Iso-seco-tanapartholides: Isolation, Synthesis and Biological Evaluation


Keywords: Natural products / Total synthesis / Terpenoids / Cleavage reaction / Inflammation

The isolation, identification and total synthesis of two plant-derived inhibitors of the NF-κB signaling pathway from the iso-seco-tanapartholide family of natural products is described. A key step in the efficient reaction sequence is a late-stage oxidative cleavage reaction that was carried out in the absence of protecting groups to give the natural products directly. A detailed comparison of the synthetic material with samples of the natural products proved informative. Biological studies on synthetic material confirmed that these compounds act late in the NF-kB signaling pathway.

Introduction

The search continues for bioactive compounds that can act as leads for drug discovery or as tools for biological studies. Whilst the use of large chemical libraries is of importance in this research area,[1] approaches based on natural products remain powerful.[2] Here we describe the total synthesis of plant-derived inhibitors of the NF-κB signaling pathway. This pathway plays a key role in inflammation, immunology and cancer. Compounds that modulate its activity have been linked to the treatment of several diseases.[3]

In the initial phase, bioactive plant-derived natural-product extracts were identified by using an NF-κB-based reporter-gene assay.[4] Bioactivity-guided fractionation of one of the extracts resulted in the isolation of what was believed to be the relevant components. Spectroscopic analysis led to structural assignment of the compounds as members of a natural-product family that includes the seco[5] and iso-seco-tanapartholides (Figure 1).[6] These interesting structures are believed to be biosynthesised by a Diels–Alder reaction of sesquiterpene lactone derived dienes with oxygen followed by fragmentation of the resulting cyclic peroxides to form the seco or iso-seco structures.[5,7]

Figure 1. Members of the iso-seco-tanapartholide family of natural products isolated from plants of the genus Artemisia and Achillea.

The total synthesis of members of this natural-product family was then carried out to confirm their structure and biological activity. A key step in our short and efficient sequence was a late-stage oxidative cleavage reaction that was carried out in the absence of protecting groups to give the natural products directly. A comparison of the previously reported spectroscopic data associated with the natural products proved interesting.

This study was completed by showing that the synthetic samples of the iso-seco-tanapartholides are indeed inhibitors of the NF-kB signaling pathway. We also present experiments that have enabled us to propose a mode of action for these inhibitors.

Results and Discussion

High-throughput screening of plant-derived extracts was carried out by using an HeLa57A cell line that expresses an NF-κB-dependent reporter.[4,8] Failure to trigger this re-
porter on addition of phorbol 12-myristate 13-acetate indicated that the tested extract inhibited the NF-κB signaling pathway. Whilst several extracts were active, follow-up studies focused on an extract from the plant Tanacetum parthenium. This extract was selected due to the robust nature of the bioactivity observed despite concerns that natural products with the same biological activity have been isolated previously from this plant, including the much studied parthenolide. Three rounds of bioactivity-guided fractionation resulted in the isolation of an enriched fraction that retained the desired activity. Detailed spectroscopic analysis of this fraction led to its assignment as the natural product, iso-seco-tanapartholide (reported structure 1), and our plans then focused on how an authentic sample of 1 might be prepared. However, in contrast to previous reports, our analysis showed that the sample of 1 we had isolated was in fact a mixture of two compounds. Figure 2a shows the observed signals assigned to the C3 proton and one each of the C2 and C13 protons and clearly shows that two signals are present for each proton. The two compounds present in the purified extract from Tanacetum parthenium were tentatively assigned the epimeric structures 1 and 2 (Figure 1), and our plans then focused on how an authentic sample of 1 might be prepared. However, in contrast to previous reports, our analysis showed that the sample of 1 we had isolated was in fact a mixture of two compounds. Figure 2a shows the observed signals assigned to the C3 proton and one each of the C2 and C13 protons and clearly shows that two signals are present for each proton. The two compounds present in the purified extract from Tanacetum parthenium were tentatively assigned the epimeric structures 1 and 2 (Figure 1).

Spectroscopic methods alone could not be used to assign the stereochemistry at C3 in the two proposed structures. The remote nature of the C3 stereocenter and the flexibility inherent in this structure contribute to the analysis challenges, as does the fact that it was not possible in our hands to obtain pure samples of the two epimers at this stage. This raised additional concerns about the existing structural assignments within the iso-seco-tanapartholide family and provided a further motivation for preparing authentic samples of both 1 and 2.

Retrosynthetically, we envisaged accessing the seco structure in 1 by oxidative cleavage of the C1,C10 syn-diol functionality in 7 in the absence of protection of the C3 alcohol and αβ-unsaturated lactone functional groups (Figure 3). We reasoned that if the oxidative cleavage reaction was carried out at an earlier stage in the synthesis, a considerable increase in the total number of steps would be required. It was therefore planned that the syn-diol unit in 7 would be introduced by dihydroxylation of the γ,δ-double bond in 8. This dihydroxylation was expected to proceed with high selectivity for the β-face of 8 providing the β-syn-diol 9 (Figure 3). This assumption was based on other work from our laboratory in which epoxidation of 8 with m-CPBA furnished almost exclusively the β-epoxide. With β-syn-diol 9 in hand it was expected that diastereoselective reduction of the C3 ketone and installation of the C11–C13 exo-methylene group would lead to substrates suitable for oxidative cleavage studies.

![Figure 3. Retrosynthetic analysis of iso-seco-tanapartholide.](image-url)

Our initial plan involved the use of 9 directly in the C3 carbonyl reduction reaction (Scheme 1). However, for practical reasons, we were forced to revise this by carrying out the reduction on a protected version of 9, the β-syn-diol 10. Reduction of the C3 carbonyl group in 10 was expected to occur under steric control with approach of the reducing agent occurring from the opposite face to the two protected alcohols giving 7 after removal of the silyl groups.
As pure samples of both 1 and 2 were required, it was proposed that epimerisation at C3 by using Mitsonobu protocols would be possible late in the synthesis; however, this proved unnecessary. It was planned to access the 5,7,5 guai- 
ocorrection

cols would be possible late in the synthesis; however, this posed that epimerisation at C3 by using Mitsonobu proto-

tipeudepsanolide (–)-

eudesmanolide (–)- epimerisation. However, protection of 

oxygen atoms moving from their ground state positions to 
torsional angles are 45°, 74° and 55°, respectively. The rela-
tively rigid nature of this framework presumably stops the 
system adopts conformations in which the hydroxy 
groups are positioned such that the O1–C1–C10–O10 tor-


dihydroxylation of the C3 carbonyl group. 

In practice, acid-mediated elimination of acetic acid from 

13 obtained from 11,[12] provided 8 in accordance with 
literature precedent. Subsequent dihydroxylation of 8 gave 
syn-diol 9 as the major component in a 3:1 mixture with 

13, with the reaction occurring exclusively at the γ,δ-double 
bond in 8. For ease of handling, the diol functionality in both 

9 and 13 was protected before reduction of the C3 carbonyl group. Attempts to protect the diol unit in 9 and 

13 as acetonides proved unsuccessful returning only the 

starting materials. This lack of reactivity could be rational-
ised when the X-ray structures of subsequent intermediates were obtained.[14] In the cases of 15–17, the seven-mem-
ered ring adopts conformations in which the hydroxy 
groups are positioned such that the O1–C1–C10–O10 torsional angles are 45°, 74° and 55°, respectively. The rela-
tively rigid nature of this framework presumably stops the 
oxidation atoms moving from their ground state positions to 
adopt the required conformation for successful acetonation. However, protection of 9 and 13 was achieved by 
treatment with TMSOTf yielding a 3:1 mixture of bis(TMS) ethers 10 and 14. Reaction of 10 and 14 with sodium 
borohydride provided two major isomers 15 and 16 that, 


despite being difficult to separate from each other, were 
readily separable from minor quantities of the other two possible isomers 17 and 18 (ratio 15+16/17+18 = 85:15; 
95% yield). When this purification was repeated on a larger 

scale, pure samples of 15, 16 and 17 were isolated and their structures assigned by X-ray analysis. Alcohol 15, the major 
product from the reduction of 10, and likewise alcohol 16, the major product from the reduction of 14, result from 
hydride addition to the less hindered face of the respective 
ketones, as expected. As 15 and 16 could not be separated easily, reaction of the mixture with TBSOTf resulted in tri-
sily ethers 19 and 20 that were readily separable. The pure 
diastereoisomers 19 and 20 were then carried through the 
subsequent steps independently.

The next challenge was to introduce the exocyclic α-

methylene group. Reaction of the lithium enolate generated 

from 19 with diphenyl diselenide gave exclusively 21 
(Scheme 2A).[15] The stereochemical outcome of this reac-
tion results from pseudoaxial approach of the electrophile and was confirmed by nOe studies on 21.[16] Oxidation of the selenide in 21 with concomitant elimination gave 22 as the only product due to the anti relationship between the 
C11 selenoxide and the C7 proton preventing formation of the 
C7–C11 double bond (Scheme 2A). Global desilylation by using TBAB gave triol 7 in excellent yield, and – pleasingly – 7 was then converted into 1 in 86% yield on treatment with lead tetracetate at 0 °C in dichloromethane. Interestingly, attempts to carry out this transformation by using sodium periodate returned only starting material, probably due to the fact that again the O1–C1–C10–O10 torsional angle in 7 is too large and inflexible to enable formation of the required cyclic periodate ester. Whilst lead tetracetate is known to cleave diols that cannot form the 
corresponding cyclic ester intermediate, the rates of these reactions are extremely slow.[16] Therefore, as cleavage of 7 
is complete within just 20 min, this reaction probably occurs via the cyclic intermediate that can be formed in this

Scheme 1. Dihydroxylation and reduction of the C3 carbonyl group.

Scheme 2. Installation of the exocyclic double bond, global deprotection and oxidative cleavage.
case due to the larger size of lead compared to iodine.\textsuperscript{[16]} When the reaction of \textit{7} with lead tetraacetate was run for extended times or at higher temperatures over-oxidation of \textit{1} was observed.\textsuperscript{[17]}

In an analogous sequence of reactions, the trisilyl-protected triol \textit{20} (Scheme 1) was transformed via the selenide \textit{23}\textsuperscript{[8]} into \textit{24}, a compound that contained the required exocyclic \textit{α}-methylene group (Scheme 2B). Subsequent desilylation of \textit{24} gave \textit{25} which, in an analogous manner to the conversion of \textit{7} into \textit{1}, underwent rapid and clean oxidative cleavage to yield \textit{2} (72\% over 4 steps). Comparison of the \textit{1}H and \textit{13}C NMR spectra of synthetic \textit{1} and \textit{2} with that reported for iso-\textit{sec-o}tanapartholide isolated from plants of the genus \textit{Artemisia}\textsuperscript{[6]} proved interesting. The spectra for both \textit{1} and \textit{2} correlated very closely with those reported for the natural product. In the absence of authentic material from these sources, further insight was gained by comparing the optical rotation of the natural material\textsuperscript{[6]} with those of synthetic \textit{1} and \textit{2}.\textsuperscript{[18]} Despite the small values involved, it appears that the relative and absolute stereochemistry assigned to iso-\textit{sec-o}tanapartholide from \textit{Artemisia} was correct (as in \textit{1}).

However, the situation was complicated when a comparison of synthetic \textit{1} and \textit{2} with our material from \textit{T. parthenium} and a sample from a plant of the genus \textit{Achillea}\textsuperscript{[19]} was carried out. NMR studies (Figure 2) showed that the bioactive material we had isolated from \textit{T. parthenium} was indeed a mixture of the two epimers \textit{1} and \textit{2} with the major isomer isolated from this plant having the same relative stereochemistry as \textit{2} [compare Figure 2a and c (full line)]. In addition, comparison of our authentic samples of \textit{1} and \textit{2}, with material isolated from a plant of the genus \textit{Achillea}\textsuperscript{[19]} gave an analogous result, again confirming that the major isomer present in this sample was \textit{epi-iso-sec-o}tanapartholide \textit{2}. Further optical rotation comparisons\textsuperscript{[8,18]} also supported our conclusion that the major isomer isolated from \textit{Achillea} had the same absolute stereochemistry as \textit{2}. For a more detailed comparison of synthetic \textit{1} and \textit{2} with the previous literature reports of their isolation, see Supporting Information.

Biological analysis\textsuperscript{[4,20]} of synthetic \textit{1} and \textit{2} confirmed that they inhibited the TNF\textsubscript{α} activation of NF-κB [\textit{IC}_{50}(1) = 7.7 \pm 0.6 \mu M; \textit{IC}_{50}(2) = 4.7 \pm 0.2 \mu M; Figure 4A] over a concentration range similar to that observed for our isolated material.\textsuperscript{[8]} To establish the mode of inhibition, cells were treated with the NF-κB activator TNF\textsubscript{α} and levels of the NF-κB inhibitor IκB\textsubscript{α} determined by western blotting. In response to TNF\textsubscript{α}, IκB\textsubscript{α} is rapidly degraded and then resynthesized as the IκB\textsubscript{α} gene is NF-κB-dependent (Figure 4B; DMSO vector). However, in the presence of \textit{1} and \textit{2}, IκB\textsubscript{α} was degraded, but resynthesis of IκB\textsubscript{α} was not observed. This indicates that \textit{1} and \textit{2} do not prevent IκB\textsubscript{α} degradation but block the transcriptional activity of NF-κB. Immunofluorescence studies were also consistent with this assumption.\textsuperscript{[8]} In addition, we determined the effect of \textit{1} and \textit{2} on the DNA binding activity of NF-κB. Recombinant-purified DNA-binding domains of the p50 and p65 subunits of NF-κB were incubated with \textit{1} and \textit{2} and a gel electrophoresis DNA-binding assay performed.\textsuperscript{[8]} Both \textit{1} and \textit{2} inhibited NF-κB-DNA binding in a dose-dependent fashion only when a thiol-based reducing agent was absent from the assay. The observed loss of biological activity of \textit{1} and \textit{2} in the presence of a thiol is consistent with a mode of action for these compounds in which covalent modification of cysteine residues in either the p50 or p65 subunit of NF-κB occurs. In addition, a close analogue of \textit{1} in which the exocyclic \textit{α}-methylene group was replaced by a methyl group [C11-(S)] did not inhibit TNF\textsubscript{α} activation of NF-κB, again suggestive of a role for the exocyclic \textit{α}-methylene group in \textit{1} as an electrophile. Interestingly, covalent modification of cysteine by inhibitors of NF-κB activation, including the sesquiterpene lactones, has been proposed previously.\textsuperscript{[4b,21]} In particular, a computational model of the proposed covalent binding mode of the sesquiterpene lactone, helenanin, to Cys38 and Cys120 of the p65 subunit of NF-κB has been described.\textsuperscript{[22]} Attempts to overlap the structure of the iso-\textit{sec-o}tanapartholides onto this model suggest that the C3 functional group would be expected to point away from the protein. If correct, this would provide an explanation for the observed disruption of the NF-κB-DNA inter-

![Figure 4](image-url) Bioactivity of \textit{1} and \textit{2}: (a) NF-κB-dependent reporter gene assay; (b) IκB\textsubscript{α} western blot analysis after TNF\textsubscript{α} activation.
action by both 1 and 2, in a manner that is independent of the C3 stereochemistry.

Conclusions

The bioactivity-guided fractionation of an extract from Tanacetum parthenium by using an NF-κB-dependent luciferase reporter gene assay is described. The purified extract was shown to contain two natural products from the iso-seco-tanapartholide family. Synthesis of authentic samples of these two natural products by using an efficient oxidative cleavage reaction clarified the structures of the compounds isolated from a series of plants. This resulted in the isolation and synthesis of the natural product epi-iso-seco-tanapartholide (2) as well as iso-seco-tanapartholide (1). Biological studies on synthetic material confirmed that these compounds act late in the NF-κB signaling pathway.

Supporting Information (see footnote on the first page of this article): Details of the Strathclyde natural-product extract collection, screening results, bioactivity-guided purification, structural assignment, experimental procedures, characterisation data for all new compounds, comparison studies of synthetic 1 and 2 with previously reported natural products and material isolated from T. parthenium and Achilles.

Acknowledgments

We would like to thank The Scottish Funding Council, SULSA and The Royal Society (NJW) for funding and Professor Mike Ferguson and Drs. C. Botting, A. McCarthy, and S. Fujihara for advice.


[9] It was also confirmed that the active fraction did not target the reporter protein, firefly luciferase, by using an in vitro assay (data not shown).
[11] As expected, this led to the C3 epimer of 7 as the major product.
[14] CCDC-734445 (for 15), -734446 (for 16), -734447 (for 17), and 734449 (for S1)$[iii]$ contain the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.
[17] It cannot be ruled out that lead tetraacetate rapidly cleaves 7 and 25 without forming a cyclic ester intermediate and that the speed of the reaction results from release of steric congestion on cleavage of the C1–C10 bond. We thank a reviewer for raising this possibility.
[18] $[ii][ii]α$D的功效(I) = +2.9 (c = 0.8, CHCl3); $[ii][ii]α$D的功效(2) = −6.5 (c = 0.2 CHCl3).

Received: September 6, 2009
Published Online: October 7, 2009