chroman ring.

Such anti-oxidant activity is not confined to tocopherol in this area: the analogous, commercially available acid, trolox 2, also obtained from trimethylhydroquinone but together with methacrylic acid and formaldehyde, displays similar anti-oxidant properties and is differentiated from the very lipophilic vitamin E components by reason of its highly polar carboxylic acid group.⁴ Contrasting applications as a preservative as well as a precursor to the tocopherols mark this compound out as an important member of this series, especially at it can be readily resolved.

![Figure 2. Trolox, a non-natural but powerful anti-oxidant](image)

Synthetic activity in this area has employed many of the known methods for the asymmetric formation of C-O bonds. Once achieved, many earlier syntheses utilised the ‘Cohen’ approach outlined in Scheme 1.⁵ Thus, a phenol or hydroquinone 3 carrying a suitably functionalised side chain, either protected or unprotected as shown, is oxidized to the corresponding quinone level and cyclised by mild acid treatment to the acetal 4, subsequent reduction of which then leads to the target chroman-2-methanol 5. There are many versions of this strategy, in which the order of events may be altered or an intermediate related to acetal 4 not isolated. Clearly, there are then many possibilities for the elaboration of a suitable side chain.⁶
Unsurprisingly, the Sharpless epoxidation\textsuperscript{2} and AD-mix methods\textsuperscript{8} have both been employed in the asymmetric synthesis of tocopherol. More modern approaches have usually focussed instead on formation of the chroman C-O bond from similar chiral, non-racemic precursors, often by intramolecular attack of the phenol group onto an activated intermediate derived from diols 3 or by a similar attack but onto an allylic alcohol or related structure, activated by a chiral catalyst, which is often palladium-based\textsuperscript{3,10} but can also be a gold\textsuperscript{11} or ruthenium species.\textsuperscript{12} Various enzymatic resolutions and related methods have also proven useful in preparing enantiomerically enriched samples of the tocopherols,\textsuperscript{13} which have also been obtained using a biomimetic approach.\textsuperscript{14}

RESULTS AND DISCUSSIONS
The setting for our work in this area was the discovery that lateral deprotonation of the 1-aminobenzotriazole derivative 6 provides a quantitative conversion into the dianion 7, which could then be reacted with a range of electrophiles to give generally excellent yields of a range of analogues.\textsuperscript{15} In the light of the classic work by Rees’ group,\textsuperscript{16} this then provided an approach to 3-substituted benzenes, following N-deprotection and amine oxidation. One particular finding was that such benzenes could be effectively trapped in an intramolecular fashion by a pendant hydroxyl group (cf. 8) to give iodo-chromans 9, as well as other bicyclic derivatives (Scheme 2).\textsuperscript{17}

![Scheme 2. Lateral metallation - benzyne trapping by an alcohol as an approach to chromans](image)

In general, alcohols have not been used in this way previously,\textsuperscript{18} probably because benzyne generation was achieved from a halobenzene or a 1,2-dihalobenzene using halogen-metal exchange, which inevitably meant that the hydroxyl group would be deprotonated; it seems that such alkoxides, being ‘hard’ nucleophiles, are incompatible with the much softer benzenes. In the final oxidation step of the present method (Scheme 2), the alcohol is not ionized: the original Rees methods used either lead(IV) acetate or N-bromosuccinimide for this purpose; we found that N-iodosuccinimide (NIS) gave enhanced yields.\textsuperscript{12} Of course, a potential bonus of this methodology is the incorporation of the iodine atom, which lends itself to additional homologation by a plethora of coupling and other reactions.
Hence, this method, overall, could be regarded a having some generality for the synthesis of 1,2,3,4-tetrasubstituted benzenes.

We wished to demonstrate that it was possible to apply this methodology to the synthesis of the tocopherol precursor alcohol 10, with the particular aim of demonstrating that such a benzyne-based approach could be successfully used to synthesise a more highly substituted target. In the light of the foregoing results, the structure of alcohol 10 leads back to two, seemingly readily available, starting materials. (Scheme 3). Trapping of the key benzyne 12 was expected to be regiospecific in favour of chroman formation and also result in incorporation of a halogen atom (X = Br or I) into the chroman 11, which should allow addition of the final methyl group. The key benzyne 12 could then be generated from 1-aminobenzotriazole 13, or its regioisomer, itself a product of a Sonogashira coupling between the fully-substituted benzene 14 and the alkyne-diol 15, followed by a double hydrogenation of both the nitro and alkyne functions and triazole formation. We chose not to use our dianion methodology (Scheme 2), as trapping of such an intermediate (cf. dianion 7) with saturated alkyl halides or epoxides was not generally very efficient and it was also likely to be a synthetic challenge to form a single regioisomer of a suitably substituted aminobenzotriazole. In addition, other model reactions suggested that there would be a lack of regioselectivity in metallations of more highly substituted N-Boc-aminobenzotriazoles. The need for and the nature of the protecting groups in the alkyne diols 15 and subsequent derivatives were also far from clear.

![Scheme 3. Retrosynthetic analysis for a possible benzyne-based approach to chroman-2-methanol 10](image)

Synthesis of the fully-substituted benzene 18 began with an efficient nitration of commercial 2,6-dimethylanisole, given that the temperature was always kept below 70 °C (see Experimental), followed by reduction of the resulting 4-nitro derivative 16a to the corresponding aniline 16b using transfer hydrogenation and acetylation to give acetanilide 16c in excellent overall yields (Scheme 4).
Subsequent iodination using $N$-iodosuccinimide in acetic acid\textsuperscript{21} worked well and the sequence was completed by a slow but efficient nitration of the resulting iodide 17 to install all the required functionality. Any variation in the order of this sequence gave (much) lower returns, as expected.

The second component, the alkyne-diol 20a was obtained from enyne 19, readily obtained by acid-catalysed dehydration of the acetylene-acetone adduct,\textsuperscript{22} by asymmetric dihydroxylation using AD-mix β.\textsuperscript{22} While we were unable to achieve useful separations of the diol enantiomers using a variety of GC and HPLC methods, the derived benzoate 20b proved readily separable by GC; unhappily, the enantiomeric enrichment was below 10\% (Scheme 5). This is perhaps not too surprising, as terminal alkenes are often not good substrates for this reaction and an alkyne group is hardly a stericly dominant substituent. However, having secured a supply of the diol 20a, we chose to continue with this essentially racemic material, to hopefully establish suitable conditions to enable completion of the projected synthesis.

The first of these steps was Sonogashira coupling between the iodo-anisole 18 and the alkyne-diol 20a. This proved to be quite slow and required relatively large amounts of the catalysts to achieve even this rate of reaction, despite the activating effect of the nitro group, which we chose to retain for this very reason, and, to a lesser extent, the $N$-acetyl group. Happily, after a 24 h reflux in THF, a 77\% isolated yield of the desired coupled product 21 was obtained (Scheme 6). Similarly, some experimentation
was required to establish optimised conditions for the simultaneous reduction of both the nitro and alkyne groups: eventually, it was found that prolonged exposure to Pearlman’s catalyst (20% palladium hydroxide on carbon) at ambient temperature secured an essentially quantitative yield of the desired aniline 22. Under more vigorous conditions, especially when a reduction was heated, we also noticed the formation of a by-product, which remained unidentified at this stage.

The same unidentified by-product was again evident in initial attempts to form the triazole ring by diazotization of aniline 22; eventually, its formation was much reduced by using three equivalents of sodium nitrite to carry out the diazotization-cyclisation step, which then delivered >70% yields of the benzotriazole 23a from which the acetyl group was readily removed by mild, base-catalysed hydrolysis to give the free benzotriazole 23b. The excellent yield was only realised when a non-aqueous work-up was used, due to the water solubility of this polar product. Unexpectedly, its NMR spectra, when obtained for a solution in CD$_3$OD, showed the presence of essentially a single tautomer, whereas in CDCl$_3$, two forms were observed, in a 2:1 ratio. Possibly, the tautomer shown, 23b, may be favoured by hydrogen bonding between the new amino group and the pendant hydroxyls. A small amount of the by-product was also isolated during this stage: a small sample showed that it did not contain a carbonyl group (IR), but did have a $^{13}$C resonance for a quaternary carbon at ~ 155 ppm, together with five methyl singlets in its $^1$H NMR spectrum, but in slightly differing positions than related derivatives 21-23a (Scheme 6). MS data (APCI) indicated its formation by loss of water: a probable molecular ion at m/z 293 (M + H) contrasted with that for the N-acetyl benzotriazole 23a at m/z 311 (M + H).

We deduced that this was the benzimidazole 24, formed by intramolecular cyclization of the N-acetyl benzotriazole 23a, either when more vigorous hydrogenation conditions were used or when diazotization was relatively slow, allowing such an (acid-catalysed) cyclisation to compete with
benzotriazole formation (Figure 3). Hence, relatively mild conditions or excess reagents were essential to secure high yields from these two steps [b) and c), Scheme 6].

The next step was electrophilic N-amination.\textsuperscript{23} In previous work,\textsuperscript{15} we had optimised the original Campbell and Rees method,\textsuperscript{24} which used hydroxylamine-O-sulfonic acid as the source of an electrophilic amino group. However, the relatively delicate nature of benzotriazole 23b meant that the method failed in this case. Fortunately, a mild alternative was the series of N-alkoxycarbonyloxaziridines, which can deliver an alkoxycarbonylamino group to a free amine function, thereby forming an N-N bond.\textsuperscript{25,26} A range of such derivatives have been assessed, all of which carry electron-withdrawing groups; one of the best appeared to be the chloral derivative 25.\textsuperscript{27} We were delighted to find that simply stirring this compound with the benzotriazole 23b in dichloromethane at ambient temperature delivered an essentially quantitative yield of the desired aminated product 26 (Scheme 7).

Rather than risk difficulties with the isolation of the doubtless polar free amine 27, we chose to deprotect using trifluoroacetic acid and, after basification, treat the residue directly with N-iodosuccinimide. We were delighted to then isolate the desired iodo-chroman 28a, in 63\% yield for the two steps, which was converted into the corresponding acetate 28b, simply to more clearly observe the AB-system due to the new CH\textsubscript{2}OR group, which overlapped with the methoxyl resonance in the initial
product 28a (Scheme 7). A modified Stille methylation, with the addition of copper(I) iodide, was then used to complete the sequence, giving a 59% yield of the target chroman, (+)-10.

A final problem was therefore to find a method suitable for obtaining the alkyne-diol 20a, or a related protected form, in optically enriched form. We thought to add considerable bulk to the alkyne group in diol 19 in the form of a large silicon group. However, exposure of such a derivative 29 to AD-mix-β once again failed to give significant chiral induction: HPLC analysis of the derived acetate 30b showed a ee of only 12.7% (Scheme 8). Hydrolytic kinetic resolution of the derived epoxy-alkyne 31 also failed to give significant chiral induction.

We therefore turned to the known if chemically more distant chiral, non-racemic diol 33. This time, AD-mix-α was used to obtain the required diol 33 with chemical and optical yields both of 91%, in agreement with the initial report of its synthesis (Scheme 9). Protection as the corresponding dioxolane 34 and selective Dibal-H reduction then proceeded smoothly to give the aldehyde 35, which was converted into the alkyne 36 by addition of a trichloromethyl group to the aldehyde, acetylation, elimination of HCl using zinc and further HCl elimination from the resulting dichloroalkene using butyl lithium. The method was readily scalable and routinely gave overall yields of around 60%. Alternatives based on Wittig chemistry (Seyferth-Gilbert or Ohira-Bestmann methods) were either low yielding or unsuccessful.

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Sadly, attempts to use the dioxolane alkyne 36 in the Sonogashira coupling with iodo-anisole 18 (see Scheme 6) were only partly successful as, unexpectedly, around half the product was simply the deiodinated arene. Attempts to hydrolyse the dioxolane group were also not viable in our hands, using Bronsted or Lewis acid-based methods, due to the extreme sensitivity of the diol 20a to dehydration. Basic deprotection conditions were therefore required and so silyl protection was next investigated. Silylation of the dihydroxyamide 33 was slow but uneventful while Dibal-H reduction of the resulting bis-silyl derivative 37 was insufficiently selective, giving the expected aldehyde 38 along with the corresponding alcohol and desilylation at the tertiary site (Scheme 10). By contrast, low temperature reduction using ethereal LiAlH₄ was completely selective and gave an 88% isolated yield, lowered no doubt by the high volatility of the aldehyde 38. Corey-Fuchs homologation then delivered the alkyne 40 by way of the dibromoalkene 39 and thence the (S)-alkyne-diol 20a.

Using the same method as above (Scheme 6), the (S)-alkyne-diol 20a was successfully coupled to the iodo-anisole 18 to give the expected (S)(+)-alkynyl anisole 21 (Figure 4). We were also able to hydrolyse the acetyl group in iodo-anisole 18 to give the corresponding aniline 14; this too coupled well with the (S)-alkyne-diol 20a leading to the homologated aniline (S)(+)-41, thereby completing a formal asymmetric synthesis of tocopherol 1.

In any synthetic scheme that encounters problems; simply altering the order of steps can often obviate these. As a small caveat to the foregoing synthesis, we have shown that this might be possible in this
case, by first incorporating an alkene into a suitable benzotriazole and then carrying out the asymmetric dihydroxylation step (Scheme 11).

Thus, alkylation of the dianion 7 (Scheme 2)\textsuperscript{15} derived from 1-aminobenzotriazole, by both (E)-crotyl and (E)-cinnamyl halides gave excellent yields of the unsaturated homologues 42\textsubscript{a,b}. Asymmetric dihydroxylation then worked well with both substrates to give the diols 43\textsubscript{a,b}, the optical purities of which were 76\% and 82\% respectively according to hplc analysis (see 30\textsubscript{b} in Experimental). Both then underwent sequential deprotection and regiospecific cyclisation, as described above, to provide reasonable but unoptimised yields of the iodochromans 44\textsubscript{a,b}. Hence, this alternative approach may have some potential in this and related areas for the elaboration of a wide variety of substituted chromans.

We have thus reached our goal of synthesising a tocopherol precursor using intramolecular benzyne trapping by an alcohol group. Clearly, the approach overall is quite lengthy but does demonstrate the viability of this key step and could probably be shortened in a number of ways. Additionally, the use of the oxaziridine 25 should be of considerable use in this and related areas in future, when N-amination is required. The final feature of note in this synthesis is the incorporation of an iodine atom at a late stage, which suggests that this chemistry could be useful for the synthesis of radio-labelled products, for example using iodine 131, which has a half-life of around eight days.

**EXPERIMENTAL**

Infrared spectra were recorded using a Perkin Elmer 1600 series FTIR instrument as thin films or KBr disks. \textsuperscript{1}H NMR spectra were all recorded for dilute solutions in deuteriochloroform, unless otherwise stated, using a Bruker AM 400 MHz instrument with chloroform as internal reference ($\delta = 7.27$). \textsuperscript{13}C NMR spectra were all recorded in deuteriochloroform using a Bruker AM 400 MHz instrument, operating at 100 MHz with chloroform as internal reference ($\delta = 77.30$). Low resolution mass spectra
were obtained using a Fisons VG Platform II instrument and high resolution spectra were measured by the EPSRC Service, Swansea University. All extracts were dried over dried magnesium sulfate. Column chromatography was performed using Silica 60A, particle size 35-70 micron, from Fisher Scientific.

2,6-Dimethyl-4-nitroanisole (16a): To an ice-cold, stirred solution of 2,6-dimethylanisole 15 (40.00 g, 294 mmol) in glacial acetic acid (60 mL) was added dropwise 70% nitric acid (60 mL). After the addition was complete and gas evolution began to subside, the solution was slowly heated to 65 °C resulting in the formation of a pale yellow solution. [CAUTION: It is essential to keep the reaction temperature below 70 °C at all times as, above this, the mixture on occasions detonated with extreme violence; protection by a blast screen is recommended in any event]. The solution was then allowed to cool to ambient temperature and diluted with water (300 mL). Nitrogen was bubbled through the resulting deep brown solution for ca. 0.5 h to remove most of the nitrogen oxides present. The resulting yellow precipitate was separated by vacuum filtration and washed with copious water then crystallised from ethanol to give the nitroanisole 16a (32.6 g, 61%), mp 91.5-93.5 °C (lit.18 mp 89-91 °C); 1H NMR δ 2.28 (s, 6H, 2 x Me), 3.71 (s, 3H, OMe), 7.83 (s, 2H, ArH); 13C NMR δ 16.8 (2 x Me), 60.3 (OMe), 124.6 (2 x ArCH), 132.7 (2 x ArC), 143.8 (ArC), 162.8 (ArC); m/z (APCI) 152 (M-29, 100%).

4-Amino-2,6-dimethylanisole (16b): To a suspension of 10% Pd-C (0.65 g) in ethanol (130 mL) at rt was added portionwise the foregoing nitroanisole 16a (10.00 g, 55 mmol) followed by cyclohexene (32 mL, 330 mmol). The mixture was then heated under reflux for 14 h, cooled to rt and filtered through celite. The filter cake was washed with ethanol and the combined filtrates evaporated to leave the sensitive aniline 16b (7.56 g, 91%) as a brown solid which was sufficiently pure for further use and which showed mp 62-63 °C (lit.31 mp 60-61 °C), IR (KBr) 3772, 3612, 3397, 1605, 1487, 1340, 1219, 1150, 1016 cm⁻¹; 1H NMR δ 2.24 (s, 6H, 2 x Me), 3.47-3.51 (br s, 2H, NH₂), 3.69 (s, 3H, OMe), 6.39 (2, 2H, 2 x ArH); 13C NMR δ 16.5 (2 x Me), 60.4 (OMe), 115.7 (2 x ArCH), 131.9 (2 x ArC), 142.5 (ArC), 150.0 (ArC); m/z (APCI) 152 (M + 1, 100%).

4-Acetamido-2,6-dimethylanisole (16c): A stirred solution of the foregoing aniline 16b (7.60 g, 50 mmol) in dry THF (100 mL) containing triethylamine (7.4 mL, 53.2 mmol) was cooled in an ice-bath before the addition of acetyl chloride (4.0 mL, 54.1 mmol). The resulting solution was stirred overnight without further cooling then quenched by the sequential addition of saturated aqueous ammonium chloride (47 mL) and 2M hydrochloric acid (47 mL). The resulting mixture was extracted with ether (3 x 75 mL) and the combined extracts washed with water (75 mL) and brine (75 mL) then dried and evaporated. Crystallisation of the residue from ethanol left the acetamide 16c (8.60 g, 89%); mp 136-137 °C [lit.34 mp 136-138 °C]; IR (KBr) 3315, 1659, 1612, 1558, 1462, 1221, 1038, 1010 cm⁻¹;
\[\text{H NMR } \delta 2.10 (s, 3\text{H, Ac}), 2.22 (s, 6\text{H, 2 x Me}), 3.64 (s, 3\text{H, OMe}), 6.94 (br s, 1\text{H, NH}), 7.08 (s, 2\text{H, 2 x ArH}); \]
\[\text{\textsuperscript{13}C NMR } \delta 16.5 (2 \text{ x Me}), 24.8 (\text{MeCO}), 60.2 (\text{Ome}), 121.1 (2 \text{ x ArCH}), 131.7 (2 \text{ x ArC}), 133.8 (\text{ArC}), 153.9 (\text{ArC}), 168.9 (\text{C=O}); m/z (APCI) 194 (M^+ , 100\%).\]

4-Acetamido-2,6-dimethyl-5-iodoanisole (17): A suspension of the foregoing acetamide 16c (7.4 g, 38.1 mmol) and N-iodosuccinimide (12.9 g, 57.2 mmol) in glacial acetic acid (120 mL) was heated to reflux for 1.5 h. The resulting purple solution was cooled to rt and neutralised with 2M aqueous sodium hydroxide to give a suspension which was vacuum filtered. The solid was washed with water then crystallised from ethyl acetate to give the iodo-anisole 17 (8.5 g, 70\%); mp 189–191 °C; IR (KBr) 3272, 1651, 1527, 1461, 1377, 1228, 1160, 1006 cm\(^{-1}\); \[\text{\textsuperscript{1}H NMR } \delta 2.08 (s, 3\text{H, Ac}), 2.12 (s, 3\text{H, Me}), 2.28 (s, 3\text{H, Me}), 3.52 (s, 3\text{H, OMe}), 7.15–7.25 (br s, 1\text{H, NH}), 7.61 (s, 1\text{H, 3-H}); \]
\[\text{\textsuperscript{13}C NMR } \delta 16.4 (\text{Me}), 22.9 (\text{Me}), 24.8 (\text{Me}), 60.4 (\text{OMe}), 96.4 (\text{C-I}), 123.0 (\text{3-CH}), 131.7 (\text{ArC}), 134.4 (\text{ArC}), 135.1 (\text{ArC}), 153.9 (\text{ArC}), 168.5 (\text{C=O}); m/z (APCI) 320 (M^+ + H, 100\%).\]

Anal. Calcd for C\(_{11}\)H\(_{14}\)INO\(_2\): C, 41.38; H, 4.42; N, 4.39. Found: C, 41.60; H, 4.35; N, 4.36.

4-Acetamido-2,6-dimethyl-5-iodo-3-nitroanisole (18): The foregoing iodo-anisole 17 (1.00 g, 3.1 mmol) was stirred in glacial acetic acid (10 mL) at rt as 70\% nitric acid (2.0 mL) was added dropwise. The resulting solution was then stirred at 65 °C for 5 h, cooled to rt, diluted with water (18 mL) and the resulting precipitate collected by filtration and crystallised from aqueous ethanol to give the nitro-iodide 18 (0.85 g, 76\%); mp 226–227 °C; IR (KBr) 3617, 3212, 1667, 1519, 1455, 1376, 1272, 1221, 1039, 989 cm\(^{-1}\); \[\text{\textsuperscript{1}H NMR } \delta 2.17 (s, 3\text{H, Ac}), 2.22 (s, 3\text{H, Me}), 2.49 (s, 3\text{H, Me}), 3.72 (s, 3\text{H, OMe}), 7.06–7.17 (br s, 1\text{H, NH}); \]
\[\text{\textsuperscript{13}C NMR } \delta 12.2 (\text{Me}), 23.7 (\text{Me}), 23.9 (\text{Me}), 61.0 (\text{OMe}), 106.3 (\text{Cl}), 125.2 (\text{ArC}), 126.3 (\text{ArC}), 127.3 (\text{ArC}), 139.9 (\text{ArC}), 156.2 (\text{ArC}), 169.3 (\text{C=O}); m/z (APCI) 365 (M^+ + H, 100\%).\]

Anal. Calcd for C\(_{11}\)H\(_{13}\)IN\(_2\)O\(_4\): C, 36.28; H, 3.60; N, 7.69. Found: C, 36.43; H, 3.81; N, 7.56.

2-Methyl-1-buten-3-yne (19): 2-Methylbut-3-yn-2-ol (50 g) and p-toluenesulfonic acid (50 g) were stirred together and heated to 90 °C under a distillation apparatus. After 3 h, the distillate was redistilled (bath temp. 50 °C) to give the enyne 19 (11.4 g, 29\%); bp 34 °C; IR (film) 3300, 3100, 1614, 1455, 1374, 1266, 1214, 1171, 1013, 961, 904 cm\(^{-1}\); \[\text{\textsuperscript{1}H NMR } \delta 1.71 (s, 3\text{H, Me}), 2.71 (s, 1\text{H, 4-H}), 5.11 (\text{app. s, 1\text{H, 1-H}}); \]
\[\text{\textsuperscript{2}H NMR } \delta 5.21 (\text{app. s, 1\text{H, 1-H}}); \]

3,4-Dihydroxy-3-methyl-1-butyne (20a): AD-mix-β (63.70 g) was added to a stirred mixture of t-butanol (225 mL) and water (225 mL) followed by methanesulfonylamine (4.30 g). Once the mixture became clear, the foregoing enyne 19 (3.00 g, 45.5 mmol) was added and the resulting mixture stirred at rt for 24 h. Sodium sulfite (68 g) was then added and stirring continued for 1 h. The mixture was then extracted with dichloromethane (4 x 100 mL) and the combined extracts washed with 2M aqueous potassium hydroxide (2 x 50 mL) then dried and evaporated. Column chromatography over silica gel eluted with ether-hexanes (2:1) then separated the diol 20a (3.1 g, 67\%) as a pale yellow oil.
IR (film) 3430, 2113, 1511, 1376, 1244, 1060, 953, 891 cm\(^{-1}\); \(^1\)HNMR \(\delta 1.37\) (s, 3H, Me), 2.39 (s, 1H, 1-H), 3.44 (d, \(J = 7.6\) Hz, 1H, 4-H\(_a\)), 3.60 (d, \(J = 7.6\) Hz, 1H, 4-H\(_b\)); \(^1\)HCNMR \(\delta 25.6\) (Me), 68.6 (3-C), 70.7 (4-CH\(_2\)), 72.6 (1-CH), 86.3 (2-C); \(m/z\) (APCI) 100 (M\(^+\), 100%).

(\pm\)-2-Hydroxy-2-methyl-3-butyn-1-yl benzoate (20b): Benzoyl chloride (26 \(\mu\)L, 0.22 mmol) was added to a stirred solution of the diol 20a (20 mg, 0.2 mmol) in dry dichloromethane containing triethylamine (30 \(\mu\)L, 0.22 mmol). After stirring overnight at rt, the solution was diluted with dichloromethane (5 mL) and washed with water (3 mL), 1M hydrochloric acid (2 mL) and brine then dried and evaporated. Short-column chromatography (SiO\(_2\); petrol-ether 4:1) separated the benzoate 20b (29 mg, 71%) as a colourless oil; IR (film) 3456, 3284, 2360, 1722, 1452, 1371, 1274, 1116, 710; \(^1\)HNMR \(\delta 1.53\) (s, 3H, 2-Me), 2.43 (s, 1H, 4-H), 2.92 (br s, 1H, OH), 4.23 (d, \(J = 11.1\) Hz, 1H, 1-H\(_a\)), 4.36 (d, \(J = 11.1\) Hz, 1H, 1-H\(_b\)), 7.36 (t, \(J = 7.6\) Hz, 2H), 7.49 (t, \(J = 7.6\) Hz, 1H), 8.00 (dd, \(J = 7.6, 1.0\) Hz, 2H); \(^1\)3CNMR \(\delta 26.4\) (Me), 67.3 (CH\(_2\)), 71.6 (2-C), 73.1 (4-CH), 85.1 (3-C), 128.9 (2 x ArCH), 130.0 (ArC), 130.2 ArCH), 133.7 (2 x ArCH), 166.8 (C=O); \(m/z\) (APCI) 205 (M+H, 100%).

GC analysis using a chiral CDX-\(\beta\) column at \(150^\circ\)C (detector: 300 \(^\circ\)C; injection: 250 \(^\circ\)C]; column head pressure: 20 psi, \(R_t\) ~ 14.2 min. showed < 10% ee against a racemic sample.

Sonogashira coupling; General Procedure:\(^{35}\) The aryl iodide (3.5 mmol) was stirred in degassed THF (40 mL) then triethylamine (14 mL) and tetrakis(triphenylphosphine)palladium(0) (0.80 g, 0.7 mmol) was added followed by the 1-alkyne (7.0 mmol). The resulting solution was degassed again by passage of dry nitrogen for 40 minutes before copper(I) iodide (133 mg, 0.7 mmol) was added and the mixture stirred at reflux for 24 h then cooled to rt and the bulk of the THF evaporated. The residue was diluted with water (35 mL) and extracted with dichloromethane (3 x 50 mL). The combined extracts were dried and evaporated and the product(s) separated by column chromatography over silica gel, typically using 5% MeOH-CHCl\(_3\) as eluant.

(\pm\)-4-Acetamido-2,6-dimethyl-5-[3’,4’-dihydroxy-3’-methylbut-1’-yn-1’-yl]-3-nitroanisole (21): Following the general Sonogashira procedure, the nitro-iodide 18 (1.30 g, 3.5 mmol) was coupled with the diol 20a (0.70 g, 7.0 mmol) using (Ph\(_3\))\(_4\)Pd (0.80 g, 0.7 mmol) to give the alkynediol 21 (1.00 g, 86%) as a pale yellow, amorphous solid, mp 178-182 \(^\circ\)C (MeOH); IR 3822, 3580, 3280, 1667, 1504 cm\(^{-1}\); \(^1\)HNMR \(\delta 1.46\) (s, 3H, 3’-Me), 2.13 (s, 3H, Ac), 2.23 (s, 3H, ArMe), 2.36 (s, 3H, ArMe), 3.04 (br s, 2H, 2 x OH), 3.43-3.50 (m, 2H, 4’-CH\(_2\)), 3.65 (S, 3H, OMe), 7.54 (BR S, 1H, NH); \(^1\)3CNMR (CD\(_3\)OD) \(\delta 12.1\) (Me), 15.5 (Me), 23.0 (Me), 26.6 (Me), 61.4 (OMe), 70.2 (CH\(_2\)), 71.4 (C), 78.7 (C), 103.9 (1’-C), 124.3 (ArC), 126.2 (ArC), 127.7 (ArC), 128.3 (ArC), 138.6 (ArC), 157.8 (ArC), 173.4 (C=O); \(m/z\) (APCI) 319 (M\(^+\)-OH, 100%). HR-MS (FAB). Calcd for C\(_{16}\)H\(_{21}\)N\(_2\)O\(_6\) (M+H): 337.1396. Found: \(m/z\) 337.1400.
(-)-4-Acetamido-3-amino-2,6-dimethyl-5-[3’,4’-dihydroxy-3’-methylbut-1’-yl]anisole (22): The alkyne-diol 21 (0.22 g, 0.66 mmol) was added to a suspension of 20% palladium hydroxide on carbon (Pearlman’s catalyst) (44 mg) in methanol (40 mL) and the mixture stirred under an atmosphere of hydrogen for 48 h then filtered through celite. The solid was washed with warm methanol and the combined filtrates were dried and the extracts were taken up into water (1.2 mL) maintained at 0°C and the hydrolysis was followed by tlc. Upon completion (ca. 4 h), the bulk of the methanol was evaporated and the residue was washed with methanol and the combined filtrates evaporated to leave the aniline 22 (0.202 g, 99%) as a colourless solid, mp 250°C. HR MS (FAB). Calcd for C_{16}H_{27}N_{2}O_{4} (M+H): 311 (M^+H, 100%). Found: m/z 311.1653.

(±)-1-Acetyl-7-(3’,4’-dihydroxy-3’-methylbutan-1’-yl)-4,6-dimethyl-5-methoxybenzotriazole (23a): To a solution of the foregoing aniline 22 (0.34 g, 1.1 mmol) in methanol (3 mL) at rt was added a solution of sodium nitrite (0.22 g, 3.3 mmol) in water (0.6 mL). The resulting solution was added slowly to a stirred solution of 10M hydrochloric acid (1.0 mL) in water (1.2 mL) maintained at 0°C. Stirring at this temperature was continued for 0.5 h then water (12 mL) was added and the mixture extracted with dichloromethane (3 x 20 mL). The combined extracts were dried and evaporated to leave the N-acetyl-benzotriazole 23a (0.266 g, 73%) as a colourless solid, mp 99-102°C (CHCl_{3}-petrol); IR (KBr) 3270, 1745, 1459, 1366, 1309, 1266, 1130, 1094 cm^{-1}; ^{1}H NMR δ 1.24 (s, 3H, 3’-Me), 1.72 (t, J = 8.8 Hz, 2H, 2’-CH_2), 2.38 (s, 3H, Ac), 2.49-2.54 (br res., 2H, 2 x OH), 2.66 (s, 3H, ArMe), 2.96 (s, 3H, ArMe), 2.99-3.15 (m, 2H, 1’-CH_2), 3.41 (dd, J = 11.3, 5.9 Hz, 1H, 4’-HA), 3.54 (dd, J = 11.3, 5.9 Hz, 1H, 4’-HB), 3.70 (s, 3H, OMe); ^{13}C NMR δ 9.6 (Me), 12.0 (Me), 22.1 (Me), 24.3 (Me), 24.9 (2’-CH_2), 37.3 (1’-CH_2), 59.5 (OMe), 68.2 (4’-CH_2), 71.8 (3’-C), 118.9 (ArC), 123.4 (ArC), 126.8 (ArC), 133.6 (ArC), 145.4 (ArC), 154.4 (ArC), 169.9 (C=O); m/z (APCI) 322 (M^+H, 100%). HR-MS (FAB). Calcd for C_{16}H_{25}N_{2}O_{4} (M+H): 322.1761. Found: m/z 322.1763.

(±)-7-(3’,4’-Dihydroxy-3’-methylbut-1’-yl)-4,6-dimethyl-5-methoxybenzotriazole (23b): Potassium carbonate (0.14 g, 1.00 mmol) and water (0.5 mL) were added to a stirred solution of the foregoing N-acetyl-benzotriazole 23a (0.10 g, 0.32 mmol) in methanol (2 mL) at rt and the hydrolysis followed by tlc. Upon completion (ca. 4 h), the bulk of the methanol was evaporated and the residue taken up into dichloromethane (20 mL), which was then dried, filtered and evaporated. The solid residue was washed with methanol and the combined filtrates evaporated to leave the benzotriazole 23b (0.09 g, 99%) as a colourless solid, mp 250-252°C (MeOH); IR (KBr) 3376, 1621, 1434, 1140, 1071, 998, 880 cm^{-1}; ^{1}H NMR (CD_{3}OD) δ 1.17 (s, 3H, 3’-Me), 1.64-1.73 (m, 2H, 2’-CH_2), 2.23 (s, 3H,
Ac), 2.45 (s, 3H, ArMe), 2.97-3.03 (m, 2H, 1'-CH\textsubscript{2}), 3.36 (d, J = 11.3 Hz, 1H, 4'-H\textsubscript{a}), 3.41 (d, J = 11.3 Hz, 1H, 4'-H\textsubscript{b}), 3.62 (s, 3H, OMe); \textsuperscript{13}C NMR (CD\textsubscript{3}OD) δ 11.8 (Me), 12.6 (Me), 24.4 (Me), 24.6 (2'-CH\textsubscript{2}), 39.6 (1'-CH\textsubscript{2}), 61.3 (OMe), 70.3 (4'-CH\textsubscript{2}), 74.4 (3'-C), 115.3 (ArC), 124.6 (ArC), 127.8 (ArC), 142.7 (ArC), 145.0 (ArC), 153.9 (ArC); m/z (APCI) 280 (M\textsuperscript{+}+H, 100%). HR-MS (FAB). Calcd for C\textsubscript{14}H\textsubscript{22}N\textsubscript{3}O\textsubscript{3} (M+H): 280.1656. Found: m/z 280.1652.

(±)-1-(tert-Butylcarbonylamino)-7-(3',4'-dihydroxy-3'-methylbut-1'-yl)-4,6-dimethyl-5-methoxybenzotriazole (26): To a stirred solution of the foregoing benzotriazole 23b (0.28 g, 1 mmol) in dry dichloromethane (8 mL) at rt was added the N-Boc-oxaziridine 25 (0.53 g, 2.0 mmol)\textsuperscript{22} and the resulting solution stirred overnight. The solvent was then evaporated and the residue separated by column chromatography (5% MeOH-CHCl\textsubscript{3}) to give the N-Boc-amino-benzotriazole 26 (0.40 g, 94%) as a yellow oil; IR (film) 3274, 1739, 1452, 1362, 1248, 1158, 1110, 1050, 906, 834, 732 cm\textsuperscript{-1}; \textsuperscript{1}H NMR δ 1.23 (s, 3H, 3'-Me), 1.32-1.75 (br s, 9H, Boc), 1.78 (t, J = 8.6 Hz, 2H, 2'-CH\textsubscript{2}), 2.30 (s, 3H, ArMe), 2.56 (s, 3H, ArMe), 2.99-3.02 (m, 2H, 1'-CH\textsubscript{2}), 3.54 (d, J = 11.3 Hz, 1H, 4'-H\textsubscript{a}), 3.65 (d, J = 11.3 Hz, 1H, 4'-H\textsubscript{b}), 3.69 (s, 3H, OMe); \textsuperscript{13}C NMR δ 7.9 (Me), 10.9 (Me), 21.9 (Me), 24.9 (2'-CH\textsubscript{2}), 27.1 (3 x Me), 37.1 (1'-CH\textsubscript{2}), 59.5 (4'-CH\textsubscript{2}), 59.6 (OMe), 72.1 (3'-C), 82.4 (CMe\textsubscript{3}), 109.0 (ArC), 124.9 (ArC), 126.0 (ArC), 129.5 (ArC), 139.4 (ArC), 152.7 (ArC), 156.7 (C=O); m/z (APCI) 395 (M\textsuperscript{+}+H, 100%). HR-MS (FAB). Calcd for C\textsubscript{19}H\textsubscript{31}N\textsubscript{4}O\textsubscript{3} (M+H): 395.2289. Found: m/z 395.2295.

(±)-8-Iodo-6-methoxy-2,5,7-trimethylchroman-2-methanol (28a): The N-Boc-amino-benzotriazole 26 (0.22 g, 0.56 mmol) was dissolved in a stirred solution of trifluoroacetic acid (1.1 mL) in dichloromethane (5 mL) at rt. The deprotection was followed by TLC and when complete (ca. 0.75 h), the volatiles were evaporated, finally under high vacuum. The residue was then dissolved in dichloromethane (5 mL) and stirred with anhydrous potassium carbonate (1.0 g) for 0.25 h and filtered. The solid was washed with warm dichloromethane and the combined filtrates concentrated to a volume of ca. 5 mL and protected from light before N-iodosuccinimide (0.38 g, 1.7 mmol) was added in one portion. The resulting purple solution was stirred at rt for a further 1 h then diluted with dichloromethane (15 mL) and the solution washed with saturated aqueous sodium thiosulfate (5 mL) and then dried and evaporated. Silica gel column chromatography (petrol-EtOAc, 2:1) separated the iodochroman 28a (0.129 g, 63%) as a yellow semi-solid; IR (film) 3390, 1739, 1362, 1248, 1158, 1110, 1058, 882 cm\textsuperscript{-1}; \textsuperscript{1}H NMR δ 1.21 (s, 3H, 2-Me), 1.67 (ddd, J = 13.6, 6.0, 4.8, 3-H\textsubscript{a}), 1.80-1.90 (m, 1H, 3-H\textsubscript{b}), 2.08 (s, 3H, ArMe), 2.35 (s, 3H, ArMe), 2.56-2.60 (m, 2H, 4-CH\textsubscript{2}), 3.57 (m, 5H, C\textsubscript{6}H\textsubscript{4}OH and OMe); \textsuperscript{13}C NMR δ 11.0 (2-Me), 19.2 (ArMe), 19.4 (3-CH\textsubscript{2}), 20.9 (ArMe), 26.9 (4-CH\textsubscript{2}), 59.5 (OMe), 68.2 (CH\textsubscript{2}OH), 77.2 (2-C), 90.0 (C-I), 117.8 (ArC), 128.4 (ArC), 132.3 (ArC), 146.8 (ArC), 149.1 (ArC); m/z (APCI) 345 (M\textsuperscript{+}-OH, 100%). HR-MS (NH\textsubscript{4}-Cl). Calcd for C\textsubscript{14}H\textsubscript{23}INO\textsubscript{3} (M+NH\textsubscript{4}): 380.0717. Found: m/z 380.0714.
(±)-2-Acetoxymethyl-8-iodo-6-methoxy-2,5,7-trimethylchroman (28b): Iodochroman 28a (30 mg, 0.08 mmol) was stirred in dry pyridine (1.5 mL) containing acetyl chloride (0.02 mL, 0.24 mmol) and 4-dimethylaminopyridine (2 mg) overnight at rt. The solution was then diluted with dichloromethane (5 mL) and washed with water (2 x 2 mL), saturated aqueous copper sulphate (2 mL) and saturated aqueous potassium carbonate (2 mL) then dried and evaporated. Silica gel column chromatography (ether-hexane 1:3) separated the acetoxoy-chroman 28b (23 mg, 75%) as a yellow oil; IR (film) 1744, 1450, 1390, 1260, 1057, 800 cm⁻¹; ¹H NMR δ 1.28 (s, 3H, 2-Me), 1.72-1.88 (m, 2H, 3-CH₂), 2.05 (s, 3H, ArMe), 2.06 (s, 3H, ArMe), 2.34 (s, 3H, Ac), 2.53 (app. t, J = 6.9 Hz, 2H, 4-CH₂), 3.56 (s, 3H, OMe), 4.02 (d, J = 11.3 Hz, 1H, 1'-H₆), 4.09 (d, J = 11.3 Hz, 1H, 1'-H₆); ¹³C NMR δ 11.0 (2-Me), 19.4 (ArMe), 20.0 (ArMe), 20.1 (3-CH₂), 20.9 (MeCO), 28.7 (4-CH₂), 59.5 (OMe), 67.2 (CH₂OH), 74.3 (2-C), 89.1 (C-I), 117.2 (ArC), 128.3 (ArC), 133.0 (ArC), 147.1 (ArC), 149.0 (ArC), 169.9 (C=O); m/z (APCI) 277 (M⁺-I, 100%). HR-MS (EI). Calcd for C₁₆H₂₁I₄O₄ (M⁺): 404.0485. Found: m/z 404.0483.

(±)-6-Methoxy-2,5,7,8-tetramethyl-2-methanol (10): To a solution of the iodochroman 28a (0.153 g, 0.75 mmol) in 1-methyl-2-pyrrolidinone (NMP; 3 mL) at rt was added triphenylphosphine (40 mg), tris-(dibenzylideneacetone)dipalladium(0).CHCl₃ (50 mg) and copper(I) iodide (15 mg). The stirred mixture was then degassed for 0.5 h before the addition of diethylamine (0.5 mL) and tetramethyltin (0.75 g, 4.5 mmol) in NMP (1 mL) and then heated at 100 °C for 24 h. The cooled suspension was then diluted with 10% aqueous sodium sulfite (40 mL) and extracted with ether (3 x 25 mL). The combined extracts were washed with 10% aqueous potassium fluoride (15 mL) then dried and evaporated. Column chromatography (SiO₂; petrol-EtOAc 2:1) of the residue separated the chroman-methanol (±)-10 (0.111 g, 59%) as a thick, colourless oil, which slowly solidified, mp 49-52 °C; IR 3310, 1450, 1233, 1040 cm⁻¹; ¹H NMR δ 1.23 (s, 3H, 2-Me), 1.74 (ddd, J = 13.6, 6.1, 4.8, 3-H₃), 1.80-1.90 (m, 1H, 3-H₃b), 2.08 (s, 3H, ArMe), 2.16 (s, 3H, ArMe), 2.30 (s, 3H, ArMe), 2.56-2.67 (m, 2H, 4-CH₂), 3.54 (2 x d (partly obscured), J_AB = 11.1 Hz, CH₃OH), 3.56 (s, 3H, OMe); ¹³C NMR δ 11.2 (Me), 11.4 (Me), 12.4 (Me), 19.9 (Me), 20.3 (Me), 27.4 (4-CH₂), 60.0 (OMe), 68.8 (CH₂OH), 75.2 (2-C), 117.3 (ArC), 122.4 (ArC), 125.1 (ArC), 128.0 (C), 146.9 (ArC), 149.3 (ArC); m/z (APCI) 251 (M⁺+ H, 100%). HR-MS (FAB). Calcd for C₁₅H₂₂O₃ (M⁺): 250.1569. Found: m/z 250.1567. The spectroscopic data were identical to those recorded previously for a while solid, no mp quoted.

4-(tert-Butyldiphenylsilyl)-2-hydroxy-2-methylbut-3-yn-1-yl acetate (30b): i) 4-(tert-Butyldiphenylsilyl)-2-methyl-1-buten-3-yne (29): Butyl lithium (24 mL of a 2.5M solution in hexanes, 60 mmol) was added dropwise to a stirred solution of 2-methyl-1-buten-3-yne 19 (4.00 g, 60 mmol) in dry THF (45 mL), cooled in an ice bath. After 0.25 h, t-butyldichlorodiphenylsilane (14.0 g, 60 mmol) was added dropwise and stirring continued for 2 h before the bulk of the THF was evaporated. The
combined extracts were dried and evaporated and the residue separated by column chromatography (ether:hexane 9:1) to give the silyle 30a (0.44 g, 70%); IR (film) 3429, 2156, 1589, 1471, 1428, 1362, 1237, 1110, 1048, 968, 921 cm⁻¹; ¹H NMR δ 1.00 (s, 9H, 1J), 1.59 (s, 9H, 2Me), 2.06 (s, 3H, MeCO), 2.54 (br s, 1H, OH), 4.02 (d, J = 11.1 Hz, 1Hδ), 4.28 (d, J = 11.1 Hz, 1Hδ), 7.29-7.37 (m, 6H), 7.69 (d, J = 6.4 Hz, 4H); ¹³C NMR δ 18.5 (CMe₃), 25.3 (3 Me), 26.0 (2Me), 27.0 (MeCO), 67.2 (2-C), 71.1 (CH₂), 84.2 (4-C), 110.6 (3-C), 127.8 (ArCH), 129.7 (ArCH), 132.9 (ArC), 135.5 (ArCH).

Chiral hplc analysis using an 5μ OD column at 0.8 mL min⁻¹ flow of 0.7% IPA in hexanes, detector set at 254 nm, gave a retention time of ~ 25 min.

2-[(tert-Butyldiphenylsilyl)ethynyl-1-yl-2-methyloxirane (31): To a well-stirred solution of the silyl butyne 29 (0.60 g, 1.73 mmol) in dichloromethane (9 mL) was added m-chloroperoxybenzoic acid (0.60 g, 3.46 mmol). The resulting suspension was stirred for 2 h at rt before saturated aqueous sodium bicarbonate (10 mL) was added and the mixture extracted with ether (3 x 15 mL). The combined extracts were dried and evaporated and the residue separated by column chromatography (ether:hexane 9:1) to give the oxirane 31 (0.59 g, 95%) as a yellow oil; IR (film) 3429, 2156, 1589, 1471, 1428, 1362, 1193, 1111, 1009, 907, 820 cm⁻¹; ¹H NMR δ 0.99 (s, 9H), 1.57 (s, 3H, 2Me), 2.71 (d, J = 5.6 Hz, 3-Hδ), 3.03 (d, J = 5.6 Hz, 3-Hδ), 7.26-7.35 (m, 6H), 7.62-7.69 (m, 4H); ¹³C NMR δ 19.0 (CMe₃), 23.3
(+)-(R)-2,3-Dihydroxy-2,N-dimethyl-N-methoxy propanamide (33): AD-mix-α directed dihydroxylation of the Weinreb amide derivative 32 (6.10 g, 47.1 mmol) of methacrylic acid as described previously[30] gave the expected dihydroxyamide 33 (7.00 g, 91%), which showed [α]D26 = +4.3 (c 2.0 g/100 mL MeOH) [lit.[30] [α]D25 = +4.7 (c 1.80 g/100 mL MeOH)] and otherwise identical spectroscopic and analytical data.

(R)-N-Methoxy-N-methyl-2,2,4-trimethyl-1,3-dioxolane-4-carboxamide (34): A solution of the dihydroxyamide 33 (3.25 g, 19.9 mmol), 2,2-dimethoxypropane (12.3 mL, 99.5 mmol) and p-toluenesulfonic acid monohydrate (76 mg) in toluene (60 mL) was heated under reflux for 2 h then cooled and evaporated. Silica gel column chromatography (EtOAc-petrol 1:1) separated the dioxolane 34 (3.60 g, 87%) as a colourless oil: IR 1658, 1456, 1372, 1247, 1211 cm⁻¹; 1H NMR δ 1.16 (s, 3H, 5-Me), 1.25 (s, 3H, 2-Me), 1.31 (s, 3H, 2-Me), 3.10 (br. s, 3H, NMe), 3.53 (s, 3H, OMe), 3.56 (d, J = 8.3 Hz, 4-Ha), 4.37 (d, J = 8.3 Hz, 4-Hb); 13C NMR δ 23.4 (5-Me), 26.1 (2-Me), 27.3 (2-Me), 60.7 ~(NMe), 61.1 (OME), 73.0 (4-CH₂), 77.7 (5-C), 110.7 (2-C), 171.7 (C=O); m/z (APCI) 204 (M+H, 100%).

(S)-4-Ethynyl-2,2,4-trimethyl-1,3-dioxolane (36): i) (R)-2,2,4-Trimethyl-1,3-dioxolane-4-carboxaldehyde (35): Dibal-H (53.2 mL of a 1M solution in THF, 53.2 mmol) was added dropwise to a stirred solution of the acetal 34 (3.60 g, 17.7 mmol) in THF cooled in an ice-water bath. After 1 h at this temperature, the reaction mixture was poured into 5% hydrochloric acid in ethanol (50 mL) and the resulting mixture partitioned between brine (20 mL) and ether (100 mL). The separated organic solution was washed with water (15 mL) then dried and evaporated to leave the aldehyde 35 (2.80 g, 100%); IR (film) 3424, 1725, 1458, 1373 cm⁻¹; 1H NMR δ 1.29 (s, 3H, Me), 1.39 (br s, 6H, 2 x Me), 3.66 (s, 2H, 4-CH₂), 9.59 (s, 1H, CHO); 13C NMR δ 18.2 (Me), 25.4 (Me), 25.8 (Me), 69.8 (4-CH₂), 76.3 (C), 83.6 (C), 201.4 (C=O); m/z (APCI) 145 (M+H, 100%), which was used immediately in the next step. ii) (S)-4-(2,2-Dichloroethen-1-yl)-2,2,4-trimethyl-1,3-dioxolane: Sodium trichloroacetate (4.70 g, 25.4 mmol) was added in portions to a stirred solution of the foregoing aldehyde 35 (2.27 g, 15.9 mmol) and trichloroacetic acid (4.20 g, 25.4 mmol) in DMF (12 mL); an ice-water bath was used to maintain the temperature below 35 °C during the addition. The mixture was then stirred without further cooling for 4 h as CO₂ was continually evolved before cooling to 5 °C and the addition of acetic anhydride (3.0 mL, 31.8 mmol), which caused increased CO₂ evolution. After a further 1 h at rt, the mixture was diluted with acetic acid (15 mL) and cooled in an ice-water bath. Zinc powder (2.10 g, 31.8 mmol) was then added in one portion and the mixture heated to 60 °C for 1 h. After cooling to rt, water (10 mL) was added and the mixture extracted with hexanes (3 x 30 mL). The combined extracts were washed with water (10 mL) and brine (10 mL) then dried and evaporated to leave the crude
dichloroalkene (1.89 g, 57%) as a pale yellow oil; IR (film) 1613, 1451, 1372, 1210, 1111, 991, 883, 810 cm⁻¹; ¹H NMR δ 1.30 (s, 3H, Me), 1.36 (s, 3H, Me), 1.42 (s, 3H, Me), 3.88 (d, J = 8.6 Hz, 4-Ha), 4.09 (d, J = 8.6 Hz, 4-Hb), 6.20 (s, 1H, :CH); ¹³C NMR δ 24.3 (Me), 26.0 (Me), 27.2 (Me), 73.9 (4-CH₂), 80.5 (C), 109.5 (C), 120.2 (:C), 136.0 (:CH). iii) (S)-4-Ethynyl-2,2,4-trimethyl-1,3-dioxolane (36): To a stirred solution of the foregoing dichloroalkene (1.64 g, 7.8 mmol) in THF (10 mL) maintained at -30 °C was added dropwise butyl lithium (9.4 mL of a 2.5M solution in hexanes, 23.5 mmol). The resulting solution was stirred for 1 h without further cooling then quenched by the addition of saturated aqueous ammonium chloride (10 mL) and diluted with ether (30 mL). The separated aqueous phase was extracted with ether (3 x 30 mL) and the combined organic solutions washed with water (10 mL) and brine (10 mL) then dried, filtered through a pad of silica gel and the filtrate and washings evaporated to leave the alkyne 36 (1.10 g, 99%) as a pale yellow oil; IR (film) 3430, 2140, 1460, 1369, 1270, 1200 cm⁻¹; ¹H NMR δ 1.28 (s, 3H, 5-Me), 1.39 (s, 3H, Me), 1.43 (s, 3H, 2-Me), 2.35 (s, 1H, 2'-H), 3.65 (d, J = 8.2 Hz, 4-Ha), 4.05 (d, J = 8.2 Hz, 4-Hb); m/z (APCI) 141 (M + H, 100%).

(+)-(R)-2,3-Di-(tert-butyldimethylsilyloxy)-2,N-dimethyl-N-methoxy propanamide (37): t-Butyldimethylsilyl chloride (23.0 g, 152 mmol) was added to a stirred solution of the dihydroxyamide 33 (10.30 g, 63.3 mmol) and imidazole (21.50 g, 316.5 mmol) in dry DMF (40 mL), which was then stirred at 35 °C for 18 h before being cooled to rt, diluted with water (10 mL) and extracted with dichloromethane (3 x 50 mL). The combined extracts were washed with water (20 mL) and brine (20 mL) and then dried and evaporated. Silica gel column chromatography (EtOAc-petrol 95:5) then separated the bis-silyl ether 37 (23.2 g, 94%) as a colourless oil; [α]D²⁶ = + 3.2 (c 2.0 g/100 mL CHCl₃); IR (film) 1670, 1472, 1387, 1361, 1251, 1214, 1033, 778, 723 cm⁻¹; ¹H NMR δ 0.00 (s, 3H), 0.01 (s, 3H), 0.08 (s, 3H), 0.11 (s, 3H), 0.83 (s, 9H), 0.84 (s, 9H), 0.01 (s, 3H), 0.08 (s, 3H), 0.11 (s, 3H), 0.83 (s, 9H), 0.84 (s, 9H), 0.85 (s, 9H).

(R)-2,3-Di-(tert-butyldimethylsilyloxy)-2-methylpropanal (38): Lithium aluminium hydride (80 mL of a 1M solution in ether, 77.7 mmol) was added dropwise to a stirred solution of the bis-silyl ether 37 (10.1 g, 25.9 mmol) in ether (260 mL) maintained at – 78 °C. Stirring was continued at this temperature for 40 mins. and then an aqueous solution of 2M sodium hydroxide (40 mL) was slowly added. After warming to rt, the resulting mixture was separated and the aqueous layer extracted with ether (3 x 100 mL). The combined organic solutions were dried and carefully evaporated to leave the aldehyde 38 (7.50 g, 88%) as a pale yellow oil, which was sufficiently pure for the next step and which showed: IR (film) 2858, 1740, 1472, 1388, 1362, 1255, 1156, 1108 cm⁻¹; ¹H NMR δ -0.08 (s, 3H, MeSi), -0.06 (s, 3H, MeSi), 0.01 (s, 3H, MeSi), 0.02 (s, 3H, MeSi), 0.80 (s, 9H), 0.82 (s, 9H), 1.12 (s, 3H, 3-Me), 3.51-3.55 (m, 2H), 9.51 (s, 1H, CHO); ¹³C NMR δ -5.8 (MeSi), -5.7 (MeSi), -3.6 (MeSi), -
3.2 (MeSi), 15.2 (2-Me), 18.1 (CSi), 18.2 (CSi), 25.7 (6 x Me), 68.4 (3-CH₂), 80.7 (2-C), 204.6 (CHO); m/z (APCI) 333 (M + H, 100%).

(+)-(S)-1,1-Dibromo-3,4-di-(tert-butyldimethylsilyloxy)-3-methylbut-1-ene (39): Triphenylphosphine (44.0 g, 167.7 mmol) was added to a stirred solution of carbon tetrabromide (27.8 g, 83.3 mmol) in dichloromethane (200 mL) followed by the foregoing aldehyde 38 (13.9 g, 41.9 mmol). The resulting solution was stirred at rt for 4 h then washed with water (30 mL), the solvent evaporated and the residue separated by silica gel column chromatography (pentane-ether 9:1) to give the dibromoalkene 39 (15.2 g, 74%) as a colourless oil; [α]₀²⁵ = + 2.7 (c 1.9 g/100 mL CHCl₃); IR (film) 1607, 1472, 1389, 1361, 1255, 1108, 1036, 836 cm⁻¹; ¹H NMR δ -0.01 (s, 9H), 1.38 (s, 3H, 3 MeSi), 0.04 (s, 3H, MeSi), 0.05 (s, 3H, MeSi), 0.81 (s, 9H), 1.38 (s, 3H, 3-Me), 3.42 (d, J = 9.1 Hz, 4-H₆), 3.52 (d, J = 9.1 Hz, 4-H₆), 6.64 (s, 1H, 2-H); ¹³C NMR δ -5.6 (MeSi), -3.0 (MeSi), -2.3 (MeSi), -2.2 (MeSi), 18.2 (CSi), 18.3 (CSi), 22.6 (3-Me), 25.8 (3 x Me), 25.9 (3 x Me), 69.7 (4-CH₂), 87.3 (3-C), 143.5 (2-CH), 204.5 (1-C); m/z (APCI) 279 (M – TBS, Me and Br +H).

(+)-(S)-3,4-Di-(tert-butyldimethylsilyloxy)-3-methylbut-1-yne (40): To a stirred solution of the dibromoalkene 39 (5.25 g, 10.8 mmol) in THF (175 mL) maintained at -30 °C was slowly added butyl lithium (13 mL of a 2.5M solution in hexanes, 32.5 mmol) and the resulting solution stirred without additional cooling for 1 h then treated with saturated aqueous ammonium chloride (10 mL) and ether (50 mL). The separated aqueous layer was extracted with ether (3 x 50 mL) and the combined organic solutions washed with water (20 mL) and brine (20 mL) then dried and evaporated. Silica gel column chromatography then separated the alkyne 40 (2.60 g, 59%) as a colourless oil; [α]₀²⁶ = + 3.5 (c 1.0 g/100 mL CHCl₃); IR (film) 3311, 1472, 1389, 1361, 1254, 1190, 1119, 1032, 939, 836, 777 cm⁻¹; ¹H NMR δ 0.00 (s, 9H, MeSi), 0.01 (s, 3H, MeSi), 0.10 (s, 3H, MeSi), 0.11 (s, 3H, MeSi), 0.80 (s, 9H), 0.84 (s, 9H), 1.36 (s, 3H, 3-Me), 2.32 (s, 1H, 1-H), 3.35 (d, J = 9.5 Hz, 4-H₆), 3.47 (d, J = 9.5 Hz, 4-H₆); ¹³C NMR δ -5.3 (MeSi), -5.2 (MeSi), -3.0 (MeSi), -2.9 (MeSi), 18.0 (CSi), 18.3 (CSi), 25.7 (3 x Me), 25.9 (3 x Me), 27.4 (3-Me), 69.9 (2-C), 71.6 (4-CH₂), 72.3 (1-CH), 87.2 (2-C); m/z (APCI) 329 (M + H, 100%). HR-MS (APCI). Calcd for C₁₇H₃₇O₂Si₂ (M⁺): 329.2332. Found: m/z 329.2337.

(-)-(S)-3,4-Dihydroxy-3-methyl-1-butyne ((S)-(-)-20a): Tetrabutylammonium fluoride (7.6 mL of a 1M solution in THF) was added dropwise to a stirred solution of the foregoing alkyne 40 (1.00 g, 3.0 mmol) at rt. After 2 h, the solvent was evaporated and the residue separated by silica gel column chromatography (petrol-EtOAc 95:5) to give the (S)-dial 20a (0.25 g, 82%) as a pale yellow oil, [α]₀²⁶ = -0.7 (c 20.0 g/100 mL CHCl₃); all other spectroscopic and analytic data were identical to the racemate (±)-20a described above.

(+)-(S)-4-Acetamido-2,6-dimethyl-5-[3′,4′-dihydroxy-3′-methylbut-1′-yn-1′-yl]-3-nitroanisole (21): Following the general Sonogashira procedure, the nitro-iodide 18 (0.364 g, 1.0 mmol) was
coupled with the (S)-diol 20a (0.20 g, 2.0 mmol) using (Ph₃P)₄Pd (0.578 g, 0.5 mmol), copper(I) iodide (0.095 g) and triethylamine (3.6 mL, 26 mmol) to give the (S)-(+-)alkyne-diol 21 (0.26 g, 77%) as a pale yellow solid, mp 173-174.5 °C (MeOH); [α]D²⁶ = +3.5 (c 1.9 g/100 mL MeOH); all other spectroscopic and analytic data were identical to the racemate (±)-21 described above.

4-Amino-2,6-dimethyl-5-iodo-3-nitroanisole (14): The nitro-iodide 18 (0.32 g, 0.88 mmol) was dissolved in methanol (5 mL) before concentrated sulfuric acid (0.18 mL) was added. The resulting solution was heated to reflux for 16 h, then cooled and poured onto ice (~20 g) and the mixture basified using potassium carbonate. The product was extracted into dichloromethane (3 x 10 mL) and the combined extracts dried and evaporated. Silica gel column chromatography then separated the nitroanisole 14 (0.18 g, 64%) as a bright yellow solid, mp 214-215 °C; IR (KBr) 3375, 1602, 1455, 1259, 1047, 799, 650 cm⁻¹; ¹H NMR δ 2.22 (s, 3H), 2.40 (s, 3H), 3.58 (s, 3H, OMe), 5.25 (br s, 2H, NH₂); m/z (EI) 322 (M⁺, 100%). Anal. Caled for C₉H₁₁N₂O₃: C, 33.56; H, 3.44; N, 8.70. Found: C, 33.78; H, 3.51; N, 8.55.

(+)-(S)-4-Amino-2,6-dimethyl-5-[3',4'-dihydroxy-3'-methylbut-1'-yn-1'-yl]-3-nitroanisole (41): By the general Sonogashira procedure, coupling between the nitroanisole 14 (0.18 g, 0.55 mmol) and the (S)-(+-)diol 21 (0.11 g, 1.1 mmol) using (Ph₃P)₄Pd (0.324 g, 0.28 mmol), copper(I) iodide (0.053 g) and triethylamine (2.0 mL, 14.3 mmol) in THF (6 mL) gave the unprotected (S)-alkyne-diol 41 (0.127 g, 79%) as a yellow solid, mp 163-164 °C; [α]D²⁵ = +6.8 (c 1.7 g/100 mL MeOH); IR 3377, 2221, 1603, 1509, 1452, 1376, 1330, 1260, 1198, 1153, 1116, 1054, 998, 912, 873, 767, 733 cm⁻¹; ¹H NMR δ 1.50 (s, 3H, 3'-Me), 2.26 (s, 3H, ArMe), 2.28 (s, 3H, ArMe), 3.05 (br res, 2H, 2 x OH), 3.55 (s, 3H, OMe), 3.56 (d, J = 11.0 Hz, 3'-H₃), 3.69 (d, J = 11.0 Hz, 4'-H₆), 5.62 (br res, 2H, NH₂); ¹³C NMR δ 13.9 (Me), 15.8 (Me), 25.9 (Me), 60.9 (OMe), 69.7 (CH₂), 71.2 (C), 78.5 (C), 102.5 (C), 108.8 (C), 128.9 (C), 134.8 (C), 140.2 (C), 141.0 (C), 148.1 (C); m/z (APCI 277 (M⁻ - OH, 100%). HR-MS (APCI). Caled for C₁₄H₁₅N₂O₄ (M⁺ - OH): 277.1188. Found: m/z 277.1197.

(E)-1-(tert-Butyloxycarbonylamino)-7-(3'-penten-1'-yl)benzotriazole (42a): Following the previously reported procedure for metalation and homologation,¹⁵ treatment of the dianion 7 derived from 1-(tert-butyloxycarbonylamino)-7-methylbenzotriazole 6 (0.30 g, 1.2 mmol) using BuLi-TMEDA with crotyl chloride (0.13 mL, 1.32 mmol) gave the (E)-pentene 42a (0.35 g, 97%) as a thick yellow oil; IR (film) 3166, 1750, 1606, 1455, 1394, 1370, 1252, 1160, 1050, 966, 910, 749 cm⁻¹; ¹H NMR δ 1.30-1.53 (br res, 9H, t-Bu), 1.54-1.58 (m, 3H, 5'-Me), 2.24-2.38 (m, 2H, 2'-CH₂), 2.94 (t, J = 8.3 Hz, 1'-CH₂), 5.39-5.41 (m, 2H, 3'- and 4'-H), 7.20-7.23 (m, 2H), 7.75-7.79 (m, 1H), 8.59 (br s, 1H, NH); ¹³C NMR δ 11.9 (5'-Me), 27.0 (3 x Me), 29.3 (CH₃), 32.7 (CH₂), 82.7 (C), 117.0 (ArCH), 123.7 (ArCH), 124.3 (CH), 125.2 (CH), 127.9 (ArCH), 129.8 (7-C), 143.7 (C-N), 152.5 (C-N), 157.2
(C=O); m/z (APCI) 302 (M+, 100%). HR-MS (FAB). Calcd for C₁₆H₂₅N₄O₂ (M⁺ + H): 303.1816. Found: m/z 303.1816.

(3'R, 4'R)-1-(tert-Butyloxycarbonylamino)-7-(3',4'-dihydroxypent-1'-yl)benzotriazole (43a): Following the procedure detailed above for the preparation of diol 20a, reaction of the foregoing pentene 42a (0.273 g, 0.9 mmol) with AD-mix-β (1.26 g) at rt for 42 h gave the diol 43a (0.196 g, 65%) as a thick, pale yellow oil; IR (film) 3252, 1749, 1508, 1456, 1370, 1254, 1159, 912, 733 cm⁻¹; ¹H NMR (rotameric) δ 1.08 (d, J = 6.3 Hz, 3H, 5'-Me), 1.22-1.52 (br s, 9H, 3 x Me), 1.71-1.80 (m, 2H), 2.75 (br s, 1H, OH), 3.00-3.13 (m, 2H), 3.13 (br s, 1H, OH), 3.35-3.48 (m, 1H, 3'-H), 3.56 (qd, J = 6.3, 6.0 Hz, 1H, 4'-H), 7.22-7.25 (m, 2H), 7.81-7.85 (m, 1H), 9.16 (br s, 1H, NH); ¹³C NMR δ 18.5 (5'-Me), 26.9 (3 x Me), 33.8 (2'-CH₂), 42.2 (1'-CH₂), 69.4 (CH), 73.8 (CH), 82.4 (C), 117.0 (CH), 123.6 (CH), 124.2 (7-C), 127.8 (CH), 129.8 (C-N), 143.7 (C-N), 152.9 (C=O); m/z (APCI) 336 (M⁺, 100%). HR-MS (FAB). Calcd for C₁₆H₂₅N₄O₄ (M⁺ + H): 337.1868. Found: m/z 337.1868. [ee: ~ 76%]

(1'R, 2R)-2-(1-Hydroxyethyl)-8-iodochroman (44a): Using the same procedure for benzene generation and trapping that gave the iodochroman 28a, treatment of the foregoing diol 43a (0.074 g, 0.22 mmol) with trifluoroacetic acid (0.44 mL) and N-iodosuccinimide (0.124 g, 0.55 mmol) gave the iodochroman 44a (0.046 g, 68%) as a thick, yellow oil; [α]D²⁹ = - 7.4 (c 6.25 g/100 mL CHCl₃); IR (film) 3367, 1454, 1261 cm⁻¹; ¹H NMR δ 1.26 (d, J = 6.0 Hz, 3H, 1'-Me); 1.45-1.61 (br res, 1H, OH), 1.67-1.78 (m, 1H, 3-H₃), 1.93-1.98 (m, 1H, 3-H₃), 2.67 (ddd, J = 16.3, 5.4, 2.7 Hz, 1H, 4-H₃), 2.77 (ddd, J = 16.3, 12.1, 5.9 Hz, 1H, 4-H₃), 3.78-3.84 (m, 2H, 2- and 1'-H), 6.56 (t, J = 7.7 Hz, 1H, 6-H), 7.02 (d, J = 7.7 Hz, 1H, 7-H), 7.49 (d, J = 7.7 Hz, 1H, 5-H); ¹³C NMR δ 18.5 (Me), 23.9 (CH₂), 24.8 (CH₂), 70.0 (CH), 81.6 (CH), 85.8 (C-I), 122.2 (CH), 123.1 (4a-C), 130.3 (CH), 136.9 (CH), 142.9 (8a-C); m/z (APCI) 286 (M⁺-H₂O, 100%). HR-MS (NH₄Cl). Calcd for C₁₆H₁₇INO₂ (M⁺ + NH₄): 322.0298. Found: m/z 322.0296.

(E)-1-(tert-Butyloxycarbonylamino)-7-(4'-phenyl-3'-buten-1'-yl)benzotriazole (42b): Following the previously reported procedure for metalation and homologation,¹⁵ treatment of the dianion 7 derived from 1-(tert-butyloxycarbonylamino)-7-methylbenzotriazole 6 (0.918 g, 3.7 mmol) using BuLi-TMEDA with cinnamyl bromide (0.87 g, 4.4 mmol) gave the (E)-butene 42b (1.05 g, 78%) as a colourless oil; IR (film) 3458, 1745, 1494, 1453, 1368, 1253, 1157, 966, 750 cm⁻¹; ¹H NMR δ 1.35 (br res, 9H, t-Bu), 2.41 (td, J = 7.1, 6.7 Hz, 2H, 2'-CH₂), 3.00 (t, J = 7.1 Hz, 1'H-CH₂), 6.10 (dt, J = 15.9, 6.7 Hz, 1H, 3'-H), 6.28 (d, J = 15.9 Hz, 1H, 4'-H), 7.03-7.21 (m, 7H), 7.66-7.72 (br res, 1H, ArH), 9.32 (br s, 1H, NH); ¹³C NMR δ 28.1 (3 x Me), 30.1 (2'-CH₂), 34.0 (1'-CH₂), 83.6 (C), 118.1 (CH), 124.7 (ArC), 125.0 (CH), 126.0 (CH), 127.2 (CH), 128.5 (CH), 129.0 (CH), 137.4 (C), 144.7 (C-N), 153.9 (C-N), 154.1 (C=O); m/z (APCI) 365 (M⁺ + H, 100%). HR-MS (FAB). Calcd for C₂₃H₂₅N₄O₂ (M⁺ + H): 365.1972. Found: m/z 365.1972.
(3’R, 4’R)-1-(tert-Butyloxy carbamylamino)-7-(3’,4’-dihydroxy-4’-phenylbutan-1’-yl)benzotriazole (43b): Following the procedure detailed above for the preparation of diol 20a, reaction of the foregoing butene 42b (0.737 g, 2.0 mmol) with AD-mix-β (2.80 g) at rt for 40 h gave the diol 43b (0.65 g, 82%) as a thick, pale yellow oil; IR (film) 3498, 1735, 1453, 1368, 1253, 1157 cm\(^{-1}\); \(^1\)H NMR (rotameric) δ 1.22-1.35 (br s, 9H, 3 x Me), 1.46-1.51 (br s, 1H, OH), 1.54-1.63 (br s, 1H, OH), 2.80-2.86 (m, 2’-H\(_2\)), 2.96-3.01 (m, 1H, 2’-H\(_\beta\)), 3.53-3.57 (m, 1H, 3’-H), 3.80-3.86 (m, 2H, 1’-CH\(_2\)), 4.30 (d, \(J = 6.6\) Hz, 1H, 4’-H), 6.95 (d, \(J = 7.0\) Hz, ArH), 7.05-7.19 (m, 6H), 7.60 (d, \(J = 7.0\) Hz, 1H, ArH), 9.70 (br s, 1H, NH); \(^{13}\)C NMR δ 25.7 (2’-CH\(_2\)), 27.1 (3 x Me), 33.4 (1’-CH\(_2\)), 74.5 (CH), 77.2 (CH), 82.6 (C), 116.9 (CH), 124.8 (CH), 126.0 (C), 126.7 (CH), 127.2 (CH), 127.8 (CH), 128.5 (CH), 130.8 (C), 142.0 (C-N), 144.4 (C-N), 154.7 (C=O); m/z (APCI) 399 (M\(^+\) + H, 100%). HR-MS (FAB). Calcd for C\(_{21}\)H\(_{27}\)N\(_4\)O\(_4\) (M\(^+\) + H): 399.2027. Found: m/z 399.2025. [ee: ~ 82%].

(-)-(3’R, 2R)-2-(1-Hydroxy-1-phenylmethyl)-8-iodochroman (44b): Using the same procedure for benzyne generation and trapping that gave the iodochroman 28a, treatment of the foregoing diol 43b (0.628 g, 1.60 mmol) with trifluoroacetic acid (6.3 mL) and N-iodosuccinimide (0.90 g, 4.0 mmol) gave the iodochroman 44b (0.334 g, 57%) as a thick, yellow oil; [\(\alpha\)]\(_D\)\(^{26}\) = -19.7 (c 3.00 g/100 mL CHCl\(_3\)); IR (film) 3564, 1562, 1494; 1448, 1237, 1194, 1093, 1045, 905, 887, 761, 701 cm\(^{-1}\); \(^1\)H NMR δ 1.53-1.70 (m, 2H, 3-CH\(_2\)), 2.60-2.65 (m, 2H, 4-CH\(_2\)), 3.12 (br s, 1H, OH), 4.01 (ddd, \(J = 10.5, 7.8, 2.7\) Hz, 1H, 2-H), 4.65 (d, \(J = 7.8\) Hz, 1H, 1’-H), 6.54 (t, \(J = 7.6\) Hz, 1H, ArH), 6.91 (d, \(J = 7.6\) Hz, 1H, ArH), 7.24-7.39 (m, 5H), 7.49 (d, \(J = 7.6\) Hz, 1H, ArH); \(^{13}\)C NMR δ 23.5 (3-CH\(_2\)), 24.7 (4-CH\(_2\)), 77.5 (2-CH), 81.4 (1’-CH), 85.8 (C-I), 122.4 (CH), 123.2 (4a-C), 127.4 (CH), 128.5 (CH), 128.6 (CH), 129.7 (CH), 137.0 (CH), 139.0 (C), 152.4 (C); m/z (APCI) 366 (M\(^+\), 100%). HR-MS (NH\(_4\)-Cl). Calcd for C\(_{18}\)H\(_{19}\)INO\(_2\) (M\(^+\) + NH\(_4\)): 384.0455. Found: m/z 384.0458.

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REFERENCES


