
Impact of Additives and ZnO Nanoparticles on the Sonodegradation of Methylene Blue

ABSTRACT

Sonochemical method for the degradation of toxic dyestuffs is now considered as a promising technique. An important dye namely Methylene Blue (MB) was degraded using ultrasound irradiation at different pH. The impact of various additives like CCl_4 , H_2O_2 , NaCl and Na_2SO_4 with different doses for MB degradation was observed. The degradation efficiency of ZnO nanoparticles was also measured. The sonolysis was conducted for 0, 5, 15, 25 minutes and the extent of degradation was measured by UV-vis spectrophotometer. Acidic media was more efficient for degradation compared to basic media. 200 $\mu\text{M/L}$ doses H_2O_2 of 20 mM concentration showed highest efficiency (41.98%) to degrade the MB molecules. On the other hand CCl_4 , Na_2SO_4 and ZnO nanoparticles also showed maximum 35.63%, 34.88% and 41.72% degradation efficiency respectively. The efficiency increases in the presence of additives and ZnO nanoparticles in comparison to the efficiency (18.88%) obtained without these additives.

Keywords: Additives; azo dye; degradation; methylene blue; nanoparticles.

1. INTRODUCTION

Organic dyes are used in a wide range of sectors, including optical discs, food, cosmetics, medicines, solar cells, and traditional textile manufacturing. To ensure clean and safe water sources, it is critical to identify a method for extracting and discarding the organic dye chemicals used on a regular basis (Chu 2001, Khorramfar et al. 2010). Although there are numerous types of dyes, azo dyes which are organic compounds bearing the functional group R-N=N-R' , represent for 70% of total dye production globally (Garcia-Segura et al. 2013). These compounds are distinguished by the azo bond chromophore group ($-\text{N}=\text{N}-$), which is related with aromatic structures and functional groups such as $-\text{NH}_2$ and $-\text{SO}_3\text{H}$ (Forgacs et al. 2004, Martínez et al. 2009). These dyes are widely used in the food, textile, paper, leather, and pharmaceutical industries because of their exceptional properties and low cost (Chang et al. 2001). The effects of various dye compounds on human health vary. When azo dyes and their intermediates degrade, they can cause damage, mutagenesis, and carcinogenesis, posing a major risk to human health (Rai et al. 2005). For example, MB is a major synthetic dye used in large quantity in water testing, sulfur analysis, medicine, and biology (Kim 2014). It also

functions as a peroxide producer and redox indicator. However, they generally disturb ecosystems and are toxic, mutagenic, and carcinogenic to both aquatic and human life (Moraisi et al. 1999, Santhy 2006, Entezari et al. 2003). It is important to develop a technology to remove this dye from water. Numerous chemical, physical, and biological approaches have been developed to remove these compounds (Entezari et al. 2003, Zhu et al. 2011, Zhou et al. 2010). Advanced oxidation processes (AOPs) have received a lot of attention due to their ability to totally mineralize organic contaminants to CO_2 and H_2O . Ultrasonography has lately been used in alternative operating procedures for wastewater treatment (Adewuyi 2001, Chowdhury 2009, Hua et al. 1997). The two main concepts of sonochemical reactions include pyrolysis reactions that occur at and inside hot regions during collapsing bubbles, as well as radical reactions by OH and H radicals produced by water pyrolysis (Misik et al. 1995).



Several research have looked at the sonochemical degradation of phenolic chemicals and dyes (Entezari et al. 2003, Okitsu et al. 2005, Inoue et al. 2006, Merouani 2010, Kobayashi et al. 2012, Kobayashi et al. 2014,

Kruus et al. 1997, Petrier et al. 1997). The development of efficient additives to accelerate the decomposition of organic molecules has been the subject of extensive investigation. Recent research investigated how adding CCl_4 or C_6F_{14} impacts sonochemical degradation (Petrier et al. 1997, Chowdhury 2009). Although CCl_4 is a toxic chemical and hence heavily controlled in use, it is an effective addition for speeding up the sonochemical breakdown of the target organic molecules. Sponza et al. found that $19 \text{ mg L}^{-1} \text{C}_6\text{F}_{14}$ boosted the rate of phenol removal in olive mill wastewater (Sponza 2014). According to Zeng et al., the sonochemical degradation of phenol rose from 0.014 to 0.031 min^{-1} or from 0.014 to 0.032 min^{-1} in the presence of 150 M CCl_4 or $1.5 \text{ M C}_6\text{F}_{14}$ (Zheng et al. 2005). According to research the action of CCl_4 or C_6F_{14} as a H atom scavenger is responsible for these positive benefits (Sponza 2014, Zheng et al. 2005). Furthermore, it has been observed that the usage of specific chemicals can affect how quickly colors deteriorate (Monira et al. 2022, Uddin et al. 2018, Hossain et al. 2013). Metal oxide semiconductors, on the other hand, have been extensively studied for their potential as photocatalysts for the removal of organic pollutants from water and air (Nagpal et al. 2019, Wawrzkiwicz et al. 2017, Danish et al. 2020, Huang et al. 2019). When the metal oxides are activated by the appropriate quantity of photon energy, active sites of electron-hole pairs form, increasing catalytic activity on the metal oxide surfaces (Lee et al. 2016). TiO_2 is considered the oldest, most common, and best material among these metal oxides due to its outstanding stability and relatively simple fabrication technique (Hashimoto et al. 2005, Dastan et al. 2016, Dastan et al. 2016, Dastan 2017). Zinc oxide (ZnO), a metal oxide with an exciton binding energy of 60 m eV and a rather broad band gap of 3.37 eV , has received substantial research as a photocatalyst for the breakdown of numerous organic contaminants. Furthermore, ZnO nanostructures are recommended over TiO_2 as photocatalyst alternatives for photodegradation due to their better solar spectrum absorption,

lower cost, and nontoxicity (Azmina et al. 2017). Numerous studies have shown that ZnO nanostructures have significant photocatalytic activity for removing organic contaminants, such as organic dyes (Hariharan 2006). Therefore, in this work we have investigated the efficiency of additives like CCl_4 , H_2O_2 , NaCl , Na_2SO_4 , and ZnO nanoparticles on the sonolytic degradation of MB at certain concentration and pH.

2. EXPERIMENTAL

Materials and equipments: Methylene Blue ($\text{C}_{16}\text{H}_{18}\text{ClN}_3\text{S}$), Merck Life Science Private Limited, Godraj, Vadhroli East, Mumbai-400079, Sodium Chloride (NaCl), Ranbaxy Fine chemicals Limited A-3, New Delhi-110020 (ISO 9001: 2000 certified company), Sodium sulphate (Na_2SO_4), Merck Specialities Private Limited, Worli, Mumbai-400018, Hydrogen peroxide (H_2O_2), Carbon tetrachloride (CCl_4), Merck, D-6100 Darmstadt, FR. Germany. All the Chemicals were used as purchased. The sonicator machine used in this work is shown in Fig. 1.

Preparation of solutions: MB solutions with a concentration of 25 mg/L were prepared. In 100 ml of distilled water, solutions of the inorganic salts NaCl and Na_2SO_4 were prepared using 36 and 13.9 grams of each salt, respectively, based on their maximum saturation points. One liter of distilled water was mixed with 1.02 mg and 2.04 ml of H_2O_2 to make 10 mM/L and 20 mM/L H_2O_2 solutions, respectively. A little syringe is used to prepare the 200 , 400 , and $600 \mu\text{M/L}$ CCl_4 . ZnO nanoparticle was prepared using sol gel technique using ZnNO_3 and KOH . Pure ZnO nanoparticle was used as catalyst with 12.5 mg dose. Water soluble dyes can absorb and reflect light, reducing the water's clarity naturally. Normally, dye effluent contains between 10 and 50 mg/L , although 1.0 mg/L dye solutions are visible and may be considered as pollutants (Hamida et al. 2022, Cheng et al. 2009).



Fig. 1. Sonicator bath used in sonochemical degradation

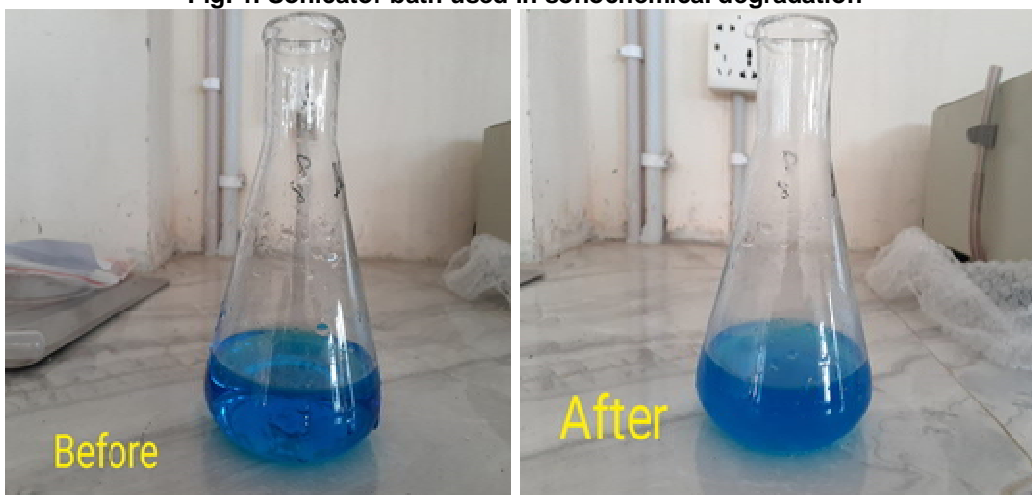


Fig. 2. Methylene blue solution

3. RESULTS AND DISCUSSION

Degradation percentage of MB at various pH:

The degradation percentages of MB without additives used in sonochemical irradiation process under various situations are presented and discussed below. The degradation percentages were calculated using the following equation.

$$\% \text{ of degradation} = (C_0 - C) / C_0 \times 100 \quad (2)$$

Where, C represents the final concentration obtained after sonication at different time and C_0 represents the initial concentration of MB solution. Fig. 3 depicts the degradation efficiency of MB at different pH. NaOH and H_2SO_4 solutions were used to change the solution's pH.

From the graph it is found that the efficiency at pH 3 is 56.33% at pH 7 is 15.06% and at pH 8 is 23.82% where the efficiency for raw solution

without changing the pH is 18.88% i. e acidic media is more efficient for degradation than basic media. There is a considerable chance of enhancing degradation efficiency using additives

that maintain an acidic environment during sonochemical breakdown. Previous research has shown that sonochemical degradation efficiency is higher in an acidic environment.

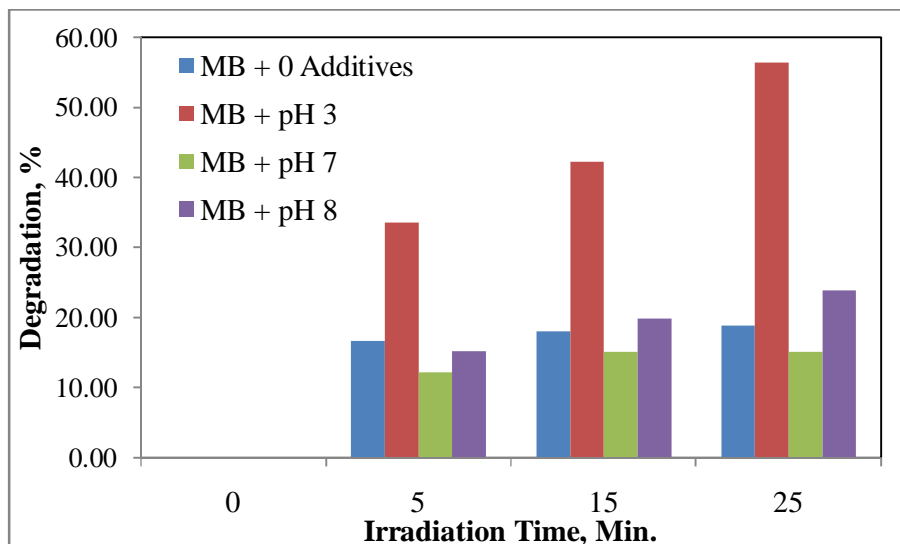


Fig. 3. Degradation efficiency for pure MB at various pH.

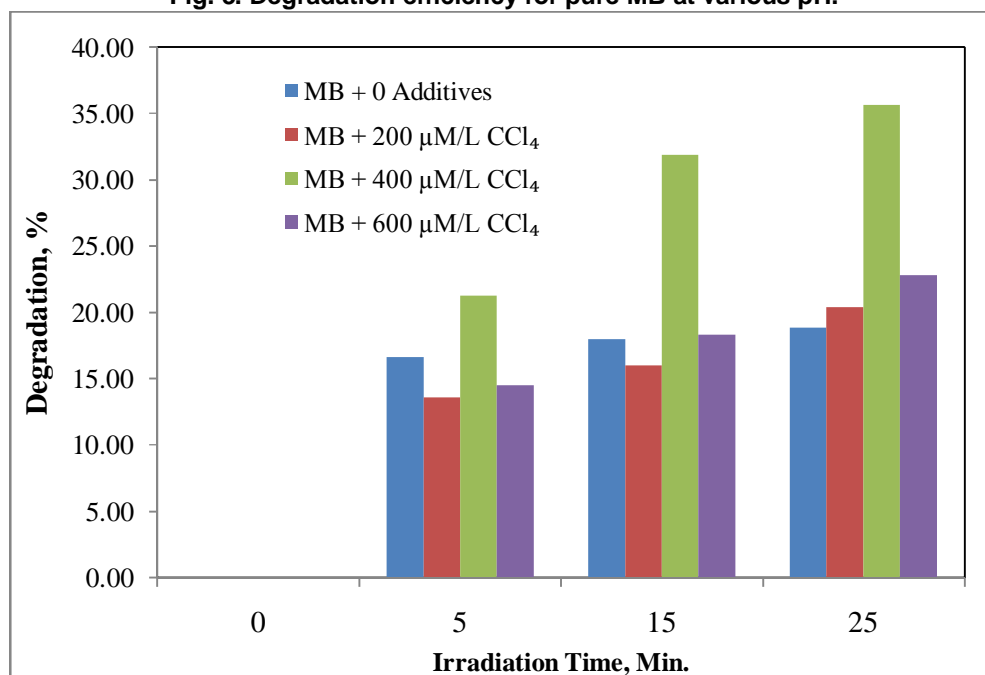


Fig. 4. Efficiency of CCl₄ for the degradation of MB

Efficiency of CCl₄ for sonolytic degradation of MB: The efficiency of CCl₄ at 200, 400, and 600 μM/L concentrations on the sonochemical degradation of MB is represented in Fig. 4.

In presence of CCl₄, the highest efficiency of MB degradation is 35.63% when 400 μM/L CCl₄ was present. In the presence of 600 μM/L CCl₄ the efficiency is 22.80% and at 200μM/L the

efficiency of degradation is 20.39%. We suggest that active Cl radicals or related species are formed during CCl_4 sonolysis, and that these radicals or species could be beneficial in the breakdown of MB, though some researchers' claims that CCl_4 could act as a H atom scavenger (Zheng et al. 2005). According to previous research (Petrier 1997, Suslick et al. 1997, Francony 1996, Hua 1996, Okitsu et al. 2008), the following responses might occur:



The radicals that form, which contain chlorine (Cl , CCl_3 , CCl_2 , and Cl_2) and so on, may speed up the breakdown of MB in an aqueous solution. Merouani et al. 2010 reported that 200 mg L⁻¹ of CCl_4 increased the sonochemical degradation rate of rhodamine B at 300 kHz by 21 times. According to Okitsu et al., adding 100 ppm CCl_4 increased the sonochemical breakdown rate of MO at 200 kHz by 41 times (Dastan 2017). The effectiveness of CCl_4 for MB degradation discovered in this investigation is consistent with these findings. Another study found that the presence of CCl_4 improves the sonolytic breakdown. From the above discussion, though CCl_4 has high toxicity (Recknagel et al. 1989) it can be used as an additive due its high degradation efficiency.

Efficiency of H_2O_2 for MB degradation:

Sonolysis of MB with 20, 100 and 200 μL doses of H_2O_2 at two different concentration 10 mM and 20 mM were carried out and the degradation efficiency is demonstrated in figure 5. It is observed from the figure that highest degradation efficiency showed by 200 μL of 10 mM H_2O_2 is 25.97% (Fig. 5 a) and 200 μL of 20 mM H_2O_2 is 41.98% (fig. 5 b) after 25 minutes irradiation i. e. higher doses and higher irradiation time increases the efficiency of degradation. The generation of free radicals as active species explains the H_2O_2 reaction process for dye discolouration.



According to research, the US/UV/ H_2O_2 technique is the most successful in decomposing

malachite green because it incorporates ultrasound to facilitate H_2O_2 scission in addition to photolysis (Banatet al. 1996). Another recent study assessed the impact of introducing hydrogen peroxide at concentrations of 0.05, 0.10, and 0.15 M. The treatment with 0.10 M H_2O_2 and 45 W was effective and energy-efficient, removing 62.9% of the water's absorbance (Miah n.d.). Sonolysis with 400 μL of H_2O_2 has been shown to improve methyl orange elimination efficiency (Monira et al. 2022). When compared to previous research, it is clear that a variety of uses for hydrogen peroxide (H_2O_2) have emerged in the field of water treatment due to its relative safety and convenience of use.

Efficiency of NaCl and Na_2SO_4 for the degradation of MB:

10 ml and 20 ml NaCl and Na_2SO_4 were added in to the MB solution in order to find the efficiency of the inorganic salts on dye degradation. Figure 6 and 7 represents the observed result. It is seen in the figure that highest degradation (34.88%) obtained by 20 ml Na_2SO_4 indicates the addition of Na_2SO_4 improved the degradation efficiency for MB. On the other hand the addition of NaCl caused it to decrease as compared to the degradation of pure MB due to the Cl^- ion produced. Previous research has demonstrated that in synthetic dye solutions containing a variety of Na_2SO_4 and NaCl mixer, Na_2SO_4 had a more modest effect on decolourization efficiency than NaCl. A higher concentration of Na_2SO_4 did not impede the decolorization process and even boosted the effectiveness of reactive bright red K-2BP in dye solutions with comparable salt or Na^+ concentrations (Zollinger Ed). Furthermore, the degradation rate has accelerated due to the increased salt concentration. Following comparison, it can be concluded that Na_2SO_4 outperforms NaCl in terms of MB degradation. Uddin et al. published a study in 2016 on the sonochemical degradation of 4-chlorophenol, phenol, catechol, and resorcinol in the absence and presence of Na_2SO_4 or NaCl (Uddin et al. 2016). The rate of phenolic compound decomposition adhered to a pseudo-first order rate constant (Bafana et al. 2011). Another study conducted by Monira et al. found that the effectiveness of methyl orange degradation increased in the presence of a 20 ml Na_2SO_4 solution.

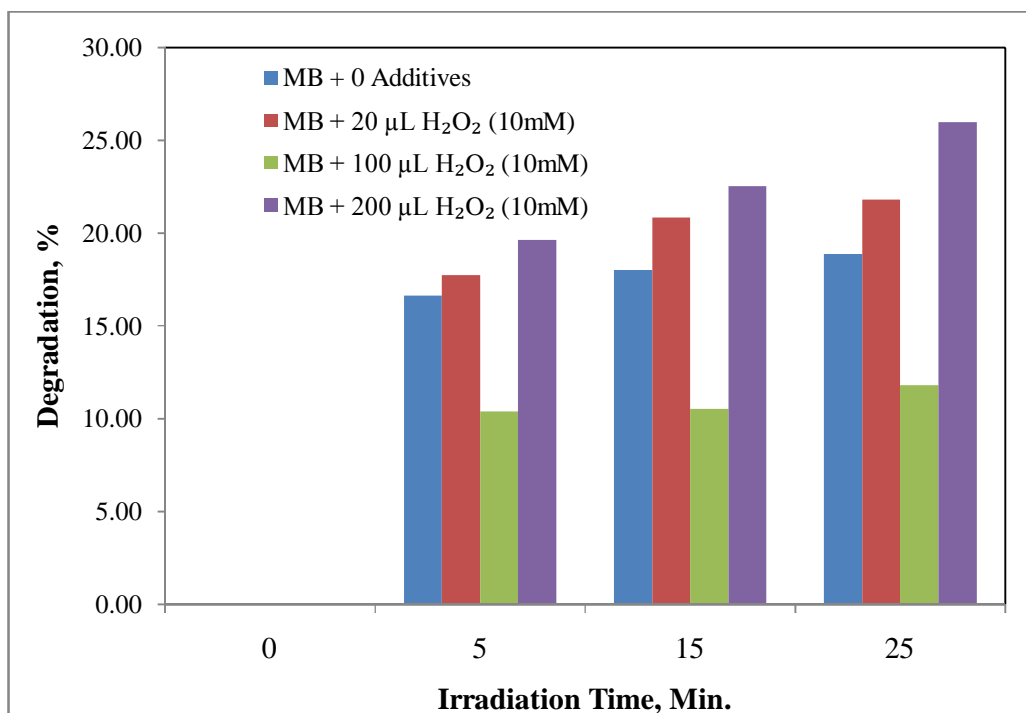


Fig. 5a. Efficiency of H₂O₂ (10mM) for the degradation of MB

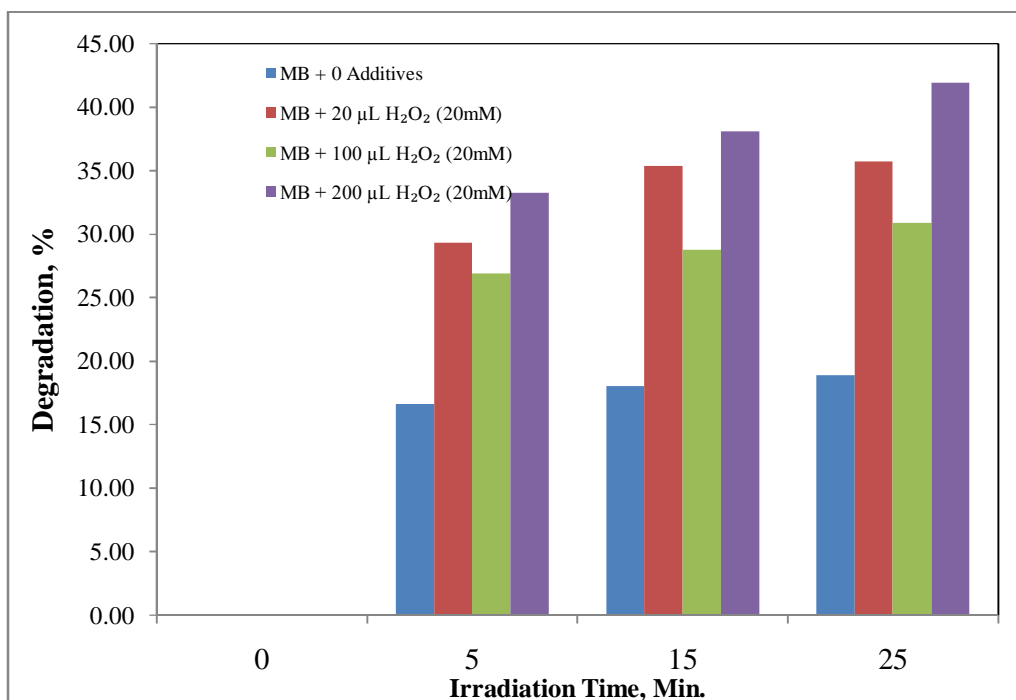


Fig. 5b. Efficiency of H₂O₂ (20mM) for the degradation of MB

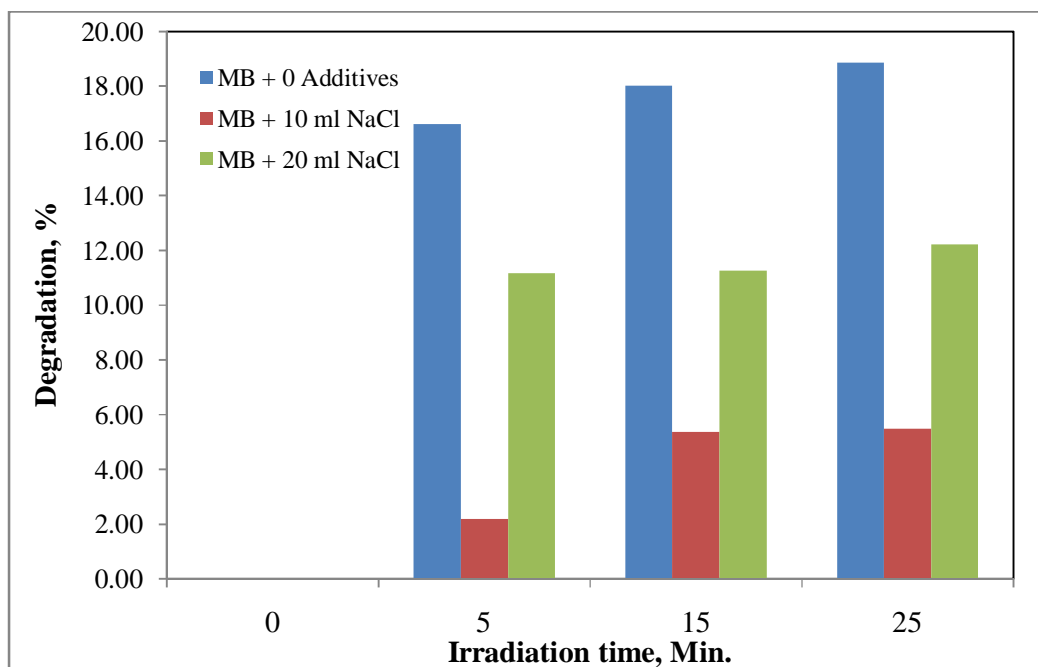


Fig. 6. Effect of NaCl on the degradation of MB

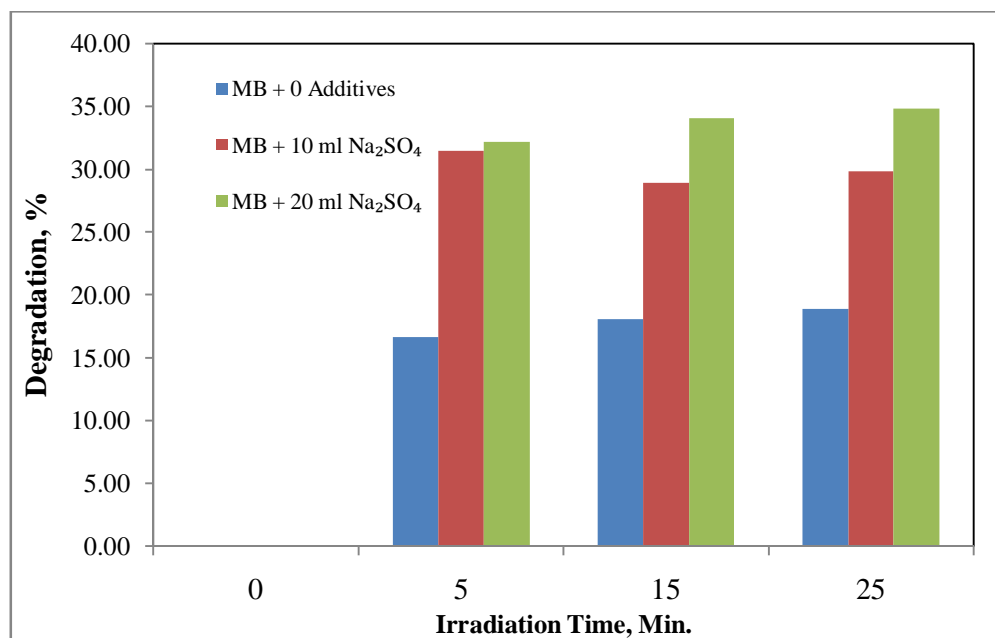


Fig. 7. Effect of Na₂SO₄ on the degradation of MB

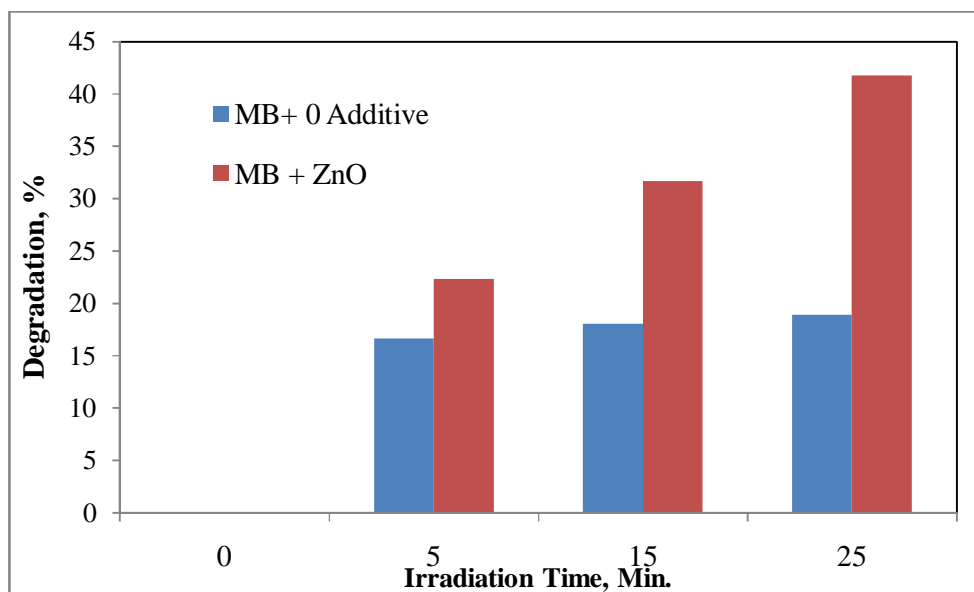


Fig. 8. Efficiency of ZnO on the degradation of MB

Effect of ZnO nanoparticle on MB degradation: Figure 8 displays the ultrasonic degradation efficiency of ZnO nanoparticle for MB. The efficiency of degradation is upto 41.73% after 25 minutes sonication. Due to advantageous spectrum absorption, porous nature and crystalline structure ZnO nanoparticle increases the degradation efficiency by producing OH radical. Maedeh Asgharian et al. observed the maximum efficiency to degrade MB in presence of rGO/ZnO/Cu compound in presence of 25 mg photocatalyst doses (Bafana et al. 2019). Kamaraj et al. showed that 99% BPA degradation caused by ZnO nanoparticles when exposed to sunshine (Kamaraj et al. 2020). Barik et al. was also showed the increased degradation efficiency for 2, 4-DCP by ZnO nanoparticle (Barik and Gogate 2017). ZnO nanoparticle is also showed one of the highest degradation efficiency of MB in this work.

4. CONCLUSIONS

The ultrasonic degradation efficiency of CCl_4 , H_2O_2 , NaCl, Na_2SO_4 and ZnO nanoparticle for MB degradation was studied at different pH, doses and concentration. The efficiency of degradation was higher at low pH compared to higher pH. 400 $\mu\text{M/L}$ dose of CCl_4 is more efficient to degrade the MB compared to other doses. On the other hand highest degradation efficiency obtained by H_2O_2 at 200 μL doses with 20 mM concentration. H_2O_2 also showed the highest degradation efficiency than the other

additives used in this study. Furthermore, Na_2SO_4 increases the degradation percentages of MB. But NaCl decreases the degradation percentages in compared to the degradation occurred for pure MB. ZnO nanoparticle enhanced the degradation efficiency considerably. Therefore, all these additives and nanoparticles except NaCl are appropriate for sonolytic degradation of methylene blue in water.

DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc.) and text-to-image generators have been used during the writing or editing of this manuscript.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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