

**Microwave- and Ultrasound-Accelerated Green Permanganate Oxidation of Thioethers**

Luu, Thi Xuan Thi; Le, Huu Tan ; Le, Thach Ngoc; Duus, Fritz

*Published in:*  
Organic Chemistry Current Research

*DOI:*  
[10.4172/2161-0401.1000129](https://doi.org/10.4172/2161-0401.1000129)

*Publication date:*  
2015

*Document Version*  
Publisher's PDF, also known as Version of record

*Citation for published version (APA):*  
Luu, T. X. T., Le, H. T., Le, T. N., & Duus, F. (2015). Microwave- and Ultrasound-Accelerated Green Permanganate Oxidation of Thioethers. *Organic Chemistry Current Research*, 4(1), 1-5.  
<https://doi.org/10.4172/2161-0401.1000129>

**General rights**

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain.
- You may freely distribute the URL identifying the publication in the public portal.

**Take down policy**

If you believe that this document breaches copyright please contact [rucforsk@kb.dk](mailto:rucforsk@kb.dk) providing details, and we will remove access to the work immediately and investigate your claim.

# Microwave- and Ultrasound-Accelerated Green Permanganate Oxidation of Thioethers

Thi Xuan Thi Luu,<sup>1,2\*</sup> Huu Tan Le,<sup>2</sup> Thach Ngoc Le,<sup>2</sup> and Fritz Duus<sup>1</sup>

<sup>1</sup>Department of Science, Systems, and Models, Roskilde University, P.O. Box 260, DK-4000, Roskilde, Denmark.

<sup>2</sup>Department of Organic Chemistry, University of Science, 227 Nguyen Van Cu, Ho Chi Minh City, Vietnam.

## Abstract

Product formation from the solvent-free oxidation of aliphatic, cyclic or aromatic thioethers by permanganate supported on copper sulfate pentahydrate has been investigated in detail with respect to the importance of the nature of thioethers, the molar ratio of potassium permanganate absorbed on copper sulfate pentahydrate and reaction activation methods. While microwave irradiation affects appropriately on the fast formation of the sulfones within 3-13 minutes to get the yield more than 74%, ultrasound irradiation activates oxidation of cyclic thioethers mildly to form the corresponding cyclic sulfoxides with the high yields in the range of 83-96%.

**Keywords:** Potassium permanganate; Oxidation; Thioethers; Microwave irradiation; Ultrasound irradiation

## Introduction

Sulfones and sulfoxides are important commercial compounds, intermediates in organic synthesis and biologically active compounds. [1] They have been mainly produced from the oxidation of thioethers. [2].

In recent decades, with the advancement of science, green chemistry has changed our life style day-to-day [3,4]. Solvent-free reaction plays an important role in the green chemistry due to prevent from solvent contamination, solvent vapour, the elimination of volatile organic solvent, hazards, toxicity and to reduce cost procedure [4]. In solvent-free oxidation reaction, permanganate ion is preferred to be used as "green" oxidation reagent because manganese dioxide formed is easily isolated and treated [3-5]. Moreover, potassium permanganate is a powerful, commercially available and un-expensive oxidation reagent [6]. The permanganate oxidation of thioethers has been investigated under the homogeneous or heterogeneous conditions by using phase-transfer catalysts supported aqueous permanganate solution, [7-9] permanganate complex, [10,11] ground permanganate, [12-14] or inorganic salts/solid supports assisted permanganate ion. [15-19] Furthermore, the efficiency of potassium permanganate on the solvent-free oxidation of thioethers has been shown clearly when it was absorbed on montmorillonite K-10, [20] manganese dioxide, [21,22] Rexyn 101 H, [23] alumina, [24] or inorganic salts [25,26].

Since ultrasound and microwave irradiation were discovered, they have contributed greatly to the green chemistry, specially on the solvent-free reaction aspects [4,27,28]. The presence of ultrasound irradiation has a good effect on the solvent-free oxidation of thioether by potassium permanganate supported on manganese dioxide, [21] besides microwave irradiation has been studied primarily on the solvent-free oxidation of dibenzyl thioether by potassium permanganate supported on copper sulfate pentahydrate [25]. In the continuation of our works on the oxidation by using potassium permanganate absorbed on copper sulfate pentahydrate called xPP/yCSP (x molar amount of potassium permanganate absorbed on y molar amount of copper sulfate pentahydrate), [29] the formation of sulfoxides or sulfones in the solvent-free oxidation of thioethers has inspired us to study under the acceleration of ultrasound and microwave irradiation (Figure 1).

## Materials and Methods

### General

Ultrasound irradiations were performed by means of a Branson 1210E-MT ultrasonic bath, operating at 47 kHz. Microwave irradiations were performed by means of a CEM MDS 200 batch microwave oven. GC/MS analyses were performed on a Hewlett Packard 5890 GC 5971A MS apparatus equipped with a J&W DB-5MS capillary column (30 m, 0.25 mm i.d., 0.25  $\mu$ m film thickness) and a Hewlett Packard 7673A autosampler. NMR spectra were recorded on a Varian Mercury 300 NMR spectrometer.

### Oxidation of thioethers into corresponding sulfoxides or sulfones by xPP/yCSP under solvent-free reaction conditions (Method A)

A suitable quantity of finely ground xPP/yCSP was added to a 5-mL round-bottom flask containing the thioether (following the molar ratio as in Table 1) and six glass balls (d=2 mm). The flask was fitted to a shaking machine and shaken at the speed of 280 rpm for a specific period of time (Table 1). Then, the reaction mixture was extracted with 45-50 mL of dichloromethane and filtered through celite layer (2 cm). The extract was removed solvent by rotational evaporation, and then, the remaining crude product was analysed by GC/MS and NMR spectroscopy.

### Oxidation of thioethers into corresponding sulfoxides or sulfones by xPP/yCSP under solvent-free reaction conditions assisted by ultrasound irradiation (Method B)

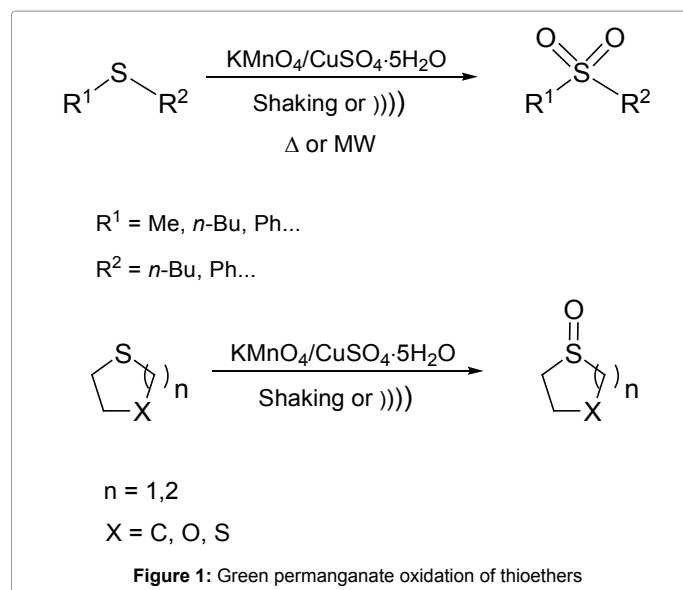
A suitable quantity of finely ground xPP/yCSP was added to a test tube (h=20.0 cm, d=3.0 cm) containing the thioether (following the

\*Corresponding author: Thi Xuan Thi Luu. Department of Science, Systems, and Models, Roskilde University, P.O. Box 260, DK-4000, Roskilde, Denmark; Tel.: +84-838-353-659; Fax: +84-838-350-096; E-mail: [lxthi@hcmus.edu.vn](mailto:lxthi@hcmus.edu.vn)

Received April 01, 2014; Accepted December 25, 2014; Published January 01, 2015

Citation: Luu TXT, Le HT, Le TN, Duus F (2015) Microwave- and Ultrasound-Accelerated Green Permanganate Oxidation of Thioethers. Organic Chem Curr Res 1:129. doi:10.4172/2161-0401.1000129

Copyright: © 2015 Luu TXT et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.



molar ratio as in Table 1). The test tube was placed into an ultrasound bath where the mixture of reactants was exposed to ultrasound irradiation for a specific period of time (Table 1). Subsequently, the reaction mixture was worked up as described in method A.

#### Oxidation of thioethers into corresponding sulfones by xPP/yCSP under solvent-free reaction conditions assisted by microwave irradiation (Method C)

A suitable quantity of finely ground xPP/yCSP was added to a test tube (h=20.0 cm, d=3.0 cm) containing the thioether (following the molar ratio as in Table 2). The test tube was placed into a beaker equipped to adhere test tubes in the CEM (Matthews, North Carolina, USA) oven. For each of the thioethers, an irradiation programme was applied to determine the most efficient reaction conditions, see Table 2. For every experiment, the temperature of the reaction mixture was measured immediately after reaction stop in order to compare with experiments performed under conventional heating. After being cooled, the reaction mixture was worked up as described in method A.

#### Oxidation of thioethers into corresponding sulfones by xPP/yCSP under solvent-free reaction conditions assisted by conventional heating (Method D)

A test tube (h=20.0 cm, d=3.0 cm) containing a suitable quantity of finely ground xPP/yCSP and the thioethers (following the molar ratio as in Table 2) were placed in an oil bath heated to the temperature measured at reaction stop of the parallel reaction run under microwave irradiation. The test tube was kept in the oil bath for a period of time corresponding exactly to that found at optimum in Method C. After being cooled, the reaction mixture was worked up as described in Method A.

#### <sup>1</sup>H- and <sup>13</sup>C-NMR spectroscopic data

The identity and purity of all products reported were ensured by NMR as well as by gas chromatography/mass spectrometry (GC/MS). Most of products are well known and spectroscopically well-characterized already, except the <sup>1</sup>H NMR as well as the <sup>13</sup>C NMR spectra of thianthrene-5,5-dioxide and thianthrene-5,5,10,10-tetroxide are described below

Thianthrene-5,5-dioxide (Entry 5, Table 1) white solid, M.p = 165.5-167.5 °C. <sup>1</sup>H-NMR (300 MHz) δ (ppm)=8.18-8.24 (m, 2H), 7.62-7.68 (m, 2H), 7.51-7.57 (m, 4H); <sup>13</sup>C-NMR (75 MHz) δ (ppm) = 135.30, 135.09, 132.04, 128.75, 127.72, 125.44.

Thianthrene-5,5,10,10-tetroxide (Entry 6, Table 1) <sup>1</sup>H NMR (300 MHz) δ (ppm)=8.31-8.37 (m, 4H), 8.02-8.09 (m, 4H). <sup>13</sup>C NMR (75 MHz) δ (ppm)=138.26, 134.94, 125.89.

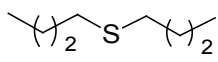
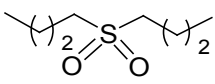
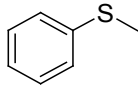
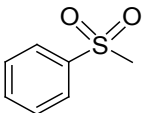
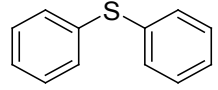
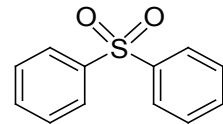
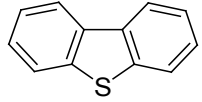
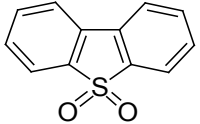
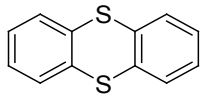
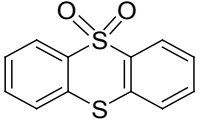
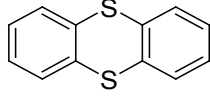
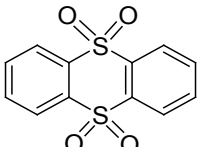

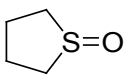
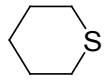
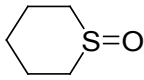
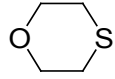
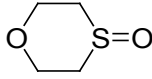
#### Results

The oxidation of aliphatic thioethers was performed on the basis of work of Shaabani and Lee who used 4 g of potassium permanganate (2 g, 12 mmol) ground with an equal amount of copper sulfate pentahydrate (2 g, 8 mmol) to convert 2 mmol dibutyl thioether into the corresponding sulfone (100%) after five-hour stirring [25]. In our work, the copper sulfate pentahydrate was dissolved in de-ionized water completely, then KMnO<sub>4</sub> was added along with a sufficient volume of de-ionized water to obtain a homogeneous solution. The solution was stirred for 10 minutes at 80°C. Subsequently, water was removed from the solution by rotational evaporation until the weight of the remaining solid mass was equal to the sum of the weights of the original ingredients. The obtained solid mass was ground in a mortar into a fine homogeneous powder.

In order to find the most efficient oxidation agent, a series of experiments was performed, where the molar ratio between KMnO<sub>4</sub> and CuSO<sub>4</sub>·5H<sub>2</sub>O was varied following the nature of thioether (aliphatic, cyclic or aromatic thioether). The most efficiency of the oxidants investigated, appeared to be PP/3CSP used for the oxidation of aliphatic thioethers and PP/5CSP used for the oxidation of cyclic or aromatic thioethers. In order to achieve the best yield of main product, we also paid attention to the molar ratio between each thioether and xPP/yCSP. The most appropriate molar ratio was also chosen in Table 1.

Altogether eleven thioethers were subjected to oxidation by xPP/yCSP under solvent-free reaction conditions, using four different methods. In the first series of oxidation reactions where the mixture of reactants was simply shaken together at room temperature (Method A, Table 1), fair to excellent yields were obtained in all cases. The next series of oxidation reactions were performed as described before but under the assistance of ultrasound irradiation (Method B, Table 1). Although the yields of the products in Method B were only slightly improved, the reaction times were shortened considerably. Obviously, ultrasound irradiation has affected well on the heterogeneous reactions (liquid/solid phase) as well as solid/solid phase reactions because cavitation collapse made the surface of solids cleaner, solid articles smaller and mass transfer easily; while above types of reactions often occur incompletely and slowly due to problem with conventional rotational mixing techniques [30-32]. In the Method A and Method B, the oxidation of aliphatic or aromatic thioethers led to the formation of sulfones (Entry 1-6, Table 1). However, due to the overlap of unhybridized *p* orbital of sulfur atom in aromatic thioethers to form a continuous ring of parallel orbitals, electron-donating of aromatic sulfur atom to oxidation reagent happened more difficulty than aliphatic sulfur atom [33-35]. Depending on structure of aromatic thioether, the oxidation did not occur (e.g. the oxidation of thiophene and benzothiophene) or occurred slowly under stronger oxidation conditions (e.g. the oxidation of dibenzothiophene and thianthrene). There was a special product formation in the oxidation of thianthrene (containing two sulfur atoms), where the product from the oxidation of one sulfur atom (thianthrene, 5,5-dioxide) was obtained after nine-

**Table 1:** Yields of products obtained by the solvent-free xPP/yCSP-promoted oxidation of thioethers by shaking method and ultrasound irradiation.

| Entry | Starting materials  | Product   | xPP/yCSP | Molar ratio <sup>a</sup> | Method A <sup>b</sup> yield <sup>c</sup> (Conv. <sup>c</sup> , time <sup>d</sup> ) | Method B <sup>b</sup> yield <sup>c</sup> (Conv. <sup>c</sup> , time <sup>d</sup> ) |
|-------|---|---|----------|--------------------------|--|--|
| 1     |    |    | PP/3CSP  | 3:6                      | 95 (100, 8)  | 98 (100, 3)  |
| 2     |    |    | PP/3CSP  | 3:6                      | 94 (100, 8)  | 96 (100, 3.5)  |
| 3     |    |    | PP/3CSP  | 1.5:3                    | 98 (100, 9)  | 99 (100, 3)  |
| 4     |  |  | PP/5CSP  | 1.5:3                    | 80 (83, 72)  | 95 (100, 8)  |
| 5     |  |  | PP/5CSP  | 1.2:4.8                  | 81 (81, 9)   | -  |
| 6     |  |  | PP/5CSP  | 1.2:4.8                  | 82 (84, 12)  | 99 (100, 5)  |
| 7     |  |  | PP/5CSP  | 3:2                      | 93 (96, 16)  | 72 (79, 7)   |
| 8     |  |  | PP/5CSP  | 3:3                      | 72 (83, 20)  | 76 (85, 3.5)   |
| 9     |  |  | PP/5CSP  | 3:4                      | 81 (92, 22)  | 82 (87, 3.5)   |

<sup>a</sup>Molar ratio = the molar ratio of thioether : xPP/yCSP. <sup>b</sup>Method A: the reactants were shaken at room temperature. Method B: the reaction was assisted by ultrasound irradiation. <sup>c</sup>Yield and Conv. (conversion yield) were based on GC/MS analysis. <sup>d</sup>Time = reaction time in hours.

**Table 2:** Yields of products obtained by the solvent-free xPP/yCSP-promoted oxidation of thioethers by microwave irradiation and conventional heating.

| Entry | Starting materials | Product | xPP/yCSP | Molar ratio <sup>a</sup> | Method C <sup>b</sup> yield <sup>c</sup> (Conv. <sup>c</sup> , time <sup>f</sup> , W <sup>g</sup> ) | Method D <sup>b</sup> yield <sup>c</sup> (Conv. <sup>c</sup> , time <sup>f</sup> , temp. <sup>h</sup> ) |
|-------|--------------------|---------|----------|--------------------------|---|---|
| 1     |                    |         | PP/3CSP  | 3:6                      | 94 (100, 1.5, 60)   | 93 (100, 1.5, 95)   |
| 2     |                    |         | PP/3CSP  | 3:6                      | 93 (100, 6, 150)  | 93 (95, 6, 92)  |
| 3     |                    |         | PP/3CSP  | 1.5:3                    | 97 (100, 3, 60)   | 97 (100, 3, 86)   |
| 4     |                    |         | PP/5CSP  | 1.5:3                    | 98 (100, 5.5, 500)  | 60 (61, 5.5, 90)  |
| 5     |                    |         | PP/5CSP  | 1.2:4.8                  | 94(100, 10.2, 630)  | 82 (90, 10.2, 98)   |
| 6     |                    |         | PP/5CSP  | 3:3                      | 74 (77, 6.5, 250)   | 46 (58, 6.5, 96)  |
| 7     |                    |         | PP/5CSP  | 3:4                      | 89 (97, 12, 350)  | 58 (65, 12, 90)   |
| 8     |                    |         | PP/5CSP  | 3:4                      | 86 (91, 12.3, 350)  | 57 (61, 12.3, 91)   |

<sup>a</sup>Molar ratio = the molar ratio of thioether : xPP/yCSP. <sup>b</sup> Method C: the reaction was assisted by microwave irradiation. Method D: the reaction mixture was heated in an oil bath at appropriate temperature (Experimental). <sup>c</sup>Yield and Conv. (conversion yield) were based on GC/MS analysis. <sup>f</sup>Time = reaction time in minutes. <sup>g</sup>W = power of microwave irradiation in Watts. <sup>h</sup>Temp. = oil bath temperature (Experimental).

hour shaking (Entry 5, Method A). Prolongation of the reaction time more than twelve-hour shaking as well as performing the reaction under ultrasound irradiation led to the formation of thianthrene, 5,5,10,10-tetraoxide. In the other cases, cyclic thioethers were oxidized into the corresponding cyclic sulfoxides without ring opening (Entry 7-9, Table 1), except the unsuccessful and ring-opening oxidation of 1,4-dithiane. It was explained that the conformation of new-born cyclic sulfoxides (transition products) were not in a plane; therefore, the next oxidation step to form cyclic sulfones took place slowly under the mild conditions.

Under heating activation, transition products could not be existed and reverted to reactants for going on to form stable products in

the next step; therefore, the oxidation of all categories of thioethers produced the corresponding sulfones in the short time. It would be of interest to check whether the drastically shortened reaction times could be effected simply by the higher reaction temperatures, so a series of experiments were performed in the presence of microwave irradiation (Method C, Table 2) in comparison with those at the same reaction times and temperatures under conventional heating (Method D, Table 2). The results of experiments (Table 2) clearly demonstrated that a conveniently increased reaction temperature could shorten drastically the formation of sulfone from the oxidation reaction. However, microwave irradiation with "dielectric heating" effects has reached the suitable reaction temperature faster and saved energy much more than conventional heating [36,37].

## Conclusion

Comprehensive experimental work has made it possible for us to introduce the oxidative reagent xPP/yCSP and emphasize the effects of activation methods on product formation. PP/3CSP and PP/5CSP are also used for fast and solvent-free oxidation of thioethers to get high yield of the products. The assistance of ultrasound or microwave irradiation efficiently shortens the reaction times compared with the shaking or conventional heating methods. Heating activations are suitable for sulfone formation through the oxidation of thioethers.

## Acknowledgments

We thank Danish International Aid Programme (DANIDA) via the Enhancement of Research Capacity (ENRECA) program and Vietnam National Foundation for Science and Technology Development (NAFOSTED) under grant number [104.01-2011.40] for financial support.

## References

- Carreno MC (1995) Applications of sulfoxides to asymmetric synthesis of biologically active compounds. *Chem. Rev.* 95: 1717-1760.
- Durst T (1979) In *Comprehensive Organic Chemistry*. Pergamon Press: Oxford, Great Britain, Vol. 3, chapter 11.6: 121-156.
- Ahluwalia VK, Kidwai M (2004) *New Trends in Green Chemistry*. Anamaya Publishers, New Delhi, India.
- Tanaka K (2003) *Solvent-Free Organic Synthesis*; Wiley-VCH, Weinheim.
- Singh N, Lee DG (2001) Permanganate: A green and versatile industrial oxidant. *Org. Proc. Res. and Develop.* 5: 599-603.
- Fatiadi AJ (1987) The classical permanganate ion: Still a novel oxidant in organic chemistry. *Synthesis*: 85-127.
- Aitken RA, Mesher STE, Ross FC, Ryan BM (1997) Effect of added benzoic acid on the Phase-Transfer Catalysed Permanganate Oxidation of Organosulfur Compounds. *Synthesis*: 787-791.
- Gokel GW, Gerdes HM, Dishong DM (1980) Sulfur heterocycles. 3. Heterogeneous, Phase-transfer, and acid-catalyzed potassium permanganate oxidation of sulfides to sulfones and a survey of their Carbon-13 nuclear Magnetic resonance spectra. *J Org Chem* 45: 3634-3639.
- Lee DG, Srinivasan NS (1982) An improved procedure for the preparation of sulfones. *Sulfur Lett.* 1: 1-3.
- Markó IE, Richardson PR, Bailey M, Maguire AR, Coughlan N (1997) Selective Manganese-mediated transformations using the combination:  $\text{KMnO}_4/\text{Me}_3\text{SiCl}$ . *Tetrahedron Lett.* 38: 2339-2342.
- Lakouraj MM, Tajbakhsh M, Tashakkorian H, Ghodrati K (2007) Fast and efficient oxidation of sulfides to sulfones with  $\text{N,N}'$ -Dibenzyl- $\text{N,N,N}'$ -tetramethyl diammonium permanganate. *Phosphorus, Sulfur, and Silicon* 182: 485-490.
- Firouzabadi H, Seddighi M (1991) Conversion of sulfides to their corresponding sulfoxides with barium permanganate  $\text{Ba}(\text{MnO}_4)_2$  under non-aqueous condition. *Synth. Commun.* 21: 211-214.
- Shaabani A, Tavasoli-Rad F, Lee DG (2005) Potassium Permanganate oxidation of organic compounds. *Synth. Commun.* 35: 571-580.
- Dishington A, Fillery S, Finlay MRV (2010) A one-pot sulfide to sulfone oxidation with *m*-chloroperoxybenzoic acid and sodium permanganate. *Tetrahedron Lett.* 51: 4211-4213.
- Lai S, Lee DG (2002) Lewis acid assisted permanganate oxidation. *Tetrahedron* 58: 9879-9887.
- Xie N, Binstead RA, Block E, Chandler WD, Lee DG, Meyer TJ, et al. (2000) Reduction of permanganate by thioanisole: Lewis acid catalysis. *J. Org. Chem.* 65: 1008-1015.
- Noureddin NA, McConnell WB, Lee DG (1984) Heterogeneous permanganate oxidations. 4. The oxidation of sulfides and selenides. *Can. J. Chem.* 62: 2113-2116.
- Wolfe S, Ingold CF (1983) Oxidation of organic compounds by zinc permanganate. *J Am Chem. Soc.* 105: 7755-7757.
- Shaabani A, Teimouri F, Lee DG (2003) Ion exchange catalysis in oxidation of organic compounds with  $\text{KMnO}_4$ . *Synth. Commun.* 33: 1057-1065.
- Shaabani A, Bazgir A, Lee DG (2004) Oxidation of organic compounds by potassium permanganate supported on Montmorillonite K10. *Synth. Commun.* 34: 3595-3607.
- Shaabani A, Mirzaei P, Naderi S, Lee DG (2004) Green oxidations. The use of potassium permanganate supported on manganese dioxide. *Tetrahedron* 60: 11415-11420.
- Shaabani A, Mirzaei P, Lee DG (2004) The beneficial effect of manganese dioxide on the oxidation of organic compounds by potassium permanganate. *Catalysis Lett.* 97: 119-123.
- Shaabani A, Lee DG (2003) Solvent free oxidations using manganese dioxide, barium manganate and potassium permanganate. *Recent Res. Devel. Organic Chem.* 7: 85-104.
- Hajipour AR, Mallakpour SE, Adibi H (2002) A facile and selective method for oxidation of sulfides and thiols to their corresponding sulfoxides and disulfides with alumina-supported potassium permanganate under solvent-free conditions. *Sulfur Lett.* 25: 155-160.
- Shaabani A, Lee DG (2001) Solvent free permanganate oxidations. *Tetrahedron Lett.* 42: 5833-5836.
- Shaabani A, Lee DG (2003) Selective oxidation of sulfides under solvent-free conditions. *Sulfur Lett.* 26: 43-46.
- Luche JL (1998) *Synthetic Organic Sonochemistry*. Plenum Press, New York.
- Hayes BL (2002) *Microwave Synthesis*. CEM Publishing, New York.
- Luu TXT, Christensen P, Duus F, Le NT (2008) Microwave- and Ultrasound-Accelerated green oxidation of alcohols by potassium permanganate absorbed on copper(II) sulfate pentahydrate. *Synth Commun* 38: 2011-2024.
- Mason TJ, Lorimer JP (1989) *Sonochemistry: Theory, applications and uses of ultrasound in chemistry*. Ellis Horwood, West Sussex.
- Gogate PR (2008) Cavitation reactors for process intensification of chemical processing applications: A critical review. *Chem Eng Process* 47: 515-527.
- Sutka VS, Gogate PR (2009) Design aspects of sonochemical reactors: Techniques for understanding cavitation activity distribution and effect of operating parameters. *Chem Eng J* 155: 26-36.
- Banerji KK (1988) Mechanism of the oxidation of organic sulfides by permanganate ion. *Tetrahedron.* 44: 2969-2975.
- Lee DG, Chen T (1991) Oxidation of organic sulfides by permanganate ion. *J. Org. Chem.* 56: 5346-5348.
- Jayaraman A, East ALL (2012) The mechanism of permanganate oxidation of sulfides and sulfoxides *J Org. Chem.* 77: 351-356.
- Kappe CO, Stadler A (2005) *Microwaves in Organic and Medicinal Chemistry*. Wiley-VCH, Weinheim, Vol. 25.
- Gole VL, Gogate PR (2013) Intensification of synthesis of biodiesel from non-edible oil using sequential combination of microwave and ultrasound. *Fuel Process Technol.* 106: 62-69.

Citation: Luu TXT, Le HT, Le TN, Duus F (2015) Microwave- and Ultrasound-Accelerated Green Permanganate Oxidation of Thioethers. *Organic Chem Curr Res* 1:129. doi:[10.4172/2161-0401.1000129](https://doi.org/10.4172/2161-0401.1000129)