



Title	Synthesis of highly condensed phospholes by the Lewis acid-assisted dehydrogenative Mallory reaction under visible light irradiation
Author(s)	Kamiyoshi, Ikki; Kojima, Yuki; Xu, Shibo et al.
Citation	Chemical Science. 2024, 15(48), p. 20413-20420
Version Type	VoR
URL	https://hdl.handle.net/11094/101968
rights	© 2024 The Author(s). Published by the Royal Society of Chemistry
Note	

The University of Osaka Institutional Knowledge Archive : OUKA

<https://ir.library.osaka-u.ac.jp/>

The University of Osaka

Cite this: *Chem. Sci.*, 2024, 15, 20413

All publication charges for this article have been paid for by the Royal Society of Chemistry

Received 23rd August 2024
Accepted 13th November 2024

DOI: 10.1039/d4sc05657d

rsc.li/chemical-science

Synthesis of highly condensed phospholes by the Lewis acid-assisted dehydrogenative Mallory reaction under visible light irradiation†

Ikki Kamiyoshi,^a Yuki Kojima,^{ID} ^a Shibo Xu,^b Kosuke Yasui,^{ab} Yuji Nishii^{ID} ^{ab} and Koji Hirano^{ID} ^{*ab}

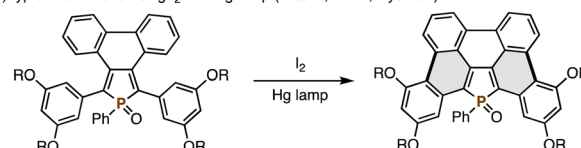
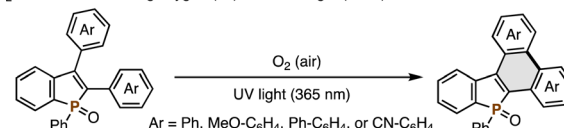
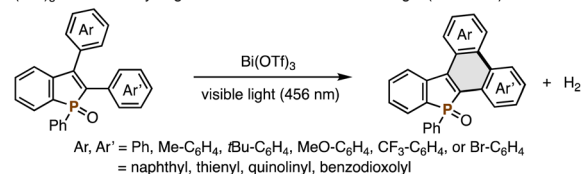
A photo-promoted oxidative cyclization, that is, the Mallory reaction of 2,3-diarylbenzophospholes has been developed. With the assistance of Bi(OTf)₃ Lewis acid, the reaction proceeds smoothly under visible light irradiation even without any external oxidants. The newly developed dehydrogenative conditions are compatible with various functional groups and substitution patterns, which enables the streamlined synthesis of highly condensed dibenzophosphole derivatives of potent interest in material chemistry. Moreover, experimental and computational studies unveil the detailed reaction mechanism. The preliminary optoelectronic properties of some newly synthesized compounds are also demonstrated.

Introduction

Highly condensed heteroaromatics are key structures in organic functional materials. In particular, phosphorus-incorporated phosphole derivatives have received significant attention because of their unique physical and optoelectronic properties in the design and synthesis of organic light-emitting diodes (OLEDs), organic photovoltaics (OPVs), and cell imaging dyes.¹ Among various reported synthetic approaches to the aforementioned important heteroaromatic core,² the photo-promoted oxidative cyclization reaction, namely, the Mallory reaction is one of the most promising candidates.³ Hissler, Réau, Nyulász, and co-workers reported successful synthesis of highly condensed dibenzophospholes by the Mallory reaction of peripherally arylated phospholes (Scheme 1a).⁴ However, a high-pressure mercury vapour lamp and I₂ external oxidant were necessary to promote the reaction. Additionally, the substituents on the phosphole nuclei are limited to strongly electron-donating dialkoxyphenyl or thienyl groups. Very recently, Zhao developed the reaction of 2,3-diarylbenzophospholes under I₂-free conditions using molecular oxygen as the oxidant, but still UV light (365 nm) was required (Scheme 1b).⁵

On the other hand, in recent years our research group serendipitously found that 3-indolylphosphole underwent

dehydrogenative cyclization even under visible light irradiation.⁶ This phenomenon was specific to a phosphole bearing the strongly electron-donating indole substituent at the C3 position, suggesting that the substrate scope can be extended by suitable electronic perturbation with the assistance of external additives. Herein, we report the significant substrate extension: the Bi(OTf)₃-assisted Mallory reaction of 2,3-diarylbenzophospholes under blue LED irradiation (456 nm) has been developed (Scheme 1c). The reaction conditions accommodate electron-donating and electron-withdrawing aryl

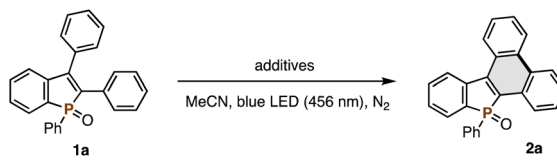
a) typical conditions using I₂ and Hg lamp (Hissler, Réau, Nyulász)b) I₂-free conditions using oxygen (air) under UV light (Zhao)c) Bi(OTf)₃-assisted dehydrogenative conditions under visible light (this work)

Scheme 1 Mallory reactions for synthesis of highly condensed phosphole derivatives: (a) typical conditions using a mercury (Hg) vapour lamp and I₂, (b) I₂-free conditions using UV light and O₂, and (c) Bi(OTf)₃-assisted dehydrogenative conditions under visible light.

^aDepartment of Applied Chemistry, Graduate School of Engineering, Osaka University, Suita, Osaka 565-0871, Japan. E-mail: k_hirano@chem.eng.osaka-u.ac.jp

^bInnovative Catalysis Science Division, Institute for Open and Transdisciplinary Research Initiatives (ICS-OTRI), Osaka University, Suita, Osaka 565-0871, Japan

† Electronic supplementary information (ESI) available. CCDC 2377855 and 2377856. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4sc05657d>

Table 1 Condition optimization for the dehydrogenative Mallory reaction of **1a** under visible light irradiation^a


Entry	Additives	Yield of 2a ^b (%)
1	Bi(OTf) ₃	33
2	Al(OTf) ₃	49
3	In(OTf) ₃	68
4	Sc(OTf) ₃	34
5	TFA	40
6	(PhO) ₂ P(O)(OH)	20
7	TfOH	0
8	PTSA	0
9	HCl	0
10	AcOH	0
11	Bi(OTf) ₃ , NaHCO ₃	(>99)
12	Bi(OTf) ₃ , Na ₂ CO ₃	<5
13	Al(OTf) ₃ , NaHCO ₃	38
14	In(OTf) ₃ , NaHCO ₃	31
15	Sc(OTf) ₃ , NaHCO ₃	8
16 ^c	Bi(OTf) ₃ , NaHCO ₃	0
17	None	0

^a Conditions: **1a** (0.050 mmol), additives (0.050 mmol), MeCN (1.0 mL), blue LED (456 nm, 40 W), ambient temperature (40–50 °C under light irradiation), 22 h, N₂. ^b Estimated by using ³¹P{¹H} NMR based on 0.050 mmol with P(O)(OEt)₃ as the internal standard. Isolated yield is in parentheses. ^c In the dark at 50 °C. Tf = trifluoromethanesulfonyl, TFA = trifluoroacetic acid, PTSA = *p*-toluenesulfonic acid, and Ac = acetyl.

groups as well as heteroaryl groups on the phosphole core. In addition, mild reaction conditions with visible light of lower energy are tolerant of a wide range of functional groups such as the Ar–Br moiety, which can be further transformed. Moreover, even in the absence of any external oxidants, the reaction proceeds smoothly with concomitant evolution of molecular hydrogen. Experimental and computational mechanistic studies and preliminary optoelectronic properties of some newly synthesized condensed phosphole derivatives are also described.

Results and discussion

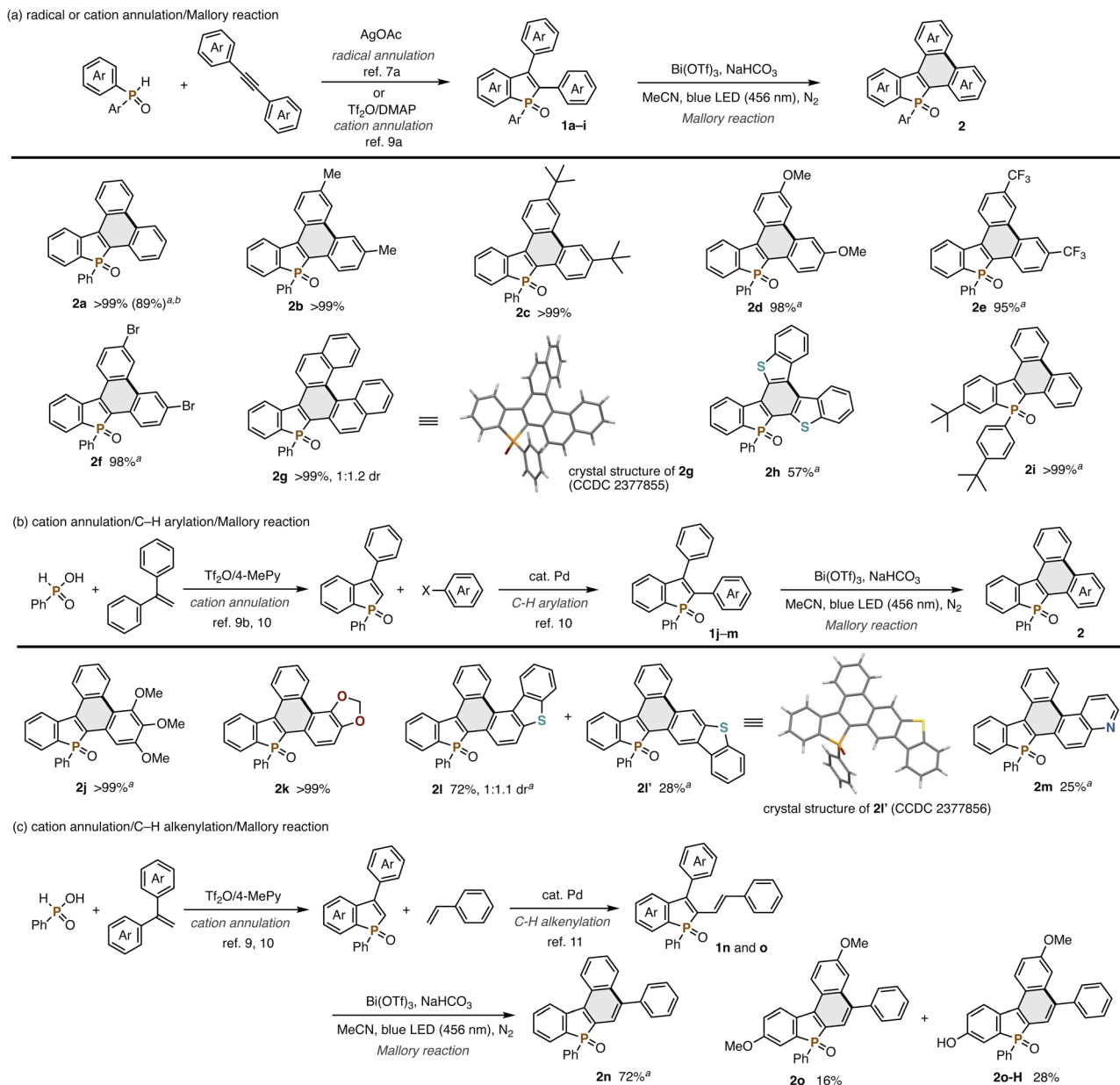
We began optimization studies with 2,3-diphenylbenzophosphole oxide **1a** (0.050 mmol) to identify suitable additives under blue LED irradiation (456 nm, 40 W, N₂ conditions, at 40–50 °C under light irradiation) in MeCN solvent (Table 1). After careful investigation, several Lewis acidic metal triflates were found to promote the reaction: Bi(OTf)₃, Al(OTf)₃, In(OTf)₃, and Sc(OTf)₃ provided the desired **2a** in 33–68% yields (entries 1–4). In addition, some Brønsted acids, TFA and (PhO)₂P(O)(OH), also showed activity (entries 5 and 6), while stronger acids including TfOH, PTSA, and HCl, gave a complicated mixture and weaker AcOH resulted in no conversion (entries 7–10). Given the

detrimental effects of TfOH (entry 7), we next tested the addition of bases to quench trace amounts of TfOH potentially generated *in situ* from metal triflates. Gratifyingly, full conversion of **1a** and isolation of **2a** in an almost quantitative yield were achieved by using a combination of Bi(OTf)₃ and NaHCO₃ (entry 11), whereas more basic additives such as Na₂CO₃ totally shut down the reaction (entry 12). In contrast, NaHCO₃ had a negative impact when combined with other metal triflates (entries 13–15). Although details still remain unclear, Lewis basic NaHCO₃ can decrease the Lewis acidity of metal triflates other than Bi(OTf)₃. Thus, the combination is crucial for satisfactory conversion of **1a**. Some control experiments were also performed: no reaction was observed in the dark even at elevated temperature (50 °C; entry 16). In the absence of additives, the starting **1a** was recovered completely (entry 17). These results clearly indicate the indispensable role of both light and Bi(OTf)₃ Lewis acid in the reaction (see the ESI† for more detailed optimization studies, including additional screening of acid additives and solvent effects).

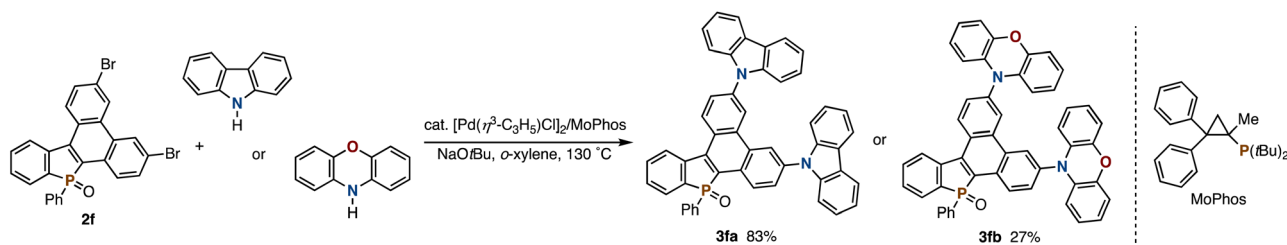
Combined with our previously developed convergent synthetic methods of 2,3-disubstituted benzophospholes, the visible-light-promoted Mallory reaction can offer a modular approach to the highly condensed dibenzophospholes with structural diversity. One is the radical annulation–Mallory reaction sequence (Scheme 2a): a variety of 2,3-diarylbenzophospholes **1a–h** readily prepared from diphenylphosphine oxide and symmetrical diaryl alkynes⁷ underwent the Mallory reaction smoothly (Scheme 2a). Electron-neutral (**2b** and **2c**), -rich (**2d**), and -deficient (**2e**) substrates were all tolerated. Owing to the visible light of relatively lower energy, the reaction conditions were compatible with the Ar–Br bond (**2f**), which is a versatile handle in downstream transformation. Actually, the Pd-catalysed double Buchwald–Hartwig amination of **2f** with carbazole and phenoxazine⁸ enabled concise synthesis of the dibenzophosphole-based donor–acceptor systems **3fa** and **3fb** in 83% and 27% yield, respectively (Scheme 3). In the case of 2-naphthyl-substituted benzophosphole **1g**, dehydrogenative cyclization occurred regioselectively to deliver the helical **2g** in an almost quantitative yield. Its structure was unambiguously confirmed by X-ray analysis (CCDC 2377855).† The heteroaromatic benzothiophene substituent could also be employed, and the corresponding P,S-doped triphenylene **2h** was obtained in an acceptable yield. Combination with the phosphorus-cation-mediated annulation of the *t*-Bu-substituted diphenyl phosphine oxide^{9a} successfully provided dibenzophosphole **2i** with the *t*-Bu-C₆H₄ substituent on the phosphorus, indicating that the diverse substitution pattern is accessible. The reaction also proceeded on a 1.0 mmol scale with synthetically useful efficacy (**2a**).

The phosphorus-dication-mediated annulation^{9b} and successive Pd-catalysed C–H arylation¹⁰ provided facile access to the benzophospholes **1j–m** with two different aromatic rings at the C2 and C3 positions, which were also amenable substrates in the Mallory reaction (Scheme 2b). The highly electron-donating trimethoxyphenyl (**2j**) and methylenedioxyphenyl (**2k**) groups at the C2 position were well tolerated under specific reaction conditions. Notably, exclusive regioselectivity was





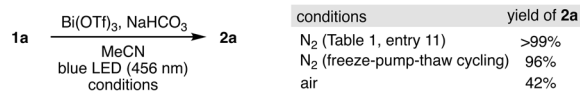
Scheme 2 Products of the Bi(OTf)₃-assisted dehydrogenative Mallory reaction of phosphole derivatives **1**. Isolated yields are shown. Conditions of the Mallory reaction: **1** (0.030–0.11 mmol), Bi(OTf)₃ (1.0 equiv.), NaHCO₃ (1.0 equiv.), MeCN (1.0 mL), blue LED (456 nm, 40 W), ambient temperature (40–50 °C under light irradiation), 22 h, N₂. ^aWith two LEDs (40 W × 2). ^bOn a 1.0 mmol scale.



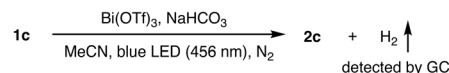
Scheme 3 Double Buchwald–Hartwig amination of dibrominated benzophosphole **2f** with carbazole and phenoxazine. See the ESI† for detailed reaction conditions.

(a) investigations of potential oxidants

- conditions screening

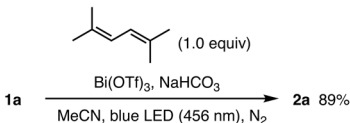


- detection of evolved H₂

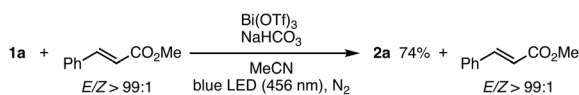


(b) investigations of excited state to be involved in cyclization process

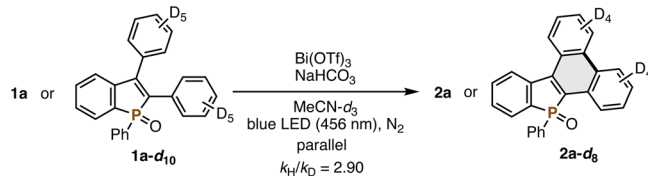
- effects of triplet quencher



- possibility of energy transfer process checked by *E* to *Z* isomerization

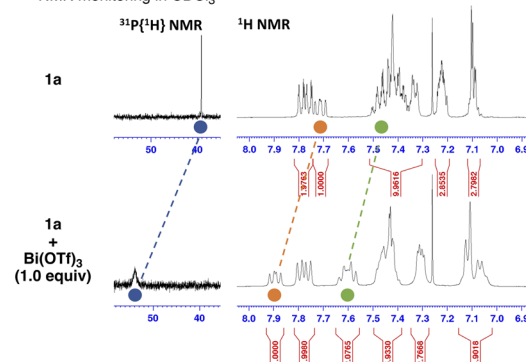


(c) deuterium-labeling experiments

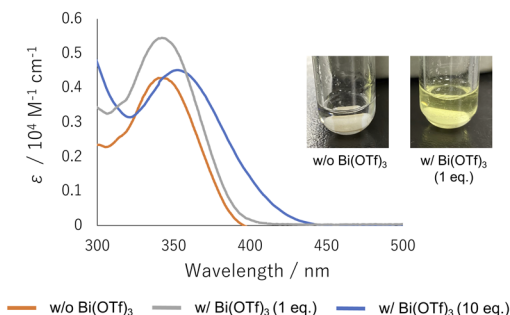


(d) NMR and UV-vis absorption spectra studies

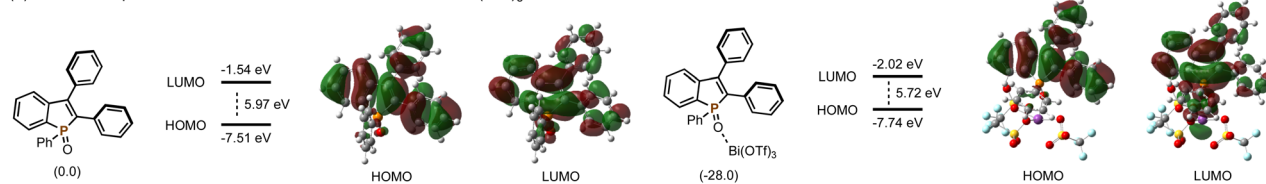
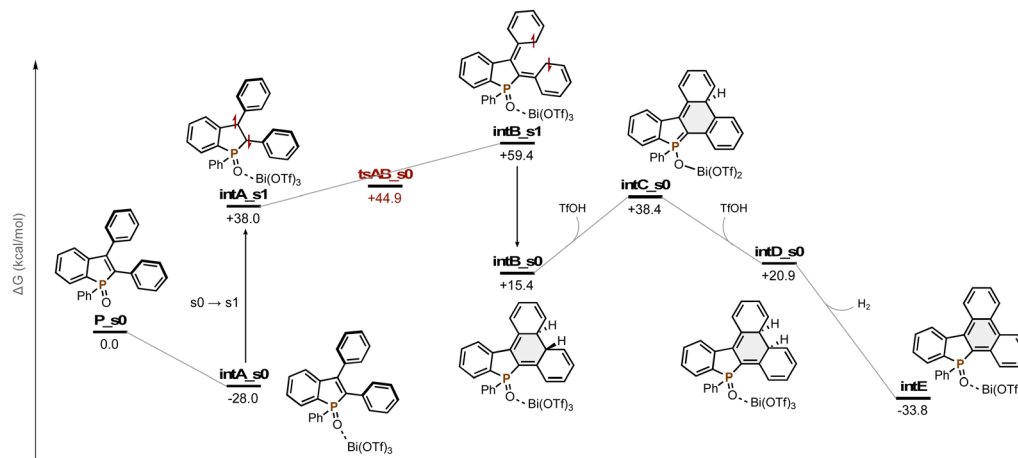
- NMR monitoring in CDCl₃



- UV-vis absorption spectra of 1a in MeCN (1.0 × 10⁻⁵ M)



Scheme 4 Experimental mechanistic studies.

(a) calculated optimal structures and HOMO/LUMO of 1a and Bi(OTf)₃-coordinated 1a(b) overall energy profile in Bi(OTf)₃-assisted dehydrogenative Mallory reaction of 1a

Scheme 5 Computational mechanistic studies. Summary of the level of theory: M06-2X/6-311+G(d,p)&SDD/PCM(MeCN)//M06-2X/6-31G(d)&LanL2DZ.



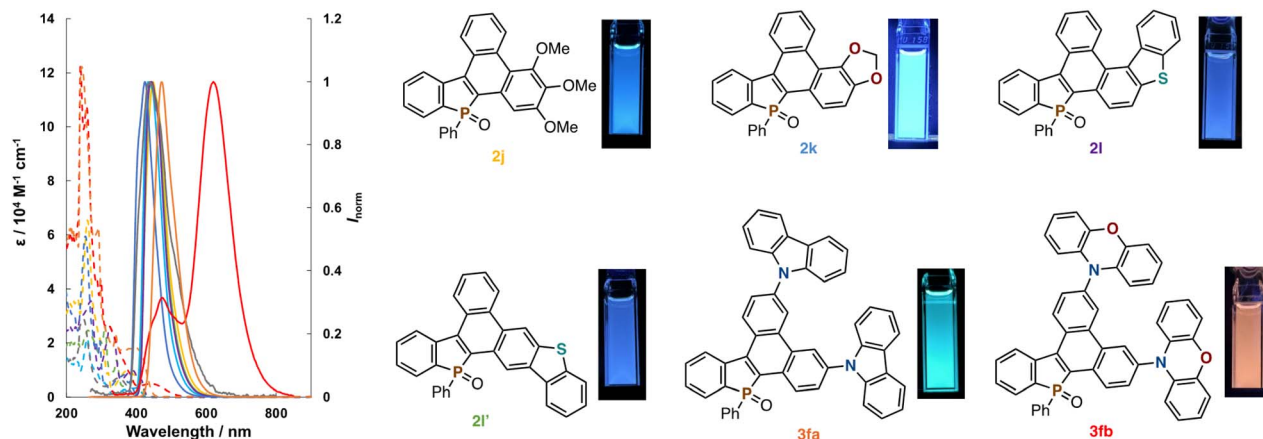


Fig. 1 UV-vis absorption (dashed lines) and fluorescence spectra (solid lines) of **1a**, **2a**, **2j**, **2k**, **2l**, **2l'**, **3fa**, and **3fb** in CHCl_3 (1.0×10^{-5} M), and fluorescence images of **2j**, **2k**, **2l**, **2l'**, **3fa**, and **3fb**.

observed in the latter case. On the other hand, the dibenzothiophene-substituted substrate **1l** afforded a regioisomeric mixture of helical **2l** and linear **2l'** (CCDC 2377856).[†] The Lewis basic quinoline-fused dibenzophosphole **2m** was also obtained albeit with a moderate yield.

The Mallory reaction was also applicable to the C2-alkenyl-C3-arylbenzophospholes **1n** and **1o**, which were also easily synthesized *via* Pd-catalysed C–H alkenylation¹¹ (Scheme 2c). In the case of **1n**, we successfully obtained the 5,7-diphenyldibenzo[*b,e*]phosphindole 7-oxide structure **2n**, which is employed as the OLED device for improvement of emission efficiency.¹² With the MeO-substituted substrate **1o**, the partial demethylation of the MeO group competitively occurred, but a good combined yield of **2o** and **2o-H** was observed.

To gain insight into the reaction mechanism, several experiments were performed next. First, we investigated potential oxidants in the present Mallory reaction (Scheme 4a). Zhao recently reported that residual oxygen in solvent is the actual external oxidant in the related UV-light-promoted Mallory reaction, which was supported by no conversion under rigorously deaerated conditions using freeze–pump–thaw cycling.⁵ Thus, our $\text{Bi}(\text{OTf})_3$ -assisted, visible-light-promoted Mallory reaction was also conducted by using the same deaeration technique. Surprisingly, the reaction proceeded smoothly without any difficulty, and **2a** was formed in 96% yield comparable with that under the standard nitrogen conditions (>99% yield, Table 1, entry 11). In addition, a lower yield was observed in air. These results suggest that oxygen in air unlikely works as the external oxidant. Finally, we successfully observed evolution of molecular hydrogen by GC after the reaction, thus indicating that the present Mallory reaction is acceptor-less, and a truly dehydrogenative reaction. To check whether the reaction is operated in the singlet or triplet excited state, we then carried out the reaction of **1a** in the presence of 2,5-dimethyl-2,4-hexadiene, which is known as a triplet quencher,¹³ but **2a** was still formed in a good yield (Scheme 4b). Even with (*E*)-methyl cinnamate, which undergoes *E* to *Z* isomerization by triplet–triplet energy transfer,¹⁴ **1a** was converted to **2a**

efficiently, and cinnamate was recovered with the maintenance of its (*E*)-configuration. Although the possibility of a triplet state intermediacy cannot be completely excluded, given the effective quenching even in the case of intramolecular cyclization,¹⁵ the aforementioned outcomes support the reaction progress *via* the singlet excited state rather than the triplet excited state. Information about the rate-determining step was obtained from the deuterium-labeling experiment with **1a-*d*₁₀** (Scheme 4c): the kinetic isotope effect (KIE) value from the parallel reaction was 2.90, thus suggesting the rate-limiting C–H cleavage. Finally, the role of $\text{Bi}(\text{OTf})_3$ was examined by using NMR and UV-vis absorption spectra (Scheme 4). Upon mixing of **1a** and $\text{Bi}(\text{OTf})_3$ in CDCl_3 , some ^1H and $^{31}\text{P}\{^1\text{H}\}$ NMR signals were shifted to the lower field. The absorption spectrum was also red-shifted, upon exposure of **1a** to $\text{Bi}(\text{OTf})_3$, to the visible light region. The colour change in the solution could also be confirmed visually. Apparently, the complexation between the phosphine oxide moiety of **1a** and $\text{Bi}(\text{OTf})_3$ enables excitation even with visible light.

Additional computational mechanistic studies were performed by using density functional theory (DFT) calculations (Scheme 5). All calculations were carried out using the Gaussian

Table 2 Optical properties of **1a**, **2a**, **2j**, **2k**, **2l**, **2l'**, **3fa**, and **3fb**^a

Compd	λ_{abs} (nm)	λ_{em} ^b (nm)	Φ_{F} ^c	$\Delta\nu$ ^d /cm ^{−1}
1a	249, 342	438	0.11	6409
2a ^e	254, 315, 364, 382	425	0.56	2649
2j	260, 379	448	0.58	4064
2k	260, 403	434	0.75	1772
2l	212, 270, 324, 392	443	0.16	2937
2l'	262, 310, 344, 391, 412	432	0.16	1124
3fa	244, 291, 341, 396	473	>0.99	4111
3fb	240, 258, 298, 317, 440	620	0.04	6598

^a Measured in 1.0×10^{-5} M solution of CHCl_3 . ^b Excited at **1a** (249 nm), **2a** (254 nm), **2j** (260 nm), **2k** (260 nm), **2l** (270 nm), **2l'** (262 nm), **3fa** (244 nm), and **3fb** (298 nm). ^c Absolute fluorescence quantum yields. ^d Stokes shifts. ^e The optical data of **2a** were taken from ref. 10.



Table 3 Absorption wavelengths, HOMO–LUMO energy gaps and differential pulse voltammetry data of compounds **2a**, **2j**, **2k**, **2l**, **2l'**, **3fa**, and **3fb**

Compd	$\lambda_{\text{onset}}^{\text{abs}}$ ^a (nm)	$E_{\text{g}}^{\text{optb}}$ (eV)	E_{ox}^{c} (V)	$E_{\text{HOMO}}^{\text{d}}$ (eV)	$E_{\text{LUMO}}^{\text{e}}$ (eV)
2a ^f	397	3.12	1.32	−6.12	−3.00
2j	422	2.94	0.91	−5.71	−2.77
2k	424	2.92	0.99	−5.79	−2.87
2l	429	2.89	1.08	−5.88	−2.99
2l'	428	2.90	1.07	−5.87	−2.97
3fa	445	2.79	0.72	−5.52	−2.73
3fb	521	2.38	0.32	−5.12	−2.74

^a Measured in CHCl_3 . ^b Determined from the onset of the absorption spectra. ^c Performed in MeCN in the presence of Bu_4NPF_6 . $\nu = 0.10 \text{ V s}^{-1}$. Values determined by DPV, versus Fc/Fc^+ . ^d The approximation for the Fc/Fc^+ level is -4.8 eV versus vacuum: $E_{\text{HOMO}} = -4.8 - E_{\text{ox}}$. ^e Estimated from E_{HOMO} and $E_{\text{g}}^{\text{opt}}$: $E_{\text{LUMO}} = E_{\text{HOMO}} + E_{\text{g}}^{\text{opt}}$. ^f The data of **2a** were taken from ref. 10.

16 program.¹⁶ The singlet ground-state (S_0) and singlet excited-state (S_1) geometries were optimized by the DFT and time-dependent density functional theory (TD-DFT) methods with the M06-2X functional and a standard 6-31G(d) basis set (LanL2DZ basis set for Bi). The M06-2X functional is a high-nonlocality functional with double the amount of nonlocal exchange (2X), with reliable performance for thermochemistry, hydrogen bonding, kinetics, and weak interactions.¹⁷ The optimized molecular structures were verified by vibrational analysis; equilibrium structures did not have imaginary frequencies and transition state structures had only one imaginary frequency. The intrinsic reaction coordinate (IRC) calculations were carried out to check the transition state leading to the reactant and the product. Single-point energies were calculated using the 6-311+G(d,p) basis set (SDD basis set for Bi), and the solvent effect of MeCN was taken into account by using the integral equation formalism PCM (IEF-PCM). We first optimized the molecular structures of **1a** and $\text{Bi}(\text{OTf})_3$ -coordinated **1a** in the S_0 state and calculated their HOMO and LUMO levels (Scheme 5a). Upon the coordination of $\text{Bi}(\text{OTf})_3$, the smaller HOMO–LUMO gap is obtained particularly by lowering the LUMO level, which is in good agreement with the NMR and UV-vis absorption spectral changes observed in Scheme 4d.¹⁸ In addition, the $\text{Bi}(\text{OTf})_3$ coordination is an energetically favoured, exothermic process ($\Delta G = -28.0 \text{ kcal mol}^{-1}$). Thus, $\text{Bi}(\text{OTf})_3$ -coordinated **1a** (**intA_s0**) is a rational starting point in the present Mallory reaction (Scheme 5b). Consistent with no conversion under dark conditions (Table 1, entry 16), the activation barrier for direct electrocyclozation in the ground state (**tsAB_s0**) is $+72.9 \text{ kcal mol}^{-1}$, which can exclude the thermal reaction pathway. On the other hand, the S_1 intermediate **intA_s1** undergoes conrotatory electrocyclozation to form **intB_s0** via **intB_s1** with a reasonable energy change ($+38.0$ to $+59.4$ to $+15.4 \text{ kcal mol}^{-1}$). The direct elimination of *vic*-hydrogens in **intB_s0** is difficult because they are *anti* to each other. Thus, enolization-induced epimerization occurs through **intC_s0** to afford the corresponding *syn*-isomer **intD_s0**. The process is uphill, but subsequent spontaneous elimination of

molecular hydrogen into the gas phase and rearomatization into **intE** are strong driving forces for the reaction process. The overall energy profile reveals that the rate-limiting step is the C–H cleaved enolization from **intB_s0** to **intC_s0**, which is also in accordance with the primary KIE value observed in Scheme 4c.

The optical properties of some newly obtained compounds in Schemes 2 and 3 (**2j**, **2k**, **2l**, **2l'**, **3fa**, and **3fb**) were preliminary surveyed in a solution state ($1.0 \times 10^{-5} \text{ M CHCl}_3$ solution). The data of absorption/emission properties ($\lambda_{\text{abs}}/\lambda_{\text{em}}$) and fluorescence quantum yields (Φ_{F}), also involving those of **1a** and **2a** as the references, are summarized in Fig. 1 and Table 2. Compared to the uncyclized **1a**, the condensed **2a–2l'** all showed bathochromic shifts of their λ_{abs} values (379–412 nm) probably because of effective π -extension. On the other hand, impacts on the emission maxima λ_{em} were little or negligible. Relatively high quantum yields were observed in the push–pull-type alkoxy-substituted **2j** and **2k**, while higher fused **2l** and **2l'** containing the dibenzothiophene core showed lower emission efficiency. In general, the smaller Stokes shifts were obtained from all compounds owing to their rigid planar structures. The carbazole-incorporated dibenzophosphole **3fa** exhibited both absorption and emission maxima in the longer wavelength regions, which is reflected by the strongly electron-donating nature of the carbazole unit. Additionally, a very high quantum yield was obtained. The introduction of phenoxazine induced much larger red shifts of λ_{abs} and λ_{em} albeit with poor emission efficiency (**3fb**).

We also examined the electrochemical properties by cyclic voltammetry (CV) and differential pulse voltammetry (DPV) in MeCN with tetrabutylammonium hexafluorophosphate (Bu_4NPF_6) as an electrolyte versus ferrocene/ferrocenium ions (Fc/Fc^+) (Fig. S14–S19†), and their HOMO and LUMO levels were estimated according to the first oxidation potentials and the optical band gaps ($E_{\text{g}}^{\text{opt}}$). The data are summarized in Table 3. The CV of all compounds showed irreversible oxidation waves, and thus E_{ox} values were determined by DPV. The electron-donating alkoxy (**2j** and **2k**) and amino (**3fa** and **3fb**) substituents largely shifted E_{ox} values in a negative direction from those of the parent **2a**. Notably, compared to the carbazole-substituted **3fa**, the phenoxazine-substituted **3fb** exhibited a higher-lying HOMO level but an almost identical LUMO level, thus suggesting its larger intramolecular charge transfer ability. On the other hand, the structural isomers **2l** and **2l'** showed almost the same values for all parameters: their electrochemical properties are less dependent on the orientation of fused dibenzothiophene.

Conclusions

We have revisited the classical Mallory reaction of 2,3-diarylbenzophosphole derivatives and developed visible-light-promoted conditions with the assistance of $\text{Bi}(\text{OTf})_3$ Lewis acid. The use of visible light with lower energy enables the concise synthesis of highly condensed dibenzophospholes bearing a variety of functional groups, which are of potent interest in organic material chemistry. Additionally noteworthy is the evolution of molecular hydrogen: the reaction proceeds



even under external oxidant-free conditions and is thus truly dehydrogenative.¹⁹ Moreover, several experimental and computational studies uncover the detailed reaction mechanism. Development of catalytic conditions and application to other (hetero)aromatic systems as well as preparation of more condensed helically chiral molecules are ongoing in our laboratory.

Data availability

All experimental procedures and spectroscopic data can be found in the ESI.†

Author contributions

K. H. conceived the idea. I. K. and S. X. performed all experiments. Y. K. conducted computational studies with DFT. K. Y. and Y. N. assisted with X-ray analysis. K. H. supervised the project and wrote the manuscript. All the authors discussed the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by JSPS KAKENHI Grant No. JP 24KJ1578 (Grant-in-Aid for a JSPS Research Fellow, to Y. K.) and JP 23K23345 (Grant-in-Aid for Scientific Research(B), to K. H.) as well as by the JST FOREST Program, Grant Number JPMJFR 211X to K. H.

Notes and references

- Reviews and books: (a) T. Baumgartner and R. Réau, *Chem. Rev.*, 2006, **106**, 4681; (b) Y. Matano and H. Imahori, *Org. Biomol. Chem.*, 2009, **7**, 1258; (c) M. P. Duffy, W. Delaunay, P.-A. Bouit and M. Hissler, *Chem. Soc. Rev.*, 2016, **45**, 5296; (d) T. Baumgartner and F. Jäkle, *Main Group Strategies towards Functional Hybrid Materials*, Wiley, 2018. Selected examples: (e) H. Tsuji, K. Sato, L. Ilies, Y. Itoh, Y. Sato and E. Nakamura, *Org. Lett.*, 2008, **10**, 2263; (f) A. Fukazawa, H. Yamada and S. Yamaguchi, *Angew. Chem., Int. Ed.*, 2008, **47**, 5582; (g) H. Tsuji, K. Sato, Y. Sato and E. Nakamura, *J. Mater. Chem.*, 2009, **19**, 3364; (h) H. Tsuji, K. Sato, Y. Sato and E. Nakamura, *Chem.-Asian J.*, 2010, **5**, 1294; (i) C. Wang, M. Taki, Y. Sato, A. Fukazawa, T. Higashiyama and S. Yamaguchi, *J. Am. Chem. Soc.*, 2017, **139**, 10374.
- For recent reviews, see: (a) B. Wu and N. Yoshikai, *Org. Biomol. Chem.*, 2016, **14**, 5402; (b) H. Hattori, K. Ishida and N. Sakai, *Synthesis*, 2024, **56**, 193.
- Reviews: (a) F. B. Mallory and C. W. Mallory, *Org. React.*, 1984, **30**, 1; (b) K. B. Jørgensen, *Molecules*, 2010, **15**, 4334; (c) A. G. Lvov, *J. Org. Chem.*, 2020, **85**, 8749.
- (a) O. Fadhel, D. Szieberth, V. Deborde, C. Lescop, L. Nyulászi, M. Hissler and R. Réau, *Chem.-Eur. J.*, 2009, **15**, 4914; (b) P.-A. Bouit, A. Escande, R. Szűcs, D. Szieberth, C. Lescop, L. Nyulászi, M. Hissler and R. Réau, *J. Am. Chem. Soc.*, 2012, **134**, 6524; (c) F. Riobé, R. Szűcs, P.-A. Bouit, D. Tondelier, B. Geffroy, F. Aparicio, J. Buendía, L. Sánchez, R. Réau, L. Nyulászi and M. Hissler, *Chem.-Eur. J.*, 2015, **21**, 6547.
- J. Li, Z. Zhuang, J. Guo, X. Dong, J. Gong, B. Z. Tang and Z. Zhao, *Adv. Sci.*, 2023, **10**, 2305516.
- K. Nishimura, S. Xu, Y. Nishii and K. Hirano, *Org. Lett.*, 2023, **25**, 1503.
- For our protocol using AgOAc, see: (a) Y. Unoh, K. Hirano, T. Satoh and M. Miura, *Angew. Chem., Int. Ed.*, 2013, **52**, 12975. For related reactions using other promoters, see: ; (b) Y.-R. Chen and W.-L. Duan, *J. Am. Chem. Soc.*, 2013, **135**, 16754; (c) W. Ma and L. Ackermann, *Synthesis*, 2014, **2014**, 2297; (d) P. Zhang, Y. Gao, L. Zhang, Z. Li, Y. Liu, G. Tang and Y. Zhao, *Adv. Synth. Catal.*, 2016, **358**, 138; (e) D. M. Ma, W. Z. Chen, G. B. Hu, Y. Zhang, Y. X. Gao, Y. W. Yin and Y. F. Zhao, *Green Chem.*, 2016, **18**, 3522–3526; (f) V. Quint, F. Morlet-Savary, J. F. Lohier, J. Lalevee, A. C. Gaumont and S. Lakhdar, *J. Am. Chem. Soc.*, 2016, **138**, 7436.
- (a) K. Suzuki, Y. Hori and T. Kobayashi, *Adv. Synth. Catal.*, 2008, **350**, 652; (b) Y. Itai, Y. Nishii, P. Stachelek, P. Data, Y. Takeda, S. Minakata and M. Miura, *J. Org. Chem.*, 2018, **83**, 10289.
- (a) K. Nishimura, Y. Unoh, K. Hirano and M. Miura, *Chem.-Eur. J.*, 2018, **24**, 13089; (b) K. Nishimura, K. Hirano and M. Miura, *Org. Lett.*, 2022, **22**, 3185.
- S. Xu, K. Nishimura, K. Saito, K. Hirano and M. Miura, *Chem. Sci.*, 2022, **13**, 10950.
- Y. Tokura, S. Xu, Y. Kojima, M. Miura and K. Hirano, *Chem. Commun.*, 2022, **58**, 12208.
- M. Y. Kang,, S. G. Hong, J. S. Ha, D. H. Lee, T. Y. Park, B. J. Jang and S. D. Seo, Organic light emitting device comprising light efficiency improvement layer comprising novel dibenzo compounds, KR2016027940A, 2016-03-10.
- (a) M. Zhu, X.-L. Huang, S. Sun, C. Zhenga and S.-L. You, *J. Am. Chem. Soc.*, 2021, **143**, 13441; (b) M. Zhu, H. Xu, X. Zhang, C. Zheng and S.-L. You, *Angew. Chem., Int. Ed.*, 2021, **60**, 7036.
- (a) T. Nevesely, M. Wienhold, J. J. Molloy and R. Gilmour, *Chem. Rev.*, 2022, **122**, 2650; (b) J. Corpas, P. Mauleón, R. G. Arrayás and J. C. Carretero, *Adv. Synth. Catal.*, 2022, **364**, 1348.
- S.-Z. Zhang, S.-S. Zhang, J.-L. Li, S. Shen, X.-L. Yang and X. Niu, *J. Org. Chem.*, 2023, **88**, 9094.
- 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.
- 17 Y. Zhao and D. G. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215.
- 18 We also confirmed that the S_0 to S_1 transition arises from the HOMO–LUMO transition by TD-DFT calculation. Oscillator strengths of **1a** and Bi(OTf)₃-coordinated **1a** are 0.2727 and 0.1982, respectively. See the ESI† for more details.
- 19 For related reviews and perspective on photochemical cross-coupling with hydrogen evolution, see: (a) H. Wang, X. Gao, Z. Lv, T. Abdelilah and A. Lei, *Chem. Rev.*, 2019, **119**, 6769; (b) S. Tang, L. Zhang and A. Lei, *J. Am. Chem. Soc.*, 2018, **140**, 13128.

