Natural Products

Aruncin B: Synthetic Studies, Structural Reassignment and Biological Evaluation

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Abstract: A ring-closing alkene metathesis (RCM)/ oxyselenolation-selenoxide elimination sequence was established to the sodium salts E- and Z-25 of the originally proposed structure for the recently isolated cytotoxin aruncin B (1), as well as to the sodium salt Z-34 of a related ethyl ether regioisomer; however, none of their corresponding free acids could be obtained. Their acid sensitivity, together with detailed analysis of the spectroscopic data indicated that profound structural revision was necessary. This led to reassignment of aruncin B as a Z-γ-alkylidenebutenolide Z-36. Although a related RCM/ oxyselenolation-selenoxide elimination sequence was used to confirm the γ-alkylidenebutenolide motif, a β-iodo Morita-Baylis–Hillman reaction/ Sonogashira cross-coupling-5-exo-dig lactonisation sequence was subsequently developed, due to its brevity and flexibility for diversification. Aruncin B (36), together with 14 γ-alkylidenebutenolide analogues, were generated for biological evaluation.

Introduction

Inhibitors of antiapoptotic proteins from the Bcl-2 family are currently considered to be promising leads for drug discovery programs, especially for the treatment of drug-resistant tumours. [1] In 2011, Woo and co-workers reported the isolation of several previously unknown cytotoxins from the aerial parts extract of Aruncus dioicus var. kamtschaticus, a plant found on the Korean island Ulleungdo. [2] Among these, aruncin B (1, Figure 1) stood out: early biological studies showed inhibitory properties towards Bcl-2 family proteins, with some selectivity for malignant over human cells. [3] The structure of aruncin B (1) was originally assigned using spectroscopic methods, as a monoterpeneoid containing a dihydropyranone core. [4] It features an unusual cross-conjugated exocyclic enol ether and, as originally reported, a potentially acid-labile tertiary allylic ethyl ether and carboxylic acid functionality within the same structure. The olefin geometry and absolute configuration remained undetermined. Its unusual structural features and promising biological activity made aruncin B (1) an interesting target for synthetic studies. In this Full Paper, we detail our synthetic studies towards the originally reported structure of aruncin B (1), as well as two closely related isomeric structures, its structural revision and synthetic routes leading to the correct structure. [5] The conciseness and convergent nature of the latter enabled access to a library of analogues for biological evaluation.

Retrosynthetically, aruncin B (1) could be derived from a dihydropyranone core (A, Scheme 1), with installation of the key enol ether being achieved through an ethoxyselenolation-selenoxide elimination sequence (C→B→A). [6] Dieno C would be obtained by ring-closing alkene metathesis (RCM) of an appropriate precursor triene (D). [7]

Results and Discussion

Synthesis of a model dihydropyranone 8

So as to establish the viability of the above strategy (Scheme 1), we first examined a simplified system that lacked the α-side-chain (i.e., R=H). α-Alkylation [7] of the bis allylic alcohol 2 (Scheme 2, obtained by addition of vinylmagnesium bromide to 3-methyl-2-butenal), [8] using the α-bromo Weinreb amide 3 (prepared from bromoacetyl bromide and NO-dime-thylhydroxylamine), [9] proceeded in modest yield. Subsequent addition of vinylmagnesium bromide gave the acid-labile enone 4. [10] RCM using Grubbs II cat. (GII) produced an inseparable 3:1 mixture of desired dihydropyranone 5 and terminal...
olefin-containing dihydropyranone 6; the latter likely being generated from enone 4 by competing initiation at the gem-dimethyl-substituted olefin. The less reactive Grubbs I cat. (GI) prevented metathetical involvement of the more sterically demanding gem-dimethyl-substituted olefin, and delivered exclusively the desired dihydropyranone 5.

Ethoxyselenylation of dihydropyranone 5 using PhSeBr/ EtOH in the presence of Et3N[11] delivered ethoxyselenide 7, isolated as a single diastereoisomer. The same level and sense of diastereoselectivity (typically >1,3 JOC, 1.5 Hz, Scheme 3) was also found in the ethoxyselenylation of all dihydropyranones examined later in this work, including selenide 23 (Scheme 7) where the relative configuration was established by single crystal X-ray diffraction studies.[12] Selenation is likely to occur from a conformer (E, Scheme 3) of dihydropyranone 5, where the isobutyl group is pseudo-equatorial, and allylic strain is minimized. The cyclic ether oxygen atom may direct approach of the electrophilic selenium species by coordination[12a] (as shown in E); this coordination would also enhance the e-ao acceptor ability of the ring C–O bond, thus increasing the relative e-ao donor ability of the allylic ring C–C bond (vs. the ring C–O bond), which already preferentially directs the attack anti to itself. This diastereoselectivity is also found in methoxyselenogation of 5- and 6-membered cyclic ethers.[12]

Hydrogen peroxide-induced oxidation[13] of ethoxyselenide 7, followed by syn-elimination of the in situ formed selenoxide, gave enol ether 8 as a separable mixture of E/Z-isomers (Scheme 2). Starting from the single diastereoisomer of ethoxyselenide 7, this sequence was expected to deliver only the E-isomer (Scheme 3). TLC monitoring of the reaction showed initial formation of the E-isomer, which isomerises under the reaction conditions and on standing.[14] One possible mode of isomerisation is via reversible conjugate addition to the δ-position of the diene. The configurational lability of the E-isomer could be explained by a[14] strain: C(S)H (vs. ring O) interaction with CMesOEt (see E-8, Scheme 3). Configurational lability of the exocyclic olefin was also found in the other γ-alkylidenedihydropyran substrates subsequently synthesised. E/Z-Configurations were assigned by comparison of the C(S)H chemical shifts (>8.0 ppm and 7.0–7.5 ppm, respectively).

The above model study showed that a RCM/ ethoxyselenation-selenoxide elimination strategy could be used to form the cross-conjugated pyranone motif. The ethoxyselenation-selenoxide elimination sequence proved to be a mild method to access the sensitive enol ether functionality. In addition, the configurational lability of the E-isomer suggested that the natural product was more likely to possess a Z-configured exocyclic enol ether.

Studies towards the originally reported structure

Synthesis of dihydropyranone 14

The viability of the above model study encouraged investigation of a system bearing an α-side-chain appropriate for aruncin B. tert-Butyl α-bromoacetate (10, Scheme 4) proved a more effective partner than the Weinreb amide derivative 3 (Scheme 2) for the O-alkylation of 3-methyl-2-butenyl-derived crude alcohol 2; the latter being achieved under mild phase-transfer catalysis (Scheme 4).[15] After conversion of the resulting ester 11 to its corresponding β-keto phosphonate 12[16] Villiéras modification[17] of the Horner–Wadsworth–Emmons reaction provided the RCM precursor, triene 13, while installing the required extra carbon atom at the α-position (missing in model triene 4). Triene 13 also possessed a greater (acid) stability than triene 4. This sequence could be carried out with either diethyl or dimethyl methylphosphonate in similar yields (50-60% over 2 steps); however, dimethyl methylphosphonate was preferred due to its...
higher water solubility, enabling removal of excess during aqueous workup, rather than by distillation. In contrast to the model study, RCM of triene 13 using GI did not give hydroxydihydropyranone 14 in useful yield (25%), probably due to the slightly more sterically demanding nature of the α-hydroxymethyl-bearing olefin partner. G II proved more promising, giving dihydropyranone 14 along with dihydropyranone 15 (ca. 9:1 mixture, inseparable); however, the latter could be converted to 14 by subsequent addition of excess 2-methyl-2-butene.

Dihydropyranone 14 possesses all the carbon atoms of the originally assigned core structure of aruncin B (1).

Original late-stage function group interconversion (FGI) attempt

The ethoxyselenation-selenoxide elimination sequence previously described (Scheme 2) was carried out on dihydropyranone 14 (Scheme 5) to give, via ethoxyselenide 16, a (separable) mixture of Z/E-enol ethers 17. Unfortunately, these intermediates undergo rapid decomposition on standing and could not be used in a subsequent (oxidation) step. However, oxidation of the ethoxyselenide 16 using excess Dess–Martin periodinane (DMP) resulted in simultaneous selenide oxidation-selenoxide elimination and oxidation of the primary alcohol to the corresponding aldehyde, resulting in the more stable Z-ketoaldehyde 18. Oxidation to the ketoacid 19, proved very challenging, proceeding at best in only 4% yield using CrO3 in pyridine; Luche reduction of ketoacid 19 failed to deliver aruncin B (1).

Redox interchange and FGI endgame

The above results suggested that it might be necessary to have the correct ring and α-side-chain oxidation levels in place prior to installation of the sensitive enol ether functionality; an initial redox interchange was therefore considered. 1,2-Reduction of dihydropyranone 14 using DIBAL-H, followed by selective TEMPO-catalysed oxidation of the primary alcohol delivered hydroxyaldehyde 20 (3:1 d.r., Scheme 6). For ease of spectral analysis diastereisomer separation was carried out at this point and the rest of the sequences presented were carried out on the major diastereoisomer 20 mj. Relative stereochemistry was originally assigned by comparison of 3J values of the major contributing half-chair conformer of each diastereoisomer. In both diastereoisomers, the isobutenyl group is expected to be pseudo-equatorial; for the major diastereoisomer, the higher 3J values (5 Hz) observed are characteristic for the oxygen substituent residing pseudo-equatorial, corresponding to the trans isomer 20 mj. For the minor diastereoisomer 20 mn, lower 3J values (<3 Hz) are characteristic for the oxygen substituent being pseudo-axial. This preference for the formation of the trans isomer has also been found in the 1,2-reduction of a 4-substituted α-(hydroxymethyl)cyclohexanone. Oxidation of the hydroxyaldehyde 20 mj using alkaline AgNO3 delivered the hydroxycacid 21. Reaction of the latter with PhSeBr in the presence of an inorganic base was used to obtain ethoxyselenide acid 22. Various oxidation conditions (H2O2, NaIO4, mCPBA/ iPr2NH or O3) were examined with acid 22 to lastly install the enol ether, but all failed to generate aruncin B (1); only decomposition was observed.

The efficiency and chemoselectivity of the above aldehyde to acid oxidation (20 mj — 21) encouraged
consideration of leaving such an oxidation to the last step. Initial ethoxyselenation of hydroxylaldehyde 20mj proceeded again diastereoselectively (Scheme 7). The relative stereochemistry was confirmed by single crystal X-ray diffraction studies on the 3,5-dinitrobenzoate derivative of ethoxyselenide 23[11] (see Supporting Information). This analysis served not only to confirm the ethoxyselenation diastereoselectivity rationalised earlier (Scheme 3), but also the diastereoselectivity in the DIBAL-H-induced 1,2-reduction of dihydropyranone 14 (Scheme 6). Subsequent oxidation of selenide 23 using NaIO₄ resulted in the Z-enol ether 24. Oxidation to the acid, using silver(I) oxide in the presence of NaOH[27] gave the sodium salt 25 which corresponded to the conjugate base of the originally reported structure of aruncin B (1). Various mild acidic conditions were then examined (workup using pH 6 buffer solution, low temperature addition (≤ 78°C) of accurately measured 1.0 equiv of HCl, carboxylic acid-supported ion-exchange resin (amberlite IR-50)), but only led to decomposition.

Since the geometry of the enol ether was not unambiguously assigned in the original isolation report of aruncin B (1, Figure 1), we were interested in establishing whether the E-isomer would present a higher stability towards acidic conditions. As noted earlier, the E→Z-isomerisation (Scheme 2) was suspected to occur in part by reversible conjugate addition to the δ-position of the dienone 17, 18 and dienal 24; we therefore considered a slightly less electron-withdrawing system, such as a dienoate, might retard the rate of isomerisation and enable access to the E-enol ether configuration.

TMSCHN₂-induced esterification of acid 21 gave hydroxyester 26 (Scheme 8). Ethoxyselenation, followed by oxidation resulted in the formation of the E-enol ether 27, which showed a much lower rate of isomerisation compared with dienal 24. Subsequent saponification gave the sodium salt E-25, which however, similarly to its Z-isomer, could not be acidified.

The above results (attempted conversion of 19, 22, Z-25 and E-25 to 1) suggested that structure 1 could not exist (as a free acid) nor correspond to aruncin B: it presents intrinsic elements of instability. Although we were unable to isolate any decomposition product(s) that may have helped to clarify the decomposition pathway(s), we consider the carboxylic acid functionality likely initiates decomposition via ethoxy protonation leading to facile loss of EtOH from the tertiary allylic position assisted by the enol ether moiety.

**Synthesis of regioisomeric ether**

Comparison of 13C NMR data for Z-25 and E-25 with reported values for aruncin B (1) indicated notable inconsistencies in the C(3), C(8), and C(10) chemical shifts (Figure 2), and we therefore considered that the site of etherification could be misassigned.

Further detailed comparison with 13C NMR data of other known tertiary allylic ethyl ethers (28[28] and (E)-O-ethylsubereinol (29)[29] Figure 2) and alcohols (30[30] and 31[31]), suggested that while the C(8) and C(10) chemical shifts observed in Na salts Z- and E-25 are not anomalous,[28,29] the shift observed and originally assigned in the natural isolate is more consistent with the tertiary allylic position being an alcohol.[28,30]
In addition to the above analytical considerations, the presence of a free hydroxy group at the tertiary allylic position might provide a lower acid sensitivity, due to its lower basicity compared with the ethoxy group, thus limiting initial protonation of the oxygen atom, suspected to trigger decomposition by loss of H₂O or EtOH, respectively. We therefore focused on synthesising an alternative regioisomic structure (34, Scheme 9) featuring the ethyl ether at C(3).

![Scheme 9. Synthesis of the regioisomeric structure 34.](image)

O-Ethylation of hydroxyaldehyde 20mj in neat EtI in the presence of Ag₂O[31] and 4 Å MS, was accompanied by oxidation of the aldehyde to the corresponding acid and subsequent esterification to give the diethyl ether ester 32. In a similar fashion to ethoxyselenation described previously, hydroxyselenylation of ester 32 could be carried out using PhSeBr in MeCN/H₂O[32], followed by oxidation with hydrogen peroxide to give Z-enol ether 33. Obtention of the Z-isomer contrasts with the results observed with the dienoate[27]. ¹H NMR analysis of the crude mixture following hydroxyselenylation-oxidative elimination of ester 32 indicated traces of E-33: chemical shifts for C(5)H (and C(7)H) 8.18 (and 5.51 ppm) (CDCl₃), respectively; these values are higher than those seen for dienoate E-27 (8.03 (and 5.35 ppm) (CDCl₃), respectively). This difference may indicate slightly higher electron deficiency in dienoate E-33 compared with E-27, which might account for the greater E → Z-isomerisation of dienoate 33. Saponification of dienoate 33 delivered the corresponding sodium salt 34, which however featured similar acid sensitivity to sodium salts 25.

The acid instability of the dihydropyranolsodium salt derivatives and the inconsistencies observed in the analytical data led us to conclude that the originally assigned structure of aruncin B or the O-ethyl variant could not be correct, and a more profound structural revision was necessary.

### Studies towards a newly postulated structure

**The Z-γ-alkylidenebutenolide hypothesis**

Reported UV data for aruncin B are consistent with an γ-alkoxy-αβγδ-unsaturated carboxyl motif being present in the molecule.[33,34] However, acid sensitivity of the tertiary allylic ether (alcohol) motif cast doubt on its coexistence with the carboxylic acid functionality, and we therefore considered that the carboxyl motif could be esterified. The ¹³C NMR shift for the ring CH₂ (in the original assignment) suggested a better match for a primary alcohol rather than an ether (Figure 3).[30]

As discussed earlier (Figure 2), the tertiary allylic position most likely corresponds to a free alcohol, with the ethyl group being borne by a secondary alcohol and the reported ¹³C shift for C(3) is more consistent for an ether rather than a free alcohol (Figure 3).

These considerations led us to postulate a γ-alkylidenebutenolide Z-36 as an alternative structure for aruncin B (Figure 4), with the ester being part of the ring and the dioxygenated moiety being a side-chain featuring a free primary alcohol and an ethyl ether of a secondary alcohol.

![Figure 3. Comparison of selected ¹³C NMR data between 1 and eupaglehnin D (35).](image)

![Figure 4. Alternative structure 36.](image)

The γ-alkylidenebutenolide motif is found in various previously reported natural products (37–44, Figures 5 and 6) including the more complex novaxenicid D (43)[46] and an oxidised tirucallane triterpenoid 44.[41] Comparison of ¹H NMR shifts with butenolide 37[36] is indicative of a Z-enol ether. Moreover, ¹³C NMR values of the secondary alcohol (ether) centre observed in compounds 38[37] and 39[38] are also consistent with the site of etherification being at the secondary alcohol. Comparison between 43 and 44, is also consistent with the tertiary allylic position being an alcohol. The revised assignment of aruncin B as 36 was only contradicted by IR data, with an anticipated C=O stretch value for this type of butenolide of 1755 cm⁻¹[40] and a reported value[20] of 1724 cm⁻¹ (KBr disc).

However, as we were informed that the original IR spectrum could not be located and none of the natural isolate was available for further analysis,[42] synthetic studies towards the revised structure were undertaken.

*Chem. Eur. J.* 2017, 23, 16525 – 16534
Although there are many known methods to γ-alkylidenebutenolides,[43] we were interested in studying whether a RCM/hydroxyselenation-selenoxide elimination sequence, which proved successful to γ-ethylidenedihydropyranoate compounds, would also enable access to the γ-ethylidenebutenolide motif. We first examined a synthesis of the model butenolide 52 (Scheme 10), lacking the α-dioxygenated side chain. Addition of vinylmagnesium bromide to acrolein,[44] followed by trapping of the in situ generated magnesium alcoholate by acryloyl chloride gave the RCM precursor, triene 47 (Scheme 10). RCM-cross-metathesis (CM) cascade using GII in the presence of 2-methyl-2-butene delivered the desired lactone 48, as an inseparable (10:1) mixture with lactone 49.[46, 18] This RCM-CM strategy avoids the intermediacy of triene 50 (from 3-methyl-2-butenal and vinylmagnesium bromide, as described earlier), which would be required in a classical RCM approach and proved to be prone to allylic rearrangement (during aqueous workup and chromatographic purification). As observed with the 6-membered ring system (Scheme 3), hydroxyselenolactone 48 proceeded with complete diastereoselectivity; the relative stereochemistry of hydroxyselenide 51 was assigned by analogy with the 6-membered ring system, on the basis of similar $J_{SOH-SeCH}$ values (≈1.5 Hz). The hydroxyselenide derived from lactone 49 was removed during chromatographic purification of hydroxyselenide 51. Finally, oxidation-selenoxide elimination of 51 delivered the Z- and E-isomers of 52. As in the dihydropyranone series, initial E-isomer formation and isomerisation over the time was observed, which again indicates a thermodynamic preference for Z-geometry. $^1$H NMR $\beta$-CH data (8.20 and 7.34 ppm for E- and Z-52, respectively; 7.47 ppm for aruncin B (1)) indicates aruncin B should possess Z-geometry.

**First-generation synthesis of butenolide 36**

With the viability of a RCM/hydroxyselenolactone-selenoxide elimination sequence to the γ-alkylidene butenolide motif established (Scheme 10), the synthesis of the butenolide 36 via α-functionalisation of lactone 48 was examined (Scheme 11), using a Morita–Baylis–Hillman (MBH) reaction, followed by appropriate FGI and hydroxyselenolactone-selenoxide elimination.

Remarkably, MBH reaction of lactone 48 with aldehyde 53,[47] using a MgI$_2$/TMEDA complex catalytic system delivered butenolide 54 as an inseparable E/Z-mixture. The concomitant and fortuitous allylic oxidation presumably occurred via hydroxy atom abstraction at the bis-allylic position; this aerobic oxidation was not observed on the 6-membered ring system, and allylic hydroxy atom abstraction is probably favoured on the 5-membered ring system by generation of the aromatic oxyfuran radical (show in parenthesis in Scheme 11). This unexpected oxidation avoids carrying out the hydroxyselenolactone-selenoxide elimination steps. Reaction of butenolide 54 in neat ethyl iodide in the presence of Ag$_2$O, facilitated by the extraordinary reactivity of the allylic hydroxy group, delivered the natural occurring γ-alkylidenebutenolides, with selected NMR data compared against related reported values for aruncin B (in parentheses).

**Figure 5.** Naturally occurring γ-alkylidenebutenolides, with selected NMR data compared against related reported values for aruncin B (in parentheses).
ciliated O-alkylation of the secondary alcohol, to give ether 55 as a single (Z)-geometric isomer. Fluoride-induced removal of the TBS group gave the free primary alcohol 36. Surprisingly, the presence of fluoride apparently promotes loss of geometrical integrity of the exocyclic unsaturation, to give the alcohol 36 as a (separable) 3:1 E/Z-isomeric mixture; the major product being the anticipated thermodynamically disfavored isomer. Analytical data recorded for alcohol Z-36 corresponded to those reported for aruncin B (see Supporting Information for details), except for the IR C=O stretch which had the value as discussed and anticipated earlier (1755 cm⁻¹, film or KBr disc).

Although the synthetic route shown in Scheme 11 delivered sufficient material to confirm the true structure of aruncin B, it was impaired by two major issues: 1) transformation of butenolide 48 to ether 54 was low yielding and poorly reproducible, and 2) loss of geometrical integrity during fluoride-induced de-silylation.

**Second-generation synthesis**

So as to address the above issues, we evaluated construction of the butenolide using a one-pot Sonogashira cross-coupling-S-exo-dig lactonisation[49] (Scheme 12). The dioxgenated side-chain and the Z-vinyl iodide functionality necessary for the final cross-coupling were installed simultaneously, by way of a β-iodo-MBH[50] between commercially available methyl propiolate and aldehyde 53 using in situ prepared MgI₂ as a source of iodide.[51] O-Ethylation of the resulting iodoacrylate 56 was carried out as previously described (Scheme 11) to give ethyl ether 57 (Scheme 12), followed by hydrolysis to give the hydroxyacid 58. Saponification was accompanied by desilylation, thus avoiding the potentially problematic use of fluoride in a subsequent step. The final step with 2-methyl-3-butyn-2-ol (59) was carried out on up to 1 mmol scale, with a low catalyst loading, and provided sufficient aruncin B (Z-36) (≈ 200 mg) for biological evaluation.

![Scheme 12. Synthesis of aruncin B (Z-36) via β-iodo MBH/ Sonogashira cross-coupling-S-exo-dig lactonisation sequence.](image)

**Possible origin of aruncin B**

Previously,[2] we postulated that the low specific rotation of natural aruncin B compared with synthetic enantiopure samples,[52] together with the unusual presence of the ethoxy group at the β'-position of the true structure of aruncin B, indicated that the latter might not be an actual natural product. Rather, we suggested that aruncin B was an artefact of the isolation process, resulting from a poorly diastereoselective incorporation of EtOH (the solvent used for the extraction)[53] to an enantioenriched natural product precursor (Scheme 13). We originally mooted that the natural precursor of aruncin B was likely to be a α-(2-hydroxyethylidene)-γ-lactone.[49] Following our initial disclosure of the structural revision of aruncin B, Zidorn and co-workers, reported the isolation of natural products from the European plant *Aruncus dioicus* var. *vulgaris*.[53] Some of the isolated monoterpenoids from the latter were identical to those isolated from *Aruncus dioicus* var. *kamtschaticus*. Based on our synthetic studies, Zidorn and co-workers revised the structure of three of them. They also reported the isolation of γ-lactone 60 and its aglycon 61 (Scheme 13); we consider the latter to be a reasonable precursor to aruncin B, through conjugate addition of EtOH/ loss of H₂O, followed by allylic oxidation—the latter resembling the oxidation observed in the generation of butenolide 54 (Scheme 11), also occurring in EtOH.

**Synthesis of analogues and biological evaluation**

**Synthesis of butenolide analogues of aruncin B**

Taking advantage of the brevity and functional group tolerance of the second-generation synthetic route to the correct structure of aruncin B (Scheme 12), this was applied to the synthesis of a range of analogues by varying different reaction components (Scheme 14). Analogues of aruncin B at the α-side-chain were obtained using a variety of aldehyde during the β'-iodo-MBH step. Higher yields were observed with non-enolisable aldehydes (Table 1, entries 3–4). The rest of the sequence was followed as for aruncin B using EtI and alkynol 59, except with R¹ = ΜBu (Table 1, entry 3), for which the hydrolysis step was carried out using TMSOK in refluxing THF.[54]
O-Alkyl-substituent analogues were obtained starting with iodo-acrylate 56 using different alkyl halides in the presence of Ag\textsubscript{2}O (Table 2). The rest of the sequence was carried out as presented previously.

| Entry | R\textsuperscript{1} | I. Yield [%] | II. Yield [%] | III. Yield [%] | IV. Yield [%] | Butenolide |
|-------|----------------------|--------------|---------------|---------------|---------------|------------|
| 1     | CH\textsubscript{3}OBn | 52           | 59            | 71            | 53            | 62         |
| 2     | Me                   | 41           | 44            | 71            | 54            | 63         |
| 3     | iPr                  | 85           | 20            | 58            | 58            | 64         |
| 4     | Ph                   | 78           | quant.        | 75            | 52            | 65         |
| 5     | nC\textsubscript{8}H\textsubscript{17} | 51          | 62            | 65            | 54            | 66         |

Table 3. O-Alkyl analogues.

| Entry | R\textsuperscript{2}Hal | I. Yield [%] | II. Yield [%] | III. Yield [%] | IV. Yield [%] | Butenolide |
|-------|--------------------------|--------------|---------------|---------------|---------------|------------|
| 1     | (R\textsuperscript{2} = H) | -            | 60            | 42            | 67            |            |
| 2     | Me                       | 77           | 98            | 67            | 68            |            |
| 3     | iPr                      | 30           | 86            | 39            | 69            |            |
| 4     | BnCl                     | 46           | 64            | 52            | 70            |            |

Derivatives at the δ-position were obtained by using different alkynes in the final cross-coupling step with hydroxy acid 58 (Table 3).

Table 4. IC\textsubscript{50} [μM] values for aruncin B analogues\textsuperscript{(56)}

| Entry | Butenolide | Derivatisation | Jurkat T cells | Human T cells |
|-------|------------|----------------|----------------|---------------|
| 1     | 62         | R\textsuperscript{1} = CH\textsubscript{3}OBn | 22.5            | 15.3          |
| 2     | 63         | R\textsuperscript{1} = Me        | > 50            | > 50          |
| 3     | 64         | R\textsuperscript{1} = iPr       | > 50            | > 50          |
| 4     | 65         | R\textsuperscript{1} = Ph        | 11.6            | 9.4           |
| 5     | 66         | R\textsuperscript{1} = nC\textsubscript{8}H\textsubscript{17} | 2.9             | 8.2           |
| 6     | 67         | R\textsuperscript{1} = H         | 26.3            | 23.5          |
| 7     | 68         | R\textsuperscript{1} = Me        | 12.5            | 16.5          |
| 8     | 69         | R\textsuperscript{1} = iPr       | 23.0            | > 50          |
| 9     | 70         | R\textsuperscript{1} = Bn        | 5.6             | 9.1           |
| 10    | 71         | R\textsuperscript{1} = CMe\textsubscript{2}OMe | > 50            | > 50          |
| 11    | 72         | R\textsuperscript{1} = iPr       | > 50            | > 50          |
| 12    | 73         | R\textsuperscript{1} = CH\textsubscript{3}OH | 3.6             | 1.2           |
| 13    | 74         | R\textsuperscript{1} = Ph        | > 50            | > 50          |
| 14    | 75         | R\textsuperscript{1} = nC\textsubscript{8}H\textsubscript{17} | > 50            | > 50          |

assays were carried out with a high number of cells (5×10\textsuperscript{3}) over 20 h.\textsuperscript{(55)} This short assay duration could potentially compromise detection of biological activity. Therefore, we tested over a longer time period (3 cell cycles), reducing the cell number (1.5–5×10\textsuperscript{3}) to avoid confluence in control wells. The most potent compounds with our assays were also tested using the method described in the original publication, so as to enable comparison between the two methods.

rac-Z-Aruncin B, and (+)- and (−)-Z-aruncin B\textsuperscript{(52)} as well as rac-E-aruncin B, were first tested against Jurkat T cells. These aruncin B isomers showed similar activity (IC\textsubscript{50} values [μM] of 16.0, 9.6, 9.7 and 9.1, respectively)\textsuperscript{(56)} and compared favourably with that originally reported for aruncin B (70 μM). The similar activity suggests that the side-chain stereochemistry is probably not directly influential in the binding/interaction with the target(s). Similar activity observed with the E-isomer could be explained by isomerisation, once the compound is present in the cell medium.

Activity of the analogues against Jurkat T cells and human T cells was then tested. For the α-side-chain derivatives (Table 4, entries 1–5), although a clear trend is not discernable, modifications at this moiety strongly affect the activity. Aryl butenolide 65 (entry 4) showed good activity against Jurkat T cells but is poorly selective, whereas n-octyl butenolide 66 (entry 5), showed greater activity against Jurkat T cells and a better se-

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lectivity towards the latter. Variation at the O-ether substituent (Table 4, entries 6–9) does not seem to significantly affect activity or selectivity. Changes at the δ-alkylidene substituent (Table 4, entries 10–14) appeared to be crucial; only butenolide 73 (entry 12), featuring a free alcohol showed activity. Compounds featuring a similar steric character to aruncin B on the δ-alkylidene moiety, such as butenolides 71 and 72 (entries 10, 11), did not show significant activities. Those observations suggest that the presence of a hydrogen bond donor substituent is critical for the activity. Other substituents such as an aromatic ring or a long aliphatic chain, 74 and 75, respectively (entries 13,14), did not show activity.

To summarise the above observations, it appears that the exact nature of the α-side-chain is critical for the activity of the butenolide, although changes in the absolute configuration or the nature of the O-alkyl group of the ether moiety are inconsequential. The nature of the δ-alkylidene substituent proved crucial to the activity of the butenolide, with the presence of a free hydroxy group apparently necessary; but the exocyclic olefin geometry does not seem important (likely due to in situ isomerisation).

The most potent compounds were then retested, following the protocol described in the original aruncin B publications[5,6] (Table 5, method B). As in the original work, auraptene was used as a positive control for cytotoxicity. For all synthetic butenolides tested, method B gave much higher IC_{50} values than the protocol used in the current work (method A).

### Table 5. Comparison of IC_{50} (µM) values between assay protocol in the current work (A) and that used in the original isolation study (B).[a]

| Entry | Compound | Jurkat T cells | Human T cells |
|-------|----------|----------------|---------------|
|       |          | Batch 1 | Batch 2 | Batch 1 | Batch 2 |
| 1     | auraptene | 83     | 57     | >50    | 205   | 19.4   | 73    |
| 2     | aruncin B | 70     | (lit.) | >245   | (lit.) |         |       |
| 3     | rac-aruncin B (Z-36) | 16.0 | >50 | NT | NT | 18.2 | >50 |
| 4     | 65       | 11.6   | 32     | 13.3   | NT    | 2.0    | 46    |
| 5     | 66       | 2.9    | 11     | 3.3    | NT    | 1.1    | 14    |
| 6     | 68       | 12.5   | >50    | 10.2   | >50   | 12.8   | NT    |

[a] NT = not tested.

### Conclusion

The Z- and E-Na salts of the postulated structure of aruncin B were synthesised through an approach relying on formation of a 6-membered ring intermediate by RCM followed by an oxy-selenolation-selenoxide elimination sequence to install the key enol ether functionality. However, the free acids could not be obtained from these Na salts, suggesting intrinsic instability. A regioisomeric ether was first envisaged as an alternative structure for aruncin B, and its Na salt was synthesised using a related RCM/ hydroxyselenolation-selenoxide elimination sequence, but featured similar acid sensitivity. Instability together with inconsistencies in the analytical data collected from the various dihydroxyran compounds suggested that aruncin B was another example of a compound of mistaken identity.[7] A butenolide was proposed as a viable alternative structure for aruncin B, and was first synthesised using a related RCM/ hydroxyselenolation-selenoxide elimination sequence, demonstrating the versatility of the strategy for the synthesis of different ring size systems, and confirming the true structure of aruncin B. Unfortunately, installation of the α-side-chain proved difficult and a preferred approach was developed involving a Sonogashira cross-coupling–5-exo-dig lactonisation strategy. This latter method also proved flexible, enabling the synthesis of 14 analogues for biological evaluation. The cytotoxic activity originally reported for aruncin B was confirmed, and some of the analogues (66 and 70) showed improved activity against malignant Jurkat T cells vs. normal T cells.

### Acknowledgements

This work was supported by the People Programme (Marie Curie Actions) of the European Union’s Seventh Framework Programme (FP7/2007-2013) under REA grant agreement 316955, and by Cancer Research UK (CRUK, C38302/13012) through the Preclinical Validation Lab, CRUK Oxford Cancer Research Centre, as well as the National Institute for Health Research Oxford Biomedical Research Centre. We also thank Prof. M. H. Woo (Catholic University of Daegu, Republic of Korea) for copies of the original spectra of 1, Prof. V. M. Macauley (Oxford) for her interest in and support of this work, and Prof. M. D. Smith and P. Gerken (Oxford) for the use of semiprep HPLC.

### Conflict of interest

The authors declare no conflict of interest.

### Keywords:
anticancer · butenolide · cytotoxic · natural products · structure elucidation

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