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A highly diastereoselective synthesis of trifluoromethylated 1,3dioxanes is described. The reaction proceeds by an addition/oxa-Michael sequence and works efficiently under mild reaction conditions, with a good substrate scope and acceptable to good yields.

Trifluoromethyl-containing heterocycles and particularly oxygenated heterocyclic systems have attracted growing attention in the last few years.¹ They are currently investigated because of their interesting applications in medicine, agriculture and material science, among others.² In spite of the continuous growth in this field, the synthesis of 1,3-dioxanes remains unexplored. Consequently, the development of new and efficient synthetic methods for that kind of heterocyclic systems is at the same time a challenge and very desirable.

Since the seminal work published by Evans and Prunet in 1993,³ the addition/oxa-Michael cascade has demonstrated to be useful in the construction of protected *syn* 1,3-diols⁴ (Scheme 1). The reaction substrates are homoallylic alcohols functionalised by a Michael acceptor. The most usual reagent is benzaldehyde but there are several examples in literature where *p*-methoxy benzaldehyde has been used successfully.⁵ Concerning the Michael acceptor and the substitution pattern the reaction has been explored with esters and amides,³ sulfones⁶ and sulfoxides,⁷ and in very few cases with substituents α to the Michael acceptor.⁸

Inspired by those previous results and by the need of a simple method to synthesise CF_3 -containing dioxanes, we decided to explore the intramolecular oxa-Michael reaction using trifluoroacetophenone and a homoallylic alcohol functionalised with an ester as substrates (Scheme 1).

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During our studies, a similar transformation on cyclic dienones in the presence of triethylamine was described by Wang et al.⁹ Even if the yields were good, the selectivity was moderate in most cases.

With our substrates, if the reaction is under thermodynamic control as it is the case with benzaldehyde,³ the major diastereomer should have all the large substituents in equatorial positions (the two alkyl chains and presumably the trifluoromethyl group).

We started our investigation with the simple known ester **1a**¹⁰ and explored the reaction under the conditions described by Wang et al. (Table 1, entry 1). After 24 hours no reaction was observed, and the starting material was recovered quantitatively. The use of a larger quantity of base and ketone or the use of another weak base such as potassium acetate did not afford better results (entries 2 and 3); therefore, we decided to turn our attention to bases and reaction conditions more familiar to us.¹¹

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Table 1 Optimisation of the reaction conditions





With tBuOLi and 2 equivalents of ketone (entry 4), the reaction proceeded and the desired dioxane 2a was isolated in 49% yield with a moderate selectivity, the major product has the same stereochemistry in all cases (see below). The change of counterion to potassium provided much better results and in 6 hours the product was isolated in 84 % yield with good selectivity (entry 5). Typically an intramolecular conjugate addition reaction of this kind is performed using successive addition of base and carbonyl compound; it was demonstrated that this serves to improve the selectivity and to increase the reaction yield.³ Hence, we used three additions of 1.1 equivalents of ketone and 0.1 equivalents of base, but after 2 hours the reaction was not complete and the product was only isolated in 70 % yield (80 % brsm, entry 6); however after 6 hours the starting material was completely consumed and the product isolated in 79% yield (entry 7). In both cases only one diastereomer was detected in the ¹H NMR spectrum of the crude reaction mixture and after purification of the product.

The stereochemistry of the major diastereomer remained uncertain. We were confident about the *cis* stereochemistry of the substituents in positions 4 and 6, but for position 2 the configuration will be dependent on the relative steric hindrance of Ph and CF_3 . Fortunately, compound **2a** was crystalline and X-ray crystallographic analysis provided its structure (Figure 1), showing the CF_3 is indeed in the equatorial position.

With these promising results in hand, we started the study of the reaction scope. All the starting materials **1** were obtained by cross metathesis reaction between the corresponding homoallylic alcohol and methyl acrylate using Grubbs II catalyst (2.5 % mol). These homoallylic alcohols were easily prepared in one step from commercially available aldehydes and allyl bromide using an aqueous Barbier reaction.¹² Then we applied the optimised conditions to all substrates **1** and we found in all cases excellent diastereoselectivity (see Table 2).



Figure 1 X-ray structure of product 2a

The use of simple aliphatic substrates afforded the best results (entries 1 to 4); there is no noticeable effect induced by the chain extension, or by the presence of aromatic rings in the aliphatic chain (compare entries 1 and 2). The use of branched R groups is also acceptable and there is no influence on the reaction diastereroselectivity; in fact, as these groups are in equatorial position we indeed expected better selectivities with these bulkier R groups. The reaction is also useful with aromatic R substituents (entries 6 to 8); however, yields are somehow lower when electron-withdrawing groups are inserted (entry 8 vs. entry 7).

The reaction mechanism is shown in Scheme 2. We postulated that this reaction is under thermodynamic control as it is the case with benzaldehyde,³ with a reversible oxa-Michael reaction. So, it is to be expected that the use of a more electrophilic carbonyl compound (trifluoroacetophenone) will cause a better reaction with the alkoxide **3**, favouring the formation of hemiketal alkoxide **4**. We hypothesise that this intermediate is stabilised by the inductive effect of the CF₃ group, facilitating the reverse oxa-Michael addition and in consequence improving the selectivity of the whole process.

Table 2 Reaction scope.^a

он R	1	0 / tBuOK (PhCOCF ₃ THI	0.3 equiv.) (3.3 equiv.) =, 0°C	$\begin{array}{c} Ph, CF_{3} \\ 0 & 0 \\ R \\ 2 \end{array}$	
-	Entry	R	Product	Yield (%) ^b	
-	1	PhCH ₂ CH ₂	2a	79	
	2	C_6H_{13}	2b	80	
	3	<i>Iso</i> propyl	2c	81	
	4	Chx	2d	82	
	5	CH₃CHOBn ^c	2e	70	
	6	Ph	2f	67 ^d	
	7	<i>p</i> -MeOPh	2g	70	
	8	<i>p</i> -ClPh	2h	63	

^aThe reaction was performed making successive additions of the base (0.1 equiv.) and the ketone (1.1 equiv.) every 15 min. ^bYields are reported for pure compounds; the ¹H NMR spectra of the crude reaction mixtures showed in all cases only one diastereomer. ^cThis product was obtained from enantiopure lactic acid and the *anti* diastereomer from the cross methatesis reaction was isolated (see supporting information). ^d80 % brsm. Journal Name

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The next step was the investigation of the effect of a substituent in the α position to the electron-withdrawing group⁸ (results in Table 3); for this purpose, we synthesised several α -substituted substrates **6** (see supporting information for the reaction sequence). When the substituent in the α position was a methyl group (entry 1), the outcome of the reaction was disappointing. The product was observed in the 1H NMR spectrum along with the intermediate hemiketal and the elimination product. We were able to isolate the desired ketal with acceptable selectivity but in low yield. On the other hand, the reaction with product 6b works smoothly and the product was isolated with excellent selectivity and in acceptable yield (entry 2); curiously the reaction was not complete after 12 hours and some starting material was recovered. The introduction of a methyl group y to the ester did not have significant effect on the selectivity (entry 3); in this case the reaction was pushed to completion with 4 additions of base and ketone to furnish 7c in 90 % yield. It has to be noted that the reaction on a substrate similar to 6c (with R = iPr) with benzaldehyde as the electrophilic reagent furnished the corresponding product in 35% yield (53% brsm) with a 67:33 selectivity.8d

Table 3 Reaction with α -substituted substrates.

$\begin{array}{c} OH & O\\ R & \overbrace{\vdots}\\ R_1 & R_2 \end{array} OR_3 \\ 6 \end{array}$			<i>t</i> BuOK (0.3 equiv.) <u>PhCOCF₃ (3.3 equiv.)</u> THF, 0°C R R R 7		F_3 O $Of_3 OR_3R_2$	
Entry	R	R ₁	R ₂	R₃	Product, Yield	Selectivity
1	PhCH ₂ CH ₂	н	Me	Et	7a , <27% [°]	82:18
2	$PhCH_2CH_2$	н	OCb^b	Et	7b , 54 [°] %	>98:2
3 ^{<i>d</i>}	$PhCH_2CH_2$	Me	OCb	Et	7c , 90 %	>98:2
л ^е	PhCH CH	ц	NUCUO	Mo	74 95 %	07.10

 a This value was calculated by analysis of the ^{1}H NMR spectrum; the crude mixture also contained the intermediate hemiketal and the elimination product; a sample was purified for analytical purpose. $^{b}Cb = CON(iPr)_{2}$ $^{c}83$ % brsm. d The reaction was performed using 4 additions of base and ketone. ^e The starting material was treated with 1.1 equivalents of base and ketone, then consecutive additions of 0.1 equivalents of base and 1.1 equivalent of ketones were made.



Figure 2 X-ray structure of product 7c.

We were also curious about the reactivity of a substrate with a nitrogen substituent in the α position. Depending on the substitution pattern of the nitrogen atom, the compound can have an enamine character or the N-H group can be acidic enough to be deprotonated before the alcohol.

We were able to synthesize compound 6d by a Horner-Wadswoth-Emmons reaction.¹³ As expected an extra equivalent of base was needed in order to completely deprotonate the NH group (entry 4). It is remarkable that this reaction proceeded in very good yield and with good selectivity, even if the deprotonation of the nitrogen can induce a nucleophilic character to the carbon in the β position to the ester. It is worth noting that in all the cases described in table 3, the reported selectivity corresponds to the position of R2; the reaction proceeds with complete selectivity for the oxygen addition.

Fortunately, the stereochemistry of the substituent in the α position could be determined by X-ray crystallographic analysis of compound 7d as shown in Figure 2. Comparison of NMR signals and coupling constants for the H-5 proton in 7d and 7b allowed us to define the configuration at C5 for compound 7b as shown in scheme 3. The stereochemistry at C5 of 7c M was assumed to be the same as that of 7b.

Some additional experiments were carried out in order to illustrate the generality of our method. First, other Michael acceptors such as Weinreb amides or sulfones were explored. The cited literature³⁻⁸ shows that the reactivity of the Michael acceptors does not have a great influence on the yield of the oxa-Michael reaction. The main difference is the diastereoselectivity.



Scheme 3 Stereochemistry for major (M) and minor (m) diastereomers.

ŌCb

7b M, J₄₋₅ = 3.3 Hz

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7c M, $J_{4-5} = 1.6$ Hz

OMe

OFt

ŌCb

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Scheme 4 Weinreb amide and sulfone as Michael acceptors

In consequence, we were curious to see if in our case Weinreb amides or sulfones are desirable substrates and if the selectivity is comparable to that obtained with esters.

As shown in scheme 4 the results are very good, the reaction yields are comparable to those obtained with simple esters and in both cases, scheme 4a and 4b, there was only one diastereosisomer visible by ¹H NMR of the crude reaction mixture and isolated products **9** and **11**. The stereochemistry of these compounds was assigned by comparison with that of the esters.

Finally, we decided to use another trifluoromethyl ketone; we speculated that, if the ketone was more electrophilic than trifluoroacetophenone, the hemiketal alkoxide **4** would be less nucleophilic and so the reaction more difficult. Thus, if the reaction worked with a ketone substituted in the aromatic ring by a strong electron withdrawing group, then it would be predictable than the reaction could be extended to the use of almost any trifluoromethyl ketone. Scheme 5 shows the result of the reaction using the *m*-nitro-trifluorocetophenone as the electrophilic reagent. In this case the reaction proved to be more difficult, four additions of base and 16 hours were needed to complete the reaction; however, product **13** was isolated in good yield and the reaction proceeded with complete diastereoselectivity.

In summary, the additional experiments let us to conclude that different Michael acceptors and other trifluoromethyl ketones can be used, thus making the method very general.

Finally, computational studies oriented to propose an explanation for the difference in reactivity and selectivity between the substrates, as well as additional experiments to increase even more the reaction scope are currently under investigation in our laboratory. The results will be reported in due course.



Scheme 5 Reaction using *m*-nitrotrifluoroacetophenone.

In conclusion, we have described a useful methodology for the synthesis of CF₃-containing 1,3-dioxanes with excellent diastereoselectivity in most cases. We also reported our preliminary conclusions about the reaction mechanism based on literature reports and experimental observations. The stereochemistry of the products was established by X-ray crystallographic analysis and extrapolation.

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Notes and references

- (a) F. M. D. Ismail, J. Fluorine Chem., 2002, 118, 27; (b) J.-A. Ma; D. Cahard, Chem. Rev., 2004, 104, 6119; (c) J. Nie; H.-C. Guo; D. Cahard; J.-A. Ma, Chem. Rev., 2011, 111, 455; (d) Y.-J. Lin; L.-N. Du; T.-R. Kang; Q.-Z. Liu; Z.-Q. Chen; L. He, Chem. Eur. J., 2015, 21, 11773.
- 2 (a) F. Li; J. Nie; L. Sun; Y. Zheng; J.-A. Ma, Angew. Chem., Int. Ed., 2013, 52, 6255; (b) J. Wang; W.-G. Kong; F. Li; J. Liu; Q. Shen; L. Liu; W.-X. Zhao, Org. Biomol. Chem., 2015, 13, 5399; (c) Y. Zhu; Z. Dong; X. Cheng; X. Zhong; X. Liu; L. Lin; Z. Shen; P. Yang; Y. Li; H. Wang; W. Yan; K. Wang; R. Wang, Org. Lett., 2016, 18, 3546.
- 3 D. A. Evans; J. A. Gauchet-Prunet, J. Org. Chem., 1993, 58, 2446.
- For a few selected examples, see: (a) D. T. Hung; J. B. Nerenberg; S. L. Schreiber, J. Am. Chem. Soc., 1996, 118, 11054; (b) T. J. Hunter; G. A. O'Doherty, Org. Lett., 2001, 3, 2777; (c) T. A. Dineen; W. R. Roush, Org. Lett., 2004, 6, 2043; (d) A. Vincent; J. Prunet, Synlett, 2006, 2269; (e) E. de Lemos; F.-H. Porée; A. Bourin; J. Barbion; E. Agouridas; M.-I. Lannou; A. Commerçon; J.-F. Betzer; A. Pancrazi; J. Ardisson, Chem. Eur. J., 2008, 14, 11092; (f) S. S. Palimkar; J. Uenishi, Org. Lett., 2010, 12, 4160; (g) A. M. Albury; M. P. Jennings, J. Org. Chem., 2012, 77, 6929; (h) R. W. Bates; T. G. Lek, Synthesis, 2014, 1731; (i) T. J. Hunter; Y. Wang; J. Zheng; G. A. O'Doherty, Synthesis, 2016, 48, 1700.
- 5 D. A. Evans; P. Nagorny; D. J. Reynolds; K. J. McRae, *Angew. Chem., Int. Ed.,* 2007, **46**, 541.
- 6 (a) R. Oriez; J. Prunet, *Tetrahedron Lett.*, 2010, **51**, 256; (b) L. Grimaud; D. Rotulo; R. Ros-Perez; L. Guitry-Azam; J. Prunet, *Tetrahedron Lett.*, 2002, **43**, 7477.
- 7 D. Gamba-Sanchez; J. Prunet, J. Org. Chem., 2010, 75, 3129.
- 8 (a) L. Grimaud; R. de Mesmay; J. Prunet, *Org. Lett.*, 2002, 4, 419; (b) D. Rotulo-Sims; J. Prunet, *Org. Lett.*, 2007, 9, 4147; (c) R. Aouzal; J. Prunet, *Org. Biomol. Chem.*, 2009, 7, 3594. (d) R. Oriez, Master Thesis, Ecole Polytechnique (Palaiseau), 2006.
- 9 F. Li; J. Wang; M. Xu; X. Zhao; X. Zhou; W.-X. Zhao; L. Liu, Org. Biomol. Chem., 2016, 14, 3981.
- 10 J. A. Gazaille; T. Sammakia, Org. Lett., 2012, 14, 2678.
- 11 In our previous studies the typical base employed was tBuOK and we performed successive additions of base and carbonyl compound.
- 12 A. Chattopadhyay, J. Org. Chem., 1996, 61, 6104.
- 13 P. A. Alexander; S. P. Marsden; D. M. Muñoz Subtil; J. C. Reader, Org. Lett., 2005, 7, 5433.

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Diastereoselective synthesis of trifluoromethylated 1,3-dioxanes by intramolecular oxa-Michael reaction

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A rapid and simple method to access trifluoromethylated 1,3-dioxanes by a sequence addition/oxa-Michael reaction has been developed; the reaction proceeds smoothly with excellent stereo selectivity.