Allylic and benzylic oxidation reactions with sodium chlorite

Samuel M. Silvestre and Jorge A. R. Salvador*

Laboratório de Química Farmacêutica, Faculdade de Farmácia, Universidade de Coimbra, Rua do Norte, 3000-295 Coimbra, Portugal

Received 31 October 2006; revised 29 December 2006; accepted 8 January 2007
Available online 12 January 2007

Abstract—Various allylic and benzylic substrates were selectively oxidized to the corresponding enones in good yields using sodium chlorite, either in combination with tert-butyl hydroperoxide in stoichiometric conditions, or associated with N-hydroxyphthalimide as catalyst. These oxidation reactions were effectively and economically performed under mild, transition-metal free conditions and therefore the dual challenge of cost effectiveness and benign nature of the processes was met.

© 2007 Elsevier Ltd. All rights reserved.

1. Introduction

Allylic and benzylic oxidations are industrially important synthetic processes due to their wide variety of applications in the synthesis of pharmaceuticals and fine chemicals.1 Relevant examples are the oxidation of Δ5-steroids to the corresponding biologically interesting Δ5-7-ketone derivatives2 and the benzylic oxidation of xanthene to xanthone.3

Traditionally, chromium(VI)-based reagents such as CrO3–pyridine complex,4 CrO3 and 3,5-dimethylpyrazole,5 pyridinium chlorochromate (PCC),6 pyridinium dichromate (PDC),7 PDC–tert-butyl hydroperoxide,8 sodium chromate,9 sodium dichromate in acetic acid,10 pyridinium fluorochromate,11 3,5-dimethylpyrazolium fluorochromate (VI)12 and a combination of an N-hydroxydicarboxylic acid imide with a chromium containing oxidant13 have been used to perform these oxidations. However, the use of large excess quantities of both reagents and volume of solvent in most of these procedures, along with the difficult work-up of the environmentally hazardous chromium residues makes such procedures inconvenient on a commercial scale.

Other classical, stoichiometric oxidative processes involve the use of manganese dioxide, potassium permanganate or selenium dioxide.14 A variety of catalytic methods for allylic and benzylic oxidations have been reported and generally peroxide-based oxidants have been the reagents of choice. Despite the fact that alkyl hydroperoxides are the most used peroxides in these reactions, the use of hydrogen peroxide associated with metal catalysts was also reported.15 Due to its importance, it is also worth to mention the use of molecular oxygen, combined with transition-metal catalysts,16 to perform these oxidation reactions.

Of particular environmental and economical interest is the use of tert-butyl hydroperoxide combined with different types of metal catalysts.17 Despite the good yields reported with CrO3,18 2,4-dimethylpentane-2,4-diol cyclic chromate,19 PDC,20 Cr(CO)6,21 RuCl2(PPh3)322 and RuCl3,23 the toxicity of these chromium compounds and the high cost of the ruthenium catalysts motivated the search for cheaper and more environmentally acceptable methods based on the use of copper24 and cobalt25 catalysts. Recently, the use of other metal catalysts such as dirhodium(II) caprolactamate,26 manganese(II) complexes27 and manganese(III) acetate28 in combination with tert-butyl hydroperoxide has also been reported.

A common difficulty associated with the homogeneous procedures reported so far is the separation step required for the removal of the catalysts, which cannot be easily recovered and reused.

The immobilization of inorganic reagents and catalysts useful in organic reactions on heterogeneous supports is a very important area in clean technology29 and led researchers to report the use of chromium-based heterogeneous catalysts in combination with tert-butyl hydroperoxide to perform these oxidations.30 We have recently reported the use of cobalt(II), copper(II), manganese(II) and vanadium(II) catalysts in heterogeneous forms for the allylic oxidation

Keywords: Sodium chlorite; Allylic oxidation; Benzylic oxidation; Transition-metal free.

* Corresponding author. Tel.: +351 239859950; fax: +351 239827126; e-mail: salvador@ci.uc.pt

0040–4020/$ - see front matter © 2007 Elsevier Ltd. All rights reserved.
doi:10.1016/j.tet.2007.01.012
of unsaturated steroids. Jurado-Gonzalez et al. reported the allylic and benzylic oxidations using tert-butyl hydroperoxide and catalytic amounts of cobalt(II) alkyl phosphonate modified silica.

The increasing concern about environment and the need for green reagents have prompted research on the use of bismuth compounds as catalysts combined with tert-butyl hydroperoxide for allylic and benzylic oxidation reactions.

The development of metal-free ecofriendly synthetic transformations is an area of current interest. These methods avoid the use of toxic and expensive metals and seem to be especially attractive for the preparation of compounds that do not tolerate metal contamination, such as pharmaceutical products. The use of oxygen in the presence of N-hydroxyphthalimide (NHPI) and an organic free-radical initiator such as dibenzoyl peroxide, acetaldehyde, azobisisobutyronitrile, anthraquinones and acridine yellow/Br2 has been reported to be suitable for these oxidations. Recently, a procedure using sodium hypochlorite (household laundry bleach) in combination with aqueous tert-butyl hydroperoxide (70% or less) has also been reported to perform allylic and benzylic oxidations.

Sodium chlorite is a very cheap oxidant and has been extensively used in water treatment and as a bleaching agent in paper and textile industries. In the field of synthetic organic chemistry, the most known use of sodium chlorite is in the efficient chemoselective oxidation of aldehydes to the corresponding carboxylic acids. More recently increasing attention has been focused on its use in organic synthesis, for example, in the oxidation of primary alcohols and epoxidation of olefins.

In this work, we report the use of sodium chlorite either in combination with tert-butyl hydroperoxide, in stoichiometric conditions, or with N-hydroxyphthalimide as catalyst for mild, transition-metal free, allylic and benzylic oxidation reactions.

## 2. Results and discussion

### 2.1. Allylic and benzylic oxidations mediated by NaClO2/tert-butyl hydroperoxide

The recent growing interest in the use of sodium chlorite as a cheap oxidizing agent in organic chemistry motivated us to study its utility in allylic and benzylic oxidation reactions.

Firstly, we explored its use in combination with tert-butyl hydroperoxide and this association revealed to be practical and useful to accomplish the referred oxidations under mild conditions. Thus, using this system it was possible to effectively perform the transition-metal free allylic oxidation of Δ^2-steroids and (+)-valencene and the benzylic oxidation of substrates (Schemes 1 and 2 and Tables 1 and 2). In general, the reactions required 1.2 equiv of NaClO2 and 5 equiv of tert-butyl hydroperoxide, relative to the substrate, and could be performed using a homogeneous solvent mixture of CH3CN and water, at 50 °C.

In a first set of experiments, dehydroepiandrosterone acetate was used as a model substrate under various experimental conditions. Blank experiments revealed that NaClO2 or tert-butyl hydroperoxide as the sole oxidant led only to traces of products after 50 h of reaction. Among the screened solvents, the best results were obtained with mixtures of CH3CN and water or with ethyl acetate (Table 1, entries 2–4 and 6). Using CH3CN or 1,4-dioxane/H2O (3:1, v/v) as solvent, longer reaction times were observed (Table 1, entry 5). Decreasing the amount of tert-butyl hydroperoxide to 1 equiv (per mmol of substrate) led to a longer reaction time (Table 1, entry 7). Reaction temperature was found to be a critical parameter. At 25 °C the relative rate
of conversion was very slow (Table 1, entry 8) in comparison to the reactions performed at 50 °C and therefore we considered the use of this temperature for the study of the process. In the presence of ethylenediamine tetracetic acid (EDTA), a metal ion sequestering agent, no significant changes were seen in the reaction indicating that this allylic oxidation was not being catalyzed by the presence of adventitious metal ions in the solvent (Table 1, entry 9).

Due to solubility reasons, the oxidation of substrate 2 was performed using CH₃CN as the solvent and the 7-ketone derivative 7 was obtained in good yield (Table 2, entry 1). When applied to substrate 3, this new oxidative system afforded product 8 in 76% yield (Table 2, entry 2).

The chemoselective allylic oxidation in the presence of a secondary hydroxyl group was studied using dehydroepiandrosterone 4 as the substrate and the corresponding 7-ketone derivative 9 was obtained with good chemoselectivity (65% isolated yield) (Table 2, entry 3). A similar oxidation was performed on (+)-valencene 5 under these conditions and the sesquiterpenoid nootkatone 10 was the major reaction product (38% yield) (Table 2, entry 4). In agreement with these results, when performed on a larger scale, only a small decrease in the reaction rate was observed (Table 2, entry 5).

In spite of longer reaction times, benzylc compounds, xanthene 11, fluorene 12 and diphenylmethane 13 were oxidized to the corresponding benzyl ketones, xanthone 15, fluorenone 16 and benzophenone 17 (Scheme 2) in high yields (Table 2, entries 6–8).

The mechanism of these oxidation reactions is worth considering. An experiment was conducted under the same reaction conditions (Section 4.2; Table 1, entry 2) but in the presence of a radical inhibitor, butylated hydroxy toluene (BHT, 20 mol %), and it was observed that the oxidation was inhibited until BHT was consumed. This information implied that this process occurred via free-radical species.

Recently, Geng et al. demonstrated the formation of chlorine dioxide radical (ClO₂) by heating NaClO₂ at 55–65 °C in CH₃CN/H₂O (4:1). As our conditions are similar, we also considered the formation of that compound. Thus, the mixture of NaClO₂ (1.2 mmol) in CH₃CN/H₂O (3:1) (12 mL) was heated at 50 °C during 15 h, and the UV spectrum of the resulting cooled solution showed the characteristic absorption peak of ClO₂ ($\lambda_{max}=359$ nm), in agreement with the previously reported results. Furthermore, the presence of ClO₂ radical was confirmed by Electron Spin Resonance (ESR) experiments.

As this reaction was not catalyzed by metal ions (Table 1, entry 9) and did not occur using only NaClO₂ or tert-butyl hydroperoxide (blank experiments), the process was most likely initiated by the homolytic cleavage of tert-butyl hydroperoxide by ClO₂ radicals, originating tert-butylperoxy radical. This reactive intermediate initiates the allylic (and benzylic) hydrogen abstraction leading to the formation of the olefin radical, which is further oxidized via a radical chain mechanism to the corresponding enone.
2.2. Allylic and benzylic oxidations mediated by 
NaClO₂, catalyzed by N-hydroxysuccinimide (NHPI) 

Due to the interesting results obtained with the system 
NaClO₂ tert-butyl hydroperoxide, we decided to study the 
use of NaClO₂ as the sole oxidant in a catalytic process. 
Knowing the fact that chlorine dioxide is a reasonable 
disproportionate product of NaClO₂ when heated and an 
inherent radical species itself,⁴⁶ although not enough active 
to perform allylic oxidation, we considered the possible 
association of NaClO₂ with a radical catalyst such as NHPI as 
a n efficient oxidizing system. Actually, this combination 
resulted in a powerful transition-metal free catalytic oxida-
tive system for these allylic and benzylic oxidations. Thus, 
the allylic oxidation of \( \Delta^5 \)-steroids \( 1-4 \) and \( (+) \)-valencene \( 5 \) and the benzylic oxidation of substrates \( 11-14 \) (Schemes 
1 and 2 and Tables 3 and 4) have been readily performed 
under mild conditions.

In general, the reactions were very fast and required 
1.5 equiv of NaClO₂ and 0.1 equiv of NHPI, relative to the 
substrate, and could be performed using a homogeneous 
 solvent mixture of CH₃CN and water, at 50°C.

The catalyst NHPI in the absence of NaClO₂ was not active, 
as revealed by the blank experiments.

The best solvent for this system was the mixture of CH₃CN 
and water (usually 2:1 or 3:1, v/v) (Table 3, entries 1 and 2), 
however, dioxane/H₂O (3:1, v/v) could also be used, 
although a longer reaction time has been observed (Table 3, 
entry 3). The use of N-hydroxysuccinimide (NHSI) in place 
of NHPI led to an increase in the reaction time and lower 
yield (Table 3, entry 4). 2,2,6,6-Tetramethylpiperidine-
1-oxyl (TEMPO) has no catalytic activity under these con-
ditions (Table 3, entry 5). The reduction of the amount of 
NHPI to 0.05 equiv led to a significant reduction in the reac-
tion rate (Table 3, entry 6). The study of the effect of temper-
ature in this reaction allowed us to observe that significantly 
shortened times were required at 50°C (Table 3, entries 2 
and 7) and for this reason we studied the process using this 
temperature.

The presence of EDTA in the reaction system did not change 
the results significantly, indicating that this reaction occurs 
without the influence of traces of metal ions in the solvent 
(Table 3, entry 8). This reaction is also not influenced by 
the molecular oxygen present in air, as the results were not 
changed when the reaction was performed under nitrogen 
atmosphere (Table 3, entry 9).

The use of NaOCl (5% aqueous solution; household laundry 
bleach) instead of NaClO₂ within the same reaction con-
ditions led to some reactivity, however, low selectivity was 
observed.

When applied to substrates 2 and 3, this new oxidative 
system afforded products 7 and 8 in 60 and 76% yields, 
respectively (Table 4, entries 1 and 2).

The reaction of dehydroepiandrosterone 4 led to the cor-
responding 7-ketone derivative 9 in moderate yield.

### Table 3. Allylic oxidation of dehydroepiandrosterone acetate 1 by NaClO₂/organic catalyst

<table>
<thead>
<tr>
<th>Entry</th>
<th>Substrate/mmol</th>
<th>NaClO₂/mmol</th>
<th>Catalyst/mmol</th>
<th>Solvent (v/v)</th>
<th>Temp/°C</th>
<th>Time/h</th>
<th>Product</th>
<th>Isolated yield b/%</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>6</td>
<td>6</td>
<td>89</td>
</tr>
<tr>
<td>2</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>50</td>
<td>6</td>
<td>6</td>
<td>75</td>
</tr>
<tr>
<td>3</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>1.4-Dioxane/H₂O (3:1)</td>
<td>50</td>
<td>14</td>
<td>6</td>
<td>84</td>
</tr>
<tr>
<td>4</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>8</td>
<td>6</td>
<td>72</td>
</tr>
<tr>
<td>5</td>
<td>1/0.25</td>
<td>0.375</td>
<td>TEMPO/0.025</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>40</td>
<td>6</td>
<td>16</td>
</tr>
<tr>
<td>6</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>24</td>
<td>6</td>
<td>76</td>
</tr>
<tr>
<td>7</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>25</td>
<td>144</td>
<td>6</td>
<td>78</td>
</tr>
<tr>
<td>8</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>7</td>
<td>6</td>
<td>77</td>
</tr>
<tr>
<td>9</td>
<td>1/0.25</td>
<td>0.375</td>
<td>NHPI/0.025</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>6</td>
<td>6</td>
<td>82</td>
</tr>
</tbody>
</table>

a Solid, 80% (Aldrich).

b EDTA aqueous solution (5 x 10⁻⁴ M).

c Performed under N₂ atmosphere.

d Traces of starting material and a by-product are visible on TLC plates but not detectable in ¹H NMR spectrum (300 MHz).

### Table 4. Allylic and benzylic oxidations by NaClO₂/NHPI

<table>
<thead>
<tr>
<th>Entry</th>
<th>Substrate/mmol</th>
<th>NaClO₂/mmol</th>
<th>NHPI/mmol</th>
<th>Solvent (v/v)</th>
<th>Temp/°C</th>
<th>Time/h</th>
<th>Product</th>
<th>Isolated yield b/%</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>2/0.5</td>
<td>1.5</td>
<td>0.05</td>
<td>1.4-Dioxane/H₂O (3:1)</td>
<td>50</td>
<td>25</td>
<td>7</td>
<td>76</td>
</tr>
<tr>
<td>2</td>
<td>3/0.25</td>
<td>0.375</td>
<td>0.025</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>50</td>
<td>7</td>
<td>8</td>
<td>76</td>
</tr>
<tr>
<td>3</td>
<td>4/1</td>
<td>1.5</td>
<td>0.1</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>11</td>
<td>9</td>
<td>50⁰</td>
</tr>
<tr>
<td>4</td>
<td>5/1</td>
<td>1.5</td>
<td>0.1</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>50</td>
<td>15</td>
<td>10</td>
<td>36⁰</td>
</tr>
<tr>
<td>5</td>
<td>1/3</td>
<td>4.5</td>
<td>0.3</td>
<td>CH₃CN/H₂O (2:1)</td>
<td>50</td>
<td>8</td>
<td>6</td>
<td>76</td>
</tr>
<tr>
<td>6</td>
<td>11/0.5</td>
<td>0.75</td>
<td>0.05</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>50</td>
<td>2</td>
<td>15</td>
<td>91</td>
</tr>
<tr>
<td>7</td>
<td>12/0.5</td>
<td>0.75</td>
<td>0.05</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>50</td>
<td>1</td>
<td>16</td>
<td>90</td>
</tr>
<tr>
<td>8</td>
<td>13/0.5</td>
<td>0.75</td>
<td>0.05</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>50</td>
<td>6</td>
<td>17</td>
<td>89</td>
</tr>
<tr>
<td>9</td>
<td>14/0.5</td>
<td>0.75</td>
<td>0.05</td>
<td>CH₃CN/H₂O (3:1)</td>
<td>50</td>
<td>18</td>
<td>18</td>
<td>75</td>
</tr>
</tbody>
</table>

a Solid, 80% (Aldrich).

b Traces of starting material and a by-product are visible on TLC plates but not detectable in ¹H NMR spectrum (300 MHz).

c Recovered by flash chromatography (ethyl acetate/light petroleum, bp 40–60°C).
higher reactivity of this system is probably the reason for the moderate chemoselectivity observed in the oxidation of this substrate (Table 4, entry 3).

The sesquiterpenoid nootkatone 10 was the major product (36% yield) of the allylic oxidation of (+)-valencene 5 under these conditions (Table 4, entry 4). When performed on a larger scale using dehydroepiandrosterone 1 as substrate, this procedure allowed the obtention of product 6 in 76% yield and with only a small decrease in the reaction rate (Table 4, entry 5).

When applied to benzylic substrates such as xanthene 11, fluorene 12 and diphenylmethane 13 this procedure allowed excellent results. Thus, the corresponding benzylic ketones, xanthone 15, fluorenone 16 and benzophenone 17 (Scheme 2) were obtained in very high yields and within short reaction times (Table 4, entries 6–8). The benzylic oxidation of isochromane 14 led to 1-isochromanone 18 in 75% yield (Table 4, entry 9).

When performed under the same reaction conditions (Section 4.3; Table 3, entry 1), but in the presence of BHT (20 mol %), this process was inhibited until the radical inhibitor consumption, which indicates that these oxidations (20 mol %), this process was inhibited until the radical action times (Table 4, entries 6–8). The benzylic oxidation of isochromane 14 led to 1-isochromanone 18 in 75% yield (Table 4, entry 9).

When performed under the same reaction conditions (Section 4.3; Table 3, entry 1), but in the presence of BHT (20 mol %), this process was inhibited until the radical inhibitor consumption, which indicates that these oxidations were also mediated by free-radical species.

It is known that NHPI is a radical catalyst, mediating various oxidation reactions by O2 via the formation of the phthalimide N-oxyl (PINO) radical intermediate. This catalyst is commonly associated with metallic or organic free-radical initiators to perform oxidation reactions using O2 as the oxidant.

We demonstrated that this reaction was not catalyzed by metal ions (Table 3, entry 8) and did not occur using only NaClO2 or NHPI catalyst (blank experiments). In addition, we confirmed by UV and ESR experiments that ClO2 was generated by heating NaClO2 at 50 °C. ESR studies of the mixture of NaClO2 (0.375 mmol) and NHPI (0.025 mmol) in CH2CN/H2O (2:1) (3 mL) at room temperature revealed the simultaneous presence of ClO2 and PINO.

Therefore, considering these observations, we propose that under these reaction conditions, NaClO2 originated ClO2 radicals, which should react with NHPI leading to the formation of PINO radical. This reactive intermediate can abstract allylic (and benzylic) hydrogens originating the olefin radical, which is in turn oxidized via a radical chain mechanism to the corresponding enone.

3. Conclusions

In summary, allylic and benzylic oxidation reactions can be cheaply and effectively performed using sodium chlorite either in combination with tert-butyld perhydroperoxide in stoichiometric conditions, or with N-hydroxypythalimide as catalyst under mild, transition-metal free conditions. These very simple, economical and ecofriendly procedures led to good yields and selectivities and, therefore, should find a large use in organic synthesis.

4. Experimental

4.1. General

Substrates 1–4, 12–14, NaClO2, t-BuOOH, NHPI, NHSI, TEMPO and BHT were commercially available from Sigma–Aldrich Co. (+)-Valencene 5 and xanthene 11 were purchased from Fluka. Reaction solvents were distilled before use, according to standard procedures. Kieselgel 60HF254/Kieselgel 60G was used for TLC analysis. Melting points were determined with a Reichert microscope apparatus and were uncorrected. IR spectral analyses were performed in a JASCO FT/IR-420 spectrophotometer. 1H and 13C NMR spectra were recorded on a Bruker AMX 300 spectrometer, in CDCl3 solution with Me4Si as internal standard. UV spectral analyses were performed in a Bausch & Lomb Spectronic 2000 spectrophotometer. ESR measurements were carried out on a Bruker EMX 1144 spectrometer (X-band) with a 100 kHz field modulation.

4.2. General procedure for allylic and benzylic oxidations mediated by NaClO2/tert-butyld perhydroperoxide

In a typical reaction, to a solution of the substrate (e.g., 17-oxoandrost-5-ene-3β-yl acetate 1, 82.6 mg/0.25 mmol) in CH3CN/H2O (3:1, v/v) (3 mL), tert-butyld perhydroperoxide (0.18 mL/1.25 mmol, 70% aqueous solution, Aldrich) was added followed by the slow addition of NaClO2 (33.92 mg/0.3 mmol, 80%, Aldrich). After 18 h under magnetic stirring at 50 °C, the reaction was complete (TLC control). The reaction mixture was poured into sodium sulfate solution (10% aqueous) and extracted with diethyl ether. The extract was washed with aqueous saturated solution of NaHCO3, water, dried and evaporated to dryness to give 7,17-dioxoandrost-5-en-3β-yl acetate 6 (62.6 mg, 75% yield). Crystallization from methanol afforded 67% of pure product; mp 180–183 °C (MeOH); lit.184 75% yield. IR (ATR): 1244, 1628, 1670, 1727, 2950, 2443, 2469, 2923 cm–1; 1H NMR (CDCl3, 300 MHz): δ 0.89 (s, 3H, 18-CH3), 1.24 (s, 3H, 19-CH3), 2.06 (s, 3H, CH3CO), 4.72 (m, 1H, 3α-CH); 13C NMR (CDCl3, 75 MHz): δ 71.90 (C-3), 126.43 (C-6), 164.79 (C-5), 170.19 (CH3CO), 200.66 (C-7), 220.14 (C-17).

4.3. General procedure for allylic and benzylic oxidations mediated by NaClO2, catalyzed by NHPI

In a typical reaction, to a solution of the substrate (e.g., 17-oxoandrost-5-ene-3β-yl acetate 1, 82.6 mg/0.25 mmol) in CH3CN/H2O (2:1, v/v) (3 mL), NHPI (4.1 mg/0.025 mmol) was added followed by the slow addition of NaClO2 (33.92 mg/0.3 mmol, 80%, Aldrich). After 6 h under magnetic stirring at 50 °C, the reaction was complete (TLC control). The reaction mixture was poured into sodium sulfate solution (10% aqueous) and extracted with diethyl ether. The extract was washed with aqueous saturated solution of NaHCO3, water, dried and evaporated to dryness to give 7,17-dioxoandrost-5-ene-3β-yl acetate 6 (76.6 mg, 89% yield). Crystallization from methanol afforded 78% of pure product.
CH); 13C NMR (CDCl3, 75 MHz): 4.3.4. Compound 10. 125.7 (C-6), 166.2 (C-5), 200.7 (C-7), 219.8 (C-17).

1.09 (s, 3H, 14-CH3), 1.71 (s, 3H, 13-CH3), 4.70 (m, 2H, 4-CH2), 4.53 (t, 2H, 4-CH2), 7.38 (t, 7.9 Hz, 2H, 1-CH, 8-CH); 13C NMR (CDCl3, 75 MHz): δ 72.00 (C-3), 126.41 (C-6), 164.13 (C-5), 170.23 (CH3CO), 201.11 (C-7), 209.64 (C-20).

CH); 13C NMR (CDCl3, 75 MHz): 4.3.6. Compound 16. 156.14 (C-4a, C-10a), 177.19 (C-9).


Acknowledgements

J.A.R.S. thanks Universidade de Coimbra for financial support. S.M.S. thanks Fundação para a Ciência e Tecnologia for a grant (SFRHB/BD/11087/2002). We kindly acknowledge the Center of NMR Spectroscopy and Center for Neurosciences and Cell Biology, University of Coimbra, Portugal (Professor C. Geraldes) and Professor J. Laranjinha (Center for Neurosciences and Cell Biology, University of Coimbra, Portugal) for the ESR experiments.

References and notes

10. Amann, A.; Ourisson, G.; Loo, B. Synthesis 1987, 1002.
Various hydroaromatic and benzylic compounds can be oxidized under a normal pressure of O2, catalyzed by NHPI (10 mol%), even in the absence of transition-metal species or organic free-radical initiators, however, reactions were generally performed in benzonitrile at 100 °C; Ishii, Y.; Nakayama, K.; Takeno, M.; Sakaguchi, S.; Iwahama, T.; Nishiyama, Y. J. Org. Chem. 1995, 60, 3934.


48. ESR spectra were obtained under the following conditions: frequency 9.7 GHz, modulation amplitude 2 G and microwave power 20 mW. The characteristic quartet signal is in agreement with the spectrum attributed to ClO2 radical: Ozawa, T.; Miura, Y.; Ueda, J.-I. Free Radical Biol. Med. 1996, 20, 837.


50. ESR spectra were obtained under the conditions reported in Ref. 48. The characteristic triplet signal is in agreement with the spectrum attributed to PINO: Ishii, Y.; Iwahama, T.; Sakaguchi, S.; Nakayama, K.; Nishiyama, Y. J. Org. Chem. 1996, 61, 4520.


