

Natural Organic Acid as Green Catalyst for Xanthenones Synthesis: Methodology, Mechanism and Calcium Channel Blocking Activity

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Xanthenones were synthesized via one-pot tricomponent reaction, under solvent-free conditions, using aldehydes, phenolic and cyclic 1,3-dicarbonyl compounds. Natural organic acids (NOAs), compounds present in many living metabolisms, were used as potential green catalysts. NOA are considered to be more eco-friendly and user-friendly alternative to traditional methodologies. Optimization studies showed that oxalic acid was the best NOA catalyst for such reaction furnishing the xanthenones with up to 93% of yield. Theoretical calculations were performed to evaluate this reaction mechanism and regioselectivity. The results showed that the regiospecificity of this three-component reaction is kinetically and thermodynamically controlled by the addition of β -naphthol C2, instead of C10, to the aldehyde. Our results also disclosed two xanthenones as novel calcium channels blockers. Eco-friendly reaction conditions, easy workup procedure, short reaction times and good yields are some of the advantages of our methodology.

Keywords: xanthenones, natural organic acid, theoretical calculations, calcium channels blockers

Introduction

Xanthenes (Figure 1) are a class of oxygenated heterocycles that contain an aromatic ring fused to 4-carbonyl-pyran ring as the basic skeleton.¹ They present several biological and pharmacologic properties: antivasular, antiarrhythmic, antihypertensive, antioxidant, anti-inflammatory, anticancer and calcium channel blocking.² On the other hand, xanthenones (Figure 1), xanthone derivatives, are poorly explored regarding their biological properties, and described only for its anti-proliferative,^{3,4} antiviral⁵ and anti-microbial properties.⁶

The literature describes the synthesis of xanthenones via a multicomponent reaction using a phenolic compound, an aldehyde, and a cyclic 1,3-dicarbonyl compound.⁷ Many Lewis or Brønsted acids may be used as catalysts: calixarenes,³

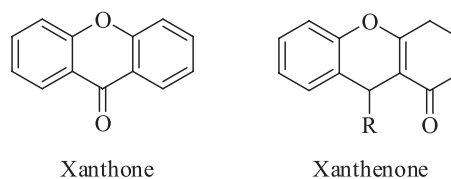


Figure 1. Xanthone and xanthenone base structure (R = aromatic or alkyl group).

ceric ammonium nitrate,⁴ polyethyleneglycol bound sulfonic acid,⁵ Hy zeolite,⁶ triflate proline,⁸ tetrabutyl ammonium fluoride,⁹ tri-methyl tetradecyl ammonium bromide,¹⁰ sulfamic acid,¹¹ *p*-toluenesulfonic acid,¹² triphenylphosphine-*m*-sulfonate/carbon tetrabromide,¹³ phosphorus(V) oxide,¹⁴ indium(III) chloride,¹⁴ scandium(II) triflate,¹⁵ boron trifluoride diethyl etherate,^{16,17} functionalized ionic liquids,¹⁸ 2,4,6-trichloro-1,3,5-triazine,¹⁹ silica-supported perchloric acid,²⁰ succinimide-*N*-sulfonic acid,²¹ ytterbium perfluorooctanoate,²² thiamine hydrochloride²³ and trityl

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chloride.²⁴ However, most of these methods do not comply with environmental-friendly protocols, presenting one or more of the drawbacks: less availability or hard catalyst preparation,^{4,5,18,21,22} long reaction time,^{8,11,16,17,19,23} cumbersome product isolation procedures.¹⁶ They also use expensive and/or detrimental metal precursors,^{14,15,22} toxic catalysts,^{11,14} and toxic organic solvents to perform the reaction or purify the product.^{13,16,17,20} A readily available and non-hazardous catalyst that promotes the formation of xanthenones in a short reaction time, under solvent-free condition, and simple work-up is highly desirable.

A more eco-friendly and user-friendly alternative employs natural organic acids (NOAs), which are biodegradable compounds found in the metabolism of many organisms.^{25,26} For example, acetic acid (the second simplest monocarboxylic acid after only to formic acid) catalyzes multicomponent reactions in the synthesis of 3-alkoxyisoindolin-1-imines,²⁷ piperidines,²⁸ bicyclic pyro,²⁹ quinazolinones and (dihydroquinazolinylphenyl) oxazolones.³⁰ NOAs bearing two carboxylic groups and used as catalysts include the oxalic, malonic and succinic acids. Oxalic acid was reported as an efficient catalyst for the synthesis of coumarins,³¹ quinoxalines,³² quinolines,³³ bis(indolyl)methanes,^{34,35} tetrahydroquinolines³⁶ and flavone.³⁷ Although less explored in multicomponent reactions, oxalic acid has also shown to be an efficient catalyst in such reactions for obtaining quinazolinones,³⁸ piperidines³⁹ and *N*-aryl-3-aminodihydropyrrol-2-one-4-carboxylates compounds.⁴⁰ Malonic acid was reported as a catalyst in polymerization reactions^{41,42} and for obtaining 1,3-dichloropropanols from glycerol;⁴³ however, its use in multicomponent reactions has not been described yet. Succinic acid was reported as a catalyst in a multicomponent reaction for the preparation of amidoalkyl naphthols⁴⁴ and α -amino phosphonates.⁴⁵ Most of the NOAs come from economically renewable resources turning them as targets for developing green multicomponent approach for substances of pharmaceutical, agricultural, cosmetic and food purposes.^{25,45-48} Despite the several described uses of acetic, oxalic, malonic and succinic acids in organic synthesis, the literature has not registered their use to furnish xanthenones.

A probable mechanistic rationalization to produce xanthenones involves an *ortho*-quinone methide intermediate, which is generated by the nucleophilic attack of the phenolic compound on the aldehyde, followed by dehydration.^{3,11,14,15,20} Although this nucleophilic addition is reasonable, there is no discussion about the regioselectivity of the intermediate. Also, to the best of our knowledge, there are no theoretical and/or experimental studies about the mechanism proposition in the literature.

We developed a green methodology to synthesize xanthenones from the one-pot condensation of benzaldehyde derivatives, β -naphthol, and dimedone, employing NOAs as organic acid catalysts. The mechanism and regioselectivity were investigated using theoretical calculations HF/6-31G(d) and LC- ω PBE/6-311++G(d,p). Moreover, due to the high biological potential of xanthenones, we also investigated this class of molecules as calcium channel blockers.

Results and Discussion

Optimization of the reaction conditions

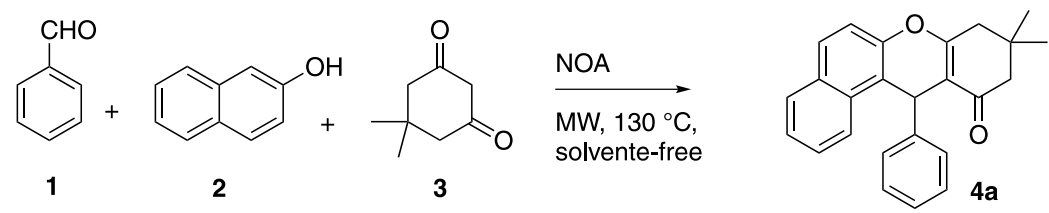
Initially, an investigation was performed to find the best catalyst and reaction condition to synthesize the xanthenone **4a** (Table 1). As a reaction model, benzaldehyde (1.0 equivalent), β -naphthol (1.2 equivalent) and dimedone (1.5 equivalent) were mixed and catalytic amounts of oxalic ($pK_{a1} = 1.2$), malonic ($pK_{a1} = 2.8$), succinic ($pK_{a1} = 4.2$), and acetic acids ($pK_a = 4.7$) (5, 10 and 20 mmol%) were added to reaction mixture. The evaluated organic acids follow a homologous series in which the number of carbons between the carboxyl groups varies; the oxalic having no carbon among them; the malonic acid has one carbon; the succinic acid has two carbons and the acetic acid is the corresponding monocarboxylic acid. The pK_{a1} value of dicarboxylic acids increases with the increase in the number of carbons between the carboxylates; a fact that can be explained by the inductive effect of the neighboring carboxyl and the stability of the corresponding conjugated base. The oxalic acid has the most stable conjugate base, because the negative charge can be stabilized by resonance and can establish an intramolecular hydrogen bond. Besides that, the other carbonyl in α , increases its acidity by inductive effect.

We chose to develop a solvent-free under microwave radiation (MW) methodology because it makes the synthesis of xanthenones simpler, cleaner and more efficient. It also provides an increment on safety, easy work-up, cost reduction, and prevents solvent waste, hazards and toxicity.⁴⁹⁻⁵¹

The catalyst quantity influences the xanthenone yields. Oxalic acid demonstrated itself to be the most efficient as a catalyst, providing similar or better yields than the other acids in all conditions.

The best condition was using 20 mmol% of oxalic acid and a reaction time of 5 min (82% yield; Table 1). This acid also showed that higher concentrations or longer reaction times lower the product yield. The behavior is opposite to the other acids, which improved the reaction yield with the

Table 1. Effect of the type and quantity of natural organic acids (NOAs) for obtaining xanthenone **4a**^a under microwave radiation (MW) and solvent-free conditions

				
entry	Natural organic acid (NOA)	NOA quantity / mmol%	time / min	Yield ^b / %
1	oxalic acid	5	5	58
2	oxalic acid	5	10	51
3	oxalic acid	10	5	62
4	oxalic acid	10	10	52
5	oxalic acid	20	5	82
6	oxalic acid	20	10	65
7	oxalic acid	25	5	66
8	malonic acid	5	5	10
9	malonic acid	5	10	26
10	malonic acid	10	5	2
11	malonic acid	10	10	51
12	malonic acid	20	5	14
13	malonic acid	20	10	55
15	succinic acid	5	5	9
15	succinic acid	5	10	35
16	succinic acid	10	5	16
17	succinic acid	10	10	21
18	succinic acid	20	5	17
19	succinic acid	20	10	54
20	acetic acid	5	5	traces
21	acetic acid	5	10	58
22	acetic acid	10	5	traces
23	acetic acid	10	10	51
25	acetic acid	20	5	51
25	acetic acid	20	10	67
26	no catalyst	—	5	16
27	no catalyst	—	10	20

^aReagents and conditions: benzaldehyde/β-naphthol/dimedone (molar ratio = 1.0:1.2:1.5); ^b yields for purified xanthenone **4a**.

increase on the reaction time. The reaction carried out in absence of catalyst furnished the desirable product **4a** in low yields (16 and 20% after 5 and 10 min of MW, respectively). Oxalic acid is the most acid catalyst herein tested. Since the formation of xanthenones involves various protonated intermediates, the stronger Brönsted-Lowry acid should be the best organic catalyst.

Most protocols describe long reaction times (30 minutes to hours) for the xanthenone synthesis.^{8,11,16,17,19,23} The

protocols using microwave radiation lasted less than 5-15 min;^{15,18,22} however, these methodologies have disadvantages such as the need to prepare the catalysts,^{18,22} the use of detrimental metals as catalysts,^{15,22} and the use of toxic solvents to purify the desirable products.¹⁵ Our protocol takes only a few minutes to complete the reaction, uses a cheap, eco-friendly and user-friendly catalyst and all the xanthenones were purified by recrystallization.

The scope of the substrates

We further investigated the scope of this methodology since the optimal conditions have been determined for the synthesis of xanthenone **4a**, using oxalic acid as the catalyst. Fifteen different xanthenones were synthesized with yields ranging 51-93% (Table 2). Three kinds of aldehydes were tested: containing electron-donating groups, electron-withdrawing groups and heteroaromatic rings. The reaction yields were not considerably affected by the presence of electron-withdrawing groups (best result using 4-nitro-benzaldehyde; 80% yield) or electron-donating groups (best result using 2-hydroxy-benzaldehyde; 93% yield) in the aldehydes. The heteroaromatic aldehyde 2-pyridinyl provided the best yield at 130 °C (85%), demonstrating a greater stability at this temperature.

Low yields were observed for 4-hydroxybenzaldehyde (35%), 4-cyano-benzaldehyde (45%), 3-methoxy-4-hydroxy-benzaldehyde (46%) and 3,4-dimethoxy-4-hydroxy-benzaldehyde (55%), using the ratio aldehyde/ β -naphthol/dimedone of 1.0:1.2:1.5. This is explained by the formation of subproducts from a side-reaction

between the dimedone and the aldehyde. In order to prevent this side-reaction, the stoichiometry of β -naphthol and dimedone was inverted, for a ratio of aldehyde/ β -naphthol/dimedone of 1.0:1.5:1.2, and we achieved better yields (Table 2, entries 7 and 10-12).

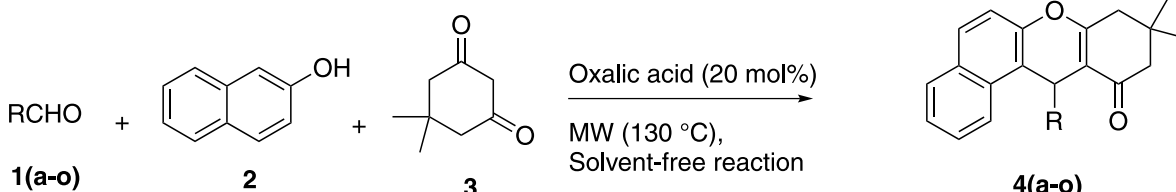
In all the cases, the pure product was readily isolated by recrystallization from methanol. The xanthenones products were characterized by melting point, infrared (IR), nuclear magnetic resonance (NMR) spectroscopy and an electrospray ionization high-resolution mass spectrometry (ESI-HRMS). Additional information was obtained from X-ray diffraction spectroscopy.

X-Ray

The crystal structure of **4e** was determined and analyzed in a diffractometer Enraf-Nonius Kappa-CCD. The ORTEP representation of **4e** is shown in Figure 2 and data related to the collection and refinement of its crystal structure are in the Table S6 of the Supplementary Information (SI).

Compound **4e** crystallized in centrosymmetric space groups, as expected from their achiral synthesis. The four

Table 2. Synthesis of xanthenones catalyzed by oxalic acid under solvent-free condition

		
entry	Aldehyde	Yield / %
1	C ₆ H ₅	4a (82) ^{a,c}
2	4-O ₂ NC ₆ H ₄	4b (80) ^{a,d}
3	4-FC ₆ H ₄	4c (72) ^{a,d}
4	4-ClC ₆ H ₄	4d (81) ^{a,d}
5	2-OHC ₆ H ₄	4e (93) ^{a,c}
6	3-OHC ₆ H ₄	4f (60) ^{a,c}
7	4-OHC ₆ H ₄	4g (68) ^{b,d}
8	4-SMeC ₆ H ₄	4h (86) ^{a,d}
9	4-OMeC ₆ H ₄	4i (70) ^{a,d}
10	4-CNC ₆ H ₄	4j (67) ^{b,d}
11	3-OMe, 4-OHC ₆ H ₃	4k (81) ^{b,d}
12	3,5-OMe, 4-OHC ₆ H ₂	4l (73) ^{b,d}
13	2-thienyl	4m (58) ^{b,d,e}
14	2-pyridinyl	4n (85) ^{a,c}
15	5-nitro-2-furyl	4o (51) ^{b,c,e}

Molar ratio of the reagents used: ^aaldehyde/ β -naphthol/dimedone (1.0:1.2:1.5) or ^baldehyde/ β -naphthol/dimedone (1.0:1.5:1.2). Yield: for the pure xanthenone after ^c5 min or ^d10 min of reaction. ^eThe temperature reaction used was 100 °C instead of 130 °C.

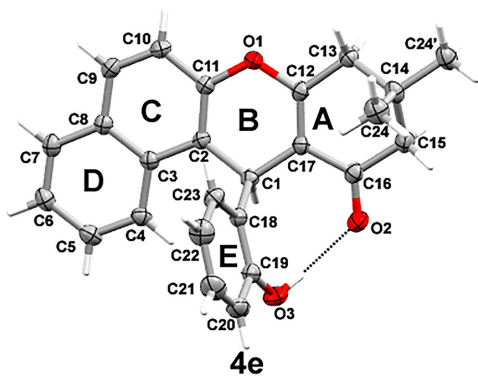


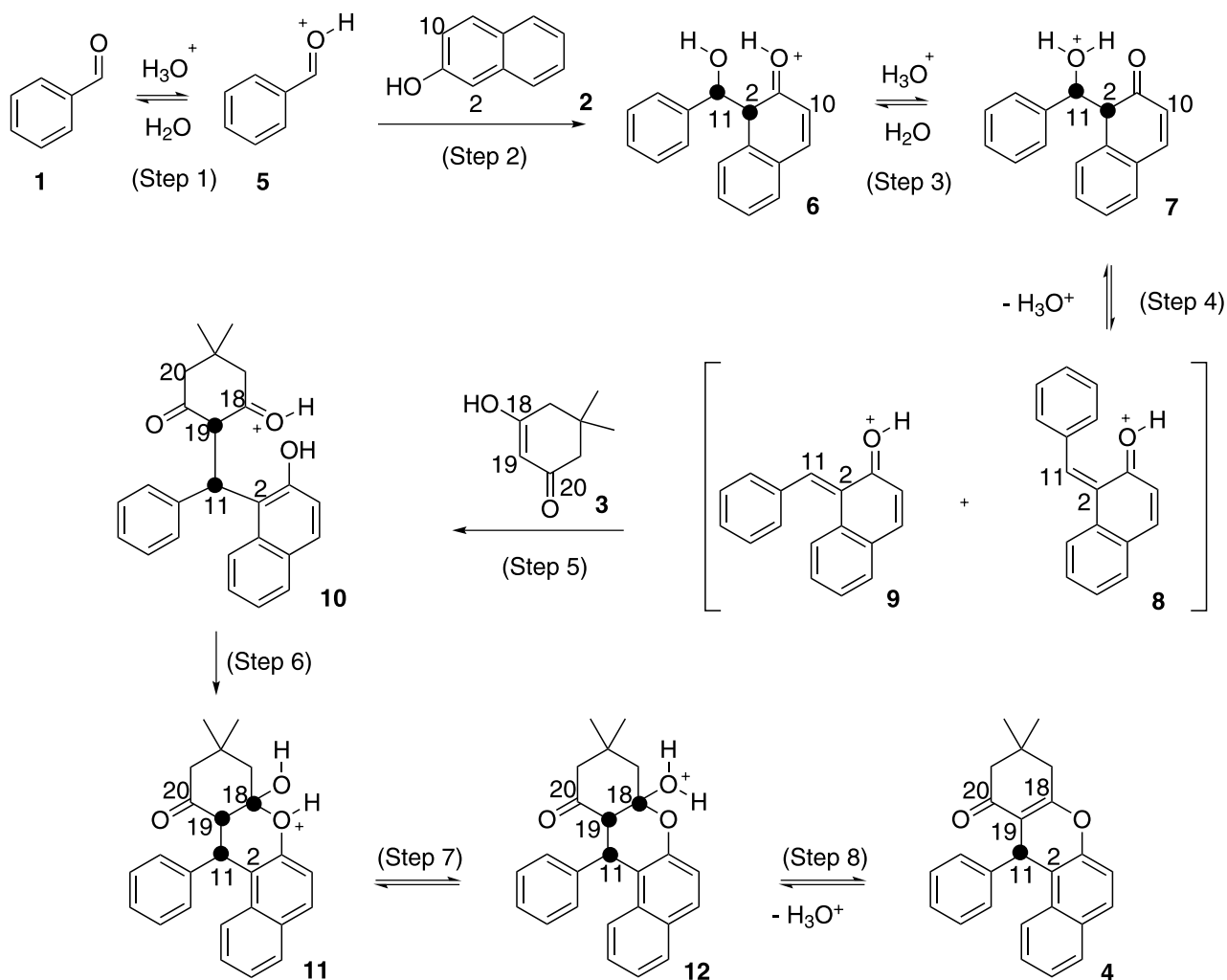
Figure 2. ORTEP image of **4e**. Non-hydrogen atoms are shown with their 30% probability ellipsoids. Rings and non-hydrogen atoms are arbitrarily labeled.

fused rings formed a nearly planar xanthene core except for ring A, which adopted a half-chair conformation with the C14 carbon removed from the least-squares plane, calculated through the other five coplanar atoms, by 0.6451(15) Å in **4e** (root-mean-square deviation (r.m.s.d.).

of the C15-C16-C17-C12-C13 fitted atoms is 0.0194 Å in **4e**). Ring B is distorted in with the C1 carbon deviating by 0.1857(14) Å in **4e** from the mean plane calculated through the C2-C11-O1-C12-C17 atoms (r.m.s.d. of the fitted atoms is 0.0245 Å in **4e**). In the structure, the C1 and C14 are on the same side of the substituted phenyl ring that is perpendicular to the xanthene mean plane. The mean plane calculated through ring E forms ca. 90° angle 81.06(4)° in **4e** with the least-squares plane encompassing the non-hydrogen atoms of the four fused rings A-D, except for C1 and C14. In addition, there is a $S_1^1(8)$ motif in compound **4e** due to the formation of an intramolecular hydrogen bonded cycle.

Theoretical investigation of the reaction mechanism

A probable mechanism to obtain xanthenones is via initial nucleophilic attack of β -naphthol (**2**) C2 on the carbonyl carbon of the benzaldehyde (**1**) (Scheme 1).^{3,11,14,15,20,24} An alternative mechanism would involve a nucleophilic attack



Scheme 1. Probable mechanism for the synthesis of xanthene (**4**) using β -naphthol (**2**) via C2 nucleophilic on the carbonyl carbon of the protonated benzaldehyde (**5**).

of β -naphthol C10 (Scheme 2); however, this pathway leads to the xanthenone **20** that is not observed for this reaction. In order to verify this regioselectivity, theoretical calculations were performed.

The energy variation of each reaction step was calculated using HF/6-31G(d) level of theory. The electronic-nuclear energies, Gibbs free energies, and entropy values for all compounds in Schemes 1 and 2 are available in the Supplementary Information (Tables S1-S4).

Comparing both schemes and the energy step-by-step, we concluded that step 2 is crucial to explain the regioselectivity. In C2 pathway, this step is more favored by energy ($\Delta G = -1.72$ kcal mol⁻¹); however, in C10 pathway, this step is thermodynamically disfavored ($\Delta G = 10.51$ kcal mol⁻¹, Tables S3 and S4).

In order to have a better regioselectivity understanding, a mechanistic study was also performed employing an improved level of theory (LC- ω PBE/6-311++G(d,p)). Scheme 3 shows the step 2 of both C2 and C10 pathways with their respective transition states (**TS1** and **TS2**). All

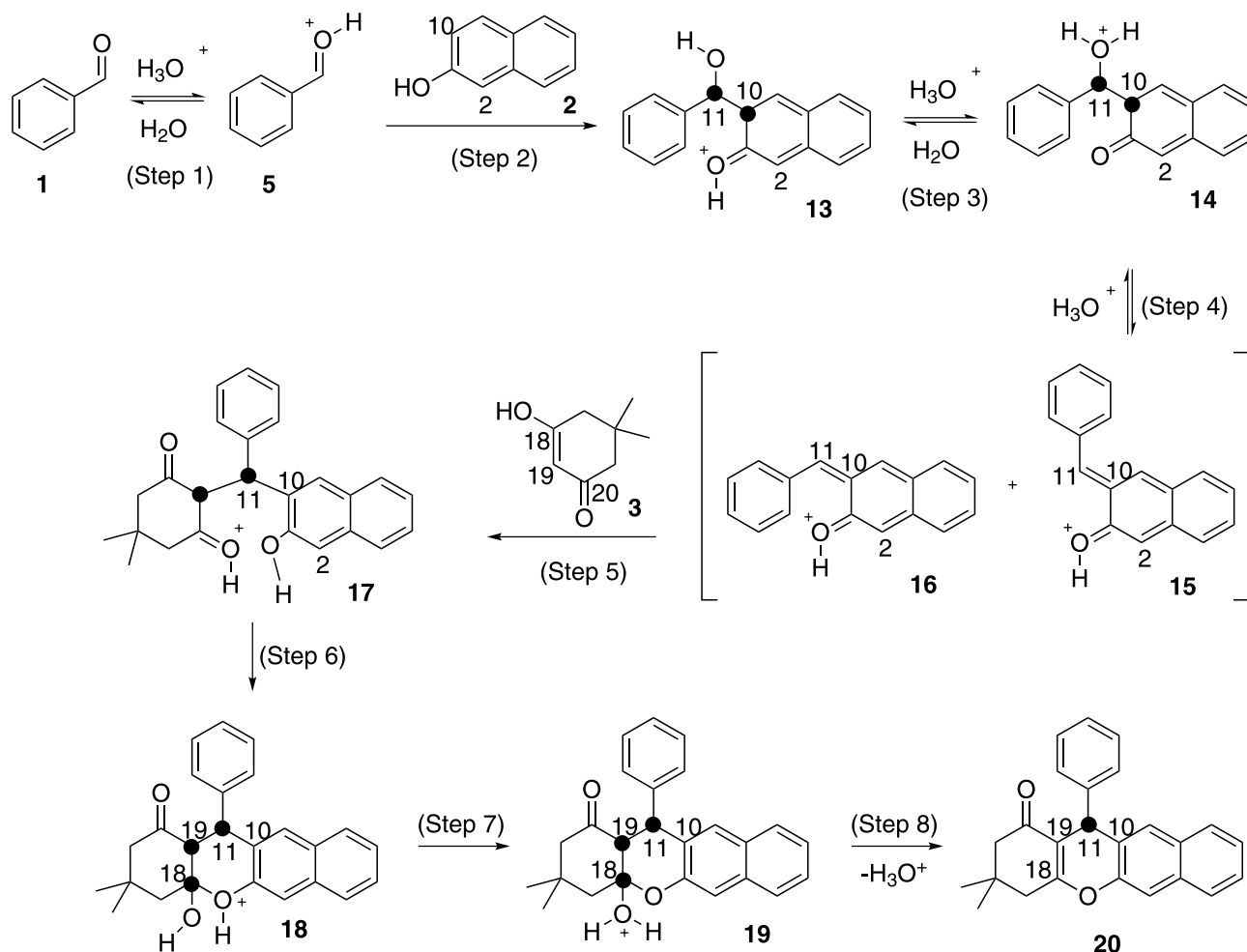
optimized structures and their energies are available in the Supplementary Information (Table S5).

The theoretical study at 130 °C showed that the activation energy (process **5** + **2** $\rightarrow$ **TS**, Table 3) to **TS1** formation ($\Delta G = 6.84$ kcal mol⁻¹) is lower than **TS2** formation ($\Delta G = 15.45$ kcal mol⁻¹). This demonstrates that the formation of **6** is thermodynamically and kinetically favorable, while the formation of **13** is not favorable (Figure 3).

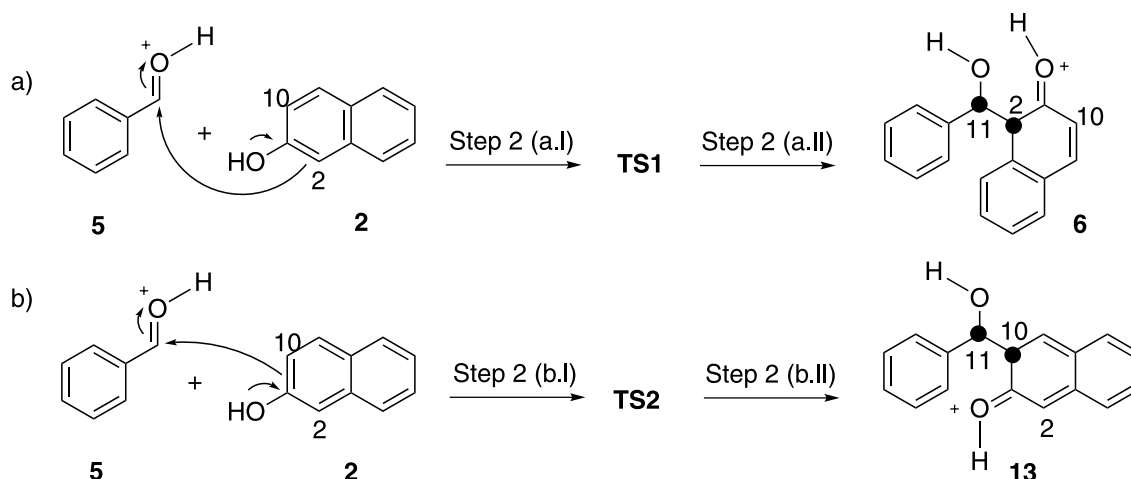
The rate constants k of the reaction were estimated according to Eyring's equation (equation 1):

$$k(T) = \frac{k_B T}{h} e^{-\Delta G^\ddagger/RT} \quad (1)$$

In equation 1, T is temperature, k_B is the Boltzmann's constant, h is the Planck's constant and R is the universal gas constant. The values were 6×10^7 and 3×10^1 s⁻¹ at 130 °C for steps 2(a) and 2(b), respectively. This means that the formation of **6** is one billion times faster than the formation of **13**. This study justifies the reason product **4**



Scheme 2. Probable mechanism for the synthesis of xanthenone (**20**) using β -naphthol (**2**) via C10 nucleophilic on the carbonyl carbon of the protonated benzaldehyde (**5**).



Scheme 3. Addition of β -naphthol (**2**) to protonated benzaldehyde (**5**) through nucleophilic attack of (a) C2 and (b) C10.

Table 3. Variations of electronic energy (ΔE_{elect}), entropy (ΔS) and Gibbs free energy (ΔG_{total}) to the studied steps at 130 °C

Process	$\Delta E_{elect} /$ (kcal mol ⁻¹)	$\Delta S /$ (cal mol ⁻¹ K ⁻¹)	$\Delta G_{total} /$ (kcal mol ⁻¹)
Step 2(a) ^a			
5 + 2 → TS1	-11.97	4.65	6.84
TS1 → 6	-11.54	1.35	-10.03
5 + 2 → 6	-23.51	6.00	-3.19
Step 2(b) ^a			
5 + 2 → TS2	-2.73	2.05	15.45
TS2 → 13	-8.61	0.40	-8.52
5 + 2 → 13	-11.34	2.45	6.93

^aThe step 2 is completely sketched in Scheme 3.

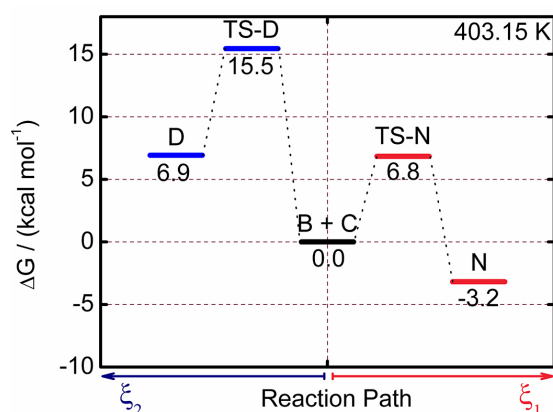


Figure 3. Energy profiles along the pathway for step 2(a) (data in red) and step 2(b) (data in blue) at 130 °C.

is obtained at the synthetic procedure while the product **20** is not observed.

The protonated benzaldehyde (**5**) suffers a nucleophilic attack receiving the charge density in its lowest unoccupied molecular orbital (LUMO) of the carbonyl carbon atom (Figure 4). The β -naphthol acts as a nucleophile donating

charge density from its highest occupied molecular orbital (HOMO) passing through C2 and leading to **6**. The formation of **13** might occur by the nucleophilic attack of β -naphthol through its energetic nearest occupied orbital to the HOMO (i.e. HOMO – 1 orbital), which presents charge density passing through C10. This attack is less favorable than the first one because the HOMO – 1 is less available than HOMO, and 19 kcal mol⁻¹ higher in energy. Furthermore, the LUMO of **5** is energetically closer to the HOMO than to HOMO – 1 of β -naphthol. This corroborates with the thermodynamic and kinetics observations. The frontier orbitals of all studied species at LC- ω PBE/6-311++G(d,p) level of theory are available in the Supplementary Information.

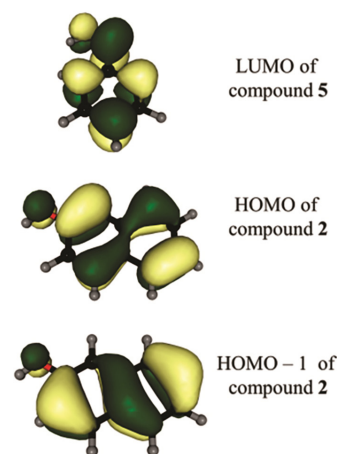


Figure 4. Frontier orbitals LUMO of protonated benzaldehyde (**5**), HOMO and HOMO – 1 of β -naphthol (**2**). Atom legends: C (black), O (red) and H (grey).

Calcium influx assay

Previously presented data by Wang *et al.*,⁵² suggested the activity of several xanthenes on blocking calcium channels

(Ca²⁺) and also beta adrenergic receptors. As xanthenones are chemically related to xanthenones, we decided to verify the potential of the synthesized xanthenones as calcium channel blockers, and for that we performed an *in vitro* calcium influx assay using cortical brain synaptosomes. All xanthenones were evaluated to determine their potential to inhibit cytosolic Ca²⁺ entry when synaptosomes were depolarized by potassium chloride (KCl) 33 mM. The changes in cytosolic Ca²⁺ may reflect the Ca²⁺ influx and were monitored continuously in fura-2-loaded synaptosomes (as described in Experimental section). Figure 5 shows the results of the most promising xanthenones in inhibiting Ca²⁺ influx induced by high potassium. Depolarization by KCl led to an increase in intracellular calcium concentration [Ca²⁺]_i, which was efficiently inhibited (ranging from 60 to 80% inhibition) by xanthenones (**4c**, **4d**, **4e**, **4g**, **4h**, **4j**, all at 200 mM) and also partially inhibited by a selective L-type Ca²⁺ channels blocker, nifedipine (5 μM). As described in literature,⁵³ the synaptosomes are commonly used to study synaptic transmission since they hold the molecular machinery necessary for the uptake, storage, and release of neurotransmitters. In addition to that, they have become common tools for drug testing.^{53,54} According to our data, nifedipine blockage did not completely abolish the increase in Ca²⁺ influx observed in synaptosomes. This data corroborates literature evidences which show that in synaptosomes just around 10% of Ca²⁺ channels are L-type and mostly are N-, P/Q-type or R-type channels.^{55,56} Furthermore, xanthenones displayed at Figure 5 were able to reduce Ca²⁺ influx to lower levels than nifedipine, suggesting that other Ca²⁺ channels might be blocked by those drugs. To clarify which channels are specifically blocked by those xanthenones detailed electrophysiological experiments would be required to test this hypothesis, which were beyond the scope of the present study.

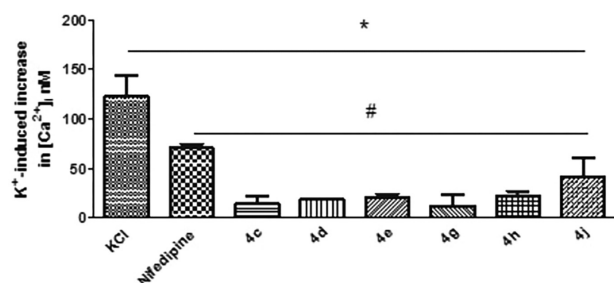


Figure 5. Effect of several xanthenones on the KCl induced cytosolic Ca²⁺ entry. The results are expressed as the mean ± S.E.M. of three independent experiments. *Indicates a significant difference compared to KCl induced Ca²⁺ entry ($p < 0.01$). #Indicates a significant difference compared to nifedipine ($p < 0.01$).

As several xanthenones were able to block Ca²⁺ influx in synaptosomes, we decided to test their effects on a model

where L-type Ca²⁺ channels are mostly expressed.⁵⁷ Thus, we performed an assay to evaluate vascular contraction induced by KCl. KCl induces vascular contraction due to vascular smooth muscle cell membrane depolarization by changing the potassium equilibrium potential and clamping the membrane potential at some value above the resting level.⁵⁸ Membrane depolarization increases [Ca²⁺]_c (cytosolic calcium) through L-type Ca²⁺ channels, activating Ca²⁺/calmodulin-dependent protein kinase II (CaMKII), which induces myosin light chain kinase phosphorylation and, consequently, vascular contraction.⁵⁹ An important component of blood pressure (BP) regulation is dependent on the contractile state of vascular smooth muscle cells (VSMCs), which is controlled to a large extent by the activities of KCNQ (subtype of potassium channel from K_v7 family) and also L-type Ca²⁺ channels.⁶⁰ Indeed, the clinical importance of L-type Ca²⁺ channels blockers is sustained by its involvement in BP regulation.

Our results show that after all xanthenones from Figure 5 were tested, the pre-treatment with xanthenones **4c** or **4e** was able to reduce the maximum contraction effect (E_{max}) induced by KCl in endothelium-denuded aortic rings (Figure 6 and Table 4) compared to the untreated control group. These results were in accordance to vascular effect induced by nifedipine which is a selective L-type Ca²⁺ channels blocker with most initial studies being performed in the early 1970s. Mikkelsen *et al.* (1979),⁶¹ as well as Pedersen *et al.* (1978),⁶² verified that nifedipine induced almost completely relaxation of arteries and veins pre-contracted by potassium. In the same way, Mikkelsen *et al.* (1978)⁶³ showed that nifedipine depressed the potassium contracture of human vessels. In vessels, vasoconstriction induced by KCl was inhibited by calciseptine (L-type Ca²⁺ blocker) or mibefradil (not selective T-type Ca²⁺ blocker) indicating

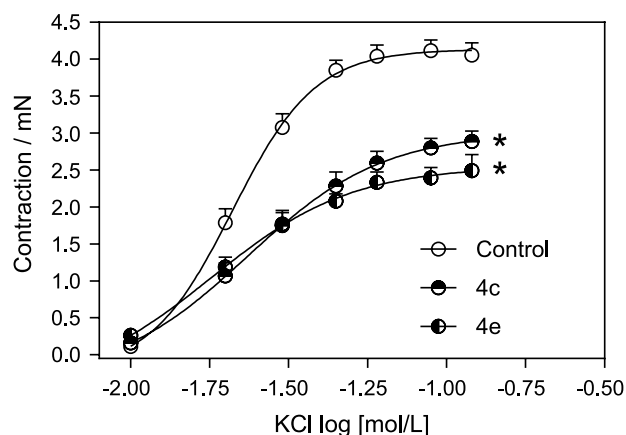


Figure 6. Effect of xanthenones on KCl-induced contractile response in endothelium-denuded aortic rings. The concentration-response curves for KCl were determined in the absence of control or after incubation with **4c** or **4e** (100 μmol L⁻¹).

that in this vascular preparation, these calcium channels are necessary for vasoconstriction.⁶⁴ Based in all these informations, our results suggest that xanthenones **4c** and **4e** could be partially blocking L-type Ca^{2+} channels, that are one of the main subtypes present in vascular tissue.^{61,62} More experiments to determine if those xanthenones may block another specific subtype of Ca^{2+} channel or may act in receptor-operated Ca^{2+} channels are needed to be studied to elucidate this point. We believed this data may be interesting in the identification of hypotensive and vasorelaxing activities of synthetic xanthenones.

Table 4. Effect of xanthene **4c** and **4e** on the E_{max} for KCl

Xanthene (100 $\mu\text{mol L}^{-1}$)	E_{max} / mN
Control	4.1 ± 0.2 (8)
4c	3.2 ± 0.4^a (4)
4e	2.9 ± 0.1^a (5)

^aCompared to respective control group ($p < 0.05$, ANOVA followed by Newman-Keuls). The number within parentheses indicates the number of animals. Values are mean $\pm$ S.E.M.

Conclusions

The present work described an alternative green protocol for the preparation of xanthenones using natural organic acids as catalysts, under solvent-free and microwave irradiation conditions. The oxalic acid was the best catalyst and furnished the xanthenones in moderate to excellent yields (51-93%). This protocol presented itself as being more eco-friendly and user-friendly alternative to traditional methodologies. The unprecedented theoretical study showed that the regioselectivity observed for such reaction is a consequence of the kinetic and thermodynamic control of the addition of β -naphthol C2, instead of the C10 atom, to the aldehyde. Additionally, our results have disclosed the xanthenones **4c** and **4e** as novel calcium channels blockers since these compounds were able to reduce the maximum contraction effect induced by KCl in endothelium-denuded aortic rings and also partially blocking L-type Ca^{2+} channels.

Experimental

General techniques

All starting materials were obtained from commercially available sources with high-grade purity and used without further purification. ^1H and ^{13}C NMR spectra at 200 and 400 MHz were obtained on a Bruker AVANCE DPX 200 and a Bruker AVANCE DRX 400 spectrometer, respectively. The chemical shifts (δ) are expressed in parts *per million* (ppm)

and are referenced to signals from tetramethylsilane (TMS) in the ^1H NMR spectra and deuterated solvent in the ^{13}C NMR spectra. ^1H NMR data are presented in the following order: chemical shift in ppm (multiplicity, integration, coupling constant (J) in hertz (Hz)). Melting points (uncorrected) were obtained on a Mettler FP 80 HT apparatus. Infrared spectra were recorded on a PerkinElmer Spectrum One spectrophotometer. The high-resolution mass spectra were obtained using a mass spectrometer with an electrospray ionization source (ESI-MS) model Shimadzu LC-ITTOF.

Well-shaped single crystals of the xanthene derivatives were chosen, mounted on a k-goniostat and exposed to graphite-monochromated X-ray beam (Mo $\text{K}\alpha$, $\lambda = 0.71073 \text{ \AA}$) using an Enraf-Nonius Kappa-CCD diffractometer equipped with a CCD camera of 95 mm lens. X-Ray diffraction experiments were carried out at room temperature. The data collection strategy was calculated by setting ϕ scans and ω scans with κ offsets. No absorption correction was applied to the raw dataset due to the negligible absorption coefficient of the compounds using a Mo $\text{K}\alpha$ beam. The crystallographic software employed was as follows: COLLECT⁶⁵ (X-ray diffraction experiment monitoring), HKL Denzo-Scalepack software package⁶⁶ (indexing, integration and scaling of raw data), SIR2004⁶⁷ (structure solving), SHELXL-97⁶⁸ (structure refinement), MERCURY⁶⁹ and ORTEP-3⁷⁰ (structure analysis and graphical representations). Direct methods of phase retrieval were used to solve the crystal structures. All the non-hydrogen atoms in the asymmetric unit were located from the electronic density Fourier map. The early model was refined by the full-matrix least squares method based on F2. In the refinements, free anisotropic and fixed isotropic thermal displacement parameters were adopted for non-hydrogen and hydrogen atoms, respectively. The isotropic thermal displacement parameters of the hydrogens were either 20% (C-H hydrogens, except those in methyl moieties) or 50% (hydrogens in methyl and hydroxyl moieties) greater than the equivalent isotropic parameter of the bonded atom. For the positions of the hydrogens, bond distances were stereochemically constrained according to the riding model.

General procedure for the xanthene synthesis

For the xanthene synthesis, aldehyde (1.0 mmol), β -naphthol (1.2 mmol), dimedone (1.5 mmol) and organic acid (5, 10 or 20 mol%) were added to a 50 mL round-bottom flask under solvent-free conditions. The reaction proceeded under MW in a DISCOVER CEM[®] reactor, in a open flask mode, using the following conditions: temperature 130 $^{\circ}\text{C}$, maximum power 250 W, hold time 3 min, and run time 5 or 10 min at high stirring with an open

tube. After completion of the reaction, methanol was added to the mixture, and the solid was filtered and recrystallized from methanol to afford the xanthenones in high purity.

Theoretical methodology

Theoretical calculations were performed using the Gaussian 09 program package.⁷¹ Gas phase optimization and evaluation of the harmonic frequencies at the HF/6-31G(d,p) level of theory were performed. Geometries were characterized as true minima in the potential energy surface (PES) when all vibrational modes were real.

It was employed the LC- ω PBE/6-311++G(d,p) level of theory⁷²⁻⁷⁶ as implemented at Gaussian 09 program.⁷¹ As criteria to geometry optimization convergence it was considered 10^{-7} Hartree to the energy and 10^{-6} Hartree Bohr⁻¹ to forces. The search for the transition states (TS) was done using QST3 method.⁷⁷ Harmonic vibrational frequency calculations were conducted analytically for vibrational analysis. Intermediates of the mechanisms were species with all real harmonic frequencies in the PES, while the TSs present one imaginary frequency. Intrinsic reaction coordinate (IRC)⁷⁸ calculations were carried out from each TS.

Purification of synaptosomes

Adult Swiss mice of both sexes (25-30 g) were decapitated and had their cortices removed and homogenized in 1:10 (m/v) 0.32 M sucrose solution containing dithiothreitol (0.25 mM) and ethylenediamine tetraacetic acid (EDTA; 1 mM). Then, the homogenates were submitted to low-speed centrifugation ($1000 \times g$, 10 min), and the synaptosomes were purified from the supernatant by discontinuous Percoll-density gradient centrifugation,⁷⁹ as previously described.⁸⁰ The isolated nerve terminals were resuspended in Krebs-Ringer-HEPES solution (KRH; 124 mM sodium chloride, 4 mM KCl, 1.2 mM magnesium sulfate, 10 mM glucose, 25 mM HEPES, pH 7.4) with no added calcium chloride to a concentration of approximately 10 mg mL⁻¹ followed by division into 200 μ L aliquots that were kept on ice until loaded with Fura-2 AM (stock solution 1 mM in dimethyl sulfoxide) for 30 min. The Ethics Committee in Animal Experimentation of the Federal University of Minas Gerais (CETEA/UFGM) approved all the *in vivo* experimental procedures under protocol number 139/2013.

Measurements of cytosolic Ca²⁺ concentration [Ca²⁺]_i

Fura-2 AM measurements of the cytosolic calcium concentration [Ca²⁺]_{initial} in synaptosomes were performed on

a Biotek spectrofluorimeter according to Gryniewicz *et al.*⁸¹ Calcium chloride was added to the synaptosomal suspension at the beginning of each fluorimetric assay (1.0 mM final concentration). Synaptosomes were stirred throughout the experiment in a plate maintained at 35 °C. Xanthenones were added to the synaptosomal suspension 10 minutes prior to membrane depolarization with KCl (33 mM). Baseline (240 s) measurements were obtained prior to the addition of drugs. At the end of each experiment, sodium dodecyl sulfate (SDS) 10% (0.1% final) was added to obtain maximal fluorescence ($R_{\max}$) by 3.0 M Tris + 400.0 mM EGTA (pH 8.6) to obtain minimal fluorescence ($R_{\min}$).⁸¹

Animals, vessel preparation and isometric tension measurements

Male Balb/c mice were housed under standard laboratory conditions with free access to food and water. Male Balb/c mice 8 to 12 weeks old (28-32 g) were killed by decapitation. The thoracic aorta was quickly removed, cleaned of adherent connective tissues and cut into rings (2-3 mm length). In all the rings, the endothelium was mechanically removed by gently rolling the lumen of the vessel on a thin wire. The aortic rings were placed between two stainless-steel stirrups and connected to an isometric force transducer (Letica Scientific Instruments; Barcelona, Spain) to measure the tension in the vessels. The rings were placed in an organ chamber containing Krebs solution with the following composition (mmol L⁻¹): sodium chloride, 135.0; KCl, 5.0; potassium dihydrogen phosphate, 1.17; calcium chloride, 2.5; MgSO₄, 1.4; sodium bicarbonate, 20.0; glucose, 11.0. The solution was maintained at pH 7.4 and gassed with 95% oxygen and 5% carbon dioxide at 37 °C. The rings were initially stretched to a basal tension of 0.5 g, and then, they were allowed to equilibrate for 60 min. Endothelial denudation was assessed qualitatively by the degree of relaxation induced by acetylcholine EC₅₀ (1 μ mol L⁻¹) in the presence of the contractile tone induced by phenylephrine (0.1 μ mol L⁻¹). For these studies with the endothelium-denuded vessels, the rings were discharged when there was more than 10% relaxation. The Ethics Committee in Animal Experimentation of the Federal University of Minas Gerais (CETEA/UFGM) approved all the *in vivo* experimental procedures under protocol number 164/2009.

Vascular contraction induced by potassium chloride (KCl) in denuded mice aorta in the presence or absence of **4c** and **4e**

The aortic rings without the endothelium were incubated with and without **4c** or **4e** (100 μ mol L⁻¹) for

30 minutes. After this time, the cumulative concentration-response curves for KCl (10-120 mM) were determined.

Statistics

The maximum effect (E_{max}) was considered to be the maximal amplitude response reached in the concentration-effect curves for the relaxant agent. The data are expressed as the mean $\pm$ S.E.M. (standard error of mean). In each set of experiments, the number of animals used (n) indicates the number of tissues studied. The differences between the mean values were assessed using one-way analysis of variance (ANOVA) followed by a Newman Keuls *post-hoc* test and values of $p < 0.05$ were considered to be significant.

Supplementary Information

The experimental data, copies of ^1H NMR and ^{13}C spectra of all xanthenones synthesized, crystal data and refinement parameters of **4e** and all information of theoretical studies are available free of charge at <http://jbcs.org.br> as a PDF file.

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