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Dynamic Trapping as A Selective Route to Renewable Phthalide from Bio-based Furfuryl Alcohol

Christopher S. Lancefield[†],^[a] Bart Fölker[†], ^[a] Razvan C. Cioc,^[d] Katarina Stanciakova,^[a] Rosa E. Bulo,^[a] Martin Lutz,^[b] Marc Crockatt^[C] and Pieter C.A. Bruijnincx^{*[a,d]}

[a]	B. Fölker, Dr. C. S. Lancefield, K. Stanciakova, Dr. R. E. Bulo, Prof. P. C. A. Bruijnincx.	
	Utrecht University	
	Universiteitsweg 99, 3584 CG, Utrecht, The Netherlands	
	E-mail: p.c.a.bruijnincx@uu.nl	
[b]	Dr. M. Lutz	
	Crystal and Structural Chemistry, Bijvoet Centere for Biomolecular Research	
	Utrecht University	
	Padualaan 8, 3584 CH, Utrecht, The Netherlands	
[c]	Dr. M. Crockatt	
	Department of Sustainable Process and Energy Systems	
	TNO	
	Leeghwaterstraat 44, 2628 CA, Delft, The Netherlands	
[d]	Dr. R. C. Cioc, Prof. P. C. A. Bruijnincx.	
	Organic Chemistry and Catalysis, Debye Institute for Nanomaterials Science	
	Utrecht University	
	Universiteitsweg 99, 3584 CG, Utrecht, The Netherlands	
	[†] Authors contributed equally. Concerting information for this article is given via a link at the	a and of the decum

[†]Authors contributed equally. Supporting information for this article is given via a link at the end of the documen

Abstract: A novel route for the production of the versatile chemical building block phthalide from bio-renewable furfuryl alcohol and acrylate esters is presented. Two challenges that generally limit sustainable aromatics production via Diels-Alder (DA) aromatization, i.e. an unfavorable equilibrium position and undesired regioselectivity when using asymmetric addends, have been addressed by using a dynamic kinetic trapping strategy. Our strategy involves the use of activated acrylates to speed up both the forward and reverse DA reactions, allowing for one of the four DA adducts to undergo a selective intramolecular lactonization reaction in the presence of a weak base. This way, the adduct is removed from the equilibrium pool, pulling the system completely to the product with the now fixed, desired regiochemistry. A single 1,2-regioisomeric lactone product is now formed in up to 86% yield and the acrylate activating agent liberated for reuse. The lactone could be aromatized to give phthalide in almost quantitative yield in the presence of Ac₂O and a catalytic amount of strong acid or in 79% using only catalytic acid.

The synthesis of renewable 'drop-in' aromatics from bio-based chemical building blocks is a rapidly emerging technology aimed at reducing the dependency of the chemical industry on fossil resources.^[1] Additionally, the rise of shale gas and its potential impact on the production of certain key aromatic building blocks has increased interest in feedstock diversification in order to safeguard production.^[2] One route to aromatics that has received significant attention of late combines a Diels-Alder (DA) reaction of biomass-derived furanics with a second dehydration (aromatization) step (Scheme 1A).^[1] Notable examples include the synthesis of: p-xylene from dimethylfuran and ethylene,[3-14] acrylic acid^[15,16] or acrolein;^[17] benzene/toluene from furan and ethylene/propylene;^[18] and (substituted) phthalic anhydride from (substituted) furan and maleic anhydride.^[19-21] In general, there are two key considerations that determine how successful these reactions are: 1) the kinetic and thermodynamic efficiency of the DA reaction and 2) the ease with which the resulting DA cycloadduct can be aromatized. Either one can make or break the process and the outcome depends heavily on the diene/dienophile combination used.

As can be seen from the previously highlighted examples, biobased DA routes to aromatics typically make use of normal electron-demand cycloadditions between electron-rich (reduced and defunctionalized) furans and electron-deficient or neutral dienophiles. Furthermore, it is common for either one or both addends to be symmetrical, thereby ensuring only one regioisomer is present in the final aromatized products. Unfortunately, these requirements lead to routes with poor atom and redox economy when considering the C₅ or C₆ carbohydrate feedstocks used to generate the bio-based furanics. For example, the synthesis of furan, which is also used to synthesize maleic



Scheme 1. A) General scheme for the production of bio-based aromatics from furanics via DA and dehydration reactions. B) Intra-molecular DA reaction reported by Torosyan *et al.*^[22] that could not be reproduced. C) This work: dynamic kinetic trapping of DA adducts for the efficient synthesis of phthalide.

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anhydride, requires decarbonylation of furfural,^[23] while (di)methylfuran requires extensive reduction with hydrogen. From an efficiency point of view, it would clearly be advantageous to develop new routes utilizing feedstocks that are closer to those obtained directly from biomass. To this end, Davis et al. have reported the synthesis of terephthalic acid via DA reactions of ethylene and 5-hydroxymethylfurfural (HMF) variants, with moderate yields and selectivities,^[24] and Jerome et al. very recently demonstrated the synthesis of a mixture of ortho and meta substituted aromatics starting from activated furfural derivatives and acrylonitrile.^[25] Here, we report on a new strategy for renewable aromatics production via DA aromatization, i.e. a new route to the versatile synthon phthalide (1). This 1,2substituted aromatic building block is synthesized from biobased furfuryl alcohol (2) and acrylic acid^[23] derivatives which can be accessed in 100% carbon yield from the C5 and C6 carbohydrate fractions, via furfural^[26] and lactic acid,^[23] respectively. Alternatively, lactic acid or directly acrylic acid are available via the chemocatalytic upgrading of bioglycerol.^[27,28] By exploiting a dynamic kinetic trapping strategy, we overcome limitations concerning the regioselectivity and equilibrium position that are inherent to this particular DA reaction and prototypical of some of the general challenges faced in DA aromatization as a method for sustainable aromatics production. In this way, 1 is obtained in an overall yield of 84% from 2.

As 2 and acrylic acid are both asymmetric, their DA reaction naturally not only leads to a mixture of endo/exo stereoisomers, which are inconsequential in the preparation of aromatic compounds, but also to ortholmeta regioisomers,^[29] with no apparent strong kinetic or thermodynamic preference for a single product. Upon dehydration this would generate mixtures of 1,2and 1,3-substituted aromatics. To overcome this regioselectivity issue, we originally pursued a strategy centered around an intramolecular DA reaction that would give complete control over the regio- and relative stereo-chemical outcome of the DA reaction, thus ensuring the correct connectivity for the eventual formation of only 1,2-substituted aromatic products. This was inspired by the work of Torosyan et al. who reported that the intramolecular DA reaction of furfuryl acrylate (3) proceeded to give cycloadduct 4 in a 50% (isolated) yield,[22] which we expected could be readily aromatized to give 1 (Scheme 1B).

In our hands, the reported procedure for the preparation of 4, which required 1 to be supported on diatomaceous earth and then heated at 60 °C for 60 h, gave no detectable intra-molecular DA product, however. Indeed, no reaction was observed at all. Further, attempts to perform the reaction in solution in a range of solvents (d₆-DMSO, CDCl₃, d₆-benzene, d₈-toluene, d₄-methanol, d₆-acetone, D₂O), or in the presence of Lewis acids (e.g. ZnCl₂, Sc(OTf)₃, Hf(OTf)₄) also failed to provide any of the expected product, as judged by ¹H NMR. Given this disappointing result, and with reports of the successful intra-molecular DA reaction of closely related furfuryl fumarates^[30,31] and maleates in mind,^[32] we decided to carry out a computational investigation into the feasibility of the target reaction using Density Functional Theory method. The computational results supported our experimental observations, suggesting the targeted intra-molecular DA is not feasible (Figure 1). In line with previous suggestions from experimental studies on furfuryl fumarates,[30] we found that initial formation of the reactive conformation (rotamer) was associated with a relatively large increase in Gibbs free energy (ΔG) of approximately 10.0 or 10.8 kcal/mol for the endo or exo reaction pathways respectively. Accessing this reactive conformer involves repositioning of the acrylate group in closer proximity to the diene and a change of the ester from a s-cis to a s-trans arrangement. This is associated with a large energy penalty of 10.0 to 10.8 kcal/mol due to the loss of the stabilising anomeric effect, increased steric strain and a larger overall dipole moment (0.9 D for the lowest energy conformer versus 4.5-4.8 D for the reactive rotamers), meaning the equilibrium population of either of the reactive conformers will be very low. Furthermore, examining the transition state energies shows that the endo pathway has a significantly higher activation energy than the exo one (30.1 vs 39.8 kcal/mol), but both are suffiently high that the forward reaction would not be feasible. Additionally, the high relative energies of the products indicates a highly endergonic reaction, further ruling out the feasibility of this intra-molecular reaction, in line with our experimental results.



Figure 1. Gibbs free energy profiles of intramolecular DA reactions calculated in the gas phase at 323K using the B3LYPD3 XC functional together with the TZ2P basis set. The global minimum substrate conformer was set to zero and all other energies were calculated with respect to these zero points.

One additional important outcome of the computational analysis was that it allowed us to identify 4-exo as a potentially stable compound, depending on synthesis conditions, with an activation energy for the retro-DA reaction of 25.7 kcal/mol, leading us to consider an alternative synthetic route. We decided to employ a dynamic kinetic trapping strategy (Scheme 1C) to produce 4-exo, making use of acyclic activated acrylates. Based on the principle of rate enhancement by the proximity effect, we anticipated to be able to trap, as the lactone, a single isomer from the mixture of isomers produced from the DA reaction of 2 and a suitably activated acrylate (Scheme 2). This approach is related to one previously utilised in the DA reaction of 2 and itaconic anhydride.^[33] From surveying commercially available acrylates we identified hexafluoroisopropyl (HFIP) acrylate (5) as a potentially suitable dienophile where the electron withdrawing HFIP group serves a dual purpose to both activate the dienophile towards DA reactions and also the ester group towards lactonization. Indeed, HFIP esters have been shown to be viable alternatives to other traditional activated esters in amide bond forming reactions,^[34] although their use in DA reactions is surprisingly seldom reported.[35,36]

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Dynamic Kinetic Trapping Using Activated Acrylates



Acrylates, Furfuryl Alcohols and Lactonized Cycloadducts Examined in this Study



Scheme 2. A new route to phthalide involving an intermolecular DA reaction of furfuryl alcohols with activated acrylates followed by selective dynamic kinetic trapping of only one isomer.

The reaction of 2 and 5 (1:1 molar ratio) neat at room temperature, proceeded smoothly to give, after 96 h, an equilibrium mixture of the four expected cycloadducts (9), totalling 66%, and remaining 2 and 5. In contrast to anhydrides,^[33] no spontaneously lactonized products could be detected under these conditions. Therefore, to promote this reaction, a catalytic amount of a weak base, NaHCO3 (20 mol%), was added to the crude mixture of cycloadducts. Gratifyingly, ¹H NMR analysis (Figure S2) revealed that one of the cycloadducts, the minor 9-exo/ortho isomer, is selectively transformed into a new compound, assigned to 4-exo based on conformational modelling, 2D NMR analysis and later confirmed via single crystal X-ray structure determination (Figure S3). The observed stereochemistry is consistent with the above computational studies showing 4-exo to he the thermodynamically favoured product. Notably, the analytical data obtained for 4-exo is substantially different to that reported by Torosyan et al. (Table S2),^[22] and given that computational analysis indicates 4-endo would be an unstable compound with respect to the retro-DA reaction (Figure 1), it seems highly unlikely that the original assignment of 4 was correct.

Due to the low equilibrium yield of the key 9-exolortho cycloadduct and the relatively slow kinetics of this DA reaction at room temperature, this did not represent a practical route to 4-exo and so optimization studies were required (Table 1). Therefore, the reaction temperature was increased to 60 °C, allowing for relatively rapid equilibration of addends and adducts, thus allowing us to apply dynamic trapping of the exo/ortho cycloadduct, by irreversible lactonization as 4-exo, as an optimization strategy. At this temperature and 20 mol% NaHCO₃ loading, 4-exo was formed in 17% yield but with a disappointing selectivity of just 20% (Entry 1). Furfuryl acrylate (3) was identified as the major side product leading to the low selectivity. Two possible routes exist for the formation of 3: direct base-catalysed transesterification of 5 by 2 or via a retro-DA reaction from 4-exo. Stability testing of the (isolated) lactone (Figure S4) suggested that 3 is formed via the former route, as 4-exo showed a relatively high thermal stability up to 100 °C. Thus, optimization of the base loading and reaction temperature led to an improved yield of 64%, with an 82% selectivity (Entry 5). A small increase in yield was observed when using 1.5 eq. of 5 (Entry 7), but changing the base or running the reaction in a solvent did not further improve the yield. Changing the acrylate to methyl acrylate (6) showed that activation of the ester was critical for the success of the reaction (Entry 14). Ultimately, the best yield of 4-exo (86% at 91% selectivity), was obtained using 4-nitrophenol acrylate (8) as the activated dienophile (Entry 16,). For both 5 and 8 recovery of the activating group should be readily achievable, for example 1,1,1,3,3,3-hexafluoroisopropyl alcohol can be recovered by distillation and 4-nitrophenol by selective extraction with a mild base, allowing recycling of these groups. Furthermore, we found that the same trapping approach could be used with 2,5bis(hydroxymethyl)furan (10, Entry 17), but introduction of a deactivating electron withdrawing group on the furan ring, as in hydroxymethylfurfural (11), was not tolerated (Entry 18).

With a reliable, high yielding route to **4-exo** in hand, our focus turned to the aromatization reaction (Table 2). Using a mixture of methanesulfonic acid (MsOH) and Ac₂O, as reported by Mahmoud *et al.*,^[20,37] **4-exo** could be readily aromatized to **1** in excellent yield (Entry 1). Interestingly, in this case, we found that

Table 1. Optimization of the DA-lactonization reaction between furfuryl alcohols and activated acrylates. Yields after 22 h using $\bf 2$ as the substrate unless otherwise stated.

Entry	Acrylate	Base	mol%	T (°C)	Yield (%)	Selectivity
1	HFIP	NaHCO ₃	20	60	17	20
2	V		2	60	34	43
3			20	80	26	28
4			2	80	54	66
5			1	80	64	82
6			0.5	80	38	70
7 ^b			1	80	68	88
8		CH ₃ O ₂ Na	20	80	23	27
9			2	80	44	55
10		CHCl ₂ CO ₂ Na	20	80	45	58
11			2	80	31	50
12		NEt ₃	2	80	30	34
13ª		NaHCO ₃	2	80	43 (68) ^c	83 (74) ^c
14	Me	NaHCO ₃	20	80	2	8
15	TFE	NaHCO ₃	20	80	33	39
16	4NP	NaHCO ₃	2	80	86	91
17 ^d	HFIP		1	80	61	66
18 ^e	HFIP		1	80	0	0

[a] Reaction run in EtOAc. [b] 1.5 eq. of HFIP acrylate. [c] Yield after 216 h. [d] Using **10** as substrate. [e] Using **11** as substrate.

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only catalytic amounts of MsOH are required (Entry 2-6), with lower loadings actually proving beneficial to the yield, which was near quantitative down to 2 mol% MsOH. Following this reaction at low temperature (0-4 °C) by NMR revealed the formation of several intermediates (Figure 2 and Figure S10) which based on NMR analysis and literature precedence^[20] we assigned to di- and

 Table 2. Optimization of the aromatization reaction to give phthalides. All reactions with 4-exo unless stated otherwise.

R = H, 4-exo	R = H, 1
R = CH ₂ OH 12-exo	R = CH ₂ OAc

	R = H, 4-exo R = I R = CH ₂ OH, 12-exo R = CH ₂ '			R = H, 1 CH ₂ OAc, 1	[:] H, 1 ₂ OAc, 13		
Entry	Solvent	Catalyst	Loading (eq.)	Ac ₂ O	T (°C)ª	Yield (%) ^b	
1	Neat	MsOH	13	20 vol%	20	75	
2			0.5	4 eq.	20/80	97°	
3			0.5	4 eq.	80	98	
4			0.1	4 eq.	80	95	
5			0.02	4 eq.	80	94	
6			0.01	4 eq.	80	39	
7		TfOH	0.01	4 eq.	80	95	
8		H ₂ SO ₄	0.01	4 eq.	80	80	
9		Amberlyst 15	0.1	4 eq.	80	82	
10	EtOAc		0.1	4 eq.	80	78	
11			0.1	2 eq.	80	89 (79) ^d	
12 ^e	Neat	MsOH	0.5	8 eq.	80	60 ^f	
13	Neat	MsOH	13	0	20	66	
14	dଃ- Toluene	MsOH	0.1	0	80	66	
15		Hf(OTf) ₄	0.1	0	80	60	
16		TfOH	0.1	0	80	63	
17		TfOH	0.01	0	80	63	
18		TfOH ^g	0.01	0	80	58	
19	CDCI ₃	TfOH	0.1	0	80	79	
20	CH₃COOH	TfOH	0.1	0	80	56	

[a] acid/Ac₂O added at 0 °C and then warmed to 20 or 80 °C as specified. [b] Determined by quantitative ¹H NMR. [c] reaction mixture heated to 80 °C for 1 h after 22 h at 20 °C. [d] Yield using recycled Amberlyst catalyst. [e] Using **12-exo** as the substrate. [f] isolated yield. [g] Silica supported TfOH acid.^[38]

Figure 2. Time course for the aromatization of 4-exo corresponding to Table 2, entry 2. [a] Mixture heated at 80 °C for 1 h.



mono-acetates representing the different stages of dehydration in the presence of Ac₂O.

Further acid screening showed that trifluoromethanesulfonic acid (TfOH) (Entry 7), H₂SO₄ (Entry 8) and (recyclable), Amberlyst 15 resin (Entry 9) were all capable of catalyzing the aromatization reaction. With TfOH, the strongest acid tested, loadings as low as 1 mol% gave excellent yields of 1. The reaction could also be run in EtOAc as a solvent and with lower amounts of Ac₂O without negative effects (Entry 10-11). 12-exo could also be aromatized under these conditions (Entry 12), although the yield was lower than for 4-exo. In the absence of a strong acid catalyst (i.e. neat Ac₂O only) no conversion was observed. Interestingly, in the absence of Ac₂O but presence of strong acid the dehydration proceeded, but with somewhat lower yields of 56-79%, highlighting the important role of Ac₂O in intercepting and stabilizing reactive intermediates during the aromatization process and thereby limiting non-productive polymerization reactions. Notably, these yields are still much higher than those obtained previously in the synthesis of phthalic anhydride,[37] suggesting 4-exo is particularly amenable to dehydration. Although rigorously dried solvents are not required for the aromatization reaction, we did find that added water has a negative effect, significantly reducing both the conversion and selectivity (Table S1).

In conclusion, we have developed a new and high yielding route (84% over two steps) to bio-based phthalide (1) from furfuryl alcohol (2) and activated acrylates such as 1,1,1,3,3,3-hexafluoroisopropyl acrylate (5) and 4-nitrophenyl acrylate (8). The choice of these substrates provides excellent overall atom and redox-efficiency considering the potential parent sugars and the opportunities to recycle the activating groups. The key step in this route involves dynamic kinetic trapping of only one of the four DA cycloadducts through selective, base-assisted lactonization. This overcomes the inherent thermodynamic and regioselectivity challenges of DA reactions using asymmetric addends, providing a high yield route to phthalide (1), itself a commodity chemical and a potential precursor to bulk chemicals such as phthalic anhydride.^[39]

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Keywords: biomass • dehydration • Diels-Alder reactions • lactonization • sustainable chemistry

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Entry for the Table of Contents



A Diels-Alder trap: An intra-molecular trapping strategy allows inherent problems in the Diels-Alder reaction of bio-based furfuryl alcohol, such as regioselectivity and unfavourable equilibrium yields, to be overcome on the way to bio-based aromatics. This strategy provides a high yielding, redox and atom economical approach to produce the versatile bio-based phthalide synthon.