Chiral photochemistry of achiral molecules

Umberto Raucci¹,²,³, Hayley Weir¹,², Christoph Bannwarth¹,²,⁴, David M. Sanchez⁵,¹,²,⁵ & Todd J. Martínez¹,²✉

Chirality is a molecular property governed by the topography of the potential energy surface (PES). Thermally achiral molecules interconvert rapidly when the interconversion barrier between the two enantiomers is comparable to or lower than the thermal energy, in contrast to thermally stable chiral configurations. In principle, a change in the PES topography on the excited electronic state may diminish interconversion, leading to electronically prochiral molecules that can be converted from achiral to chiral by electronic excitation. Here we report that this is the case for two prototypical examples – cis-stilbene and cis-stiff stilbene. Both systems exhibit unidirectional photoisomerization for each enantiomer as a result of their electronic prochirality. We simulate an experiment to demonstrate this effect in cis-stilbene based on its interaction with circularly polarized light. Our results highlight the drastic change in chiral behavior upon electronic excitation, opening up the possibility for asymmetric photochemistry from an effectively nonchiral starting point.

¹Department of Chemistry and The PULSE Institute, Stanford University, Stanford, CA 94305, USA. ²SLAC National Accelerator Laboratory, 2575 Sand Hill Road, Menlo Park, CA 94025, USA. ³Present address: Italian Institute of Technology, Genova, GE, Italy. ⁴Present address: Institute of Physical Chemistry, RWTH Aachen University, Aachen, Germany. ⁵Present address: Design Physics Division, Lawrence Livermore National Laboratory, Livermore, CA, USA. ✉email: Todd.Martinez@stanford.edu
Chirality is a symmetry property widespread in nature, and examples include the molecules of life (e.g., proteins, nucleic acids). A molecule is chiral when it cannot be superimposed on its mirror image (enantiomer), and chirality can occur in asymmetric or dissymmetric systems (i.e., molecules that do not have any symmetry elements or do not have rotation-reflection axes). The effective (i.e., observable) chirality of a given molecule is governed by the topography of its potential energy surface (PES), specifically the height of the barrier separating the minima of the two enantiomers (the PES is symmetric with respect to an interconversion coordinate, Fig. 1). If the interconversion barrier is much higher than the thermal energy ($k_B T$), the two enantiomers can be isolated and easily characterized (i.e., thermally stable chiral molecules, Fig. 1a). Otherwise, rapid interconversion between the two enantiomeric configurations takes place making them inseparable (i.e., thermally achiral molecules, Fig. 1a).

By tuning the PES topography it is possible to influence and control the inherent chiral behavior of the molecule. One such way is by promoting the molecule to an electronically excited state where the new electronic configuration can induce an increase/decrease of the interconversion barrier resulting in a shift of the enantiomeric ratio. Furthermore, the electronic excitation can introduce other accessible photoreactions (e.g., photoisomerization) which quench enantiomeric interconversion (Fig. 1b). In this case, the molecule can be considered electronically prochiral: it can be converted from achiral to chiral by electronic excitation. Electronic prochirality is a special case of the non-equilibration of excited state rotamers (NEER) principle where the two rotamers are effective enantiomers$^{1-4}$. Applied mostly to photocyclization reactions (e.g., precalciferol, triene), the NEER hypothesis states that ring closure through a conical intersection occurs more rapidly than conformational changes due to an enhancement of the rotational barrier in the excited state.

Herein, we examine the concept of electronic prochirality via the relationship between enantiomer interconversion and cis-trans photoisomerization around a carbon–carbon double bond. In recent decades, this has been exploited to generate a local asymmetry in the PES resulting in unidirectional rotation$^{5-13}$. Indeed, the preferential selectivity of a particular direction of motion is the result of a nonequilibrium process in which the local topography of the PES is either sloped or characterized by a substantially lower energy barrier in that direction when compared to other possible pathways. The presence of chiral elements represents the simplest way to break the symmetry of the PES on the atomistic scale, ensuring that one rotational direction, e.g., clockwise (CW) or anticlockwise (ACW), is energetically preferred over the other.

Inspired by this principle, several generations of artificial light-driven molecular motors have been reported$^{15-13}$. In these synthetic nanomachines, the introduction of asymmetric elements (e.g. stereogenic center or chirality axis) results in unidirectional rotation (CW or ACW) around a carbon–carbon double bond. Feringa and co-workers achieved full 360° unidirectional rotation by designing sterically overcrowded alkenes$^{6,8,9}$. Here, a carbon–carbon double bond connects the stator and the rotor, and axial chirality is enforced by steric hindrance between the two halves. Bulky substituents in strategic positions around the rotational axis induce large twisting of the double bond, directing the olefinic bond rotation during the light-induced isomerization$^{6,9}$. Marchand and co-workers reported that without axial chirality a single stereocenter in the aliphatic position of the isomerizing double bond is enough to ensure significant directionality in the photoisomerization$^{10}$. Recently, Wang et al. exploited the axial chirality of cyclohexenylidene group to achieve unidirectional motion in a protonated Schiff-base$^{11}$.

Here, we extend these findings to simple, highly symmetric molecules that are thermally achiral in the electronic ground state, where rapid interconversion between the two enantiomeric configurations takes place. Our simulations suggest that for electronically prochiral molecules, chirality can be enhanced or “stiffened” when the molecule is promoted to the excited state despite thermal achirality in the ground state. We demonstrate this effect on the cis-trans photoisomerization of 1,2-diphenylethylene (stilbene) and 1–1’-bis-indanylidene (stiff-stilbene).

Results
Stilbene and stiff-stilbene represent prototypical examples of carbon–carbon double bond photoisomerization (Fig. 2). The steric hindrance between the phenyl groups prevents cis-stilbene and cis-stiff-stilbene from being completely planar in the ground electronic state ($S_0$), resulting in a reduction in symmetry from $C_{2v}$ to $C_s$, with the $C_2$ rotational axis perpendicular to the ethylene bond. Two helically twisted enantiomers arise for both cis-stilbene and cis-stiff-stilbene showing right-hand and left-hand helicity ($P$ and $M$, respectively, Fig. 2).

Although the photoisomerization of stilbene and stiff-stilbene has been the topic of numerous theoretical and experimental studies over the past century$^{14-19}$, to our knowledge the cis-trans photoisomerization dynamics have never been explored from the

Fig. 1 Schematic potential energy surfaces (PESs) for chiral and achiral molecules. a A barrier separates two minima corresponding to the two enantiomers ($E_1$ and $E_2$): if the interconversion barrier is much higher than the thermal energy ($k_B T$), the molecule is chiral, otherwise rapid interconversion between the two enantiomeric configurations takes place making them inseparable (thermally achiral molecules). b PESs for electronically prochiral molecules which can be converted from achiral to chiral by electronic excitation. The example here shows a case where the molecule is thermally achiral on the ground state, but a similar scenario is also possible where the molecule is truly achiral on the ground state.
perspective of its rotational directionality. Here, we explore the rotational directionality in the non-adiabatic dynamics of the P and M enantiomers of cis-stilbene and cis-stiff-stilbene in the gas phase using a graphical processing unit (GPU)-accelerated multiconfigurational electronic structure theory in TeraChem coupled to ab initio multiple spawning (AIMS) coupled to ab initio multiple spawning (AIMS) trajectories for both P and M enantiomers that are initiated from thirty initial conditions (positions and momenta) sampled from a 0 K harmonic Wigner distribution around their respective ground state minimum. Figure 3 shows the time evolution of the central ethylenic twist angle ($\tau$ in Fig. 2) around the photoisomerizing carbon–carbon bond for the two enantiomers (P and M) of cis-stilbene and cis-stiff-stilbene on the first singlet excited state PES, S1. We observe that the photoisomerization dynamics for each isomer are unidirectional: the evolution of the P enantiomer shows an increasing value of the $\tau$ angle until the P $S_1/S_0$ conical intersection (CI) region is reached at 90° (CW rotation), whereas the excitation of the M isomer leads to ACW rotation of $\tau$ to $-90^\circ$ at the $M S_1/S_0$ CI.

In addition to the photoisomerization around $\tau$, cis-stilbene may also relax back to the ground state via photocyclization leading to 4a,4b-dihydrophenanthrene (DHP). The cis-trans isomerization and ring closure to DHP are orthogonal pathways accessed through distinct conical intersections. As a consequence of the unidirectional dynamics, an enantioselective photocyclization is observed. Indeed, the CW rotation in (P)-cis-stilbene leads to (4aS,4bS)-4a,4b-dihydrophenanthrene (S,S-DHP), while (4aR,4bR)-4a,4b-dihydrophenanthrene (R,R-DHP) is formed from the ACW motion in (M)-cis-stilbene.

No P-M helical inversion is observed on $S_1$ for both cis-stilbene and cis-stiff-stilbene indicating the occurrence of excited state axial chirality that leads to unidirectional photoisomerization. Hence, in spite of being formally achiral at thermal equilibrium on $S_0$ due to the fast helical inversion process (inversion barriers are 1.6 and 4.9 kcal/mol for stilbene and stiff-stilbene, respectively, and transition state structures are shown in Supplementary Fig. 1), cis-stilbene and cis-stiff-stilbene behave as chiral molecules during their short life on $S_1$. Indeed, the helical inversion barrier increases on $S_1$ to 19 and 13 kcal/mol for stilbene and stiff-stilbene, respectively. As can be inferred from Supplementary Fig. 1, the helical inversion in cis-stilbene proceeds via phenyl rotation on the ground state, whereas the transition state is planar on the excited state, reached by ethylenic torsion. This behavior can be easily rationalized by inspecting the frontier molecular orbitals involved in the electronic excitation (Supplementary Fig. 2) as routinely done in organic photochemistry. Indeed, the $S_0 \rightarrow S_1$ transition corresponds to a HOMO $\rightarrow$ LUMO $\pi-\pi^*$ one-electron excitation characterized by a transfer of electron density from the central ethylenic bond to the $\pi$ orbitals of the adjacent carbon pairs. The new electronic arrangement reached on $S_1$ lowers the barrier for the rotation around the central carbon–carbon double bond and hinders the rotation of the phenyl groups necessary for the P-M inversion. The resulting unidirectional photoisomerization is thus observed as a result of a nonequilibrium process in the excited state, in which the local topography of the PES is sloped in the direction of the cis-trans isomerization compared to the helical inversion pathway (Fig. 1 and Supplementary Fig. 3).

Nevertheless, in order to achieve photo-induced net unidirectional motion, the preferential excitation of a specific enantiomer is required. Photoexcitation with non-polarized or linearly polarized light is unsuitable for this purpose because it would lead to a racemate in the excited state. In contrast, the differential absorption of circularly polarized light (CPL) results in bands with opposite signs in the electronic circular dichroism (ECD) spectrum for both enantiomers, meaning that an excess of one enantiomer is excited over the other. In Supplementary Fig. 4, we present the simulated relative absorption spectra for right circularly polarized light (r-CPL) of the (M)- and (P)-conformers of cis-stilbene. We use the quantum mechanically determined optical anisotropy Kuhn factor, $g_L$ (ratio of the dipole strength and the rotatory strength) for the $S_0 \rightarrow S_1$ excitation to evaluate the enanitomergetic excess of excited chiral species following r-CPL absorption (a more detailed discussion is reported in the SI). In the optical window 230–350 nm, the M-enantiomer preferentially absorbs the r-CPL, while the l-CPL is preferentially absorbed by...
Fig. 3 Time evolution of the twist angle (\( \tau \)) on the \( S_1 \) state for \( P \) (blue) and \( M \) (red) enantiomers. Cis-stilbene is shown in the upper panel (a) and cis-stiff-stilbene is in the lower panel (b). The bold line represents the weighted average of the total population over all the trajectory basis function (TBFs) on \( S_1 \). Each line represents a TBF on \( S_1 \), where the thickness of each line is proportional to its contribution to the weighted average.

the \( P \) conformer. As a consequence, the excitation of the ground state sample of cis-stilbene with \( \chi \)-CPL will preferentially excite the left-handed helical twisted \( M \)-enantiomer, whereas the \( P \) enantiomer can be preferentially excited with \( \lambda \)-CPL.

To supplement our findings for the unidirectional photoisomerization of cis-stilbene, we simulated a rather straightforward experiment based on the enantioselective photocyclization leading to \( S,S \)-DHP and \( R,R \)-DHP. Indeed, the excitation of cis-stilbene (existing as a racemic mixture of the \( P \) and \( M \) enantiomers at thermal equilibrium) with non-polarized UV light will lead to a racemic mixture of the chiral DHP photoproduct, whereas an excess of one enantiomer over the other is expected with the CPL excitation. Due to the fact that DHP can absorb CPL in the same optical window as \( S,S \)-stilbene, the enantiomeric excess (ee) of DHP reached at the photo-stationary state can be predicted considering the kinetic model reported in Fig. 4a. In agreement with the results of the AIMS simulations, we assume that the cis-trans isomerization and the DHP cyclization reaction, and the ground state helical inversion process, respectively. The above expression demonstrates how the ee depends on the asymmetry of the electronic excitations (via the \( g \) factors), the cis-trans photoisomerization, and the ground state helical inversion processes.

For a given set of \( g \) factors, three limiting cases arise: the cis-trans photoisomerization is faster than the ground state helical inversion (\( k_{cis\rightarrow trans} \gg k_{rac} \)), \( S_0 \) racemization is faster than the cis-trans isomerization (\( k_{rac} \gg k_{cis\rightarrow trans} \)) or these processes take place on the comparable time scale (\( k_{cis\rightarrow trans} \sim k_{rac} \)). The wavelength-dependent enantiomeric excess of DHP is reported for the three cases in Fig. 4b. According to our simulation, the maximum enantiomeric excess reached around 250 nm is \( \approx 0.23\% \) when the photoisomerization is faster than the helical inversion process (which is the case for cis-stilbene due to its sub-picosecond cis-trans photoisomerization), whereas it is \( \approx 0.25\% \) in the other two limit cases. The predicted ee is comparable with what Butchardt and co-workers reported for the asymmetric synthesis of chiral helicenes with CPL.

\[
\begin{align*}
\text{ee} & = \frac{[S,SDHP] - [R,RDHP]}{[S,SDHP] + [R,RDHP]} \\
& = \frac{k_{cis\rightarrow trans}(g_{DHP} - g_{\text{std}}) + 2k_{rac}(g_{DHP} - g_{\text{std}})}{2k_{trans}(1 - g_{\text{std}}) + 2k_{rac}(1 - g_{DHP})}
\end{align*}
\]

where \( g_{\text{std}} \) and \( g_{DHP} \) are the \( g \) factors of cis-stilbene and DHP, respectively (Supplementary Fig. 4), while \( k_{cis\rightarrow trans} \) and \( k_{rac} \) represent the kinetic constants of the cis-trans photoisomerization reaction, and the ground state helical inversion process, respectively.
Our kinetic model shows that the expected enantiomeric excess is strongly dependent on the \( g \) factors. Considering that the anisotropy factors are typically quite small (less than 0.01) a low \( ee \) should be expected\(^3\). Future improvements may include the excitation to states with larger \( g \) factors as well as shifting of the photo-stationary equilibrium, e.g., by selective removal of one of the enantiomeric photoproducts. Indeed, the optimal condition to maximize the \( ee \) is reached when one enantiomer is preferentially formed and the other destroyed by the same monochromatic CPL irradiation (corresponding to \( g \) factors with opposite sign, see Supplementary Fig. 6 for a generic \( A \rightarrow B \) photoreaction). Since the anisotropy factors are wavelength-dependent, the aim is to find the optical window that maximizes the excitation asymmetry.

In conclusion, our non-adiabatic simulations of stilbene and stiff-stilbene suggest that electronic excitation can change the...
inherent chiral behavior of molecules. We coin the term electronically prochiral to describe molecules whose enantiomer interconversion can be hindered in the excited electronic state, opening up the possibility for asymmetric photochemistry from an effectively nonchiral starting point. Exploiting such asymmetric photochemistry and its connection with unidirectional motion will represent the next step toward designing new generations of responsive smart materials.

Methods

Computational details. The excited state non-adiabatic dynamics of cis-stilbene and cis-stiff-stilbene were investigated with ab initio multiple spawning22–23 interfaced with multi-configurational electronic structure methods. GFP accelerated State-Averaged Complete Active Space Self-Consistent Field (SA-2-CASSCF(2,2)/6-311 G*) was employed to model the photochemistry of cis-stilbene. This level of theory has been shown to provide a reliable description of cis-stilbene photoisomerization17,19, agreeing well with experimentally determined lifetimes and branching ratios. For example, the computed branching ratio of 44:52:4 for the three photoproducts (cis-stilbene, trans-stilbene, and DHP)11–19 is in line with recent experiments using transient absorption spectroscopy with 318 nm excitation wavelength, which yielded 5% DHP.

Modelling the excited state landscape of cis-stiff-stilbene has been shown to be a challenging task19. Here, we employ the a-flatting Floating Occupation Molecular Orbital Complete Active Space Configuration Interaction (a-FOMO-CASCI) electronic structure method 9,44, which provides a balanced description of the twisting barriers on both the cis and trans side. The main idea behind a-FOMO-CASCI is to recover effects on the electronic state splitting arising from dynamic electron correlation that is mostly absent in the FOMO-CASCI wavefunction through an a scaling of the state-specific energy splitting while leaving the state-averaged energy untouched in the same way as a-CASSCF44. Adopting (0.8)-FOMOβ = (0.2)-CAS(2,2)Cl/6-31 G* (where a is the scaling factor, and β represents the FON temperature) with two electrons into two π orbitals, the computed ratio of the cis and trans barriers shows good agreement with experimental estimates. Critical points of the ground and excited-state potential energy surfaces are reported for cis-stiff-stilbene in Supplementary Fig. 7. The TeraChem electronic structure package24–25 was employed to perform all the electronic structure calculations.

Initial conditions generation. Thirty initial conditions (ICs) for each enantiomer (P and M) of cis-stilbene and cis-stiff-stilbene were selected out of 500 geometries sampled from a 0 K harmonic Wigner distribution corresponding to geometry and frequencies computed at B3LYP/6-31 G* and B3LYP/6-31 G** for cis-stilbene and cis-stiff-stilbene, respectively. These 500 phase space points were used to simulate the electronic absorption spectra (Supplementary Fig. 8). The absorption spectra were generated by single-point energy calculations at SA-2-CASSCF(2,2)/6-311 G* and (0.8)-FOMOβ = (0.2)-CAS(2,2)Cl/6-31 G*, for cis-stilbene and cis-stiff-stilbene, respectively. The final spectra were obtained by broadening the S0 Æ S1 excitation energies with Gaussian functions (full-width half maximum of 0.2 eV). We randomly selected 30 different ICs from the 500 phase space points used to simulate the spectra. These ICs (positions and momenta) were placed on the S1 manifold.

P-M thermal barriers. Transition states (TS) for the P-M helical inversion were evaluated by the Transition State Finder in DL-FIND through the ChemShell/ TeraChem interface. B3LYP-D3/def2-TZVP(1)-CASSCF(2,2)-CASPT2 was employed. The TS involved in the helical inversion is reported in Supplementary Fig. 1.

Simulation of circular dichroism spectra. The electronic absorption and ECD spectra (Supplementary Fig. 9) were computed by means of the simplified time-dependent density functional theory (sTD-DFT) 15,44. We used 1000 structures from the aforementioned 0 K harmonic Wigner distribution. At each of these structures, we computed Kohn-Sham density functional theory single-point energies with BHLYP/def2-TZVP(1)-CASSCF(2,2)-CASPT2, Mulliken populations and orbital energies were then used to compute all vertical excitations up to 10 eV with the sTD-DFT method as implemented in the sTDA package20. All excitations were blue-shifted by 0.2 eV and convolved with Gaussians of 0.24 eV width at 1/e maximum to match the previously reported experimental absorption (in hexane) 15. The ECD spectra and the relative r-CPL absorption spectra were obtained and shifted accordingly. The spectroscopic data for stiff-stilbene was determined in the same way. A blue shift of 0.35 eV was used to match the absorption spectrum from ref. 9.

Data availability

Data generated in this study have been deposited in Zenodo (https://zenodo.org) under the accession code https://doi.org/10.5281/zenodo.6430143.

Code availability

Results described in this manuscript have been obtained with publicly available software packages such as TeraChem (http://www.terachem.com), MolPro (http://www.molpro.net), and sTDA (https://github.com/grimme-lab/stda).

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References

1. Jacobs, H. J. C. & Havinga, E. Photochemistry of Vitamin D and its Isomers and of Simple Trienes. Adv. Photochem. 11, 305–373 (1979).
2. Vroegop, P. J., Lugtenburg, J. & Havinga, E. Conformational equilibrium and photochemistry of hexa-1,3,5-trienes. Tetrahedron 29, 1393–1398 (1973).
3. Havinga, E. & Schlatmann, J. L. M. A. Remarks on the specificities of the photochemical and thermal transformations in the vitamin D field. Tetrahedron 16, 146–152 (1961).
4. Whitesell, J. K., Minton, M. A. & Tran, V. D. The non-equilibration of excited rogers (neer) principle. Ground-state conformational bias in triene photocyclizations. J. Am. Chem. Soc. 111, 1473–1476 (1989).
5. Kottas, G. S., Clarke, I. L., Horinek, D. & Michl, J. Artificial molecular rotors. Chem. Rev. 105, 1281–1317 (2005).
6. Feringa, B. L. The art of building small: from molecular switches to molecular motors. J. Org. Chem. 72, 6635–6652 (2007).
7. Baroncini, M., Silvi, S. & Credi, A. Photo- and redox-driven artificial molecular motors. Chem. Rev. 120, 200–268 (2020).
8. Koumura, N., Zijlstra, R. W. J., van Delden, R. A., Harada, N. & Feringa, B. L. Light-driven monodirectional molecular rotor. Nature 401, 152–155 (1999).
9. Roke, D., Wezenberg, S. J. & Feringa, B. L. Molecular rotary motors: unidirectional motion around double bonds. Proc. Natl Acad. Sci. USA 115, 9423 (2018).
10. Marchand, G. et al. Directionality of double-bond photoisomerization dynamics induced by a single stereogenic Center. J. Phys. Chem. Lett. 6, 599–604 (2015).
11. Wang, J., Orgunati, B. & Durbeej, B. Light-driven rotary molecular motors without point chirality: a minimal design. Phys. Chem. Chem. Phys. 19, 6952–6956 (2017).
12. Huber, L. A. et al. Direct observation of hemithioindigo-motor unidirectionality. Angew. Chem. Int. Ed. 56, 14356–14359 (2017).
13. Guentert, M. et al. Sunlight-powered kHz rotation of a hemithioindigo-based molecular motor. Nat. Commun. 6, 8406 (2015).
14. Quick, M. et al. Photoisomerization dynamics of stiff-stilbene in solution. J. Phys. Chem. B 118, 1389–1402 (2014).
15. Fu, W., Kosmidis, C., Schmid, W. E. & Trushin, S. A. The photochemical cis-trans isomerization of free stilbene molecules follows a Hula-Twist pathway. Angew. Chem. Int. Ed. 43, 4178–4182 (2016).
16. Liu, F. & Morokuma, K. Computational study on the working mechanism of a stilbene light-driven molecular rotor: slowed minimal energy path and unidirectional nonadiabatic photoisomerization. J. Am. Chem. Soc. 134, 4861–4876 (2012).
17. Quenneville, J. & Martinez, T. J. Ab initio study of cis–trans photoisomerization in stilbene and ethylene. J. Phys. Chem. A 107, 829–837 (2003).
18. Impuro, R. & Santoro, F. Excited-state behavior of trans and cis isomers of stilbene and stiff stilbene: a TD-DFT study. J. Phys. Chem. A 109, 10058–10067 (2005).
19. Weir, H., Williams, M., Parrish, R. M., Hohenstein, E. G. & Martinez, T. J. Nonadiabatic dynamics of photoexcited cis-stilbene using Ab initio multiple spawning. J. Phys. Chem. B 124, 5476–5487 (2020).
20. Uftinsev, I. S. & Martinez, T. J. Quantum chemistry on graphical processing units. 2. Direct self-consistent-field implementation. J. Chem. Theory Comput. 5, 1004–1015 (2009).
21. Uftinsev, I. S. & Martinez, T. J. Quantum chemistry on graphical processing units. 3. Analytical energy gradients, geometry optimization, and first principles molecular dynamics. J. Chem. Theory Comput. 5, 2619–2628 (2009).
22. Uftinsev, I. S. & Martinez, T. J. Quantum chemistry on graphical processing units. 1. Strategies for two-electron integral evaluation. J. Chem. Theory Comput. 4, 222–231 (2008).
23. Ben-Nun, M. & Martinez, T. J. Ab initio quantum molecular dynamics. Adv. Chem. Phys. 121, 439–512 (2002).
24. Ben-Nun, M. & Martinez, T. J. Nonadiabatic molecular dynamics: validation of the multiple spawning method for a multidimensional problem. J. Chem. Phys. 108, 7244–7257 (1998).
25. Ben-Nun, M., Quenneville, J. & Martinez, T. J. Ab initio multiple spawning: photochemistry from first principles quantum molecular dynamics. J. Phys. Chem. A 104, 5161–5175 (2000).
26. Celani, P., Ottani, S., Olivucci, M., Bernardi, F. & Robb, M. A. What happens during the picosecond lifetime of 2A1 cyclohexa-1,3-diene? A CAS-SCF study of the cyclohexadiene/hexatriene photochemical interconversion. J. Am. Chem. Soc. 116, 10141–10151 (1994).

27. Celani, P., Bernardi, F., Robb, M. A. & Olivucci, M. Do photochemical ring-opening occur in the spectroscopic state? 1B2 pathways for the cyclohexadiene/hexatriene photochemical interconversion. J. Phys. Chem. 100, 19364–19366 (1996).

28. Bearpark, M. J. et al. Cooperating rings in cis-stilbene lead to an S0/S1 conical intersection. J. Phys. Chem. A 101, 3841–3847 (1997).

29. Zimmerman, H. E. Mechanistic organic photochemistry. Acc. Chem. Res. 8, 1–11 (1969).

30. Michl, J. Physical Basis of Qualitative MO Arguments in Organic Photochemistry. Top. Curr. Chem. 46, 1–59 (1974).

31. Kuhn, W. The physical significance of optical rotatory power. Trans. Faraday Soc. 26, 293–308 (1930).

32. Rau, H. Asymmetric photochemistry in solution. Chem. Rev. 83, 535–547 (1983).

33. Bernstein, W. J., Calvin, M. & Burchardt, O. Absolute asymmetric synthesis. I. Mechanism of the photochemical synthesis of nonracemic helicenes with circularly polarized light. Wavelength dependence of the optical yield of octahelicene. J. Am. Chem. Soc. 94, 494–498 (1972).

34. Shen, Y. & Chen, C.-F. Helicenes: synthesis and applications. Chem. Rev. 112, 1463–1535 (2012).

35. Feringa, B. L. In control of motion: from molecular switches to molecular motors. Acc. Chem. Res. 34, 504–513 (2001).

36. Feringa, B. L., van Delden, R. A., Kourum, N. & Geertsema, E. M. Chiropical molecular switches. Chem. Rev. 100, 1789–1816 (2000).

37. Huck, N. P. M., Jager, W. F., de Lange, B. & Feringa, B. L. Dynamic control and amplification of molecular chirality by circular polarized light. Science 273, 1686 (1996).

38. Kovalenko, S. A., Dobryakov, A. L., Ioffe, I. & Ernsting, N. P. Evidence for the phantom state in photoinduced cis–trans isomerization of stilbene. Chem. Phys. Lett. 493, 255–258 (2010).

39. Slavíček, P. & Martínez, T. J. Ab initio floating occupation molecular orbital-complete active space configuration interaction: an efficient approximation to CASSCF. J. Chem. Phys. 132, 234102 (2010).

40. Hollas, D., Štůlík, L., Hohenstein, E. G., Martínez, T. J. & Slavíček, P. Nonadiabatic Ab initio molecular dynamics with the floating occupation molecular orbital-complete active space configuration interaction method. J. Chem. Theory Comput. 14, 339–350 (2018).

41. Snyder, J. W., Parrish, R. M. & Martínez, T. J. α-CASSCF: An efficient, empirical correction for SA-CASSCF to closely approximate MS-CASPT2 potential energy surfaces. J. Phys. Chem. Lett. 8, 2432–2437 (2017).

42. Becke, A. D. Density-functional thermochemistry. III. The role of exact exchange. J. Chem. Phys. 98, 5648–5652 (1993).

43. Grimme, S., Antony, J., Ehrlich, S. & Krieg, H. A consistent and accurate ab initio parametrization of density functional dispersion correction (DFT-D) for the 94 elements H–Pu. J. Chem. Phys. 132, 154104 (2010).

44. Weigend, F. & Ahlrichs, R. Balanced basis sets of split valence, triple zeta potential energy surfaces. J. Phys. Chem. Lett. 3, 3297–3305 (2005).

45. Bannwarth, C. & Grimme, S. A simplified time-dependent density functional theory approach for electronic ultraviolet and circular dichroism spectra of very large molecules. Comp. Theo. Chem. 1040–1041, 45–53 (2014).

46. Grimme, S. A simplified Tamm-Dancoff density functional approach for the electronic excitation spectra of very large molecules. J. Chem. Phys. 138, 244104 (2013).

47. Becke, A. D. Density-functional exchange-energy approximation with correct asymptotic behavior. Phys. Rev. A 38, 3098–3100 (1988).

48. Lee, C., Yang, W. & Parr, R. G. Development of the Colle-Salvetti correlation-energy formula into a functional of the electron density. Phys. Rev. B 37, 785–789 (1988).

49. Becke, A. D. A new mixing of Hartree-Fock and local density-functional theories. J. Chem. Phys. 98, 1372–1377 (1993).

50. Tanigushi, M. & Lindsey, J. S. Database of absorption and fluorescence spectra of >300 common compounds for use in photochemCAD. Photochem. Photobiol. 94, 290–327 (2018).

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Competing interests
The authors declare no competing interests.

Additional information
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Correspondence and requests for materials should be addressed to Todd J. Martinez.

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