

Cite this: *Chem. Sci.*, 2024, 15, 213

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

Received 12th October 2023
Accepted 17th November 2023

DOI: 10.1039/d3sc05414d

rsc.li/chemical-science

Oxidative cleavage of ketoximes to ketones using photoexcited nitroarenes†

Lucas T. Göttemann,‡ Stefan Wiesler‡ and Richmond Sarpong *

The methoxime group has emerged as a versatile directing group for a variety of C–H functionalizations. Despite its importance as a powerful functional handle, conversion of methoximes to the parent ketone, which is often desired, usually requires harsh and functional group intolerant reaction conditions. Therefore, the application of methoximes and their subsequent conversion to the corresponding ketone in a late-stage context can be problematic. Here, we present an alternative set of conditions to achieve mild and functional group tolerant conversion of methoximes to the parent ketones using photoexcited nitroarenes. The utility of this methodology is showcased in its application in the total synthesis of cephanolide D. Furthermore, mechanistic insight into this transformation obtained using isotope labeling studies as well as the analysis of reaction byproducts is provided.

Introduction

The total synthesis of natural products continues to be an exercise that highlights unaddressed gaps in synthetic methodology.¹ In particular, the high density of functional groups in complex natural products necessitates the development of functional group tolerant methods that can effect the selective reaction of a targeted reactive group among many others.² Recently, as a part of a total synthesis study of natural products in the cephalotane family,^{3,4} we required a method that would convert a methoxime to the corresponding ketone. Over the last decade, methoximes have emerged as superior directing groups for C–H functionalization,^{5–10} leading to value-added products where a variety of groups are installed. Despite their important role in directing C–H functionalizations, methoximes, unlike *O*-acetyl oximes, are notoriously difficult to convert to the parent ketone, which is often the desired functional group. The conditions for the removal of methoximes are generally harsh and, in some instances, can result in low yields (Scheme 1).

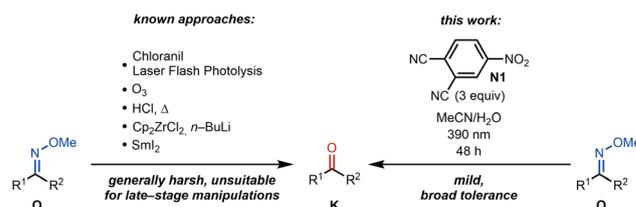
Therefore, the established conditions that are required for the conversion of methoximes to ketones are incompatible with a wide range of functional groups, making their use in a late-stage context challenging.^{11–15}

In our synthesis of cephanolide D (Scheme 2), a methoxime (1) that was prepared from a precursor ketone, had served as a directing group to effect carbomethoxylation at C15 giving rise to 2 through a palladated intermediate.¹⁶ However, myriad

attempts to convert the methoxime moiety in 2 back to the ketone group, including using strongly acidic¹⁷ or reductive¹⁸ conditions, resulted in, among many other undesired outcomes, elimination of the tertiary hydroxy group in 2 and subsequent decarboxylation of the bridging lactone. The only successful approach that was identified for removing the methoxime group in 2 was to effect ozonolysis cleavage of the oxime^{19,20} followed by a reductive workup. Even in that case, the reaction proceeded capriciously and did not lead to full conversion. Typically, reaction yields in the 35% range were realized along with a significant amount of recovered starting material (64%).

The yield of the desired product dropped precipitously upon prolonged exposure to ozone, likely due to competing oxidation at other susceptible sites on the molecule including the arene²¹ and benzylic positions.²²

While effective, ozonolysis can be impractical, especially on a small scale where it is difficult to determine the precise stoichiometry of ozone.²³ Furthermore, the inhalation and explosivity hazards of ozone and ozonides,²⁴ respectively, can make their use a significant safety risk. As a result of these considerations, we sought to identify milder conditions that could be

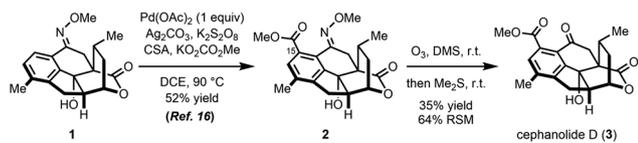


Scheme 1 Known conditions for the cleavage of methoximes to ketones and our approach.

Department of Chemistry, Latimer Hall, University of California, Berkeley, California 94720, USA. E-mail: rsarpong@berkeley.edu

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3sc05414d>

‡ These authors contributed equally.



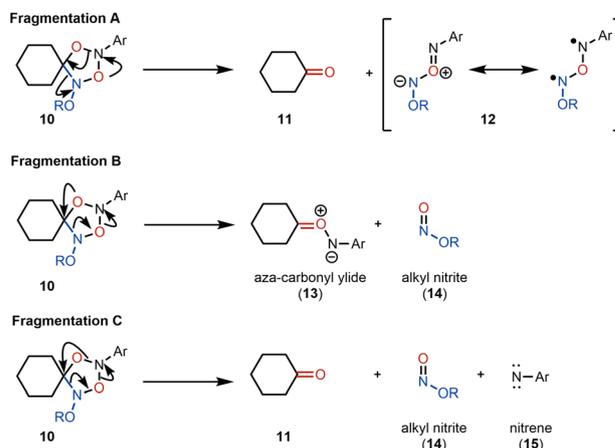
Scheme 2 Ozonolytic cleavage of the cephanolide D methoxime.

employed as an alternative for the cleavage of oximes — ideally, in a practical, safe, stoichiometrically precise and functional group tolerant fashion for applications in complex molecule synthesis.

Recently, concurrent reports from the groups of Leonori²⁵ and Parasram²⁶ described the use of photoexcited nitroarenes to cleave alkenes (Scheme 3). Notably, a range of nitroarenes were surveyed in these studies, whereby in some cases, the nature of the nitroarene significantly affected the yield of the oxidative cleavage product. Several functional groups were tolerated under these oxidative cleavage conditions including oxidation-susceptible groups such as a sulfide (see **4**).²⁷ These initial reports set the stage for additional developments such as the dihydroxylation of alkenes,²⁸ hydroxylation of aliphatic substrates,²⁹ and the oxidation of aldehydes and aldimines to carboxylic acids and amides, respectively.³⁰ Furthermore, Li recently reported a photoinduced cascade process between nitroarenes and amines to form isoxazolidine derivatives.³¹

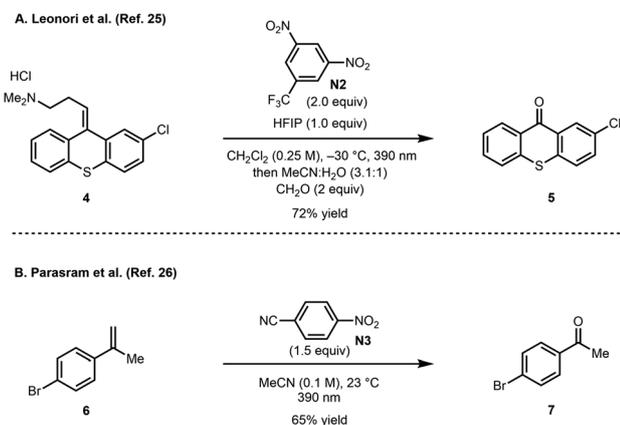
Results and discussion

On the basis of the alkene oxidative cleavage reports from Leonori and Parasram, we envisioned a cleavage of oximes using photoexcited nitroarenes that might proceed through an analogous mechanism. However, we anticipated that this transformation was likely to be challenging due to the significantly different properties of the oxime double bond compared to a C–C double bond. We envisioned that for cyclohexanone-derived ketoxime **8** (Scheme 4), reaction with a photoexcited nitroarene³² might result in the formation of an exciplex (see **9**), which upon cycloaddition would yield photoadduct **10**. Several scenarios could then be envisioned for the fate of photoadduct

Scheme 4 Possible fragmentation pathways for **10** in the oxidative cleavage.

10 (Scheme 4). In the fragmentation A pathway, the anticipated ketone product (*i.e.*, cyclohexanone, **11**) could form directly along with byproduct **12**, shown as its diradical and ylide resonance forms, which is expected to be a transient species. Alternatively, in fragmentation B, aza-carbonyl ylide **13** and alkyl nitrite **14** would be the anticipated products. Again, aza-carbonyl ylide **13** would be expected to be short-lived. In the fragmentation C pathway, direct formation of the ketone would be accompanied by alkyl nitrite **14** and aryl nitrene **15** as byproducts. Here, nitrene **15** would be expected to be fleeting. We reasoned that because a reactive intermediate would be formed in each of the fragmentation scenarios from photoadduct **10**, a trapping agent should be introduced to quench the reactive byproducts. We rationalized that water would be ideal for these purposes. In the fragmentation A pathway, **12** would be expected to hydrolyze to alkyl nitrite **14** and an aryl hydroxylamine. The formation of an aryl hydroxylamine byproduct would also be expected from the hydrolysis of aza-carbonyl ylide **13** (fragmentation B) and upon reaction of nitrene **15** with water through an OH-insertion.

On the basis of this rationalization, we embarked on a survey of conditions to effect the photoexcited nitroarene-mediated oxidative cleavage of oximes using tetralone methoxime **16** (which we envisioned to be a suitable model substrate for our cephanolide D synthesis). In addition to varying reaction conditions (*i.e.*, irradiation wavelength and duration, temperature, and solvent), we also tested several nitroarenes in our optimization studies (see the ESI† for details). Of note, in the cases where incomplete cleavage of the starting oxime was observed, a side-product observed in the mixture was the oxime diastereomer (*Z*-isomer), which presumably arises from photoisomerization under the reaction conditions.³³ With an initial



Scheme 3 Pioneering work by Leonori and Parasram.



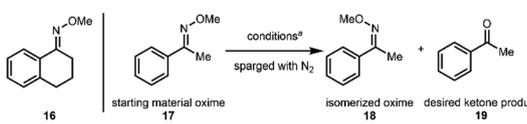
set of effective conditions in hand (see conditions in entry 1, Table 1), we began to examine the scope of this reaction.

Unfortunately, these conditions turned out not to be generalizable and we observed low yields for additional substrates that were tested such as acetophenone-derived oxime **17** (20% yield, entry 1), necessitating a second round of optimization for less reactive compounds. We chose acetophenone methoxime (**17**) as the substrate for this fine-tuning to achieve more broadly applicable conditions (see the ESI† for the full screening table). We retained the solvent mixture (4 : 1 MeCN : H₂O), which was shown to be significantly superior to other solvent mixtures used in our initial screening. We started by testing lower wavelength irradiation. LEDs centered at 390 nm showed an immediate improvement and resulted in a significant reduction of side reactions (entries 1–3). We then screened several nitroarenes and observed improved yields using nitroarenes **N2** and **N1**, which possess more electrophilic character in their excited states as compared to **N3** (entries 3–5).²⁵ Increasing the nitroarene equivalents and using a setup with Kessil lamps instead of a photoreactor³⁴ (see pictures in the ESI†) led to significantly higher yields (entries 5–7). Overall, the optimized conditions that we identified from these two surveys using tetralone methoxime (**16**) and acetophenone methoxime (**17**) are: using nitroarene **N1** (3 equiv) in a 4 : 1 mixture of acetonitrile and water with irradiation using LEDs centered at 390 nm at 40 °C (the measured temperature of our setup using a household fan for cooling) for 48 h (Table 1, entry 7). Finally, as shown in entry 8, in the absence of the nitroarene, the starting oxime is recovered in 90% yield along with 7% of the isomerized oxime.

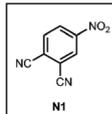
We then investigated the scope of the reaction to test the functional group tolerance of our method and its efficacy.

Importantly, we selected substrates featuring groups that are often encountered in more complex settings (*e.g.*, in total synthesis), where existing methods for converting methoximes to ketones have not been as effective. As shown in Fig. 1, a range of benzylic ketoximes are effectively cleaved under the conditions we have identified to afford the corresponding ketones.³⁵ Of note, sterically encumbered methoximes can be cleaved (see **34**) and oxidatively sensitive groups such as a thiomethyl group (**45**), which is oxidized to the sulfone under ozonolysis conditions (see the ESI†), are tolerated to some extent, emphasizing the complementarity of the current method to existing methods. Furthermore, tetralone, acetophenone, benzosuberone, and chromanone derived oximes all participate faithfully in this oxidative cleavage reaction. In addition, free oximes (see **39**), acetylated oximes (see **40**) and hydrazones (**41**) all undergo oxidative cleavage to the corresponding ketones under the conditions that we have identified. In general, we have found that substrates that proceed in low conversion, but high mass balance, can be further converted to the product over an extended reaction time, but the reaction rate decreases noticeably. In cases where poor mass balance was observed, longer wavelength irradiation (using LEDs centered at 420 nm) was beneficial to the reaction outcome. This is likely due reactivity of intermediates at the shorter wavelengths that were previously employed, which led to undefined side reactions. Mainly, the yields improved by irradiation at longer wavelengths for substrates with benzylic C–H groups that are known to undergo benzylic oxidation reactions. Albeit proceeding with lower mass balance upon irradiation at 420 nm over a prolonged period, we did not observe significant benzylic C–H hydroxylation products under these conditions.

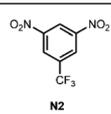
Table 1 Optimization of the reaction conditions for methoxime **17**



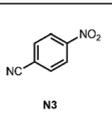
Entry	Wavelength	Nitroarene	Solvent	Time ^b	Yield 17 : 18 : 19
1	420 nm	N3 (1.5 equiv)	MeCN/H ₂ O 4 : 1	48 h	6%/70%/20%
2	405 nm	N3 (1.5 equiv)	MeCN/H ₂ O 4 : 1	48 h	7%/65%/23%
3	390 nm	N3 (1.5 equiv)	MeCN/H ₂ O 4 : 1	48 h	10%/58%/29%
4	390 nm	N2 (1.5 equiv)	MeCN/H ₂ O 4 : 1	48 h	14%/33%/47%
5	390 nm	N1 (1.5 equiv)	MeCN/H ₂ O 4 : 1	48 h	7%/21%/65%
6	390 nm	N1 (3.0 equiv)	MeCN/H ₂ O 4 : 1	48 h	<5%/17%/74%
7 ^c	390 nm	N1 (3.0 equiv)	MeCN/H₂O 4 : 1	48 h	-/trace/95%
8	390 nm	—	MeCN/H ₂ O 4 : 1	48 h	90%/7%/–



N1



N2



N3

^a A reaction time of 48 h is consistent with values reported for photochemical reactions with nitroarenes.^{25,26} ^b The reactions were carried out in a photoreactor.²⁸ A reaction time of 48 h is consistent with values reported for photochemical reactions with nitroarenes.^{25,26} ^c The reaction was carried out using Kessil lamps (distance to LED = 3 cm).



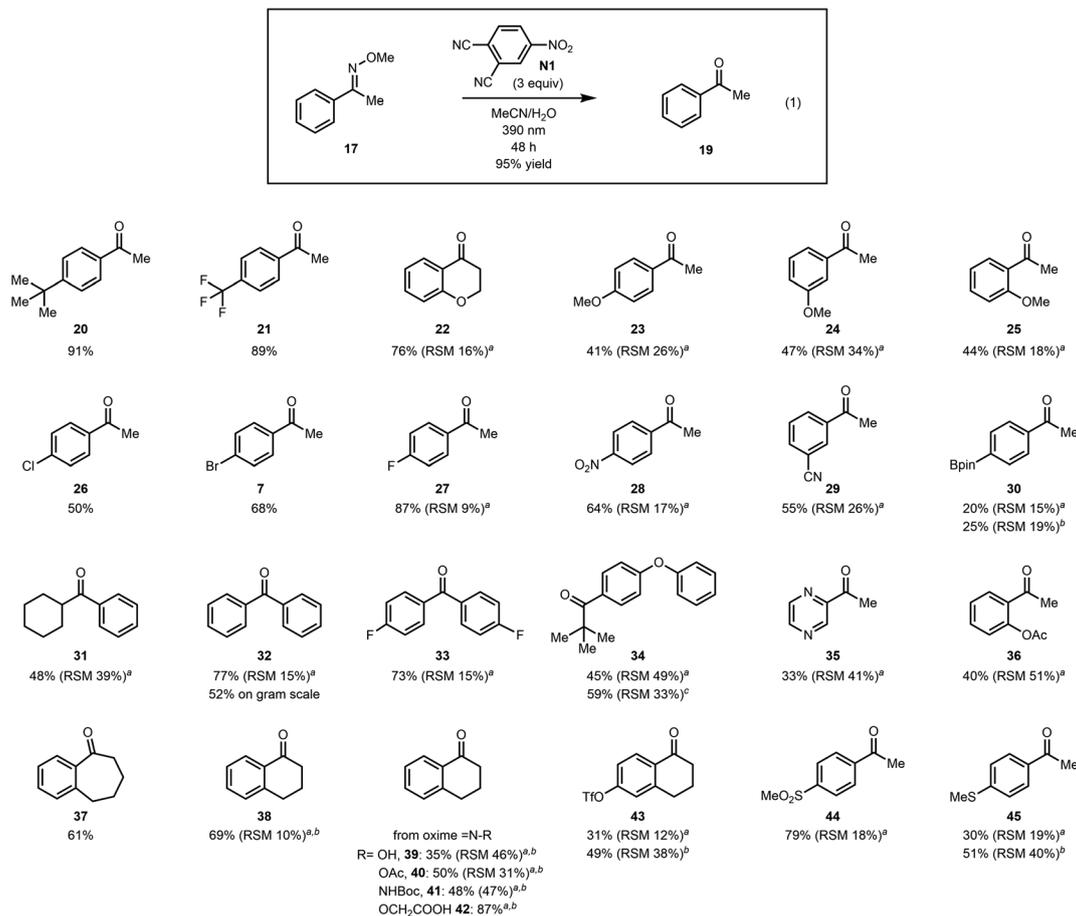


Fig. 1 Scope of benzylic methoxime cleavage. ^aRSM = recovered starting material. The RSM is a combined yield of oxime diastereomers. ^bIr-radiation was conducted at 420 nm using nitroarene **N3** for 48 h. ^cThe reaction was irradiated for 96 h.

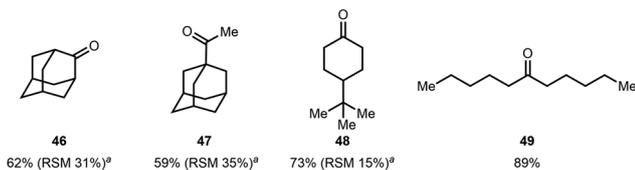
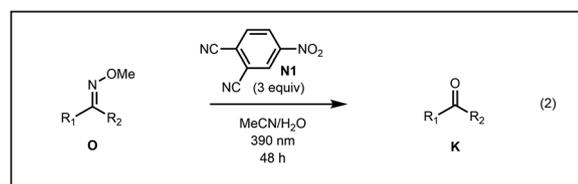
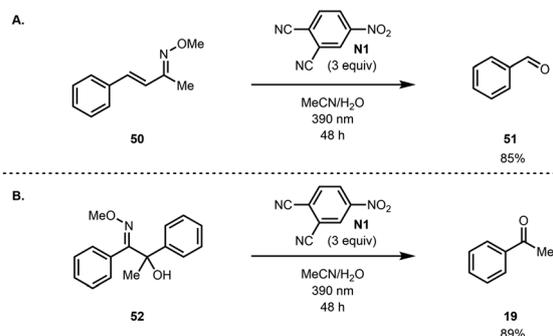


Fig. 2 Methoxime cleavage of aliphatic compounds. ^aRSM = recovered starting material. The RSM is a combined yield of oxime diastereomers.

The photoexcited nitroarene oxidative cleavage of a series of aliphatic methoximes was also investigated (Fig. 2). The reaction was found to work well for several aliphatic ketoximes including cyclic and sterically encumbered as well as acyclic substrates (see **46–49**).

However, the reaction does not tolerate alkenes, which are cleaved under the reaction conditions. For example, α,β -unsaturated oxime **50** leads to benzaldehyde (**51**) as the main detected product (Scheme 5A). Similarly, we found that acyloin derived methoxime **52** undergoes C–C bond cleavage under the reaction conditions to give acetophenone (Scheme 5B).

Of note, ozonolysis of **52** also results in the formation of acetophenone (**19**). Furthermore, we observed that substrates with free alcohol groups as well as amines were not tolerated



Scheme 5 Limitations and side reactions.



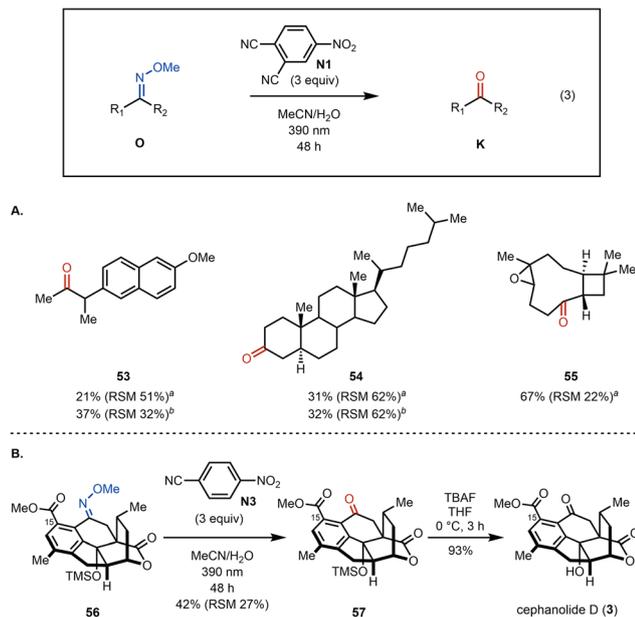


Fig. 3 (A) Application of methoxime cleavage in bioactive and complex substrates. (B) Synthesis of cephanolide D (3). ^aRSM = recovered starting material. The RSM is a combined yield of oxime diastereomers. ^bThe reaction was irradiated for 96 h.

under the reaction conditions, leading to a very low conversion of the starting material (see the ESI† for examples).

The oxime cleavage reaction has also been found to work well in several complex substrates (Fig. 3). For example, naproxen derived methylketone **53** is formed from the corresponding methoxime using our standard photoexcited nitroarene oxidative cleavage conditions in 21% yield (isolated along with 51% of the methoxime starting material). Cholestanone (**54**) is obtained in 31% yield (along with 52% of the methoxime starting material) upon subjection to the cleavage conditions. Similar to substrate **34** the yield for methylketone **53** could be improved upon irradiation for 96 h (see Fig. 3A), however cholestanone derived methoxime **54** suffered from solubility issues. On the other hand, the cleavage of the corresponding methoxime generated from kobusone (**55**) gives **55** in 67% yield (along with 22% of the methoxime starting material). Finally, in the context of our synthesis of cephanolide D, we subjected methoxime **56**³⁶ to the newly developed oxidative cleavage conditions (Fig. 3B). To our delight, our conditions were tolerated by the ester, lactone and even the TMS group, giving rise to ketone **57** — showcasing the use of photoexcited nitroarenes as an effective alternative for methoxime cleavage in compounds bearing various reactive groups.³⁷ Cephanolide D (**3**) was then obtained from **57** in 93% yield upon treatment with TBAF.

In order to test our hypothesis of a photoexcited nitroarene mediated oxidative cleavage of methoximes to the corresponding ketones, a comprehensive photophysical study was carried out. As a part of this investigation, the UV-vis data, including the absorption maxima of all the methoxime starting materials, ketone products, and nitroarenes at various concentrations were obtained experimentally. Our observations indicated that at a concentration of 50 μM, the methoximes, ketones, and

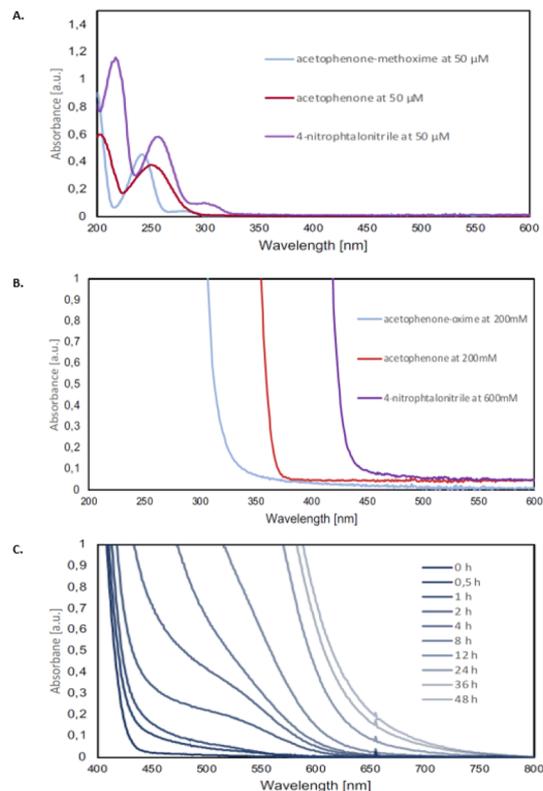
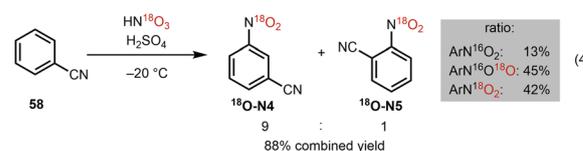


Fig. 4 (A) Absorption maximum for **17**, **19** and **N1** at 50 μM. (B) UV-vis absorption spectra for **17** and **19** at 200 mM and **N1** at 600 mM. C. UV-vis spectra for the reaction monitoring experiment indicating redshift using **17** and **N1** under standard reaction conditions and taking aliquots at the indicated time.

Table 2 ¹⁸O incorporation experiments

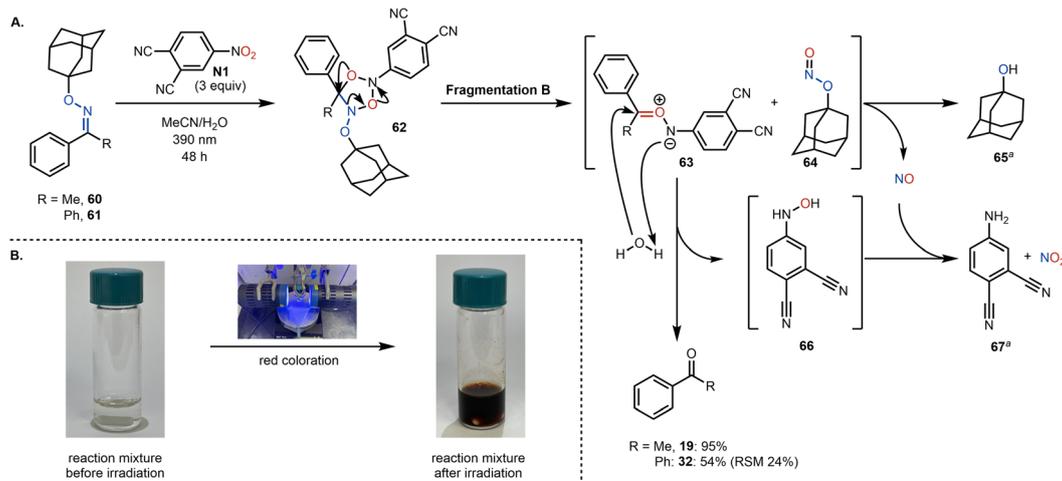
entry	wavelength	nitroarene	solvent	time	¹⁸ O incorporation
1	390 nm	¹⁸ O-N4 (1.1 equiv)	MeCN/H ₂ O 4:1	48 h	6%
2	390 nm	N4 (1.1 equiv)	MeCN/H ₂ ¹⁸ O 4:1	48 h	69%
3	390 nm	¹⁸ O-N4 (1.1 equiv)	MeCN/H ₂ ¹⁸ O 4:1	48 h	77%

nitroarenes do not absorb light centered at 390 nm. However, at higher concentrations, which reflect our reaction conditions (0.2 M), the nitroarenes strongly absorb whereas ketones can partially absorb at 390 nm (Fig. 4B). These absorption profiles strongly support the hypothesis that the reaction is mediated by



Scheme 6 Preparation of ¹⁸O labelled nitroarene ¹⁸O-N4.





Scheme 7 (A). Proposed mechanism of the oxidative cleavage. (B). Color change after 48 h reaction time. ^aByproducts were both detected by GC-MS and ¹H NMR.

photoexcited nitroarenes. Furthermore, our observations also explain the relatively long reaction time of 48 h since the ketone, formed over the course of the reaction (*i.e.*, continuously increasing concentration) partially absorbs light and reduces the efficiency of the absorption of the nitroarene and therefore reaction. This rationale is further supported by an increasing redshift over the course of the 48 h reaction duration, which suggests that an inner filter effect slows down the reaction.

To gain further insight into the proposed mechanism for this oxime cleavage method, we have carried out a series of experiments using ¹⁸O-labeled nitroarene ¹⁸O-N₄ and H₂¹⁸O (see Table 2). ¹⁸O-Labeled nitroarene ¹⁸O-N₄ was prepared by nitration of benzonitrile using ¹⁸O-labeled nitric acid and sulfuric acid analogous to the protocol of Wibaut (Scheme 6).³⁸

In our mechanistic studies, we have found that subjecting adamantanone methoxime (**59**) to our standard conditions using the ¹⁸O-labeled ¹⁸O-N₄ in a mixture of acetonitrile and ¹⁶O-labeled water leads to a 6% ¹⁸O incorporation in the ketone product (entry 1). The use of unlabeled (*i.e.*, ¹⁶O) nitroarene **N4** and ¹⁸O-labeled water leads to a 69% incorporation of ¹⁸O in the ketone product (entry 2), suggesting a fast O-atom exchange in the ketone group of adamantanone under the reaction conditions. The use of ¹⁸O-labeled nitroarene ¹⁸O-N₄ and ¹⁸O-labeled water leads to the ketone product in a slightly more isotopically enriched form (77% incorporation of ¹⁸O; entry 3) than in the case where only ¹⁸O-labeled water is used (*i.e.*, entry 2). While strict conclusions as to the source of ¹⁸O label in the observed products cannot be drawn (*i.e.*, the oxygen atom in the ketone product might arise from the nitroarene or through fast exchange with water), the slight enrichment in ¹⁸O for the product under the conditions where only ¹⁸O-N₄ is used (entry 1), and also the higher enrichment for the case where ¹⁸O-N₄ and H₂¹⁸O are used (compare entry 3 to entry 2) may support the nitroarene, in addition to the water co-solvent, as sources of the oxygen atom in the product. These preliminary labeling studies support the proposed mechanism (Scheme 4) wherein a cycloaddition between the photoexcited nitroarene and oxime is proposed (see 9).

To gain further mechanistic insight, we sought to investigate the fate of the oxime residue. Two chemical probes (**60** and **61**) bearing a bulky, non-volatile, alkyl substituent (in place of the methyl group) were synthesized and subjected to our reaction conditions (Scheme 7). Assuming the formation of cycloadduct **62**, fragmentation B pathway (*vide supra*) would give rise to aza-carbonyl ylide **63** and transient adamantyl nitrite **64**. Subsequently, aza-carbonyl ylide **63** could be trapped by water yielding the desired ketone (**19** or **32**) as well as hydroxylamine **67**, which can then be envisioned to undergo NO mediated reduction to aniline **67**.^{39,40} We assumed that the NO arises from the photochemical decomposition of alkyl nitrite **64**, resulting in the formation of 1-adamantanol (**65**) and NO.⁴¹ Both **65** and **67** were detected in the crude reaction mixture (using GC-MS and NMR; see the ESI† for more details), providing additional support for the fragmentation B pathway. Due to the reactive nature of the intermediates described, other concomitant decomposition pathways of **62** are also plausible (*e.g.*, *via* the fragmentation A pathway), which would explain a partial incorporation of ¹⁸O from the nitroarene as observed in the labeling experiments.

Conclusions

In summary, building on recent studies that describe the use of photoexcited nitroarenes to effect oxidative cleavage of alkenes, we report here the removal of oxime groups to afford the corresponding ketones under a related set of conditions. The substrate scope of the transformation is relatively broad but is not tolerant of alkenes, which are cleaved under the reaction conditions (consistent with the precedent of Leonori and Parasmam). The success of the current studies enabled the application of this new protocol to a preparation of cephalolide D from a precursor methoxime. While the mechanism of the reaction is yet to be fully elucidated, isotopic labelling studies and the detection of several by-products support both the water cosolvent and nitroarene as possible sources of the O-atom in the ketone product that is obtained. Overall, this protocol offers



an alternative to the strongly acidic or reductive conditions as well as ozonolysis cleavage conditions that are usually employed in the conversion of ketoxime groups to ketones.

Author contributions

The project was conceived by S. W. The experimental work was performed by L. G. and S.W. The initial draft was written by R. S. with input from all authors. L. G., S. W. and R. S. were involved in editing and finalizing the manuscript. R. S. supervised the project and secured funding.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by funding from the US National Science Foundation (CHE-1700982). S.W. thanks the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation, WI 5665/1-1) for a postdoctoral scholarship. The authors are grateful to Dr Hasan Celik and UC Berkeley's NMR facility in the College of Chemistry (CoC-NMR) for spectroscopic assistance. Instruments in CoC-NMR are supported in part by NIH S10OD024998. The authors thank Magan Powell (UC Berkeley) for assistance with GC-MS.

Notes and references

- 1 K. C. Nicolaou, S. Rigol and R. Yu, *CCS Chem.*, 2019, 3–37.
- 2 R. A. Shenvi, D. P. O'Malley and P. S. Baran, *Acc. Chem. Res.*, 2009, **42**, 530–541.
- 3 M. Haider, G. Sennari, A. Eggert and R. Sarpong, *J. Am. Chem. Soc.*, 2021, **143**, 2710–2715.
- 4 G. Sennari, K. E. Gardner, S. Wiesler, M. Haider, A. Eggert and R. Sarpong, *J. Am. Chem. Soc.*, 2022, **144**, 19173–19185.
- 5 Y.-F. Liang, X. Wang, Y. Yuan, Y. Liang, X. Li and N. Jiao, *ACS Catal.*, 2015, **5**, 6148–6152.
- 6 C. Ma, C.-Q. Zhao, Y.-Q. Li, L.-P. Zhang, X.-T. Xu, K. Zhang and T.-S. Mei, *Chem. Commun.*, 2017, **53**, 12189–12192.
- 7 Y. Park, S. Jee, J. G. Kim and S. Chang, *Org. Process Res. Dev.*, 2015, **19**, 1024–1029.
- 8 Y.-F. Liang, X. Li, X. Wang, Y. Yan, P. Feng and N. Jiao, *ACS Catal.*, 2015, **5**, 1956–1963.
- 9 B. S. Pilgrim, A. E. Gatland, C. H. A. Esteves, C. T. McTernan, G. R. Jones, M. R. Tatton, P. A. Procopiou and T. J. Donohoe, *Org. Biomol. Chem.*, 2016, **14**, 1065–1090.
- 10 S. Yu, B. Wan and X. Li, *Org. Lett.*, 2015, **17**, 58–61.
- 11 H. J. P. De Lijser and C.-K. Tsai, *J. Org. Chem.*, 2004, **69**, 3057–3067.
- 12 R. E. Erickson, P. J. Andrulis, J. C. Collins, M. L. Lungle and G. D. Mercer, *J. Org. Chem.*, 1969, **34**, 2961–2966.
- 13 E. Tan, M. Zanini and A. M. Echavarren, *Angew. Chem., Int. Ed.*, 2020, **59**, 10470–10473.
- 14 M. Kitamura, Y. Shintaku, D. Kudo and T. Okauchi, *Tetrahedron Lett.*, 2010, **51**, 4890–4893.
- 15 F. Huang and S. Zhang, *Org. Lett.*, 2019, **21**, 7430–7434.
- 16 Z.-Y. Li and G.-W. Wang, *Org. Lett.*, 2015, **17**, 4866–4869.
- 17 R. J. Sharpe and J. S. Johnson, *J. Am. Chem. Soc.*, 2015, **137**, 4968–4971.
- 18 M. M. Majireck, J. A. Witek and S. M. Weinreb, *Tetrahedron Lett.*, 2010, **51**, 3555–3557.
- 19 (a) Y. Ito, H. Yokoya, Y. Umehara and T. Matsuura, *Bull. Chem. Soc. Jpn.*, 1980, **53**, 2407–2408; (b) D. J. Weitz and M. D. Bednarski, *J. Org. Chem.*, 1989, **54**, 4957–4959.
- 20 K. Griesbaum, B. Övez, T. S. Huh and Y. Dong, *Liebigs Ann.*, 1995, **1995**, 1571–1574.
- 21 E. A. Skrotzki, J. K. Vandavasi and S. G. Newman, *J. Org. Chem.*, 2021, **86**, 14169–14176.
- 22 W. Gao, L. Du, W. Jiao and Y. Liu, *Chin. J. Chem. Eng.*, 2020, **28**, 808–814.
- 23 A. L. Gomez, T. L. Lewis, S. A. Wilkinson and S. A. Nizkorodov, *Environ. Sci. Technol.*, 2008, **42**, 3582–3587.
- 24 A. Lazen, *Emergency and Continuous Exposure Limits for Selected Airborne Contaminants*, Defense Technical Information Center, Fort Belvoir, VA, 1984, vol. 1.
- 25 A. Ruffoni, C. Hampton, M. Simonetti and D. Leonori, *Nature*, 2022, **610**, 81–86.
- 26 D. E. Wise, E. S. Gogarnoiu, A. D. Duke, J. M. Paolillo, T. L. Vacala, W. A. Hussain and M. Parasram, *J. Am. Chem. Soc.*, 2022, **144**, 15437–15442.
- 27 A. Maggiolo and E. A. Blair, *Ozone Chemistry and Technology*, American Chemical Society, N.W. Washington 6, D.C., 1959.
- 28 C. Hampton, M. Simonetti and D. Leonori, *Angew. Chem., Int. Ed.*, 2023, **62**, e202214508.
- 29 J. M. Paolillo, A. D. Duke, E. S. Gogarnoiu, D. E. Wise and M. Parasram, *J. Am. Chem. Soc.*, 2023, **145**, 2794–2799.
- 30 J. K. Mitchell, W. A. Hussain, A. H. Bansode, R. M. O'Connor, D. E. Wise, M. H. Choe and M. Parasram, *Org. Lett.*, 2023, **25**, 6517–6521.
- 31 X.-Z. Shao, G.-Y. Xu, W. Fan, S. Zhang and M.-B. Li, *Org. Chem. Front.*, 2023, **10**, 624–631.
- 32 R. Hurley and A. C. Testa, *J. Am. Chem. Soc.*, 1966, **88**, 4330–4332.
- 33 X. Zhang and T. Rovis, *J. Am. Chem. Soc.*, 2021, **143**, 21211–21217.
- 34 A. Penn PhD M2 was used. See ESI† for details.
- 35 Generally, caution must be taken when isolating the product ketones since in some instances their volatility must be taken into consideration as this can result in diminished yields.
- 36 As we mentioned before, free alcohols react poorly and are therefore a limitation of this method. For this reason we decided to use the TMS protected methoxime **62** instead of methoxime **10**.
- 37 Subsequently, we removed the TMS group in **63** using TBAF to yield cephanolide D (93% yield).
- 38 J. P. Wibaut and R. van Strik, *Recueil*, 1958, **77**, 316–326.
- 39 E. Damiani, L. Greci and C. Rizzoli, *J. Chem. Soc. Perkin Trans.*, 2001, **2**, 1139–1144.
- 40 H. Weber, A. Grzesiok, R. Sustmann and H.-G. Korth, *Z. Naturforsch.*, 1994, **49b**, 1041–1050.
- 41 G. R. McMillan, *J. Phys. Chem.*, 1963, **67**, 931–932.

