Reactions of Vinylcyclopropane and Bicyclopropyl Compounds With Maleic Anhydride

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Abstract. Vinylcyclopropane and bicyclopropyl C₆ compounds are studied as diene precursors in the Diels-Alder reaction. Their rearrangement under different thermal and microwave conditions leads to specific isomeric hexa-, penta- or butadienes, condensed with maleic anhydride as a dienophile. Adduct stereochemistries were compared to those of previously-synthesized model compounds, and are fully characterized with 2D NMR and GC-MS spectroscopies. Some unsaturated terpenes bearing vinylcyclopropane moieties were also condensed, and their adduct structures assigned.

Key words: Microwave, Thermal Diels-Alder Reactions, Diene Precursors, Vinylcyclopropane as a Diene

Introduction

One of the most useful reactions for building alicyclic structures is the Diels-Alder condensation (DA). Its scope and limitations, its different modes of operation, its control of adduct stereochemistry, and its mechanisms have all been widely studied, and several systematic rules have been discovered [1-3]. Several thermal, photochemical, catalytic, enzymatic or microwave (MW)-assisted, high-pressure and variable-temperature DA condensations were attempted (not always successfully) on the synthetic or biological building blocks, both dienes and dienophiles [4, 5].

The search for some diene analogs led to the design of more functional diene precursors, for example, vinyl-oxiranes, -aziridines or 1,3-diepoxides [6], or precursors. In this last case, the closest double bond alternative is the cyclopropane [7, 8]. In “conjugation” to the vinyl or allylic bonds, this ensemble – when built on, for example, a six-carbon chain with the three carbons necessary for cyclopropane formation – will offer a hexadiene-like system.

As an extension of the vinylcyclopropane (1) systems based on bicyclopropyl (2), allyl cyclopropane (3) or cyclopropyl propenes (4) – all of six carbons – could also be considered as potential diene sources. Of course, in order to react as a 1,3-conjugated diene, the cyclopropane moiety (or moieties) of the vinylcyclopropane or bicyclopropyl derivatives should undergo an opening and numerous rearrangements [9-15].

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Resumen. Vinilciclopropano y compuestos biciclopropilos C₆ se estudiaron como precursores de dienos en reacciones de Diels-Alder. La transposición de estos compuestos bajo condiciones térmicas o de irradiación de microondas conduce especificamente a hexa-, penta- o butadienos isoméricos, los cuales cicloadicionan con anhidrido maleíco como dienófilo. La estereocquímica de los aductos se estableció por comparación con aductos modelo previamente preparados y se caracterizaron por espectroscopia de RMN de dos dimensiones y CG-EM. Algunos terpenos insaturados llevando consigo el sistema vinilciclopropano también se condensaron, y fue asignada la estructura de sus aductos.

Palabras clave: Microondas, reacciones térmicas de Diels-Alder, precursores de dienos, vinilciclopropanos como dienos.

The dienes obtained as a result of previously-referenced rearrangements of cyclopropane moieties of 1 or 2, could be identified as Diels-Alder adducts with maleic anhydride, which is a high-efficiency dienophile and compared to some model adducts obtained from the pure dienes (Scheme 1).

Although Diels-Alder reactions are feasible under a wide variety of conditions, we chose to perform thermal and MW-assisted experiments because of their simplicity, and because of the proven “green” character of the latter. It is also of some interest to study the potential subtle differences between these two modes of activation of reagents entering cycloaddition, as has already been observed in our previous works (14).

It is expected that the reactions involving cyclopropane moieties will first lead to the production of conjugated six-
carbon diene systems \((C_6)\) with hexa, penta or buta-diene structures; these dienes react with maleic anhydride as per the stereochemistry revealed by the rearrangements [15]. Although some of these dienes do not lead directly to the \(s\)-\(cis\)-1,3-diene system necessary for efficient Diels-Alder addition the minor differences on the geometry of the dienes formed will be reflected by different adducts [16]. Because of the evident multistep character of the proposed synthesis, the stereochemistry of adducts produced from vinylcyclopropane cannot be predicted by Woodward-Hoffmann rules (2), however the dienes, once formed, should react with dienophile accordingly.

In this study we present the Diels-Alder reaction on the following compounds: allyl cyclopropane (3), 3-cyclopropyl-2-propene (4 \(cis\) and 4 \(trans\)), and 1-cyclopropyl propanol (5) as a compound leading via the dehydration of the previous two compounds, and the only two cyclopropane rings compound, bicyclopropyl (2). In the series of terpenes which contain vinyl and cyclopropyl moieties, we performed the reactions on 2-carene (6), and two model terpene dienes – terpinene (7) and \(\alpha\)-phellandrene (8) [17].

It is worth noting that terpenes were already used as diene precursors by Diels in 1938 for carene and thujene systems and even extended to cyclobutane-containing terpenes [18, 19]. It is also necessary to point out that 2-carene (6) has an endocyclic double bond and an exocyclic cyclopropane within its bicyclo [4,1,0] heptane skeleton. This last can either undergo rearrangement to a bicyclo[2,1,2] heptane skeleton, or open its cyclopropane bond, both of which give the seven-member ring (18) (Scheme 2). This work is also an extension of our previous observations on cubebene (10), a tricyclic sesquiterpene extracted from Solidago Canadiensis species. Cubebene contains the vinylcyclopropane system and undergoes both diene formation and epimerisation under Micro Wave-assisted conditions. Finally; in this work we try to rationalise some of those results by studying simpler diene-generating systems [14, 21].

In order to have several models of maleic anhydride adducts for \(trans,trans\)-hexadiene (11 \(tt\), \(trans,cis\)-hexadiene (11 \(ct\)), 4-methyl 1,3-pentadiene (12) and 2,3-dimethyl 1,3-butenes (13) adducts from pure isomeric six-carbon dienes were synthesized, a series of terpenes (7 and 8 and their maleic anhydride adducts) were also prepared (Scheme 2, 3 and 5).

All Diels-Alder reactions were performed in two parallel ways: thermal and MW-assisted manner for all simple diene precursors and terpene compounds. The starting dienes were also subjected to attempts at thermal and MW-assisted isomerisation without maleic anhydride in order to evaluate their possible structural alteration prior to their reaction as dienic compounds.

It must be noted that Micro Wave-assisted treatments are based upon conditions that differ from those that prevail under conventional thermal reactions. The main difference is that the susceptibility of the reagents towards microwaves may well be very different from those of the products. Hence, the interaction between microwaves and the various components is selective with a bias toward compounds characterized by a higher dielectric constant. While this may offer advantages in terms of conversion rate (yields), it has significant effect on the speed of the reaction and the selectivity (less side reactions or reduced degradation). In our examples, this behaviour can be explained easily from the reduced interactions the products have towards microwaves (reduced value of dielectric constant). This not only implies that less energy is imparted to them (thus, polymerization is reduced), but also that reaction kinetics are increased: for a given applied power, assuming constant reflection, the electrical field increases over time. Electrical field intensity is well known to be the most important factor in Micro Wave-assisted processes such as chemical synthesis [22-24].

In all cases, adducts and some diene structures were modelled in order to evaluate stability while the 500 MHz NMR studies will allow for unambiguous identification of single adducts or mixtures thereof.
Results and discussion

The diene-generating systems from different vinylcyclopropanes were of two origins: simple systems containing one or two cyclopropane moieties (all of which were six-carbon compounds leading to dienes), and terpene-originating systems, such as 6, 7 or 8. The results of their condensation with maleic anhydride are presented in two following sections. (Scheme 3 and 5)

Simple diene series

In this series of experiments, four diene precursors were synthesized. In a mono cyclopropyl series – allyl cyclopropane (3) (alternative name 3-cyclopropyl 1-propene), 1-cyclopropyl propene (4 cis or 4 trans) and as a model to previous one 1-cyclopropyl propanol (5) were synthesized. For the two cyclopropyl-containing series, the parent bicyclopropyl (2) was also prepared.

These syntheses were achieved by several methods (24-28):

- a. direct Grignard coupling reaction, by the Wurtz-Grignard approach with allyl bromide to obtain the allyl cyclopropane (3), as well as by the similar reaction on dicyclopentyl Zn (25);
- b. step-by-step Grignard reaction of cyclopropylmagnesium bromide on propionaldehyde to obtain, isolate, and separate 1-cyclopropyl propanol (5) which can lead to cis (4 cis) or trans (4 trans) 1-cyclopropyl propene by dehydration, used as obtained, in a mixture condensed with maleic anhydride or as a pure 4 trans isomer separated from this mixture;
- c. cyclopropylmagnesium bromide coupling to cyclopropyl bromide, leading to the bicyclopropyl (2);
- d. in situ synthesis of olefins from 1-cyclopropyl propanol (5), later leading to dienes as in point b.

Details of these syntheses are summarized in the Experimental section.

These cyclopropane compounds (with the exception of carbinol 5) are unstable, and difficult to synthesize. The use of closed vessels (stainless steel reactor for thermal, volatile and all-glass tube for Micro Wave) helped reduce the loss of material, and allowed the transformation of most of the diene into adducts (the procedures are described in the Experimental).

As mentioned previously, the dienes formed from these precursors were reacted with the maleic anhydride. The resulting adduct mixtures made via both thermal and MW-assisted routes were characterized, especially with help of their GC retention time, Rt (Table 1 and Experimental), GC-MS and high-resolution 1D and 2D NMR spectroscopic data obtained for four independently-synthesized adducts made from high-purity dienes. Particular attention was given to the proton high-resolution NMR, and minor differences in mass spectra of adducts (Table 2). All model diene adducts (14-17) (Scheme 3) displayed the isomass molecular ion at m/z 180, with a major fragmentation pattern leading to fragments at m/z 152, and retro-Diels-Alder fragments of various intensities at m/z 98, 70 and 82 (Table 2).

In order to position the given adducts, their Rt and the GC-MS single-ion monitoring at m/z 180 were used (Table 2). In principle, the hexadiene, methyl pentadiene and dimethyl butadiene adducts – formed from 11tt, 11ct, 12 and 13 dienes – are identified according to those parameters and confirmed by NMR.

From these results it was observed that, for a thermal reaction starting with allyl cyclopropane (3), two maleic anhydride adducts were observed, 14 and 15 (ratio 9:1), corresponding to

\[ \text{Table 1. Thermal and micro-wave assisted reaction on cyclopropane substrates} \]

<table>
<thead>
<tr>
<th>Substrate</th>
<th>Overall adducts yield (%)</th>
<th>Isomeric adducts obtained (ratio)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thermal / M-W</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>23% 45%</td>
<td>Hexadiene adducts 83%, 14 and 15 ratio 9:1 (thermal) and 4:1 (M-W)</td>
</tr>
<tr>
<td>4t and 4c (mixture of trans and cis isomers 3:1)</td>
<td>55 / 25</td>
<td>Hexadiene adducts 90%, 14 and 15 ratio 95:5 (thermal) and 94:6 (M-W)</td>
</tr>
<tr>
<td>4t (pure)</td>
<td>45 / 20 (max)</td>
<td>Adducts 14 (45%) and 15 (23%), methyl pentadiene adducts 36% (thermal)</td>
</tr>
<tr>
<td>5</td>
<td>24 / 13</td>
<td>Adducts 14 45%, 15 23% (thermal and M-W)</td>
</tr>
<tr>
<td>2</td>
<td>20-25 / 46</td>
<td>Isomeric pentadiene adducts up to 17%</td>
</tr>
</tbody>
</table>

* ratio of Diels-Alder adducts evaluated from m/z 180 ion mass chromatogram...
trans,trans-2,4-hexadiene (14) (major) and cis,trans-2,4-hexadiene (15) (minor) adducts. The dienes were formed first by the isomerisation of the double bond from 1,2 - to 2,3 - position of the diene, and by the cyclopropane opening. As witnessed by the formation of the same adducts from compounds 4t and 4c.

The alternative way – first the opening of the cyclopropane followed by the isomerisation of the double bond system to the conjugated diene could also be considered. The dienes 11tt and 11ct are both formed from 3 but product 16 cannot be formed (as it is from 4t alone or from the mixture of 4t and 4c). This alternative way was proposed by the Referee of our paper. It cannot however explain the diene 12a which gives the adduct 16 as shown in Scheme 4.

In a MW-assisted reaction the same yield (ca 35%) and 4:1 ratio of adducts were observed, however, less secondary products were formed.

For the cis and trans 1:3 mixture of compound 4, the thermal reaction gives a much higher yield of adducts than does the MW-assisted reaction (55% to 25%): the major adduct 14 comes from trans, trans-2,4-diene (11 tt) adduct, and the minor adduct 15 comes from the cis,trans diene (11 ct) (ratio 95:5 in thermal and 94:6 in MW method). The reaction was also repeated for the purified 4 t olefin, with results as described in Table 2.

Table 2. Thermal and micro-wave (M-W) assisted reactions of model dienes and maleic anhydride

<table>
<thead>
<tr>
<th>Diene</th>
<th>Adduct obtained</th>
<th>Yield (%)</th>
<th>R, **</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Thermal</td>
<td>Micro-wave</td>
</tr>
<tr>
<td>11tt</td>
<td>14</td>
<td>95</td>
<td>90</td>
</tr>
<tr>
<td>11ct</td>
<td>15</td>
<td>69</td>
<td>60</td>
</tr>
<tr>
<td>12</td>
<td>16</td>
<td>43</td>
<td>-</td>
</tr>
<tr>
<td>13</td>
<td>17</td>
<td>34</td>
<td>52</td>
</tr>
</tbody>
</table>

**relative retention time on column HP-5MS 60m of this adduct 7.30 min was normalized to 1.00

In this last case, the minor methyl pentadiene adducts were also identified – in particular, the adduct originating from the 4-methyl-1,3-pentadiene (12), identified in an MW-assisted reaction (3%). This result could be related to the position of the double bond in “conjugation” to the cycloproply moiety, which plays an activation role for the opening of either of two cyclopropane bonds, thereby leading to the diene formation (Scheme 4).

In the next diene precursor studies, 1-cyclopropyl propanol (5) was subjected to Diels-Alder reaction with maleic anhydride, giving 24% of adducts in thermal reaction and less than 15% under MW heating. The major adduct 14 corresponded to the trans,trans -diene (45%), followed by adduct 15 (13%). The lower overall yield of these reactions is probably related to the dehydration of alcohol necessary to produce the first double bond. The elimination of water from such cyclopropyl carbinol was not catalyzed by, for example, base or acid addition in either set of reaction conditions used.

The last and most challenging reaction of the bicyclopropyl (2), formed from several thermal and microwave generally laborious attempts, lead to a conclusion that the yields of adducts are much higher for the micro-wave assisted closed
vessel method, 46% than for thermal 20-25%, however even in the Micro Wave-assisted reaction, the presence of adduct 14 (50% yield) was followed by 15 (23-25%), 4-methyl 1,3-penta
diene adduct (16) (5%), and finally 2,3-dimethyl-1,3-butadi
eine adduct (17) (1%). In the thermal series, the ratio and yields of 14, 15 and methyl pentadiene adducts were similar, and butadiene adduct 17 was obtained at traces level only. These
last results indicate that the single cyclopropane ring in bicy
clopropyl (2) rearranged nonspecifically to two hexadienes by cleavage of a, a’ (or a, c’) cyclopropane bonds. It was observed also that the cleavage of cyclopropane bond followed either the already-observed cleavage of a, b’ bond, or a new cleavage of b, b’ bond, to give minor pentadiene and butadiene adducts, respectively (Scheme 4).

The most important factors in the identification of the adducts were the polynuclear 1D and 2D NMR spectroscopies. NMR analysis of these mixtures was performed using specific signals, with the signals from adduct methyls used as a starting point. For example, the distinction between the cis, trans and trans, trans, trans 2,4-hexadiene adducts 14 and 15 of two methyl proton chemical shifts (1.38 and 1.46 ppm), protons 4 and 7 as well as their coupling patterns to the maleic bridge protons and NOE effects. For example, both bridge protons and CH as well as CH₃ and vinyl protons (H-5, H-6) showed strong NOE for adduct 14 but one strong NOE effect only between H-3alpha and CH₃ alpha and also CH₃-C₄ and H-5 for adduct 15 was observed.

The results presented here led to the conclusion that, in all cases, there is an overwhelming propensity for the 11 tt diene formation from cyclopropyl propene precursors, leading to the adduct 14.

The bicyclopropyl (2) system showed the possibility of the rearrangement of the cyclopropane by several C-C bond cleavages, leading to the formation of 11 tt, but also 11 ct, along with a methyl pentadiene fraction and traces of 13. It should also be mentioned that other remaining isomeric methyl pentadienes (12, 2-methyl, all 3-methyl etc.) adducts were not synthesized separately. However, for this last series, the yields of these reactions remain low – less than 20%. For the MW-assisted reactions the polymerisation is certainly lower; but at best, these yields are in the same general range as for thermal reactions.

The addition of maleic anhydride is efficient in the presence of aprotic and anhydrous solvents. When preparing adducts, it is useful to replace the benzene or toluene solvent with ethyl acetate, to help reduce the polymerisation and opening of the anhydride to form the diacid. In this manner we also avoided the spontaneous isomerisation of 11 ct to 11 tt, observed for hexadienes under lengthy condensation (e.g. 24 hrs or longer).

**Terpene series**

In a series of more complex compounds which contain vinyl and cyclopropyl moieties, we performed the reactions on 2-carene (6) with the help of two model terpene dienes, terpinene (7) and α-phellandrene (8) (Scheme 5).

The rearrangements of the 2-carene (6) should lead, via one of three possible cyclopropane bond cleavages, to either:

a. 1,7,7-trimethyl bicyclo [2,1,2] hept-2-ene (19), a possible precursor to the α-terpinene (7) and α-phellandrene (8) after the corresponding allylic proton shift, the cyclopropane bond opening corresponded to the favoured E(pro-trans)-cyclopropane bond cleavage;

b. the unobserved iso-terpene skeleton of 1-methyl 3-isopropyl 1,3-cyclohexadiene (20) corresponding to the cleavage of the homo allylic to the double bond cyclo
propane bond; or

c. 1,5,5-trimethyl cycloheptene (18) skeleton terpene corresponded to the unfavoured Z(pro-cis)-cyclopropane bond cleavage (28). (Schemes 2 and 5).

The reaction of 2-carene (6) carried out under thermal or MW-assisted conditions with maleic anhydride led to a mixture of two adducts identified as the maleic anhydride of ter
pinene (7) and phellandrene (8) adducts, respectively; almost invariably the ratios were between 4:6 to 1:2 in favour of the phellandrene-originating product 23. The parallel synthesis of these two compounds from the addition of single-terpene dienes 7 and 8 to maleic anhydride allowed for their unambiguous identification. This result means that the 2-carene (6) is non-specifically rearranged to these two dienes, which react with the dienophile.

In order to further test this hypothesis, the 2-carene (6) is heated or irradiated under MW without maleic anhydride in several solvents (such as ethyl acetate, benzene or toluene); GC reveals that both individual terpene dienes are formed. All three terpenes show good stability under these conditions, less than 2% of 7 rearranges to 8 after 24 h of conventional heating or under MW irradiation in these solvents (18).
The NMR spectrum from terpene adducts 22 and 23 (see Experimental) display some interesting features, as confirmed by 2D experiments. The most interesting coupling observed, which confirms the proposed structures for adduct 23, is the 4J coupling between the H-9 vinylic proton and the H-10 beta proton; this confirms the proposed alpha orientation of the isopropyl group. NOESY experiments on the same system confirm this adduct’s structure by showing the strong effects between H-2 – H-10 exo and H-6 – H-11 exo as well as H-9 interaction with both isopropyl methyls. For adduct 22, however, the relevant features used for the confirmation of the structure were the dipolar correlation between H-2 and H-10 exo and H-6 and H-11 exo protons. As in the previous cases, the conformational energies for all terpene adducts were calculated, both for structures resulting from exo or endo approaches or having isopropyl group in equatorial position for adduct 22.

Conclusion

The aim of this study was to test the potential of cyclopropane-bearing precursors 2-5 to produce dienes via the rearrangements. It was observed that they are able to produce Diels-Alder adducts in both thermal and MW-assisted reactions, once their rearrangements are completed. The formation of dienes from the strained cyclopropane precursors was relatively easy, as shown by the diene adducts obtained. The same cannot be said about the synthesis of all cyclopropane compounds in this study, which was laborious and difficult: products were unstable, and yields were low. The most suitable solvent for these reactions was found to be ethyl acetate, which was preferred over chloroform, benzene, toluene, and water.

Although the results of condensations under MW-assisted conditions generally lead to cleaner mixtures of products in less time and with less solvent used, the expected differences between condensations under thermal and MW-assisted modes were not observed. The same could be concluded about the reactions in general, except that the bicyclopentyl (3) under MW-assisted conditions produces more butadiene. It is necessary to point out that the MW-assisted reactions were performed via heating through the solvent mode; reactions using focused MW should be also performed, in order to verify this conclusion.

In general however for all cyclopropene precursors 2-5 the lack of more pronounced selectivity of the rearrangements toward some specific dienes, except hexadienes, was observed under both heating conditions used. The Diels-Alder condensation and the ratio of diene adducts obtained in the sequence of reactions attempted is then only the indication of the selectivity of the multi-step rearrangement process. This also may be concluded about terpenes with more rigid structures, where the stereochemistry of the starting vinylcyclopropane could lead to more specific adduct isomers. However, many questions of mechanism of the rearrangement remain unanswered, for example of whether the vinylcyclopropane systems allow the 1,5-sigmatropic shifts through the cyclopropane ring, because of its partial double bond character, or whether the rearrangement could instead proceed through a series of 1,3-sigmatropic and allylic shifts.

As a further perspective to this study, the use of the aforementioned focused MW and photocatalytic reactions on the substrates 2-5 is an interesting alternative. It will also be interesting to build (for example) cyclohexene structure from the vinylcyclobutane (for 6-carbon-containing compounds).

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Experimental

Materials and methods

All cyclopropyl products and most of reagents were purchased from Aldrich Chemicals. The purification of volatile material was made using Buchi-Kugelrohr distillation apparatus and several flash chromatography on usual absorbents (Si-gel).

GC-MS measurements

GC/MSD Analysis: 10 µL aliquot was pipetted from each sample and diluted with 1 mL of hexane then were analyzed on a GC/MSD system consisting of an Agilent 6890N GC equipped with an Agilent 7683 automatic liquid sampler and interfaced directly to an Agilent 5973N mass selective detector controlled via Agilent’s ChemStation software. A 60-m HP-5MS capillary column (0.25 mm id, 0.25 µm) was used. Elution conditions were as follows: constant flow of 1.0 mL/min with an initial pressure of 18.54 psi; initial oven temperature of 70°C maintained for 2 min, then heated at 20°C/min up to 230°C where it was maintained for 5 min; injection volume of 0.2 µL in a split-less mode at 250°C with a purge-off time of 2.00 min.; the MSD was operated in selected ion monitoring (within m/z 100-300 range) with a transfer line kept at 280°C, quadrupole at 150°C and source at 230°C. Tuning was performed using the auto-tune feature with perfluorotributylamine (PFTBA) and the electron multiplier voltage was nominally set at 1529V.

The electron impact (EI) mass spectra are given in m/z (1%) of the most intense ion.
NMR

NMR high resolution experiments were performed using Varian INOVA spectrometer, at 500 MHz for proton and 125 MHz for carbon-13 respectively. In all cases the samples were prepared in CDCl₃ (or in different solvents as indicated) as ca 5% solution at 25 °C, chemical shifts were referred to the TMS in δ scale, are expressed in ppm and the coupling constants in Hz: s (singlet), d (doublet), t (triplet), q (quadruplet), m (multiplet).

Selected 2D experiments were run using a standard Varian package of INOVA programs for COSY and NOEs. HSQC experiments for short range couplings, less than 140 Hz, and one bond coupling were used. The long-range proton-carbon experiments (HMBC) were recorded with Varian’s program.

Other spectrometers used for more routine analysis of known products were Bruker apparatus 200 MHz, 300 MHz and 400 MHz NMR (UdeMoncton, CEA Saclay, IRSN CNRS Gif,respectively as indicated in the experimental ). The chemical shifts are expressed in δ (ppm) and coupling constants in Hz.

All the 1D and 2D detailed high resolution NMR spectra are available on request from the corresponding author (CKJ, Canada).

Synthesis of substrates 2-5

Allylcyclopropane (alternative name: 3-cyclopropyl 1-propane) (3)

Compound 3 was synthesized according to the modified procedure of Farneth (29) via Grignard-Wurtz coupling between the commercial cyclopropylmagnesium bromide and allyl bromide at room temperature. Yield of 3: 25-28%, bp 55-60°C/ 25 torr (product should be kept under Nitrogen)

1H NMR (200 MHz, CDCl₃) (δ, ppm, # of protons): 0.1-0.9 (5H), 1.95 (2H), 4.9 (1H, broad d, 10), 5.1 (1H, broad d, q, 15), 5.72 (1H, tt)

1-cyclopropyl propanol (5)

Carbinol 5 was synthesized according to modified procedure of Ferreri and Khusid (28, 30) from ethylmagnesium bromide and cyclopropenyl aldehyde at -20 to -78°C in ether or in tetrachloroethylene, with moderate (56-70%) yields. The synthesis was not optimised. When the dicyclopropyl zinc method is used with propionaldehyde as a starting aldehyde according to Reetz methodology (25), the yields of these reactions did not exceeded 25% and further improvements were abandoned because of the lack of variable temperature probe NMR facility and low temperature reactions incompatible with the microwave in use.

The analytical sample of carbinol, bp ca 135-140°C (dec.), was stored under the dry nitrogen atmosphere.

1H NMR (CDCl₃, 400 MHz): cyclopropyl signals 0.51 (2H), 0.23 (2H), 0.88-0.9 (1H), 2.82 (m, 1H), chain signals 1.60 (q, 2H), 1.01 (t, 3H);

13C NMR: 2.2, 2.7 and 10.2 (cyclopropyl), 17.5 (C3), 30.7 (C2), 75.2 (C1);

MS (EI, m/z, %): M+ 100 (2), 99 (M-1,100%), 71 (34), 57 (23).

Trans and cis 1-cyclopropyl propene (4t and 4t)

The 60-40 mixture of isomeric product 4t (major) and 4c (minor) was prepared by dehydration of 5 using catalytic amount of sulfuric acid at 80-90° with overall 55-65% yield. The compound 4t (trans isomer) was also synthesized with 30-35% by elimination in basic conditions. Both 4c and 4t isomers are relatively unstable. The pure 4t was separated by flash chromatography (b.p; 70-3°/23 torr).

4t Isomer

1H NMR (200MHz, CDCl₃): 0.1-0.9 (4H), 1.70 (1H), 4.93 (1H, qdd 1, 17, 16, H-2), 5.48 (1H, dqq, 1, 17, 16, H-1), 1.64 (3H, dd, CH₃)

4c isomer (spectral data extracted from the 300 MHz NMR spectrum of this mixture).

NMR 1H (300MHz, CDCl₃): 0.1-0.9 (m, unresolved 4H), 1.3 (1H), 4.70 (qdd, 1H, 10, 10, 1.8, H-3), 5.35 (1H, m, dqq, 1, 10, 7, H-2), 1.70 (3H, 7, 2, CH₃)

Bicyclopropyl (2)

This compound was synthesized by three methods.

a. according to the Farneth methodology (28-9) using a Simmons-Smith reaction on a tetrabromo derivative obtained from dibromocarbene addition to butadiene (31) with max. 20 % yield. Compound 2 was used as such in the Diels-Alder reactions.

b. direct coupling of cyclopropylmagnesium bromide to cyclopropylbromide at -78° using a Grignard reaction in THF solution with low yield of product 2 (10-2%). The unsaturated cyclic products and cyclopropanol derivatives were obtained instead in these reactions.

c. dicyclopropyl zinc addition to cyclopropyl bromide at -78°; yield of 2 within the 5% range.

The analytical sample of 2: bp 50-65 °C, one gas-chromatographic peak.

1H NMR (200 MHz, CDCl₃): 0.1 (4H), 0.3-0.35 (4H), 0.80-0.85 (2H).

Diels-Alder synthesis with diene precursors (2-8)

The above mentioned reactions, according to routes a-d, were performed at 50°C under thermal conditions in closed stainless steel vessels or in all-glass closed system at room-temperature under the MW irradiation using MAP technology (21-3) and a
one-pot method with the dienophile present in the vessel. The volatility of the dienes and cyclopropane containing intermediates (e.g. 5) require the reaction mixtures, after the quenching with the maleic anhydride to obtain the adducts, to be immediately extracted with ether and analysed by GC-MS and their NMR spectra were recorded. This method was developed in order to separate any polymer fraction due to diene polymerisation, which can be important especially at higher (50 °C) temperatures for all dienes and their precursors. This also minimizes the aromatisation of adducts with or without dienophile expulsion already observed.

After several unsuccessful attempts the thermal reaction temperature was decreased to 25° and then to 0°, because of the volatile and unstable character of dienes at 50° (32).

The crude reaction mixtures were analysed by GC-MS. At higher temperatures, reactions with anhydride showed, using GC-MS as a detection method, the presence of up to 40 compounds. These same reactions carried out in a micro-wave assisted mode reduced this number to 25 or 28. The major adduct diene – anhydride was always present in all these series, and its abundance was used to quantify the data (Table 2).

The use of a closed vessel “one-pot” method conforms to the practice widely applied in combinatorial chemistry for soft reaction conditions in particular. The lower temperatures helps minimizes losses in the volatile substrates and reduced their rearrangements (33).

**Synthesis of Diels–Alder adducts.**

Physical description: adducts 14-17 were obtained as oils or semisolid products decomposing easily when heated and their further crystallisation was not attempted. The NMR spectra of adducts 14-17, 22 and 23 were recorded on 500 MHz spectrometer.

**Adduct 14 (on 11t)**

Alternative name: 4a,7a-dimethyl-3a,4,7,7a-tetrahydro isobenzofuran-1,3-dione (14)

Product 14 was synthesized with a 90-95% yield.

H NMR (CDCl₃): 3.34 (H-3a and H-7a), 2.48 (H-4 and H-7), 5.77 (H-5 and H-6), 1.43 (CH₃-C₄ and CH₃-C₇)

13C: 46.15 (C3a and C7a), 30.0 (C4 and C7), 134.15 (C-5 and C-6), 16.1 (CH3-C4 and CH3-C7a), 171.5 (C=O).

**Adduct 15 (on 11ct)**

Alternative name: 4a,7a-dimethyl-3a,4,7,7a-tetrahydro isobenzofuran-1,3-dione (15)

1H NMR (CDCl₃): 1.46 (CH₃-C₄, CH₃-C₇), 5.85 and 5.80 (H-5, H-6), 3.28 and 3.41 (H-3a, H-7a), 2.34 and 2.28 (CH–CH₃)

**Adduct 16 (on 12)**

Alternative name: 4,4-dimethyl-3a,4,7,7a-isobenzofuran-1,3-dione (16)

1H NMR (CDCl₃): 5.6-5.7 (2H, H-5, H-6), 1.23 (3H, s, CH₃-C₄ ax) and 1.08 (3H, s, CH₃-C₄ eq), 3.07 (1H, ddd 8.5, 9, 4, H-7a), 2.98 (1H, d, 8.5, H-4a), 2.05 (1H, H-7eq) and 2.28 (1H, H-7ax)

**Adduct 17 (on 13)**

Alternative name: 5,6-dimethyl-3a,4,7,7a-tetrahydro isobenzofuran-1,3-dione (17)

1H NMR (CDCl₃): 3.36 (tt, 3.5, 1.0; H3a, H7a), 2.46 (td, -15, 1.0, H-4 and H-7), 2.29 (broa d, H-4' and H-7'), 1.717 (CH₂-C₅ and CH₂-C₆)

13C NMR: 40.3 (C3a and C7a), 30.4 (C4 and C7), 132.7 (C-5 and C-6), 174.46 (C1 and C3), CH3 17.6.

**Adduct on terpinene (22)**

Alternative name: 1-isopropyl-α7-methyl-α – 4-oxa –tricyclo[5,2,2,0]204,204] undec-8-e ne-3,5-dione (22)

The equimolar quantities of terpenes and maleic anhydride were dissolved in toluene and heated overnight. Adduct 22 was separated by precipitation after partial removal of the solvent.

Yield: 67%.

**Adduct of α-phellandrene (23)**

Similar procedure to adduct 22. Yield: 77%

1H NMR (CDCl₃): 3.18 (1H, ddd, 2.0, 3.5 and 6.0, H-1), 3.08 (1H, ddd, 3.5 and 8.5, H-2), 3.14 (1H, ddd 3.5 and 8.5, H-6), 2.96 (1H, dddd, 0.5, 2.0, 3.0 and 5.0, H-7), 1.32 (1H, m, 1.5, 5.5, 7.0 and 9.0, H-10), 1.79 and 1.044 (2H, ddd, 3.0, 9.0 and 3.5; ddd 3.5, 8.5 and 3.5, H-11 ax and H-11 eq), 1.78 (3H, d, 2.0, H-12), isopropyl chain 1.12 (1H, d, heptuplet, 9.5 and 7.0, H-13), 0.92 (3H, d, 7.0, CH14), 0.81 (3H,d, 7.0, CH3-15), 5.76 (1H, m, ddd, 0.5, 2.0, 6.5(H-9))

13C NMR: 35.06 (C-1), 46.07 (C-2), 173.0 (C-3), 172.7 (C-5), 44.04 (C-6), 37.72 (C-7), 142.2 (C-8), 122.3 (C-9), 44.27 (C-10), 29.7 (C-11), 20.43 (C-12), 32.86 (C-13), 20.7 (C-14), 20.12 (C-15)

**References**


13. However it is worth noticing that the sigmatropic shift was considered by Woodward-Hoffmann formalism only for a limited number of concerted reactions. The vinylcyclopropane (1) system could undergo a sigmatropic shift only after allylic rearrangement and followed by the opening of the cyclopropene ring. The allylic and/or sigmatropic shifts can then transform the vinylcyclopropane or bicyclopropyl into isomeric dienes of specific geometry. The Cope rearrangement can also be considered as a step toward the final conjugated diene formation. The vinylcyclopropane to cyclopropene rearrangement could then be seen as a 1,3-sigmatropic reaction. The accepted mechanism of this reaction proceeds via the formation of 1,5-diradicals in two isomeric configuration: Z (cis) or E (trans). The first one leads to cyclopentene and the second to dienes or other products (Scheme 4).


17. The parent of these terpenes should be considered thujene (9) with a homoanannular and endovincylcyclopropene which unfortunately is not commercially or easily available as a pure isomer.


26. Several attempts synthesis described in the literature mentioned the assumed presence of these products, their short lifespan, easy polymerisation or aromatisation. Several procedures were either incomplete or irreproducible. Many particular and non commercial reagents are used in order to obtain e.g. the bicyclopropene (29), in particular zinc-Grignard (24, 28) or photolysis reactions (27).


32. The above mentioned difficulties are related to the six carbon or lower chains of dienes involved and to the cyclopropylcarbinols. The other isomeric model C4 dienes were not commercially accessible. Some of them cannot be obtained from the cyclopropene compounds of this study and then are not potential diene precursors.

33. The small volume stainless steel vessel conditions are relatively close to those applied in using the MW methods.