RESEARCH ARTICLE

REVISED Synthesis of bismuth oxyhalide ($\text{BiOBr}_z\text{I}_{(1-z)}$) solid solutions for photodegradation of methylene blue dye

[version 2; peer review: 3 approved]

Previously titled: 'Synthesis of bismuth oxyhalide ($\text{BiOBr}_z\text{I}_{(1-z)}$) solid solutions for photodegradation of methylene dye'

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V2 First published: 23 Aug 2021, 4:43
<https://doi.org/10.12688/aasopenres.13249.1>

Latest published: 04 Feb 2022, 4:43
<https://doi.org/10.12688/aasopenres.13249.2>

Abstract

Background: The removal of textile wastes is a priority due to their mutagenic and carcinogenic properties. In this study, bismuth oxyhalide was used in the removal of methylene blue (MB) which is a textile waste. The main objective of this study was to develop and investigate the applicability of a bismuth oxyhalide ($\text{BiOBr}_z\text{I}_{(1-z)}$) solid solutions in the photodegradation of MB under solar and ultraviolet (UV) light irradiation.

Methods: Bismuth oxyhalide ($\text{BiOBr}_z\text{I}_{(1-z)}$) ($0 \leq z \leq 1$) materials were successfully prepared through the hydrothermal method. Brunauer-Emmett-Teller (BET), transmission electron microscope (TEM), X-ray diffractometer (XRD), and scanning electron microscope (SEM) were used to determine the surface area, microstructure, crystal structure, and morphology of the resultant products. The photocatalytic performance of $\text{BiOBr}_z\text{I}_{(1-z)}$ materials was examined through methylene blue (MB) degradation under UV light and solar irradiation.

Results: The XRD showed that $\text{BiOBr}_z\text{I}_{(1-z)}$ materials crystallized into a tetragonal crystal structure with (102) peak slightly shifting to lower diffraction angle with an increase in the amount of iodide (I^-). $\text{BiOBr}_{0.6}\text{I}_{0.4}$ materials showed a point of zero charge of 5.29 and presented the highest photocatalytic activity in the removal of MB with 99% and 88% efficiency under solar and UV irradiation, respectively. The kinetics studies of MB removal by $\text{BiOBr}_z\text{I}_{(1-z)}$ materials showed that the degradation process followed nonlinear pseudo-first-order model indicating that the removal of MB depends on the population of the

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version 1 23 Aug 2021	 view	 view	 view

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adsorption sites. Trapping experiments confirmed that photogenerated holes (h^+) and superoxide radicals ($\cdot O_2^-$) are the key species responsible for the degradation of MB.

Conclusions: This study shows that bismuth oxyhalide materials are very active in the degradation of methylene blue dye using sunlight and thus they have great potential in safeguarding public health and the environment from the dye's degradation standpoint. Moreover, the experimental results agree with nonlinear fitting.

Keywords

Hydrothermal, Flower-like, Bandgap, Thermal effect, Solar, Photocatalysis, Ultraviolet, Superoxide, Photogenerated holes

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Competing interests: No competing interests were disclosed.

Grant information: This work was supported by Botswana International University of Science and Technology and the department of Chemical and Forensic sciences [initiation grant project code S00130]. Cecil K. King'ondou and Anita Etale are grantees of the Future Leaders – African Independent Research (FLAIR) scheme from the African Academy of Sciences.

The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

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How to cite this article: Gembo RO, Aoyi O, Majoni S *et al.* **Synthesis of bismuth oxyhalide ($BiOBr_zI_{(1-z)}$) solid solutions for photodegradation of methylene blue dye [version 2; peer review: 3 approved]** AAS Open Research 2022, 4:43 <https://doi.org/10.12688/aasopenres.13249.2>

First published: 23 Aug 2021, 4:43 <https://doi.org/10.12688/aasopenres.13249.1>

REVISED Amendments from Version 1

There were no major changes made on the new version of this manuscript. However, a few changes that were suggested by the Peer reviewers were incorporated on the manuscript. Firstly, The title of the manuscript has been revised to capture methylene blue for clarity to the reader(s). Secondly, the degradation product(s) was not analyzed due the unavailability of LC-MC at the University. Nevertheless, the degradation process was monitored through the color reduction and the deduction was made out of that. Thirdly, the effects of methylene blue dye on human and animals has been clearly stated in the new version of the manuscript.

The photocatalyst was not tried on the real samples because there is no textile industry around the university where the study was carried out, therefore the authors resorted on synthetic textile waste. In conclusion, the new version has included the work done by Ren *et al.* on "Synthesis of the bismuth oxyhalide solid solutions with tunable band gap and photocatalytic activities" which was not included in the previous version. Finally, the location of Fig. S1, is indicated and labelling of the X-axis of Figure 6.

Any further responses from the reviewers can be found at the end of the article

Introduction

The availability of clean water is key for human and environmental health. The rise in demand for dyed commodities such as plastics, and textiles has led to an increase in the discharge of organic dyes into the environment and the deterioration of water quality. Approximately 17–20%^{1,2} of water pollution is from the dyeing and textile industries. About 7×10^5 tons^{3,4} of organic dyes are used to manufacture 3×10^7 tons of textiles annually⁵. Textile dyeing also consumes large volumes of water. However, approximately 30%, (2×10^5 tons)³, of dyes are lost as waste during the manufacturing process thereby finding their way into the environment and water bodies. Therefore, much attention has been directed towards developing efficient treatment techniques for these organic dyes in wastewater in the recent past. These techniques include conventional ones such as biological, physical, and chemical treatment⁴.

However, several limitations associated with these methods have been reported. First, treated effluent often does not meet the set standards for parameters such as color and chemical oxygen demand (COD) due to the limited effectiveness of these methods in breaking down the dyes. Second, the wastewater often contains both inorganic and organic compounds, making it hard to treat by conventional methods^{5–8}. Therefore, there is a need to employ a method that is effective, less costly, and environmentally friendly. Oxidative processes such as photocatalysis, the Fenton method, photolysis, sonolysis, sonocatalysis, sonoFenton, photo-Fenton, and ozonolysis^{5,9–17} have recently been explored. Among these, photocatalysis, which depends on *in-situ* photogenerated positively charged holes (h^+), hydroxyl radicals ($\cdot OH$), negatively charged electrons (e^-), superoxide radicals ($\cdot O_2^-$) has been demonstrated to be promising in terms of cost, toxicity, recyclability, mild reaction conditions, ease of operation, efficiency, and high

degradation ability^{18–23}. Photocatalysis is also a better option because it could potentially mineralize bio-recalcitrant compounds to produce carbon dioxide, water, and other inorganic substances hence no waste is left for secondary disposal⁸. Various semiconductor photocatalysts such as metal oxides and sulfides have been widely probed and applied in environmental remediation.

Traditionally, TiO_2 has been known to be effective in the treatment of wastewaters. However, titanium is costly and not readily available. Furthermore, TiO_2 only absorbs in the ultra-violet (UV) region, which is about 3–5% of the solar spectrum due to its wide bandgap of 3.2 eV^{17,19,22}. Instability and photo-corrosion are also drawbacks associated with metal-oxide semiconductors. Much of the inexpensive and abundant visible radiation energy is not harnessed in wastewater treatment with these types of metal oxide photocatalysts because they are only active within the UV region which constitutes a very small fraction of the solar insolation^{18,24–27}. Metal sulfides like CoS_2 , In_2S_3 , CdS , and Sb_2S_3 have been studied and found to have a proper location of conduction valence bands and high sensitivity to visible light, however, they are costly, prone to photo-corrosion, and the heavy metals involved are toxic^{2,19}.

To overcome these drawbacks, it is imperative to develop alternative photocatalytic materials that are more stable but less costly. Bismuth oxyhalide ($BiOZ$ ($Z = Cl, Br, I$)) materials are a novel class of photocatalyst due to their environmental friendliness, outstanding photocatalytic performance under both solar and UV irradiation, unique optical and electrical properties, unique layered crystal structure, high oxidation capacity, good chemical stability, and internal electric field effect (IEE)^{19,23,28–30}. $BiOZ$ compounds are tetragonal matlockite (space group $p4/nmm$) which consist of $(Bi_2O_2)^{2+}$ layer interleaved (Z)² halogen ions ($Z = Cl, Br, I$) slabs which crystallize into a layered structure, forming $[Z-Bi-O-O-Bi-Z]$ stacked slices. These slices interact with halogen ions along the *c*-axis via the Weak Van der Waals forces. Due to the layered structure, bismuth oxyhalide compounds, display unique optical, mechanical, and electrical qualities and have found application in fields including photocatalysis, organic synthesis, nitrogen fixation, and solar-driven H_2 generation. Furthermore, the electric field that forms between $(Bi_2O_2)^{2+}$ and $2Z^-$ layers enhance the separation of photoexcited e^- and h^+ , thus improving the photocatalytic activity^{30–32}. Bismuth oxyhalides have been studied as photocatalysts, interfaced with other photocatalytically active materials, and quaternary alloys³¹.

The yellow colour of $BiOBr$ and the coral red colour of $BiOI$ indicate that they strongly absorb light within the visible range, however, their performance is still low⁸. Therefore, improving the photocatalytic abilities of $BiOZ$ compounds is necessary for practical applications. Several approaches, for instance, doping with metals and/or non-metals, compositing $BiOZ$ with other materials, for example, TiO_2 , Ag , $AgCl$, $AgBr$, WO_3 , and $AgPO_4$, noble metals deposition, synthesis of different heterojunctions^{20–22} have been shown to enhance the activity of these $BiOZ$ photocatalysts in dye degradation.

Zhang *et al.*, synthesized nitrogen-doped graphene quantum dots/BiOZ (Z = Br, Cl) for the removal of rhodamine B (RhB) dye¹⁴. Lee *et al.*, prepared AgX (X = Cl, Br, I)/BiOZ for the removal of methyl orange, RhB, and MB¹⁵. Qu *et al.*, reported the treatment of methyl orange with TiO₂/CQDs/BiOZ (Z = Cl, Br, I) heterostructure. It has been shown that the preparation of solid solutions generally enhances the activity of a catalyst.

The development of solid solution has been acknowledged to change the bandgap, crystal, and electronic structures of a photocatalyst which can impact their photocatalytic properties¹⁶. Solid solutions such as BiOI_zBr_{1-z}, BiOCl_zI_{1-z}, zBiOBr-(1-z)BiOI, BiOBr_zI_{1-z}, BiOM_zR_{1-z} (M, R = Cl, Br, I), and BiO(ClBr)_{1-z}/2I_z have been synthesized and have shown improved photocatalytic activity when compared to corresponding single components^{19,23,29,31,33}. These materials have been utilized in the remediation of wastewater containing different organic dyes. Zhang *et al.*,¹⁶ fabricated BiOBr_zI_{1-z} nanoplates for the removal of RhB. Gnyem *et al.*,²⁰ prepared hierarchical nanostructured 3D flowerlike BiOCl_zBr_{1-z} for RhB degradation under visible light irradiation. Zhang *et al.*,²¹ synthesized BiOCl_zBr_{1-z} nanoplate solid solutions for the removal of RhB under the irradiation from visible light. Zhang *et al.*,²² synthesized BiOBr_zI_{1-z} solid solutions for the removal of various dyes from textile wastes.

However, to the best of the authors' knowledge, the application of BiOBr_zI_{1-z} solid solutions in the treatment of methylene blue (MB) dye and comparative studies on the photocatalytic performance of these materials under natural sunlight and UV light has not been reported. Additionally, no report on the comparison of the experimental kinetic data to both linear and non-linear fitting using Langmuir-Hinshelwood (L-H) kinetic model. Methylene blue dye is extensively used in printing and textile dyeing industries due to its intense blue colour and thus a common pollutant in industrial wastewaters.

Methylene blue dye if not monitored can adversely affect human beings and animals by causing eye burns which is a permanent health damage. Additionally, the MB can cause painful micturition, nausea, increase in heart rate, quadriplegia disorders in breathing, quadriplegia, cyanosis, mental confusion, jaundice, vomiting, tissue necrosis, and methemoglobinemia³⁴. The purpose of the present study is to enhance the absorption of visible light by BiOZ. Thus, for the improved visible-light uptake, the BiOBr_zI_{1-z} solid solutions with a suitable bandgap were prepared by doping the bromide and the iodide ions through varying the amounts of the dopants from 0 to 1 in the compound. The photocatalytic activity of BiOBr_zI_{1-z} solid solutions was evaluated by determining their efficiency in removing MB under solar and UV light exposure. Furthermore, the efficiency of the most active materials was optimized by careful adjustment of catalyst dosage, temperature, and pH, while the degradation pathway was examined by use of different scavengers.

Methods

Duration of methodology

The preparation of the photocatalysts was done between Oct–Nov 2018. After the preparation, the degradation efficacy

of the prepared samples was tested before characterization. This was carried out in Jan 2019. The XRD, SEM, TEM, BET, Raman characterization was carried out between Feb–Dec 2019. The photodegradation studies were carried out throughout 2019 and between Jan–May 2020. The XRD and Raman analysis was carried out at the Botswana International University of Science and Technology (BIUST). The SEM characterization was done at Botswana Institute for Technology Research and Innovation (BITRI), while TEM and BET analysis was done at Global Change Institute, University of the Witwatersrand, South Africa. No adjustment was done to the scanning electron microscope micrographs. All the degradation performance was carried out at BIUST. OriginPro 8.5.0 SR1 b161 software was used for data analysis and graphing. Alternatively, R and excel can be applied for analysis.

Chemicals and materials

Analytical standard bismuth nitrate (Bi(NO₃)₃·5H₂O, Cat No. 10035-06-0), potassium iodide (KI, Cat No. 290813PQ), sodium bromide (NaBr, Cat No. 27656), acetic acid (CH₃COOH, Cat No. AA539), ethylenediaminetetraacetic acid disodium (EDTA, Cat No. 1046725) salt, sodium nitrate (NaNO₃, Cat No. 030311SN), sodium hydroxide (Cat No. 051214SO), silver nitrate (Cat No. 111214SN), benzoquinone (Cat No. 606-013-00-3), tert-butanol (Cat No. 603-005-1), nitric acid (Cat No. 110815NC) and methylene blue (MB, Cat No. MB057) reagents used in this work were acquired from Sigma-Aldrich. The dye working solutions used in this investigation were made by adding a 10 mg methylene blue in 1000 mL of distilled water (DI) to make the desired concentrations.

Synthesis of BiOBr_zI_{1-z} solid solution

Bismuth oxyhalide (BiOBr_zI_{1-z}) solid solutions were synthesized via hydrothermal method at 160 °C. Typically, a 0.4123 mM Bi(NO₃)₃·5H₂O solution was prepared in CH₃COOH. A solution mixture having stoichiometric amounts of KI and NaBr was then drop wisely added into the above Bi-based solution while continuously stirring. The resultant mixture was then put into 23 mL stainless steel autoclave lined with teflon (4749 PTFE A280AC Teflon, Parr Instrument Company) and heated at 160 °C in an oven, for 24 h. When the reaction was complete, the sample was naturally cooled to room temperature, centrifuged (Heraeus Megafuge 40 centrifuge) 6 times at 6000 rpm to effectively separate the product, and washed 6 times with deionized. The product obtained was then dried at 70 °C for 24 h and used in photodegradation reactions. The molar ratios of Br and I was varied between 0 and 1 with Z = 0.0, 0.2, 0.4, 0.6, 0.8, 1.0 for BiOBr_zI_{1-z} materials and samples prepared hereafter labelled BiOI, BiOBr_{0.2}I_{0.8}, BiOBr_{0.4}I_{0.6}, BiOBr_{0.6}I_{0.4}, BiOBr_{0.8}I_{0.2}, and BiOBr, respectively.

Point of zero charge (pH_{pzc}) determination

The procedure used in this work was adopted from Tahira *et al.*,²³. The PZC values for BiOBr_zI_{1-z} samples were determined in 0.1 M solution of NaNO₃ at 298 K. Typically, 0.1 g of the samples was dispersed in NaNO₃ solution (0.1 M, 30 mL) in various reaction flasks. Adjusting the initial pH of the mixtures to 2, 3, 4, 5, 6, 7, 8, 9, 10, and 11 was done using 0.1 M HNO₃ and NaOH. Each reaction vessel was then agitated

at 130 rpm in a shaker (Stuart orbital shaker SSL1) for 24 h. The final pH of the mixtures was determined by a Basic20 pH-meter (Crisson). A graph of the difference between final and initial pH (ΔpH) values was then plotted against the initial pH values. The pH_{PZC} was taken to be the initial pH at which ΔpH is 0.

Characterization

The crystal structure of the synthesized materials was obtained by X-ray diffraction (XRD) at room temperature by a Bruker D8 Advance powder diffractometer with a Cu tube X-ray source and a LynxEye XE-T energy-dispersive strip detector. The radiation used was a Cu-K α ($\lambda = 1.54056$ nm) with increasing current and voltage of 40 kV and 40 mA, correspondingly. The patterns were acquired at a 2θ scan rate of steps 0.02 degrees with a time of 0.500 sec/step from 5 to 90°. Raman spectra were acquired by LabRAM HR800 Raman spectrophotometer, exciting the samples with 532 nm laser. FEI Tecnai G2 Spirit transmission electron microscope at an acceleration voltage of 120 kV and scanning electron microscope (SEM, Gemini SEM 500, Carl Zeiss, 5.00 KX magnification) with energy-dispersive X-ray detector (EDX) were applied to obtain the morphology and the elemental composition of the $\text{BiOBr}_{z(1-z)}$ solid solutions. Porosity and surface area investigations were performed by Brunauer-Emmett-Teller technique (Micro metrics Tristar 3000 porosity analyzer). The samples were degassed at 150°C for 5 h before sample analysis at liquid nitrogen temperature (77.350 K). Ultraviolet-visible (UV-Vis) spectrophotometer (UV201, Shimadzu) was applied to study the photodegradation of MB.

Photodegradation studies

To test for the photodegradation ability of the $\text{BiOBr}_{z(1-z)}$ solid solutions, 50 mg of the material was added in 50 mL of 10 mg^{-1} dye solution. The equilibrium (adsorption-desorption)

between the photocatalyst and the solution containing the dye was attained through the stirring of the mixture in the dark for 30 min, thereafter the photodegradation was carried out under solar irradiation and UV lamp operating at 0.16 Amps with a UV output of 254 nm (UVP UVG-54). The set up for the photodegradation under UV lamp and solar is shown in Figure S1 (Extended data³⁵). During the reaction process, 2 mL of aliquot was drawn at 30-minute intervals from the mixture to monitor the degradation of MB. The separation of the photocatalyst and the dye solution was achieved through centrifuging the aliquot for 7 min at 13000 rpm. The concentration was monitored using UV-Vis spectrophotometer within a 200–800 nm range.

The absorption of MB was examined at a maximum wavelength of ~ 661 nm, according to the Beer-Lambert law. An experiment without a sample (blank) was also carried out under UV and solar irradiation as a control. The same procedure was followed for the two conditions (solar and UV light). The active species responsible for the photodegradation process was investigated by conducting scavenging experiments. Tert-butanol (TBA), EDTA, silver nitrate, (AgNO_3), and P-benzoquinone (BQ) were applied to quench $\cdot\text{OH}$, h^+ , e^- , and $\cdot\text{O}_2^-$, respectively. The concentration of the quenchers used was 1 mM.

Results and discussion

X-ray diffraction

Figure 1 displays the XRD spectra of BiOI, $\text{BiOBr}_{0.2\text{I}_{0.8}}$, $\text{BiOBr}_{0.4\text{I}_{0.6}}$, $\text{BiOBr}_{0.6\text{I}_{0.4}}$, $\text{BiOBr}_{0.8\text{I}_{0.2}}$, and BiOBr.

It was observed that all the diffraction peaks of the spectra are sharp showing an efficacious crystallization of $\text{BiOBr}_{z(1-z)}$ materials synthesized via the hydrothermal method. The pure $\text{BiOBr}_{z(1-z)}$ materials can be attributed to tetragonal structure

Figure 1. X-ray diffraction (XRD) patterns of $\text{BiOBr}_{z(1-z)}$ materials.

BiOBr (PDF Card No. 04-002-3609) and BiOI (PDF Card No. 00-10-0445). Even though the composition of the materials is different, they possess a similar tetragonal phase. The BiOBr peaks at $2\theta = 11.0, 31.8,$ and 32.3° are assigned to the hkl values (001), (102), and (110), respectively. While the BiOI peaks at $2\theta = 29.6$ and 31.7° can be respectively assigned to the hkl values (102) and (110).

There is an observable shift in the diffraction peaks to lower diffraction angles with an increase of I content in $\text{BiOBrI}_{z(1-z)}$ compound. This observation was also reported by Xu *et al.*, and Ren *et al.*,^{19,33} who suggested that the shift is a result of bromide ions substitution (ionic radius of 0.196 nm) with the larger iodide ions (ionic radius of 0.216 nm) that leads to an expansion of the interlayer spacing. The percentage difference between the ionic radii of bromide and iodide ion is 9%, a value less than the maximum of 15% for substitution to occur. Additionally, the unlimited formation of solid solutions between BiOI and BiOBr can further be shown by the shift in diffraction peaks to lower angles as the amount of iodine increases²⁵. Cell parameters also increased gradually with an increase in I content (Extended data, Table S1³⁵)¹⁸. Figure 1 shows a perfect change from BiOBr to BiOI in the $\text{BiOBrI}_{z(1-z)}$ pattern, this has been observed before by Lei *et al.*,²⁶. There are no peaks attributed to impurities, indicating that the samples consisted purely of the desired phases.

Raman spectroscopy

To further understand the structure of the as-prepared materials, the Raman technique was used. Figure S2 (Extended data³⁵) represents the Raman spectra of $\text{BiOBrI}_{z(1-z)}$ solid solutions prepared by varying amounts of Br and I ions. To show a variation in structural features of $\text{BiOBrI}_{z(1-z)}$ solid solutions, the Raman spectra were recorded from $25 - 500 \text{ cm}^{-1}$. Bismuth oxyhalide belongs to the tetragonal PbFCl type structure with space of $p4/nm$, hence the active modes of the Raman exhibited are A_{1g} , E_g , and B_{1g} . BiOBr samples presented two major Raman peaks at 45 and 106 cm^{-1} allotted to the first-order vibration of Bi-O and A_{1g} internal Bi-Br stretching mode, respectively²⁷. The stronger bands of BiOI at 78 and 145 cm^{-1} is allotted to the A_{1g} stretching mode from internal Bi-I bonds. It can also be observed that the peak at 45 cm^{-1} diminishes while the peak 106 cm^{-1} changes to a lower value of 78 cm^{-1} as the amount of I^- ions increases. The shifting of the peaks to the lower Raman value agrees with the XRD results, which also indicates that as the amount of I^- increases the (102) plane shifts to the lower angles.

Morphological analysis by SEM and TEM

The examination of surface morphologies of $\text{BiOBrI}_{z(1-z)}$ solid solutions was done using SEM, as shown in Figure 2(a-f). The samples display plate-like morphology with interleaved nanoplates of varying sizes. The formation of the plate-like

Figure 2. SEM micrographs of (a) BiOBr, (b) $\text{BiOBr}_{0.8}\text{I}_{0.2}$, (c) $\text{BiOBr}_{0.6}\text{I}_{0.4}$, (d) $\text{BiOBr}_{0.4}\text{I}_{0.6}$, (e) $\text{BiOBr}_{0.2}\text{I}_{0.8}$, (f) BiOI. TEM images of (g) BiOBr, (h) $\text{BiOBr}_{0.6}\text{I}_{0.4}$, and (i) BiOI. No modifications have been made to the images apart from adding the scales.

morphology can be attributed to the internal structure of bismuth oxyhalide (BiOZ , $Z = \text{Cl}, \text{Br}, \text{I}$) where $(\text{Bi}_2\text{O}_2)^{2+}$ layers sandwich between the two slabs of halogen atoms resulting in the anisotropic growth of BiOZ at a c axis forming 2D structures. As shown in Figure 2a, pure BiOBr formed relatively large plates which upon iodine doping reduced in size and self-assembled in 3D flowers, Figure 2b,c. As the amount of bromide decreased and iodine increased towards pure BiOI , the size of the flakes increased forming relatively large 2D structures akin to those of pure BiOBr . TEM images were obtained to further understand the BiOBr , $\text{BiOBr}_{0.6}\text{I}_{0.4}$, and BiOI SEM morphology. The TEM micrographs, Figure 2g, h, i, confirm the plate-like morphology observed by SEM for all the $\text{BiOBr}_z\text{I}_{(1-z)}$ materials. The presence of Br, Bi, O, and I was confirmed by EDX examination. The amounts of the elements detected agree with the $\text{BiOBr}_z\text{I}_{(1-z)}$ formula, (Extended data, Figure S3³⁵).

Surface area analysis

The textural properties of BiOBr , $\text{BiOBr}_{0.6}\text{I}_{0.4}$, and BiOI materials were obtained using N_2 adsorption-desorption analysis. The isotherms displayed in Figure S4a (Extended data³⁵) can be allotted to type III based on IUPAC classification. This type of isotherm has no recognizable monolayer formed due to the relatively weak interaction between the adsorbent and the adsorbate hence leading to a clustering of adsorbed molecules around the most favourable sites on the surface of a macroporous or nonporous solid. Compared to type II isotherms, the quantity of molecules adsorbed remains finite at $p/p_0 = 1$ which is the saturation pressure²⁸. The BET surface areas of the as-prepared materials were 0.517, 3.249, and 1.890 m^2/g for BiOBr , $\text{BiOBr}_{0.6}\text{I}_{0.4}$, and BiOI , respectively. The BiOBr sample has a larger and thicker microplate, which has a much lower surface area. Figure S4b (Extended data³⁵) shows Barret-Joyner-Halenda (BJH) plots of BiOBr , $\text{BiOBr}_{0.6}\text{I}_{0.4}$,

and BiOI materials obtained from desorption isotherms. The BJH plots confirmed that the corresponding macropores peaks of the pore-diameters distributions of BiOBr , $\text{BiOBr}_{0.6}\text{I}_{0.4}$, and BiOI could be found up to 159, 170, and 153 nm, respectively. The BJH result shows that the pore diameter distributions of the materials are wide. This can be ascribed to the inter-crossing of the spaces between the microplate structure²⁹.

pH_{PZC} and optical properties

The results are displayed in Figure S5 of the Extended data³⁵. The PZC (the pH at which the surface charge of a material is 0) of BiOBr , $\text{BiOBr}_{0.8}\text{I}_{0.2}$, $\text{BiOBr}_{0.6}\text{I}_{0.4}$, $\text{BiOBr}_{0.4}\text{I}_{0.6}$, $\text{BiOBr}_{0.2}\text{I}_{0.8}$, and BiOI materials was found to be 5.27, 5.27, 5.29, 5.31, 5.38, and 5.39, respectively. At a pH lower than the PZC, the material will be positively charged whereas at a pH higher than PZC the materials acquire a negative charge³⁰. The photocatalytic performance of a semiconductor depends on the band structure of the photocatalyst. Figure 4 shows the UV/Vis spectra of the as-synthesized $\text{BiOBr}_z\text{I}_{(1-z)}$ materials.

When the composition of iodine is increased from 0 to 1, there is a significant redshift of the absorption edges from 478 to 632 nm for pure BiOBr and BiOI , respectively. Whereas the absorption edges of $\text{BiOBr}_z\text{I}_{(1-z)}$ materials whose composition lie between that of BiOBr and BiOI were found to be 546, 577, 591, and 611 nm for $Z = 0.8, 0.6, 0.4$, and 0.2 , respectively. The change is consistent with the colour of the material, from white to coral-red (Figure 3b). The spectra revealed that the absorption of BiOBr is slightly within the visible region and the absorption edge of BiOI extends towards the visible region indicating that it is highly responsive within visible light. The bandgap energy (E_g) values of the obtained $\text{BiOBr}_z\text{I}_{(1-z)}$ materials are determined using Equation 1 below:

$$\alpha h\nu = A(h\nu - E_g)n/2 \quad (1)$$

Figure 3. (a) Ultraviolet-visible absorption spectra for the $\text{BiOBr}_z\text{I}_{(1-z)}$ solid solutions (b) optical images of $\text{BiOBr}_z\text{I}_{(1-z)}$ suspension in methanol.

Figure 4. Photodegradation efficiencies under (a) solar and (b) ultraviolet (UV) light after 3 h of irradiation.

where α is absorption coefficient, h is Planck's constant, ν is frequency of light, A is absorbance, and E_g is bandgap energy⁸, and n is dependent on the transition characteristics of the semiconductor. For BiOZ, n is 4 owing to its indirect change. The corresponding bandgap energies of BiOBrI_{z(1-z)} materials were calculated and the results are shown in Table 1. The bandgap energy of the materials could be tailored from 2.59 to 1.96 eV by reducing the value of Z from 1 to 0 as shown in Table 1, demonstrating that doping I atoms into BiOBr crystal reduced the bandgap and increased the absorption range of BiOBr. This improves the photocatalytic performance of BiOZ.

The Mulliken electronegativity theory was used in calculating the conduction band (CB) and valence band (VB) electric potentials using Equation 2 and Equation 3:

$$E_{CB} = \chi - E^c - \frac{1}{2}E_g \quad (2)$$

$$E_{VB} = E_{CB} + E_g \quad (3)$$

where E_{CB} and E_{VB} are the electric potential edges, respectively. E^c is the free electrons energy on the hydrogen scale (4.5 eV), E_g is semiconductor's bandgap energy. While χ is the semiconductor electronegativity, equivalent to geometric mean electronegativity of atoms forming the compound.

From the calculations obtained from the UV-Vis results, the CB and VB of BiOBrI_{z(1-z)} materials were approximated and the results are presented in Table 1. The gradual decrease in VB edge potential from 2.98 to 2.42 with increasing iodine composition indicates a weak oxidation ability and a stronger light absorption capability³¹. The generation of $\cdot O_2^-$ radicals depends on the CB edge potential, the more positive the CB, the more difficult it is to generate $\cdot O_2^-$ radicals. Moreover, increased iodide concentration leads to a reduction in the level of VB. This in turn reduces the bandgap energy thereby

facilitating photosensitization of the catalyst. Therefore, the highest photocatalytic performance of BiOBr_{0.6}I_{0.4} material; the most active material, is ascribed to the suitable bandgap structure that may attain the equilibrium between the light absorption capacity and redox power.

Photodegradation studies

Adsorption of MB by BiOBrI_{z(1-z)}. The adsorption ability of a photocatalyst is well-known to play an important role in the photodegradation process. Figure S6 in the Extended data³⁵ displays the MB adsorption uptakes by BiOBrI_{z(1-z)} materials at constant concentration and catalyst dosage. The adsorption uptake values for the MB dye by BiOBr, BiOBr_{0.8}I_{0.2}, BiOBr_{0.6}I_{0.4}, BiOBr_{0.4}I_{0.6}, BiOBr_{0.2}I_{0.8}, and BiOI were 12, 17, 19, 16, 15, and 14%, respectively. The adsorption efficiency was found to be in the order BiOBrI_{0.6}I_{0.4} > BiOBr_{0.8}I_{0.2} > BiOBr_{0.4}I_{0.6} > BiOBr_{0.2}I_{0.8} > BiOI > BiOBr. Fitting of the adsorption data to adsorption isotherms using nonlinear model fitting procedures (Figure S7, Extended data³⁵) indicated that the process of adsorption was according to Langmuir. The Langmuir constant and the maximum adsorption capacity of BiOBr_{0.6}I_{0.4} were found to be 6.10×10^{-2} L/mg and 31.5 mg/g, respectively. This implies that at the highest concentration used in this study, the adsorption sites were not saturated with the dye.

Kinetics, scavenging experiments and recyclability. To broadly understand the photocatalytic ability of the as-prepared materials, the removal of MB was performed under irradiation from solar and UV light in the presence and absence of the photocatalyst materials. Stirring of photocatalyst and dye mixture was carried out in the dark for half an hour before illumination to attain adsorption-desorption equilibrium (Figure S6, Extended data³⁵). Figure 4a, b shows the removal of MB under the irradiation from solar and UV light, respectively. The degradation in the absence of the photocatalyst (photolysis) both under the solar and UV light showed a negligible change

Table 1. Absorption thresholds, electronegativity, bandgap energy, the conduction band (CB) and valence band (VB) edges of as-prepared BiOBr_zI_{1-z} materials.

Parameters	BiOBr _z I _{1-z} materials					
	Z = 1.0	Z = 0.8	Z = 0.6	Z = 0.4	Z = 0.2	Z = 0.0
Absorption thresholds (nm)	478	546	577	591	611	632
Electronegativity (χ) (eV)	6.18	6.14	6.08	6.04	5.99	5.94
E _g (eV)	2.59	2.27	2.15	2.10	2.03	1.96
E _{CB} (eV)	0.39	0.51	0.51	0.49	0.48	0.46
E _{VB} (eV)	2.98	2.78	2.66	2.59	2.51	2.42

indicating that MB is stable under both light sources hence photolysis can be neglected. Therefore, the removal of MB can be attributed to solar and UV light in the presence of the photocatalysts. The photodegradation efficiency was calculated using Equation 4 below:

$$\text{Degradation efficiency, } \mu = \frac{C_0 - C_t}{C_0} \times 100\% \quad (4)$$

where C_0 is the concentration of MB dye at equilibrium, C_t the concentration at the time, interval, t . Table S2 in the Extended data³⁵ shows the percent degradation of MB after 3 h under solar and UV-light irradiation using BiOBr, BiOBr_{0.8}I_{0.2}, BiOBr_{0.6}I_{0.4}, BiOBr_{0.4}I_{0.6}, BiOBr_{0.2}I_{0.8}, and BiOI. BiOI consistently displayed the least photocatalytic performance amongst all the materials despite absorbing at the longest wavelength of 632 nm. This could be due to a higher recombination rate of photoexcited e^- and h^+ pair⁸. Photocatalytic ability in most cases depends on the surface area, such that the lower the surface area, the lower the activity due to lower adsorption of the dye solution. However, despite their low surface areas, BiOBr_zI_{1-z} solid solutions in our work showed high photocatalytic activity revealing that the specific surface area was not the key reason for the improved photocatalytic performance of this material. The higher activity could be attributed to the electric field that built-in between (Bi₂O₃)²⁺ and 2Z⁻ layers enhancing the separation of photoexcited e^- and h^+ , thus improving the photocatalytic performance.

Importantly, although no obvious trend in the photocatalytic activity of the obtained materials was observed, doping seemed to improve the photocatalytic ability of the photocatalysts. The bandgap of BiOBr_zI_{1-z} solid solutions ranges from 1.96 to 2.59 eV which is generally low, hence requires lower energy for activation. After 3 h of irradiation both under solar and UV light, BiOBr_{0.6}I_{0.4} recorded the highest activity with degradation efficiency of 99% and 88% under solar and UV-light, respectively. The greatest photocatalytic performance of BiOBr_{0.6}I_{0.4} can be linked to its flower-like morphology that could lead to stronger absorption of light than the 2D structures. Finally, the optimal composition of BiOBr_{0.6}I_{0.4} promotes the photoactivity by minimizing the recombination of photoexcited electron-hole pair.

The model that was used to fit the experimental data obtained from UV-light irradiation was pseudo-first-order. The equation $\ln(C_0/C_t) = \kappa t$, was used to determine the pseudo-first-order rate constant. In the equation, $\ln(C_0/C_t) = \kappa t$, κ and t denote the rate constant (min^{-1}) and time of irradiation, respectively. C_0 is the initial concentration of MB dye, whereas C_t is the concentration at the time interval, t . As presented in Table S3 of the Extended data³⁵ the reaction rate constants of blank, BiOI, BiOBr_{0.2}I_{0.8}, BiOBr_{0.4}I_{0.6}, BiOBr_{0.6}I_{0.4}, BiOBr_{0.8}I_{0.2}, and BiOBr were 5.56×10^{-5} , 0.00432, 0.00846, 0.01188, 0.00696, 0.00505, 0.00192, and min^{-1} , respectively. The reaction rate constants results indicate that BiOBr_{0.6}I_{0.4} with the highest reaction rate of 0.01188 min^{-1} was approximated to be three and six times greater than BiOBr and BiOI, respectively.

The pseudo-first-order kinetic model, derived from the Langmuir-Hinshelwood (L-H) kinetic model under conditions of low dye concentration, has been used extensively to evaluate the reaction kinetics of photodegradation processes^{36,37}. Whereas the linear equation has been widely (almost exclusively) used during kinetic studies in photodegradation studies, it has been noted that the nonlinear model fitting gives more accurate results. This is highlighted in Figure 5, which shows a comparison of nonlinear and linear fitting for BiOBr and BiOBr_{0.6}I_{0.4} in which nonlinear model-fitting has a lower sum squared error (SSE) values compared to linear fitting. As the deviation from L-H kinetic model increases, the differences in parameters obtained from nonlinear and linear model fitting becomes significant as shown in Table 2 in which the differences in the obtained rate constants are 1.8% for BiOBr and 10% for BiOBr_{0.6}I_{0.4}. Results for nonlinear fitting of BiOBr_zI_{1-z} materials are shown in Figure S7 of the Extended data³⁵ and summarised in Table 2.

The main active species involved during photocatalysis are known to be $O_2^{\cdot-}$, e^- , h^+ , and $HO^{\cdot}$. However, the main active species during the degradation of MB was established through the quenching experiment over BiOBr_{0.6}I_{0.4} under the irradiation from the UV light. P-benzoquinone (BQ), silver nitrate ($AgNO_3$), EDTA-2Na, and tert-butanol (TBA) was added to the reaction vessels to quench $O_2^{\cdot-}$, e^- , h^+ , and $HO^{\cdot}$ respectively. When TBA and $AgNO_3$ was added, the removal efficiency was

Figure 5. Linear and nonlinear fitting of (a) BiOBr and (b) BiOBr_{0.6}I_{0.4} to L-H kinetic model.

Table 2. Summary of linear and nonlinear fitting parameters for BiOBr and BiOBr_{0.6}I_{0.4}.

	BiOBr			BiOBr _{0.6} I _{0.4}		
	Sum squared error (SSE)	R ²	k/min	Sum squared error (SSE)	R ²	k/min
Nonlinear	0.0012	0.9971	0.00432	0.028	0.9627	0.01188
Linear	0.04	0.9827	0.0044	0.14	0.9556	0.01322

insignificant compared to when no quencher was added as presented in Figure 6. This indicates that the HO[•] and e⁻ were not the main active particles in the photodegradation reaction system. However, when BQ and EDTA-2Na were added, there was an obvious reduction in degradation efficiency as a result of the suppression of h⁺ and O₂^{-•} which were the major species in the reaction system. Rashid *et al.*,³⁸ made similar conclusions for the photocatalytic mineralization of aqueous ciprofloxacin.

The recyclability/reusability of a material is key for its practical application. Therefore, the stability of the prepared materials was determined through the recyclability of BiOBr_{0.6}I_{0.4} for the removal of MB. The results are displayed in Figure 7. In this study, BiOBr_{0.6}I_{0.4} was reused five times. After every cycle, the catalyst was centrifuged, cleaned, and dried for recovery. The degradation efficiency was reduced to 82% from 88%. The slight drop in efficiency of BiOBr_{0.6}I_{0.4}, even after five cycles, signifies that the BiOBr_zI_(1-z) materials showed remarkable good stability.

Thermal contribution on the MB photodegradation under solar. It is observed that the photocatalytic efficiency was higher under solar irradiation compared to UV light. Unlike the

controlled conditions under which the removal of MB under irradiation from UV was done, the removal under solar was performed in open-air that may have introduced other factors that led to the efficiency enhancement. Therefore, an investigation, particularly on the thermal contribution, was done, with other factors like catalyst loading and dye initial concentration kept constant. BiOBr_{0.6}I_{0.4} being the best photocatalyst was chosen for the study. An experiment was performed with a set up having four different reaction vessels: two aluminium foil-covered vessels, one with catalyst, and another without and another two uncovered vessels with similar catalyst conditions (with/without) were set up. The temperatures of the four samples were recorded as 27 °C before exposure to sunlight. Table S3 in the Extended data³⁵ shows the changes in temperature with irradiation time.

For the covered vessels, the temperature increases steadily up to the 150th minute whereas for the uncovered vessels, the temperature increases up to the 120th minute, with no noticeable change thereafter. The covered vessels recorded the highest temperature due to the build-up of heat that could not escape as the vessels were covered. Further, MB degradation still occurred in the covered vessels, despite the absence of light, which may be attributed to the build-up of heat energy.

Figure 6. Effects of quenchers on the removal of methylene blue (MB).

Figure 7. Recyclability test for BiOBr_{0.6}I_{0.4} for the removal of 10 mgL⁻¹ MB after 3 h of light illumination at pH 7.0 with 1 gL⁻¹ catalyst dosage.

The significantly higher degradation efficiency afforded by the uncovered vessels compared to their covered counterparts despite the fact that the uncovered vessels were able to prevent heat build up by dissipating it provides a credible evidence that the MB degradation in our study was largely photo-driven. [Figure 8](#)

illustrates the photocatalytic change for MB both from covered and uncovered flasks.

Effects of temperature. Temperature is a relevant parameter since the rate of reaction is temperature dependent. The

Figure 8. Effects of thermal contribution towards MB degradation under solar irradiation.

influence of temperature on the degradation process of MB over $\text{BiOBr}_{0.6}\text{I}_{0.4}$ is shown for the temperature range 30 – 60°C in Figure S8a of the Extended data³⁵. Other parameters were kept constant. It was observed that a rise in temperature caused a gradual increase in degradation efficiency. An increase in temperature led to the enhancement in photooxidation as a result of the increased frequency of molecules collision. Furthermore, when the temperature increases, the reaction competition is increased thereby restraining electron-hole recombination^{38–40}. Figure S8b (Extended data³⁵) displays the Arrhenius plot of k versus T^{-1} where the activation energy (E_a) of $9.1 \times 10^{-1} \text{ J mol}^{-1}$ was obtained for the removal of MB. The lower activation energy implies that the removal of MB over $\text{BiOBr}_{0.6}\text{I}_{0.4}$ requires little energy hence making it economical.

Effect of amount of catalyst on MB removal. To determine the influence of the amount of catalyst on the MB removal, a dosage of $\text{BiOBr}_{0.6}\text{I}_{0.4}$ was increased from 0.01 to 0.07 g/L under optimal conditions (10 mg/L MB concentration, pH, 7.0, and 150 min reaction time at $30 \pm 2^\circ\text{C}$). The result is shown in the Extended data, Figure S9³⁵. The removal of MB was increased from 30% to 99% with an increasing amount of dosage from 0.01 g to 0.05 g. The removal was maximum at 0.05 g, which is the optimal catalyst loading. At the optimal dosage, there is maximum availability of the catalyst surface area plus active sites for the production of active radicals under UV-light irradiation. However, the catalyst dosage above 0.06 g results in a decrease in degradation since the screening effect and light scattering tend to become higher^{41–43}. Additionally, the illumination of key active primary oxidants in photocatalysis will be prevented when the catalyst dosage exceeds the optimum amount hence the degradation efficiency will reduce^{44,45}. Economically, therefore, optimum catalyst dosage is vital, as less energy would be required for regeneration.

Effect of pH on photocatalysis. The initial pH of a wastewater solution can significantly influence the process of waste treatment since the extent of MB removal is affected by the charge on the photocatalyst surface. The catalyst surface charge depends on the concentration of OH^- and H^+ in aqueous solutions⁴². To establish the role of pH on MB removal, the initial pH of the mixtures was adjusted to 3.0, 5.0, 7.0, 8.0, and 14.0 by adding 1 M HNO_3 or 1 M NaOH while other conditions were kept constant (0.05 g $\text{BiOBr}_{0.6}\text{I}_{0.4}$ loading and 10 mg/L MB concentration). When the pH values of the mixtures were lower than pH_{PZC} the surface of the catalyst acquires positive charge while if the pH is high, the material acquires negative charge. The results are shown in Figure S10 of the Extended data³⁵. The degradation efficiencies recorded were 47, 28, 86, 78, and 75% for pH values of 3.0, 5.0, 7.0, 8.0, and 14.0, correspondingly.

The high efficiencies at the pH of 7.0, 8.0, and 14.0 can be ascribed to negatively charged $\text{BiOBr}_{0.6}\text{I}_{0.4}$ ($\text{pH}_{\text{PZC}} = 5.3$) surface. Thus, if the pH of the mixtures were adjusted to the initial pH of 7.0, 8.0, and 14.0, adsorption of the dye on to the positively charged $\text{BiOBr}_{0.6}\text{I}_{0.4}$ surface will be greater due to electrostatic attractions, which enhances the adsorptive process thereby enhancing the degradation efficiencies. On the other hand, adjusting initial pH of the pollutant to 3.0 and 5.0 reduces the degradation efficiency because of the electrostatic repulsion between $\text{BiOBr}_{0.6}\text{I}_{0.4}$ ($\text{pH}_{\text{PZC}} = 5.29$) surface which has a positive charge and the cationic dye⁴³. Thus, lowering the degradation efficiency of the material.

Conclusion

The preparation of a series of $\text{BiOBr}_{z(1-z)}$ solid solutions via the hydrothermal technique was successful. The crystal structure, morphology, pore size, and surface area were obtained by XRD, SEM, TEM, and BET, respectively. The investigation

of the photocatalytic ability of the prepared $\text{BiOBrI}_{z(1-z)}$ materials was carried out through the removal of MB, additionally the optimization of the operational parameters was also performed. The XRD revealed that the (102) plane shift to smaller diffraction angles with increasing I composition. From the SEM analysis, it was observed that pure BiOBr was made of ultrathin nanoplates with various sizes whereas the morphology of $Z = 0.8, 0.6, 0.4, 0.2$, and 0.0 was 3-Dimension flower-like structures comprising of nanosheets. The BET results showed that the prepared photocatalysts had low surface areas. The surface characterization showed that the point of zero charge (PZC) of BiOBr, $\text{BiOBr}_{0.8}\text{I}_{0.2}$, $\text{BiOBr}_{0.6}\text{I}_{0.4}$, $\text{BiOBr}_{0.4}\text{I}_{0.6}$, $\text{BiOBr}_{0.2}\text{I}_{0.8}$, and BiOI was found to be 5.27, 5.27, 5.29, 5.31, 5.38, and 5.39, respectively. The percentage degradation efficiency over $\text{BiOBr}_{0.6}\text{I}_{0.4}$ under solar and UV light irradiation was 99 and 88%, respectively. The experimental data was fitted to the first-order kinetics and the R^2 obtained were 0.997, 0.970, 0.963, 0.926, 0.905, and 0.939 for BiOBr, $\text{BiOBr}_{0.8}\text{I}_{0.2}$, $\text{BiOBr}_{0.6}\text{I}_{0.4}$, $\text{BiOBr}_{0.4}\text{I}_{0.6}$, $\text{BiOBr}_{0.2}\text{I}_{0.8}$, and BiOI, correspondingly. The rate constant of $\text{BiOBr}_{0.6}\text{I}_{0.4}$ was three and six times greater than that of individual BiOBr and BiOI, correspondingly. Even after five runs, the material still showed outstanding efficiency and stability. Hence, this study has demonstrated that $\text{BiOBrI}_{z(1-z)}$ solid solutions synthesized by hydrothermal technique are promising for the degradation of MB in wastewaters. From the linear and non-linear forms of Langmuir-Hinshelwood (L-H) isotherms, nonlinear provided the most flexible curve fitting functionality that prevented the errors brought by various approximations resulting from

simple linear regression. The experimental results well fitted the nonlinear and pseudo-first-order model. From the trapping experiment conducted, it was established that the $\cdot\text{O}_2^-$ and h^+ radicals were the key active species responsible for the removal of MB. Therefore, the use of solid solutions has promised to be a better route for environmental remediation.

Data availability

Underlying data

Zenodo: Synthesis of Bismuth Oxyhalide ($\text{BiOBrI}_{z(1-z)}$) Solid Solutions for Photodegradation of Methylene Blue Dye. <http://doi.org/10.5281/zenodo.4987341>³⁵.

This project contains the following underlying data:

- Degradation, scavenger, pH, catalyst dosage, thermal contribution, XRD, SEM TEM, Raman, Point-of-Zero Charge Raw Data.xlsx

Extended data

Zenodo: Synthesis of Bismuth Oxyhalide ($\text{BiOBrI}_{z(1-z)}$) Solid Solutions for Photodegradation of Methylene Blue Dye. <http://doi.org/10.5281/zenodo.4987341>³⁵.

This project contains the following extended data:

- Supporting Information - Figures and Tables.docx

Data are available under the terms of the [Creative Commons Attribution 4.0 International license](#) (CC-BY 4.0).

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Current Peer Review Status:

Version 2

Reviewer Report 14 February 2022

<https://doi.org/10.21956/aasopenres.14470.r29125>

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John Mmari Onyari

Department of Chemistry, University of Nairobi, Nairobi, Kenya

Thanks. The revisions made are appropriate.

Competing Interests: No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Version 1

Reviewer Report 22 September 2021

<https://doi.org/10.21956/aasopenres.14371.r28852>

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John Mmari Onyari

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This study, involved the synthesis and characterization of Bismuth oxyhalide (BiOBr_z I (1-z)) ($0 \leq z \leq 1$) materials. Bismuth oxyhalide was used in the removal of methylene blue (MB) which is a textile waste that is discharged to the environment. The photocatalytic performance of the materials was examined by photodegradation of the MB done under solar and ultraviolet (UV) light irradiation. The presentation of results is well done and the conclusions drawn in the study were

adequately supported by the results.

The manuscript makes a significant contribution to scientific knowledge and my recommendation is **"Approved with Reservations."** The paper is to be accepted for indexing after the minor revisions to address issues summarized below.

1) Page 4: "However, to the best of the authors' knowledge, the application of BiOBrz I (1-z) solid solutions in the treatment of methylene blue (MB) dye and comparative studies on the photocatalytic performance of these materials under natural sunlight and UV light has not been reported." Include in the literature review and discussion section work done by Ren *et al.* (2013¹).

2) Page 5, Photodegradation studies: "The set up for the photodegradation under UV lamp and solar is shown in **Figure S1**." Where is Figure S1 shown?

3) Page 7: "The corresponding bandgap energies of BiOBrz I (1-z) materials were calculated and the result is shown in Table 1". [... **results are shown** ...].

4) Page 8 Fitting of the adsorption data to adsorption isotherms using nonlinear model fitting procedures (**Extended data**, Figure S7³³). See also Page 9 etc ... As presented in **Table S3** of the **Extended data**³³ Page 10 are shown in Figure S7 of the **Extended data**³³. Reference is made to "Extended data" several times. Why not just refer to the key finding and give the reference?

5) Page 10: Label the x-axis in Figure 6.

6) Page 11: Check the spelling of "forth" in Figure 7 (replace "forth" by "fourth"?).

7) Page 11 last line: "Figure S8b displays the Arrhenius plot of k" - where is Figure S8b shown?

8) Page 11 Effect of amount of catalyst on MB removal. To determine the influence of the amount of catalyst on the MB removal, a dosage of BiOBr_{0.6}I_{0.4} was increased from 0.01 to 0.07 g/L under **optimal conditions** (10 mg/L MB concentration, pH, 7.0, and 150 min reaction time).- indicate the optimal temperature used.

9) Page 13 References:

Pavithra KG, Kumar PS, Jaikumar V, **et al.**: etc. - Why is et al., used in this section? Check the Journal guidelines.

Silveira E, Marques PP, Silva SS, **et al.**:

Zhang LH, Zhu Y, Lei BR, et al.:

Drumond Chequer FM, de Oliveira GAR, Anastacio Ferraz ER, **et al.**:

Sun X, Zhang Y, Li C, **et al.**:

References

1. Ren K, Liu J, Liang J, Zhang K, et al.: Synthesis of the bismuth oxyhalide solid solutions with tunable band gap and photocatalytic activities. *Dalton Trans.* 2013; **42** (26): 9706-12 [PubMed Abstract](#) | [Publisher Full Text](#)

Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Development and Utilization of various adsorbent materials for adsorption of dyes and heavy metals

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Author Response 15 Dec 2021

Botswana International University of Science Botswana International University of Science

15th December 2021

Summary of Revisions.

Detailed Response to Reviewers' comments

All changes are highlighted in "red" in the revised manuscript. Please refer to these changes for more detailed information.

Reviewers Comments.

Reviewer #3:

General Comment: APPROVED WITH RESERVATION

This study, involved the synthesis and characterization of Bismuth oxyhalide ($\text{BiOBr}_z\text{I}_{(1-z)}$) ($0 \leq z \leq 1$) materials. Bismuth oxyhalide was used in the removal of methylene blue (MB) which is a textile waste that is discharged to the environment. The photocatalytic performance of the materials was examined by photodegradation of the MB done under solar and ultraviolet (UV) light irradiation. The presentation of results is well done and the conclusions drawn in the study were adequately supported by the results. The manuscript makes a significant contribution to scientific knowledge and my recommendation is “

Approved with Reservations. The paper is to be accepted for indexing after the minor revisions to address issues summarized below.

Comment #1: Page 4: "However, to the best of the authors' knowledge, the application of $\text{BiOBr}_z\text{I}_{(1-z)}$ solid solutions in the treatment of methylene blue (MB) dye and comparative studies on the photocatalytic performance of these materials under natural sunlight and UV light has not been reported." Include in the literature review and discussion section work done by Ren *et al.* (2013¹).

Response:

The authors are very grateful for that important suggestion and the comment.

Literature Review

$\text{BiOBr}_z\text{I}_{1-z}$ (M, R = Cl, Br, I) solid solutions (Ren *et al.*, 2013) which was the work done by Ren *et al.* has been added on the literature review part as was suggested by the reviewer. The change has been highlighted on the revised manuscript.

Discussion

*From the X-ray diffractometry data, a shift in diffraction peaks to lower diffraction angles was observed in our study. This observation was also reported by Xu *et al.*, and Ren *et al.* (Ren *et al.*, 2013). The work of the Ren *et al.* has been acknowledged as was suggested by the reviewer on the result and discussion part. One reference was added*

Reference

No. 33 Ren, K., Liu, J., Liang, J., Zhang, K., Zheng, X., Luo, H., Huang, Y., Liu, P., & Yu, X. (2013). Synthesis of the bismuth oxyhalide solid solutions with tunable band gap and photocatalytic activities. Dalton Transactions, 42(26), 9706–9712. <https://doi.org/10.1039/c3dt50498k>

Comment #2: Photodegradation studies: "The set up for the photodegradation under UV lamp and solar is shown in **Figure S1**." Where is Figure S1 shown?

Response:

The Figure S1 is shown on the Extended Data [35]. The change has been made on the revised manuscript and it is highlighted

Reference

No. 35 R. O. Gembo, O. Aoyi, S. Majoni, A. Etale, S. Odisitse, and C. K. King'onde, "Synthesis of bismuth oxyhalide ($\text{BiOBr}_z\text{I}_{(1-z)}$) solid solutions for photodegradation of methylene blue dye," AAS Open Res., vol. 4, p. 43, 2021, doi: 10.12688/aasopenres.13249.1.

Comment #3: Page 7: "The corresponding bandgap energies of $\text{BiOBr}_z\text{I}_{(1-z)}$ materials were calculated and the result is shown in Table 1". [... **results are shown** ...].

Response:

Thanks to the reviewer for being able to notice the error. The word "is" has been replaced by "are". The manuscript has been revised to capture the change.

Comment #4: Page 8 Fitting of the adsorption data to adsorption isotherms using nonlinear model fitting procedures (**Extended data**, Figure S7³⁵). See also Page 9 etc ... As presented in **Table S3** of the **Extended data**³⁵ Page 10 are shown in Figure S7 of the **Extended data**³⁵. Reference is made to "Extended data" several times. Why not just refer to the key finding and give the reference?

Response:

The suggestion is very significant and the authors appreciate it. However, if all the information is

included in the manuscript, it would be congested with a lot of figures hence some information has been kept in the Extended Data therefore the references.

Comment #5: Page 10: Label the x-axis in Figure 6.

Response:

*The comment is appreciated. The X – axis has been labelled as “**Quenching Agent**”. The change is highlighted on the figure.*

Comment #6: Page 11: Check the spelling of “forth” in Figure 7 (replace “forth” by “fourth”?).

Response:

*The comment is appreciated. The word “**Forth**” is replaced by “**Fourth**” the change has been made on figure 7*

Comment #7: Page 11 last line: "Figure S8b displays the Arrhenius plot of k" - where is Figure S8b shown?

Response:

The comment is appreciated. The Figure S8b is shown on the Extended Data [35]. The change has been made on the revised manuscript and it is highlighted

Reference

No. 35 R. O. Gembo, O. Aoyi, S. Majoni, A. Etale, S. Odisitse, and C. K. King'ondur, "Synthesis of bismuth oxyhalide ($\text{BiOBr}_{2-1-x}\text{I}_x$) solid solutions for photodegradation of methylene blue dye," AAS Open Res., vol. 4, p. 43, 2021, doi: 10.12688/aasopenres.13249.1.

Comment #8: Page 11 Effect of amount of catalyst on MB removal. To determine the influence of the amount of catalyst on the MB removal, a dosage of $\text{BiOBr}_{0.6}\text{I}_{0.4}$ was increased from 0.01 to 0.07 g/L under **optimal conditions** (10 mg/L MB concentration, pH, 7.0, and 150 min reaction time).- indicate the optimal temperature used.

Response:

*The comment is appreciated. The optimal temperature is **$30 \pm 2^\circ\text{C}$***

Comment #9: Page 13 References:

Pavithra KG, Kumar PS, Jaikumar V, **et al.:** etc. - Why is et al., used in this section? Check the Journal guidelines.

Silveira E, Marques PP, Silva SS, **et al.:**

Zhang LH, Zhu Y, Lei BR, et al.:

Drumond Chequer FM, de Oliveira GAR, Anastacio Ferraz ER, **et al.:**

Sun X, Zhang Y, Li C, **et al.:**

Response:

The authors would to commend the reviewer such an important question. Nonetheless, The original document includes all the authors in the references. However, the references have the et al because that's the AAS guideline on presenting the references.

Competing Interests: No competing interest

Reviewer Report 14 September 2021

<https://doi.org/10.21956/aasopenres.14371.r28856>

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Elizabeth N. Ndunda

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Environmental pollution is becoming a global challenge and studies on remediation techniques for these environmental pollutants are timely. The current manuscript clearly details the degradation of methylene blue dye, which is found in textile waste, using bismuth oxyhalide. The manuscript is well written, with the following minor suggestions:

1. The title needs to be revised to capture methylene blue dye as used in text.
2. The authors need to mention if they tried the catalyst on real samples.
3. Besides MB being a textile waste, the authors should further mention any effects that MB might have on humans, animals and the environment.

Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: fabrication of sensor, synthesis of molecularly imprinted polymers for

analytical applications.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 15 Dec 2021

Botswana International University of Science Botswana International University of Science

15th December 2021

Summary of Revisions.

Detailed Response to Reviewers' comments

All changes are highlighted in "red" in the revised manuscript. Please refer to these changes for more detailed information.

Reviewers Comments.

Reviewer #2:

General Comment: APPROVED

Environmental pollution is becoming a global challenge and studies on remediation techniques for these environmental pollutants are timely. The current manuscript clearly details the degradation of methylene blue dye, which is found in textile waste, using bismuth oxyhalide. The manuscript is well written, with the following minor suggestions:

Comment #1: The title needs to be revised to capture methylene blue dye as used in text.

Response: *The authors acknowledge the reviewer for such an important observation about the title "Synthesis of bismuth oxyhalide ($\text{BiOBr}_2\text{I}_{(1-z)}$) solid solutions for photodegradation of methylene dye". The title has been revised as suggested. The change has been highlighted on the revised manuscript.*

New title: *Synthesis of bismuth oxyhalide ($\text{BiOBr}_2\text{I}_{(1-z)}$) solid solutions for photodegradation of methylene blue dye*

Comment #2: The authors need to mention if they tried the catalyst on real samples.

Response: *Thanks to the reviewer for such a vital concern. However, the catalyst was not tried on the real samples because there is no textile industry around the university where the study was carried out, therefore the authors resorted for synthetic textile waste.*

Comment #3: Besides MB being a textile waste, the authors should further mention any effects that MB might have on humans, animals and the environment.

Response: *Thanks to the reviewer for that significant question on whether the dye has effect on both human and animals.*

Added paragraph on the revised manuscript: *... Methylene blue dye if not monitored can adversely affect human beings and animals by causing eye burns which is a permanent health damage. Additionally, the MB can cause painful micturition, nausea, increase in heart rate, quadriplegia disorders in breathing, quadriplegia, cyanosis, mental confusion, jaundice, vomiting, tissue necrosis, and methemoglobinemia. (Ramírez-Aparicio et al., 2021). The reference below was added in response to the comment.*

Reference

No. 34 Ramírez-Aparicio, J., Samaniego-Benítez, J. E., Murillo-Tovar, M. A., Benítez-Benítez, J. L.,

Muñoz-Sandoval, E., & García-Betancourt, M. L. (2021). Removal and surface photocatalytic degradation of methylene blue on carbon nanostructures. Diamond and Related Materials, 119(July), 108544. <https://doi.org/10.1016/j.diamond.2021.108544>.

Competing Interests: No competing interest

Reviewer Report 08 September 2021

<https://doi.org/10.21956/aasopenres.14371.r28854>

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Ndzondelelo Bingwa

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The authors describe synthesis of bismuth oxyhalides and their application in photocatalytic degradation of an industrial (textile) pollutant, methylene blue. They further studied the kinetics of adsorption using the Langmuir-based kinetic model. The results are well presented. The structure of the work is commendable. However, there are a few minor issues that need to be taken care of.

1. The product(s) of degradation must be stated in the abstract. Otherwise, the reader might think that the only process taking place is removal by adsorption.
2. The magnification on the TEM images is too low. It would be great if the images are taken at higher magnification. This will allow the determination of interplanar spacing.
3. The catalytic recyclability and reusability is a study can be reported in a better manner. Comparing conversions at the end of the reaction or at a certain reaction time might be deceiving. It is always advisable to study catalyst reusability with kinetics. In this way, it is easy to see if the catalyst is deactivating by looking at the reaction rates rather than conversion at time t.

Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Heterogeneous catalysis, material synthesis, photocatalysis, nanomaterials for catalysis

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 15 Dec 2021

Botswana International University of Science Botswana International University of Science

15th December 2021

Summary of Revisions.

Detailed Response to Reviewers' comments

All changes are highlighted in "red" in the revised manuscript. Please refer to these changes for more detailed information.

Reviewers Comments.

Reviewer #1:

General Comment: APPROVED

The authors describe synthesis of bismuth oxyhalides and their application in photocatalytic degradation of an industrial (textile) pollutant, methylene blue. They further studied the kinetics of adsorption using the Langmuir-based kinetic model. The results are well presented. The structure of the work is commendable. However, there are a few minor issues that need to be taken care of.

Comment #1: The product(s) of degradation must be stated in the abstract. Otherwise, the reader might think that the only process taking place is removal by adsorption.

Response: Thanks to the reviewer for that vital comment. Nevertheless, the degradation products were not analyzed as it requires LC-MS which is not available at our university, therefore the analysis was not possible. Nevertheless, the degradation was monitored through color reduction.

Comment #2: The magnification on the TEM images is too low. It would be great if the images are taken at higher magnification. This will allow the determination of interplanar spacing.

Response: The authors thank the reviewer for this excellent suggestion. It is true that the

magnification applied was low to reveal the interplanar spacing. However, the interplanar spacing can be determined by high resolution transmission electron microscope unlike in this work where the materials were analyzed using transmission electron microscope which only gives the micrographs.

Comment #3: The catalytic recyclability and reusability is a study can be reported in a better manner. Comparing conversions at the end of the reaction or at a certain reaction time might be deceiving. It is always advisable to study catalyst reusability with kinetics. In this way, it is easy to see if the catalyst is deactivating by looking at the reaction rates rather than conversion at time t.

Response: The authors thank the reviewer for this excellent suggestion that provide more insight and evidence on the stability of the prepared materials, unfortunately a stability with kinetics was not studied as the authors only did a study to establish the stability of the prepared photocatalysts. The cyclic voltammetry was not performed to establish whether the catalyst was deactivating or not.

Competing Interests: No competing interest