

Cite this: *Org. Biomol. Chem.*, 2024, **22**, 8285

Unveiling the reactivity of 2*H*-(thio)pyran-2-(thi)ones in cycloaddition reactions with strained alkynes through density functional theory studies†

Wei Huang,^a Kangqiao Wen,^a Scott T. Laughlin *^a and Jorge Escorihuela *^b

Over the past two decades, click chemistry transformations have revolutionized chemical and biological sciences. Among the different strain-promoted cycloadditions, the inverse electron demand Diels–Alder reaction (IEDDA) has been established as a benchmark reaction. We have theoretically investigated the IEDDA reaction of *endo*-bicyclo[6.1.0]nonyne (*endo*-BCN) with 2*H*-pyran-2-one, 2*H*-thiopyran-2-one, 2*H*-pyran-2-thione and 2*H*-thiopyran-2-thione. These 2*H*-(thio)pyran-2-(thi)ones have displayed different reactivity towards *endo*-BCN. Density functional theory (DFT) calculations show, in agreement with experiments, that *endo*-BCN reacts significantly faster with 2*H*-thiopyran-2-one compared to other 2*H*-(thio)pyran-2-(thi)one derivatives because of the lower distortion energy. Experimentally determined second-order rate constants for the reaction of a 2*H*-pyran-2-thione with different strained derivatives, including a 1-methylcyclopropene derivative and several cycloalkynes (*exo*-BCN, (1*R*,8*S*)-bicyclo[6.1.0]non-4-yne-9,9-diylo)dimethanol, dibenzocyclooctyne and a light activatable silacycloheptyne, were used to validate the computational investigations and shed light on this reaction.

Received 31st July 2024,
Accepted 16th September 2024

DOI: 10.1039/d4ob01263a

rsc.li/obc

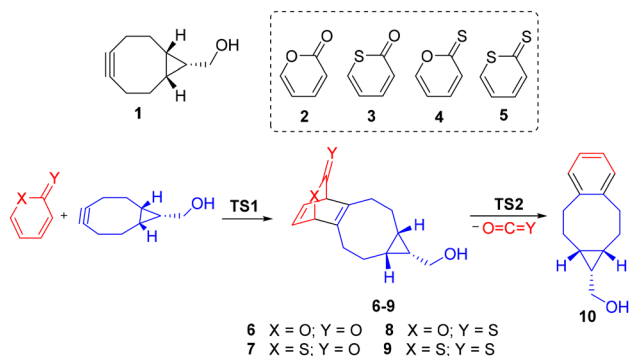
Introduction

Since the seminal 2001 work of Sharpless and coworkers,¹ click chemistry has blossomed to include new reactions optimized for diverse applications. For example, after reports showing the toxicity of Cu(I),² strain-promoted cycloadditions were developed as an alternative to the copper(I)-catalysed azide alkyne cycloaddition (CuAAC).³ Among the strain-promoted cycloadditions, the strain-promoted azide–alkyne cycloaddition (SPAAC),⁴ the inverse electron demand Diels–Alder reaction (IEDDA),⁵ and more recently, the strain-promoted oxidation-controlled quinone cycloaddition (SPOCQ),⁶ have emerged as powerful organic transformations. Due their high rates and efficiency, strain promoted cycloadditions have been successfully applied in bioconjugate functionalization,⁷ surface chemistry,⁸ and polymer science.⁹ Theoretical studies on the mechanism of SPAAC,¹⁰ IEDDA¹¹ and SPOCQ,¹² have led to a deeper understanding of these widely used organic reactions.

Among the different dienes used in IEDDA reactions, 1,2,4,5-tetrazines have been widely investigated as counterparts for reacting with alkynes/alkenes.¹³ Conversely, reactivity studies of 2*H*-pyran-2-ones and analogues, the focus of this report, are scarce. Kranjc and Kočevar reported in 2005 the Diels–Alder reaction of 2*H*-pyran-2-ones with alkynes under thermal conditions and at high pressures to afford aniline derivatives.¹⁴ In 2022, Xian and coworkers reported a click-and-release strategy based on the reaction of 2*H*-pyran-2-thiones and other analogues with a strained alkyne, (1*R*,8*S*,9*S*)-bicyclo[6.1.0]non-4-yn-9-ylmethanol (*endo*-BCN, **1**).¹⁵ In that report, computational studies based on the nucleus-independent chemical shift (NICS) were used to predict the cycloaddition efficiency of *endo*-BCN with the 2*H*-(thio)pyran-2-(thi)one analogues. In a recent work, we synthesized substituted pyrones and pyranthiones and studied their biorthogonality, demonstrating their utility in protein labelling and subcellular delivery of hydrogen sulfide in living cells.¹⁶ A deeper understanding of these reactions can help in the design of 2*H*-pyran-2-thiones and heterocyclic analogues with enhanced reactivity for potential applications in chemical biology.

Previous studies have shown that dienes 2–5 can react with strained alkynes through an IEDDA reaction to form an unstable non-aromatic intermediate 6–9 via **TS1** (Scheme 1).¹⁵ This intermediate then undergoes a retro Diels–Alder reaction through **TS2** to provide the aromatic derivative **10** and release a molecule of CO₂, COS or CS₂ as a byproduct (Scheme 1).

^aDepartment of Chemistry, Stony Brook University, 100 Nicolls Road, Stony Brook, NY 11790, USA. E-mail: scott.laughlin@stonybrook.edu^bDepartamento de Química Orgánica, Universitat de València, Avda. Vicente Andrés Estellés s/n, Burjassot, 46100 Valencia, Spain. E-mail: jorge.escorihuela@uv.es† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ob01263a>



Scheme 1 Schematic representation of the IEDDA reaction of *exo*-BCN **1** with *2H*-pyran-2-one (**2**), *2H*-thiopyran-2-one (**3**), *2H*-pyran-2-thione (**4**) and *2H*-thiopyran-2-thione (**5**).

Understanding the trends in their reactivity regarding the *2H*-(thio)pyran-2-(thio)one derivative will be key for the further development of this click reaction.

Herein, a systematic theoretical study of cycloadditions of *endo*-BCN with *2H*-pyran-2-one (**2**), *2H*-thiopyran-2-one (**3**), *2H*-pyran-2-thione (**4**) and *2H*-thiopyran-2-thione (**5**) was performed using the activation strain model. A combined experimental-computational study was conducted to evaluate reactivity of a *2H*-pyran-2-thione analogue towards different strained dienophiles including 1-methylcyclopropene (MeCp) and cycloalkynes such as *endo*-BCN, ((1*R*,8*S*,9*R*)-bicyclo[6.1.0]non-4-yn-9-yl)methanol (*exo*-BCN), ((1*R*,8*S*)-bicyclo[6.1.0]non-4-yne-9,9-diyl)dimethanol (DMBCN), dibenzocyclooctyne (DBCO) and a light activated silacycloheptyne (Si7HT).

Results and discussion

As mentioned above, oxygen and sulphur-containing heterocycles such as *2H*-pyran-2-one (**2**), *2H*-thiopyran-2-one (**3**), *2H*-pyran-2-thione (**4**) and *2H*-thiopyran-2-thione (**5**) can react with a strained cycloalkyne such as *endo*-BCN to form a non-aromatic intermediate (**6–9**) via **TS1**. Thus, we performed a computational study in order to rationalize the observed reactivity of the different oxygen and sulphur-containing heterocycles. To this end, the effect on the reactivity of the oxygen and sulphur atoms in these heterocycles was investigated by means of DFT calculations at the M06-2X/6-311+G(d,p) level of theory. Initially, a conformational search was performed to analyse the multiple conformations of *endo*-BCN (**1**) and the conformation with the lowest energy and an energy gap of 4.0 kcal mol⁻¹ towards the other conformations was selected for the mechanistic study.

The formation of the aromatic derivative **10** can be rationalized in terms of the following general reaction mechanism, which can be divided into two main steps: [4 + 2] cycloaddition and retro Diels–Alder reaction with the elimination COS, CO₂ or CS₂ depending on the oxygen or sulphur-containing heterocycle. The Gibbs energy profile for the reaction of *endo*-BCN

with the different dienes **2**, **3**, **4** and **5** is shown in Fig. 1. The first step of the mechanism involves the [4 + 2] cycloaddition and is exergonic. In line with similar Diels–Alder reactions, a pre-reaction complex with higher energy than both separated reactants was located.¹⁷ The interaction between the diene and *endo*-BCN (**1**) led to the formation of the molecular pre-reaction complex (**RC**). A decrease in the enthalpy of the reaction system of around 6 kcal mol⁻¹ was computed of this transformation accompanied with substantial reduction in the entropy (Fig. S2, ESI[†]). Therefore, Gibbs energy (ΔG) of the formation of pre-reaction complex (**RC**) has a positive value, which excluded the possibility of pre-reaction complexes as stable intermediates. As inferred from Fig. 1, *2H*-pyran-2-thione (**4**) has the lower activation Gibbs energy (21.2 kcal mol⁻¹) of the four heterocyclic derivatives (**2–5**). For *2H*-pyran-2-one (**2**), the computed activation Gibbs energy was higher (22.7 kcal mol⁻¹), whereas for *2H*-thiopyran-2-thione (**5**) the barrier was up to 25.3 kcal mol⁻¹.

Finally, *2H*-thiopyran-2-one (**3**) exhibited the highest barrier (26.5 kcal mol⁻¹) for the [4 + 2] cycloaddition among the four derivatives. Fig. 2 shows the transition state structures and activation Gibbs free energies (in kcal mol⁻¹) for the cycloaddition of BCN (**1**) with heterocyclic structures **2**, **3**, **4** and **5**. Given the lack of symmetry in the heterocyclic dienes, all cycloadditions occurred *via* non-synchronous transition state structures with C...C bond forming distances within the range of 2.13–2.47 Å. The pattern of reactivity was 3 < 5 < 2 < 4. Carbonyl derivatives, *i.e.*, *2H*-pyran-2-one (**2**) and *2H*-thiopyran-2-one (**3**), had activation Gibbs free energies of 22.7 and 26.5 kcal mol⁻¹, respectively, whereas for thiocarbonyl compounds, *2H*-pyran-2-thione (**4**) and *2H*-thiopyran-2-thione (**5**), activation Gibbs free energies of 21.2 and 25.3 kcal mol⁻¹,

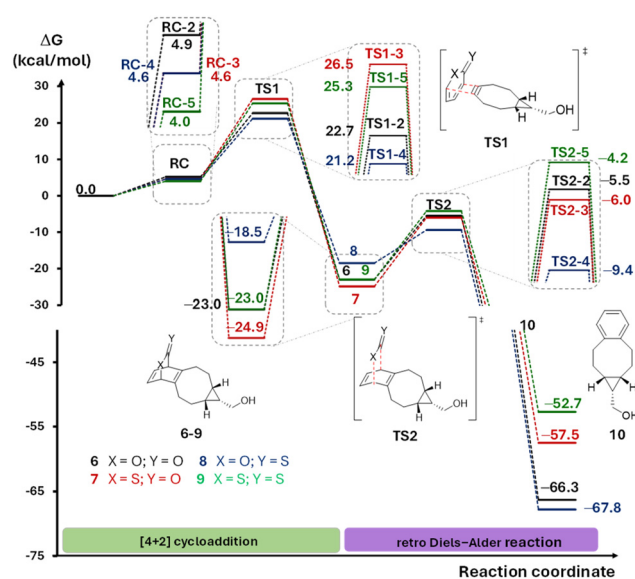


Fig. 1 Gibbs energy profile (in kcal mol⁻¹) for the cycloaddition of *endo*-BCN (**1**) with dienes **2**, **3**, **4** and **5**.



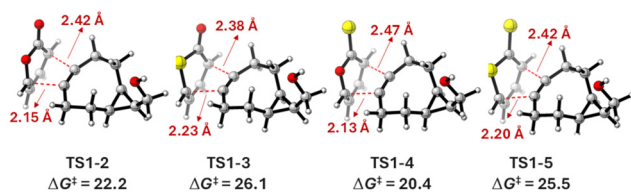


Fig. 2 Optimized transition state structures for the [4 + 2] cycloaddition. Activation Gibbs energies (ΔG^\ddagger) are given in kcal mol⁻¹, and distances are shown in Angstroms.

respectively, were computed. The analysis of transition state structures reveals that 2*H*-pyran-2-thione (4) has the longest forming bond distance (2.47 Å), and therefore the earliest early transition state leading to the lowest activation energy. The forming bond distances are almost identical for the oxygen and sulphur-containing heterocyclic structures 2, 3, 4 and 5. According to the experimental data, 2*H*-pyran-2-thione (4) reacts 3.7 times faster than 2*H*-pyran-2-one (2) in a PBS buffer (pH 7.4 with 10% DMSO).¹⁵ Computational data predict a higher reactivity for 2*H*-pyran-2-thione (4) compared to the other oxygen and sulphur-containing heterocycles.

From non-aromatic intermediates 6–9, a retro Diels–Alder reaction affords the aromatic product 10 and releases a molecule of COS for 7 and 8, CO₂ for 6 or CS₂ for 9. Fig. 3 shows the transition state structures and activation Gibbs free energies (in kcal mol⁻¹) for the retro Diels–Alder reaction to yield the final product 10. The lowest activation Gibbs energy for the retro Diels–Alder reaction was found for intermediate 8 (TS2-4) (coming from 2*H*-thiopyran-2-thione (4)), with a barrier of 13.4 kcal mol⁻¹, which delivered the final product 10 with an exergonicity of –49.3 kcal mol⁻¹. Thiopyran-derivatives 7 and 9 (coming from 2*H*-thiopyran-2-one (3) and 2*H*-thiopyran-2-thione (5)) had barriers of 18.9 and 18.8 kcal mol⁻¹, respectively; whereas intermediate 6 (TS2-2) showed a Gibbs activation barrier of 17.5 kcal mol⁻¹. After the second transition state (TS2), the formation of the aromatic product 10 is highly exergonic, acting as the thermodynamic driving force of this process. The high exergonicity observed from the retro Diels–Alder reaction is due to the recovery of the aromaticity from non-aromatic intermediates 6–9 to deliver the final aromatic

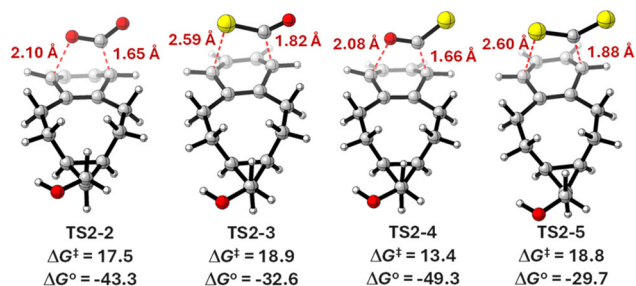


Fig. 3 Optimized transition state structures for retro Diels–Alder reaction. Activation Gibbs energies (ΔG^\ddagger) and free energies of reaction (ΔG°), and distances are shown in Angstroms. Energies are given in kcal mol⁻¹.

product 10 and the release of a molecule of COS, CO₂, or CS₂. In general, the O...C and S...C distances for the breaking bonds were around 2.1 and 2.6 Å, respectively, whereas the C...C distance for the breaking bond in the case of pyran-derivatives 6 and 8 (TS2-2 and TS2-4) was found to be shorter than that of thiopyran-derivatives 7 and 9 (TS2-3 and TS2-5). Overall, the type of atom at the heterocycle becomes a key factor with a significant effect the thermodynamic reaction feasibility (ΔG°). In this regard, those reactions involving the breaking of a C...O bond (TS2-2 and TS2-4) are more exergonic than those in which a C...S is being broken (TS2-3 and TS2-5).

Comparing the two transient steps (TS1 and TS2), the [4 + 2] cycloaddition is the rate determining step (RDS) for the IEDDA reaction between *endo*-BCN and 2*H*-(thio)pyran-2-(thio) ones, as also observed for IEDDA reactions of 1,2,4,5-tetrazines.¹⁸ According to the computed activation free energies for both transition states, and the high exergonicity of the overall process, the cycloadduct (10) would be formed after the reaction with the release of CO₂, COS or CS₂. Finally, a comparison between M06-2X/6-311+G(d,p), M06-2X/Def2TZVPP and ω B97xD/Def2TZVPP methods was performed to validate the results and each method yielded very similar results in terms of transition state barriers (ΔG^\ddagger) (see Table S3, ESI[†]).

Next, the activation strain model was used to rationalize the trend in reactivity.¹⁹ Fig. 4a shows the distortion/interaction-activation strain (D/I-AS) analysis of the transition state structures. According to this model, the optimized transition state structure is separated into two fragments (distorted BCN and distorted heterocyclic structure), and the energy difference between the distorted fragment and the corresponding optimized structure is defined as the distortion energy (ΔE_{dist}). The difference between the activation energy and the total distortion energy is the interaction energy (ΔE_{int}). This model has been successfully applied to rationalize the reactivity of cycloadditions, substitution and elimination reactions, and transition metal catalysed reactions.²⁰ As inferred from the analysis, BCN distortion varies slightly (green arrows, 2.6 to 2.9 kcal mol⁻¹), but the variation in distortion energy for each diene is more significant (blue arrows). The variation of geometrical parameters of distorted *endo*-BCN at the TS was very similar for the four structures (Table S4, ESI[†]). Small angle deviations (lower than 3°) from the *endo*-BCN reactant structure agree with the computed low BCN distortion energies. The high distortion energy for the dienes can be associated to the distorted and bended structure at the TS in comparison with the almost planar structure of the reactant. In this regard, 2*H*-pyran-2-thione (4) displayed the lowest distortion in line with the lowest activation energy. The substitution of oxygen by sulphur at the pyranthione ring (from 4 to 5) was reflected in a significant increase in the distortion energy of the heterocycle, leading to a higher activation energy. A similar effect was observed in the atomic substitution from 2 to 3.

Finally, a non-covalent interaction (NCI) analysis was performed on the transition states (Fig. 4b). Upon inspection of the NCI surfaces, it is clear for all compounds a similarly sized contour was computed, with an attractive interaction in the



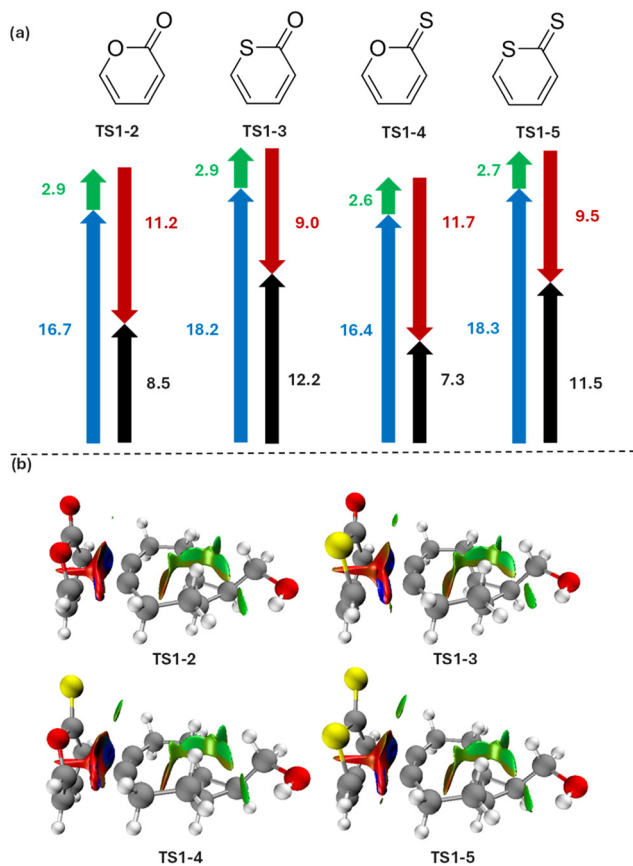


Fig. 4 (a) Distortion/interaction analysis on the transition structures (blue arrow for distortion energy of heterocyclic compound; green arrow for distortion energy of BCN; red arrow for interaction energy, and black arrow for activation potential energy). Energies are given in kcal mol⁻¹. (b) NCI surfaces for the transition state structures.

region where C–C bonds are being formed, but also with a strong repulsive interaction.

Fig. 5 shows the activation strain diagram (ASD) of the cycloaddition of *endo*-BCN with 2*H*-pyran-2-one (black line), 2*H*-thiopyran-2-one (red line), 2*H*-pyran-2-thione (blue line) and 2*H*-thiopyran-2-thione (green line). The distortion energies for reactions of 2*H*-pyran-2-one (black line) and 2*H*-pyran-2-thione (blue line) are quite similar, and significantly lower than that for 2*H*-thiopyran-2-one (red line), and 2*H*-thiopyran-2-thione (green line) along the complete reaction coordinate. This trend is also observed for the activation energy and the interaction energy, and the study suggests the reactivity is controlled by the distortion energy. An earlier transition state indicates that the transition state geometry is closer to that of the reactants, which is reflected in a lower distortion energy for 2*H*-pyran-2-thione (4). It is evident that the reactivity differences between 2*H*-pyran-2-one (2) and 2*H*-pyran-2-thione (4) with *endo*-BCN comes from the significant differences in the distortion energy between reactants along the reaction coordinate as both reactions show similar interaction energies. Furthermore, these type of reactions are inverse electron-demand Diels–Alder reactions, in which the HOMO of the die-

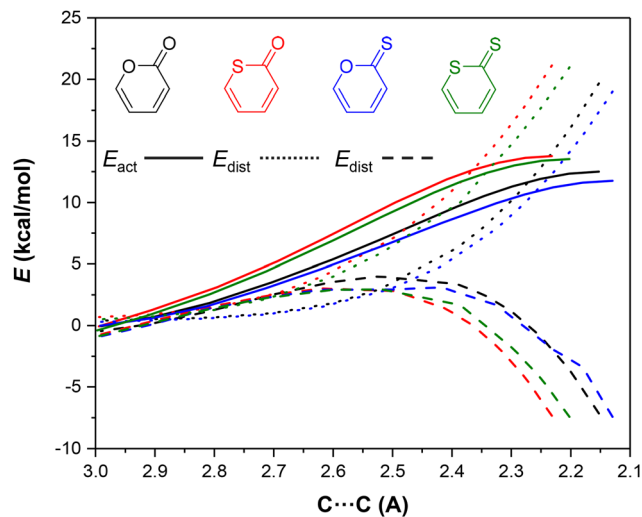


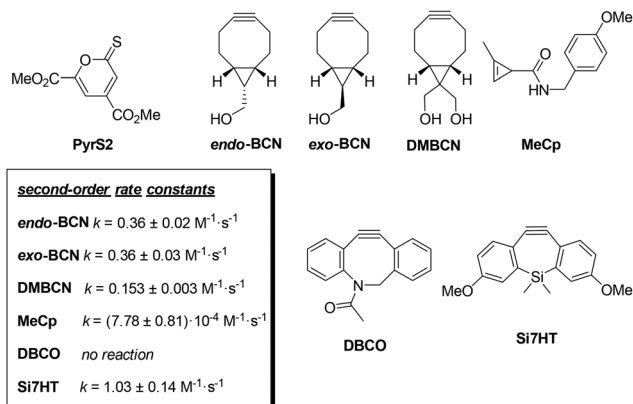
Fig. 5 Activation strain diagrams of the cycloaddition of BCN with the different heterocyclic compounds along the reaction coordinate projected onto the C...C bond stretch, computed at M06-2X/6-311+G(d,p).

nophile interacts with the LUMO of the diene. A more effective orbital overlap interaction was computed between the HOMO of *endo*-BCN and the LUMO of 2*H*-pyran-2-thione (4) compared to the other set of compounds.

Given the faster kinetics observed for 2*H*-pyran-2-thione among the different oxygen and sulphur-containing heterocycles,^{15,16} we experimentally studied the reactivity of a 2*H*-pyran-2-thione with several popular dienophiles. For this study, we choose a diester substituted 2*H*-pyran-2-thione, **Pyrs2**, the current best in class compound based on our previous studies with these reagents in protein labelling experiments.¹⁶ The dienophile reaction partners we explored were 1-methylcyclopropene (MeCp) and the cycloalkynes *exo*-BCN, DMBCN,²¹ DBCO²² and light activatable Si7HT.²³ We conducted kinetics studies for reaction of each of these dienophiles with **Pyrs2** (see ESI† for further details). Reactivity tests towards *exo*-BCN, DMBCN, DBCO and MeCp and were carried out at 25 °C in 50% CH₃CN/H₂O; whereas for Si7HT the kinetic data was determined at 25 °C in CH₃CN due to the limited stability of silacycloheptyne after UV illumination (Scheme 2). As expected, **Pyrs2** exhibited high reactivity with *exo*-BCN, resulting in a second-order rate constant of 0.36 M⁻¹ s⁻¹. Contrary to other related cycloadditions, *endo*-BCN and *exo*-BCN displayed a similar reactivity. Slightly slower kinetics was observed for DMBCN, having a second-order rate constant of 0.15 M⁻¹ s⁻¹. In the case of DBCO, no significant reaction was observed at room temperature after 24 h, whereas the methylcyclopropene derivative (MeCp) had a second-order rate constant of 8 × 10⁻⁴ M⁻¹ s⁻¹. Finally, a second-order rate constant of 1.03 M⁻¹ s⁻¹ was determined for Si7HT after light activation. In all cases, the cycloadducts were isolated after the release of COS.

To further support our experimental findings, computational investigations using DFT calculations were performed





Scheme 2 Chemical structures of 2*H*-pyran-2-thione **PyrS2**, the **PyrS2** reaction partners evaluated here, and their second-order rate constants for the reaction with 2*H*-pyran-2-thione **PyrS2**.

at the M06-2X/6-311+G(d,p) level of theory and including the implicit solvent model SMD. To this purpose, we determined the Gibbs energy barriers for the cycloaddition of 2*H*-pyran-2-thione **PyrS2** with different strained counterparts. The transition state structures and the calculated activation energy (ΔE^\ddagger), activation Gibbs energy (ΔG^\ddagger), and Gibbs energy of the reaction (ΔG_{rxn}) are displayed in Fig. 6a. It is worth mentioning that activation energy barriers (ΔE^\ddagger) are lower than Gibbs energy barriers (ΔG^\ddagger). Gibbs energy barriers take into account both enthalpy and entropy changes that occur during a chemical reaction. These differences reveal that entropic contributions on these reactions are significant. The TS structures are all asynchronous with C...C bond forming distances ranging from 2.24 to 2.34 Å. The lowest activation Gibbs energy was computed for Si7HT, in agreement with the faster experimentally determined kinetics, having an earlier transition state and a longer forming C...C bond distance than other strained counterparts. The computed barriers for the BCN derivatives *exo*-BCN and DMBCN were found to be 19.0 and 18.8 kcal mol⁻¹, respectively. In contrast, to other related cycloadditions, *endo*-BCN and *exo*-BCN displayed a similar reactivity. In the case of DBCO, the high activation Gibbs energy barrier of 25.1 kcal mol⁻¹ is in agreement with the lack of reactivity of this dibenzocyclooctyne derivative, as observed for other cycloaddition reactions.^{12a} In agreement with experimental results, the computed rate constants that predict: (1) DMBCN will react faster than *endo*-BCN and *exo*-BCN with 2*H*-pyran-2-thione **PyrS2**, (2) MeCp has a low reactivity, and that (3) Si7HT will react with **PyrS2** around 3 times faster than *endo*-BCN and *exo*-BCN. To our surprise, the kinetics observed for DMBCN and **PyrS2** were slightly slower than expected from our computational findings, which might imply other factors like solvent effects played some role in reaction kinetics.

The D/I-AS analysis of the transition state structures of 2*H*-pyran-2-thione **PyrS2** with the different studied strained counterparts is shown in Fig. 6b. In this case, the D/I-AS analysis was performed at the transition state as a defined point

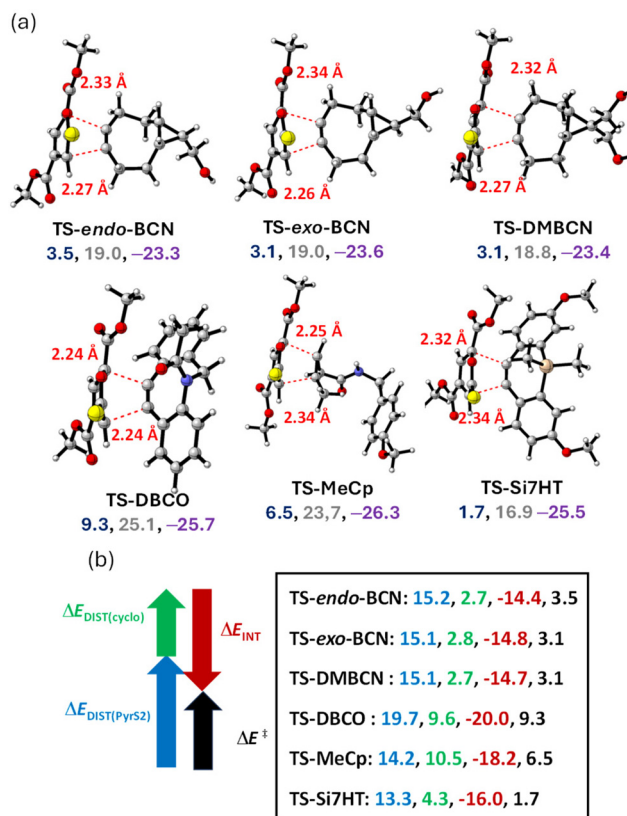


Fig. 6 (a) Optimized transition state structures at the M06-2X/6-311+G(d,p) level of theory with bond forming distances in Å. Values of activation energy (ΔE^\ddagger), activation Gibbs energy (ΔG^\ddagger), and Gibbs energy of the reaction (ΔG_{rxn}) are shown below each structure in black, green, and purple, respectively, in kcal mol⁻¹. (b) Values of distortion energy of pyrones ($E_{\text{DIST}}(\text{pyro})$, in blue), distortion energy of strained cyclooctadiene ($E_{\text{DIST}}(\text{cyclo})$, in green), interaction energy (E_{INT} , in red) and activation energy (ΔE^\ddagger , in black). Energies are given in kcal mol⁻¹.

for each reaction under study. It should be noted that performing the D/I-AS analysis at the state geometry can lead to incorrect interpretations if transition states occur at highly different bond forming distances.²⁴ However, here the forming bond lengths are very similar in all cases, warranting the comparison at the transition state. As shown in Fig. 6b, the distortion energy for **PyrS2** (blue arrow) decreased with the smaller ring sizes with relatively smaller energy for MeCp and Si-7HT (14.2 and 13.3 kcal mol⁻¹, respectively). In contrast, biphenyl flanked cyclooctyne, DBCO (19.6 kcal mol⁻¹), showed significantly increased distortion energy for **PyrS2** compared with non-fused ring cyclooctynes. The relatively larger distortion energy of **PyrS2** was possibly resulting from the large steric effect from DBCO. This was further confirmed by the distortion energy for dienophiles. Another biphenyl flanked dienophile, Si7HT showed much smaller distortion energy compared with DBCO owing to the lower steric interactions after ring size contraction. The non-phenyl flanked cyclooctyne dienophiles had even lower distortion energy, providing more support for the vital role of steric effects in dienophiles.



Interestingly, MeCp showed the highest distortion energy (10.5 kcal mol⁻¹) with **PyrS2**, implying that the interaction between alkynes and **PyrS2** were much more favorable than that with alkenes. Additionally, the interaction energy was relevant to steric effects with more favorable interactions with cyclooctynes (*exo*-BCN, *endo*-BCN, and DMBCN) with non-fused rings compared to fused ring counterparts (DBCO and Si7HT). The interactions between alkynes were also favorable compared that with alkenes.

Conclusions

In this computational study, we have studied the inverse electron-demand Diels–Alder reactions of BCN towards 2*H*-pyran-2-one, 2*H*-thiopyran-2-one, 2*H*-pyran-2-thione and 2*H*-thiopyran-2-thione. DFT calculations have been carried out to rationalize why 2*H*-pyran-2-thiones have higher reactivity compared to the other counterparts. The enhanced reactivity of 2*H*-pyran-2-thiones (**4**) with respect to the other set of derivatives results from a lower distortion energy. The reaction of 2*H*-pyran-2-thiones was further extended different strained derivatives and DFT calculations showed a good agreement with experimentally determined second-order rate constants. We envision that this mechanistic study will help in the understanding and development of more reactive 2*H*-pyran-2-thiones and related analogues in the future.

Experimental

Reactivity tests and second order rate constant determination

Reactivity tests between **PyrS2** and DBCO were carried out at 25 °C in a 50% mixture of CD₃CN/D₂O, which was monitored by ¹H nuclear magnetic resonance (NMR) spectroscopy using 1,3,5-trimethoxybenzene as the internal standard. Reactions between **PyrS2** and MeCp were monitored using HPLC and the cycloadducts were analysed by LC-MS. Second order rate constants were determined by mixing **PyrS2** with dienophiles (*exo*-BCN, DMBCN, MeCp) in a 1 : 10, 1 : 15, or 1 : 20 ratio for pseudo 1st order kinetics and monitoring the characteristic absorption decay of **PyrS2** at 412 nm using a plate reader. Uncaging tests with the Si7HT precursor **Cp-7HT** and **PyrS2** were carried out at room temperature in CH₃CN with light exposure using a mounted LED (Thorlabs, 365 nm, input power 3 W) and reaction monitoring using HPLC assay. For kinetics analysis, reaction mixtures containing the Si7HT precursor **Cp-7HT** (2 to 3 mM) in CH₃CN were uncaged for 1 min in a quartz test tube and then mixed with **PyrS2** (0.2 mM) to make final concentrations of 1 to 1.5 mM for **Cp-7HT** and 0.1 mM for **PyrS2**, respectively. After that, the reaction mixtures were aliquoted into the plate reader for absorption monitoring at 412 nm over time. All the cycloadducts were isolated from reaction mixtures after kinetics determination and characterized by ¹H NMR, cosy, and HRMS for reaction validation.

Computational details

All calculations were performed using Gaussian 16.²⁵ Geometry optimizations of the minima and transition states were carried out using the M06-2X functional²⁶ with the 6-311+G(d,p) basis set and SMD model²⁷ for mimicking the solvent (50% CH₃CN/H₂O mixture, $\chi = 55.69$ at 25 °C).²⁸ This functional has proved to give accurate energetics for cycloaddition reactions according to previously reported studies.²⁹ Vibrational frequencies were computed at the same level to verify that optimized structures are energy minima or transition structures and to evaluate zero-point vibrational energies (ZPVE) and thermal corrections at 298 K. Intrinsic reaction coordinate (IRC) calculations were performed to verify the expected connections of the first-order saddle points with the local minima found on the potential energy surface.³⁰ Finally, a comparison between M06-2X/6-311+G(d,p), M06-2X/Def2TZVPP and ω B97xD/Def2TZVPP methods was performed to validate the results and all the methods yield very similar results (see ESI† for details). Distortion/interaction analysis were done at the M06-2X/6-311+G(d,p) level. To this end, the optimized transition state structure is separated into two fragments (distorted BCN and distorted heterocyclic structure), followed by single-point energy calculations on each fragment obtained from the IRC analysis. The energy difference between the distorted fragment and corresponding optimized structure is defined as the distortion energy. The difference between the activation energy and the total distortion energy is the interaction energy. ASD was performed using Gaussian 16 by single point calculations of separate reactants from the IRC analysis. The optimized structures indicated and shown in this study are illustrated using the CYLview program.³¹ The presence of noncovalent interactions was evaluated using a reduced density gradient (RDG) analysis³² performed with Multiwfn³³ and visualized by visual molecular dynamics (VMD).³⁴ Orbital overlap was calculated with Multiwfn.³⁵

Author contributions

JE and STL conceived and planned the research. WH and KW performed the kinetic study and analysed the data, and then discussed the results with STL. JE performed DFT calculation and analysed the data. JE and STL wrote the manuscript. All authors contributed to the manuscript and have approved the final version of the manuscript.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.



Acknowledgements

The computational resources from the Servei d'Informàtica de la Universitat de València (SIUV) are gratefully acknowledged for providing access to supercomputing resources.

References

- H. C. Kolb, M. G. Finn and K. B. Sharpless, *Angew. Chem., Int. Ed.*, 2001, **40**, 2004–2021.
- (a) L. M. Gaetke and C. K. Chow, *Toxicology*, 2003, **189**, 147–163; (b) M. C. Linder, *Mutat. Res.*, 2012, **733**, 83–91.
- (a) C. W. Tornøe, C. Christensen and M. Meldal, *J. Org. Chem.*, 2002, **67**, 3057–3064; (b) V. V. Rostovtsev, L. G. Green, V. V. Fokin and K. B. Sharpless, *Angew. Chem., Int. Ed.*, 2002, **41**, 2596–2599; (c) N. J. Agard, J. A. Prescher and C. R. Bertozzi, *J. Am. Chem. Soc.*, 2004, **126**, 15046–15047.
- (a) J. M. Baskin, J. A. Prescher, S. T. Laughlin, N. J. Agard, P. V. Chang, I. A. Miller, A. Lo, J. A. Codelli and C. R. Bertozzi, *Proc. Natl. Acad. Sci. U. S. A.*, 2007, **100**, 9116–9121; (b) S. T. Laughlin, J. M. Baskin, S. L. Amacher and C. R. Bertozzi, *Science*, 2008, **320**, 664–667; J. Dommerholt, F. P. J. T. Rutjes and F. L. van Delft, *Top. Curr. Chem.*, 2016, **374**, 16.
- M. L. Blackman, M. Royzen and J. M. Fox, *J. Am. Chem. Soc.*, 2008, **130**, 13518–13519.
- (a) A. Borrmann, O. Fatunsin, J. Dommerholt, A. M. Jonker, D. W. P. M. Löwik, J. C. M. van Hest and F. L. van Delft, *Bioconjugate Chem.*, 2015, **26**, 257–261; (b) R. Sen, J. Escorihuela, F. L. van Delft and H. Zuilhof, *Angew. Chem., Int. Ed.*, 2017, **56**, 3299–3303; (c) D. Gahtory, R. Sen, A. R. Kuzmyn, J. Escorihuela and H. Zuilhof, *Angew. Chem.*, 2018, **130**, 10118–10122; (d) B. Albada, J. F. Keijzer, H. Zuilhof and F. L. van Delft, *Chem. Rev.*, 2021, **121**, 7032–7058.
- (a) M. F. Debets, S. S. van Berkel, J. Dommerholt, A. J. Dirks, F. P. J. T. Rutjes and F. L. van Delft, *Acc. Chem. Res.*, 2011, **44**, 805–815; (b) J. J. Bruins, D. Blanco-Ania, V. van der Doef, F. L. van Delft and B. Albada, *Chem. Commun.*, 2018, **54**, 7338–7341.
- J. Escorihuela, A. T. M. Marcelis and H. Zuilhof, *Adv. Mater. Interfaces*, 2015, **2**, 1500135.
- K. Li, D. Fong, E. Meichsner and A. Adronov, *Chem. – Eur. J.*, 2021, **27**, 5057–5073.
- (a) F. Schoenebeck, D. H. Ess, G. O. Jones and K. N. Houk, *J. Am. Chem. Soc.*, 2009, **131**, 8121–8133; (b) Y. Liang, J. L. Mackey, S. A. Lopez, F. Liu and K. N. Houk, *J. Am. Chem. Soc.*, 2012, **134**, 17904–17907; (c) J. Dommerholt, O. van Rooijen, A. Borrmann, C. Fonseca Guerra, F. M. Bickelhaupt and F. L. van Delft, *Nat. Commun.*, 2014, **5**, 5378.
- (a) Y.-F. Yang, Y. Liang, F. Liu and K. N. Houk, *J. Am. Chem. Soc.*, 2016, **138**, 1660–1667; (b) B. J. Levandowski, D. Svatunek, B. Sohr, H. Mikula and K. N. Houk, *J. Am. Chem. Soc.*, 2019, **141**, 2224–2227; (c) P. García-Aznar and J. Escorihuela, *Org. Biomol. Chem.*, 2022, **20**, 6400–6412.
- (a) J. Escorihuela, A. Das, W. J. E. Looijen, F. L. van Delft, A. J. A. Aquino, H. Lischka and H. Zuilhof, *J. Org. Chem.*, 2018, **83**, 244–252; (b) J. Escorihuela, W. J. E. Looijen, X. Wang, A. J. A. Aquino, H. Lischka and H. Zuilhof, *J. Org. Chem.*, 2020, **85**, 13557–13566; (c) J. A. M. Damen, J. Escorihuela, H. Zuilhof, F. L. van Delft and B. Albada, *Chem. – Eur. J.*, 2023, **29**, e202300231.
- (a) H. Wu and N. K. Devaraj, *Acc. Chem. Res.*, 2018, **51**, 1249–1259; (b) S. Mayer and K. Lang, *Synthesis*, 2017, 830–848; (c) P. Kumar, O. Zainul, F. M. Camarda, T. Jiang, J. A. Mannone, W. Huang and S. T. Laughlin, *Org. Lett.*, 2019, **21**, 3721–3725; (d) A. Jemas, Y. Xie, J. E. Pigga, J. L. Caplan, C. W. am Ende and J. M. Fox, *J. Am. Chem. Soc.*, 2022, **144**, 1647–1662; (e) L. Liu, D. Zhang, M. Johnson and N. K. Devaraj, *Nat. Chem.*, 2022, **14**, 1078–1085.
- K. Kranjc and M. Kočevar, *New J. Chem.*, 2005, **29**, 1027–1034.
- Q. Cui, T. W. Pan, M. Shieh, S. S. Kelly, S. Xu, W.-J. Qian and M. Xian, *Org. Lett.*, 2022, **24**, 7334–7338.
- W. Huang, N. Gunawardhana, Y. Zhang, J. Escorihuela and S. Laughlin, *Chem. – Eur. J.*, 2024, **30**, e202303465.
- (a) R. Jasiński, M. Kwiatkowska, V. Sharnin and A. Barański, *Monatsh. Chem.*, 2013, **144**, 327–335; (b) E. Dresler, A. Wróblewska and R. Jasiński, *Molecules*, 2023, **28**, 5289; (c) D. Loco, I. Chataigner, J. Piquemal and R. Spezia, *ChemPhysChem*, 2022, **23**, e202200349.
- (a) J. Cioslowski, J. Sauer, J. Hetzenegger, T. Karcher and T. Hierstetter, *J. Am. Chem. Soc.*, 1993, **115**, 1353–1359; (b) D. Svatunek, M. Wilkovitsch, L. Hartmann, K. N. Houk and H. Mikula, *J. Am. Chem. Soc.*, 2022, **144**, 8171–8177.
- (a) F. M. Bickelhaupt and K. N. Houk, *Angew. Chem., Int. Ed.*, 2017, **56**, 10070–10086; (b) F. Liu, Y. Liang and K. N. Houk, *Acc. Chem. Res.*, 2017, **50**, 2297–2308.
- (a) A. M. Sarotti, *Org. Biomol. Chem.*, 2014, **12**, 187–199; (b) W.-J. van Zeist and F. M. Bickelhaupt, *Org. Biomol. Chem.*, 2010, **8**, 3118–3127; (c) P. Vermeeren, T. Hansen, P. Janssen, M. Swart, T. A. Hamlin and F. M. Bickelhaupt, *Chem. – Eur. J.*, 2020, **26**, 15538–15548; (d) N. Grimblat and A. M. Sarotti, *Org. Biomol. Chem.*, 2020, **18**, 1104; (e) S. Portela and I. Fernández, *Eur. J. Org. Chem.*, 2021, 6102–6110; (f) T. Hansen, P. Vermeeren, L. de Jong, F. M. Bickelhaupt and T. A. Hamlin, *J. Org. Chem.*, 2022, **87**, 8892–8901; (g) T. Hansen, J. C. Roozee, F. M. Bickelhaupt and T. A. Hamlin, *J. Org. Chem.*, 2022, **87**, 1805–1813; (h) J. Escorihuela and L. M. Wolf, *Organometallics*, 2022, **41**, 2525–2534; (i) J. Escorihuela, *Synthesis*, 2023, 1139–1149.
- J. Dommerholt, S. Schmidt, R. Temming, L. J. A. Hendriks, F. P. J. T. Rutjes, J. C. M. van Hest, D. J. Lefeber, P. Friedl and F. L. van Delft, *Angew. Chem., Int. Ed.*, 2010, **49**, 9422–9425.
- (a) M. F. Debets, S. S. van Berkel, S. Schoffelen, F. P. J. T. Rutjes, J. C. M. van Hest and F. L. van Delft, *Chem. Commun.*, 2010, **46**, 97–99; (b) A. Kuzmin,



- A. Poloukhine, M. A. Wolfert and V. V. Popik, *Bioconjugate Chem.*, 2010, **21**, 2076–2085.
- 23 M. Martínek, L. Filipová, J. Galeta, L. Ludvíková and P. Klán, *Org. Lett.*, 2016, **18**, 4892–4895.
- 24 T. A. Hamlin, D. Svatunek, S. Yu, L. Ridder, I. Infante, L. Visscher and F. M. Bickelhaupt, *Eur. J. Org. Chem.*, 2019, 378–386.
- 25 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16, Revision B.01*, Gaussian, Inc., Wallingford CT, 2016.
- 26 Y. Zhao and D. G. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215–241.
- 27 A. V. Marenich, C. J. Cramer and D. G. Truhlar, *J. Phys. Chem. B*, 2009, **113**, 6378–6396.
- 28 L. G. Gagliardi, C. B. Castells, C. Ràfols, M. Rosés and E. Bosch, *J. Chem. Eng. Data*, 2007, **52**, 1103–1107.
- 29 (a) R. S. Paton, J. L. Mackey, W. H. Kim, J. H. Lee, S. J. Danishefsky and K. N. Houk, *J. Am. Chem. Soc.*, 2010, **132**, 9335–9340; (b) F. Liu, Y. Liang and K. N. Houk, *J. Am. Chem. Soc.*, 2014, **136**, 11483–11493.
- 30 (a) C. Gonzalez and H. B. Schlegel, *J. Phys. Chem.*, 1990, **94**, 5523–5527; (b) K. Fukui, *Acc. Chem. Res.*, 1981, **14**, 363–368.
- 31 Legault, C. Y. *CYLVIEW, 1.0b*, Université de Sherbrooke, Canada, 2009, <https://www.cylvview.org>.
- 32 E. R. Johnson, S. Keinan, P. Mori-Sánchez, J. Contreras-García, A. J. Cohen and W. Yang, *J. Am. Chem. Soc.*, 2010, **132**, 6498–6506.
- 33 T. Lu and F. Chen, *J. Comput. Chem.*, 2012, **33**, 580–592.
- 34 W. Humphrey, A. Dalke and K. Schulten, *VMD: Visual molecular dynamics*, *J. Mol. Graphics*, 1996, **14**, 33–38.
- 35 T. Lu and Q. Chen, *J. Phys. Chem. A*, 2023, **127**, 7023–7035.

