Heterostructured Nanophotocatalysts for Degradation of Organophosphate Pesticides from Aqueous Streams

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ABSTRACT. The present paper focuses on the synthesis, characterization and application of nanophotocatalyst for degradation of quinalphos and monocrotophos. Novel heterostructured ZnO/TiO$_2$ photocatalyst (Z$_9$T) was prepared and characterized with X-ray diffraction (XRD), SEM and UV–vis diffuses reflectance spectroscopy. The average crystalline size of synthesized Z$_9$T was found to be 21.48 nm. The pesticides were degraded in the presence of nanophotocatalysts i.e., TiO$_2$, ZnO, TiO$_2$/ZnO mixed in various proportions and heterostructured nanophotocatalyst synthesized by Sol-Gel method. The batch experiments were performed by adding photocatalyst to 100 ml of pesticide solution and suspension was subjected to irradiation under UV light. In case of mixed catalyst, the maximum degradation of monocrotophos and quinalphos has been observed when ZnO and TiO$_2$ were in the ratio of 7:3 and 8:2 respectively. The degradation efficiency with synthesized heterostructured nanophotocatalyst (Z$_9$T) was found to be comparable with TiO$_2$.

Key words: Nanophotocatalyst, Degradation, Quinalphos, Monocrotophos, Z$_9$T

INTRODUCTION

Organophosphate pesticides (OP’s) are relatively soluble in water and have significant potential for entering aquatic environments through surface runoff, sprays, and soil leachates. These have been detected frequently in environmental waters, and the issue of their removal from aqueous streams has gained momentum. Pesticides contamination of aqueous streams is matter of great concern to the regulation bodies as well as environmental societies. The current technologies available for water treatment are found to be insufficient for removal of the pesticides from aqueous streams. Alternately, the Advanced Oxidation Processes (AOP’s) have emerged as best available technology (BAT) as per US EPA for removal of organic pollutants.

Environment friendly semiconductor nanomaterials have found diverse applications in numerous fields such as photovoltaics,$^1$ photocatalysis,$^{2-4}$ self-cleaning coatings,$^5$ advanced oxidation,$^6$ and photoelectrocatalytic degradation of organic compounds.$^7$ Several semiconductors can act as photocatalyst for light-induced chemical transformations because of their unique electronic structure, e.g. TiO$_2$, ZnO, Fe$_2$O$_3$, WO$_3$ and CdS. Among all of these, TiO$_2$ and ZnO have been studied extensively for the removal of organic pollutants such as synthetic dyes, pesticides, phenols, chlorophenols etc. from the aqueous streams. Both, TiO$_2$ and ZnO absorb 5% of solar radiation.

Quinalphos (O,O-diethyl O-quinoxalin-2-yl phosphorothioate) and Monocrotophos (dimethyl [(E)-4-(methylamino)-4-oxobut-2-en-2-yl] phosphate) are widely used OP’s in Indian agriculture. Quinalphos (QP) is used for controlling caterpillar and scale insect on fruit trees, cotton, vegetables, peanuts and pest complex on rice.$^8,9$ Monocrotophos (MCP) is most popular and widely used OP owing to its low cost and high efficiency in controlling pests mainly on cotton crop, rice and sugarcane, and active against large variety of insects in India. World Health Organization (WHO) and Environmental Protection Agency (EPA) classified Quinalphos in class II as moderate toxic and MCP in class I – highly toxic compound. However, Monocrotophos is continually used for the control of major pests in agriculture in developing countries like India primarily due to lack of alternative replacements.$^{10}$

Many studies have been reported on degradation of OP’s (containing P=S and P–S groups), such as Methamidophos,$^{11,12}$ Malathion,$^{13,14}$ Parathion$^{15}$ etc. by heterogeneous photocatalysis. These findings have showed that OP’s degrade rapidly and non-selectively in an illuminated suspension of TiO$_2$. The feasibility of photocatalytic degradation of monocrotophos using floating TiO$_2$:SiO$_2$ photocatalyst beads,$^{16}$ synthesized La doped ZnO nanoparticles and IO$_2$: doped TiO$_2$ nanoparticles was reported.$^{17,18}$ No study has been reported for photocatalytic degradation of Quinalphos using photocatalyst.
In continuation with the work done earlier by our research group, the present paper reports (i) the degradation of Quinalphos and Monocrotophos in the presence of TiO₂ and the effects of various process parameters (catalyst loading, type of catalyst and pH) on the efficiency of photocatalytic degradation, (ii) synthesis and characterization of nanophotocatalyst named Z₉T and comparison with TiO₂, ZnO, TiO₂/ZnO in mixed ratio in the presence of UV light.

EXPERIMENT

Material
Commercial samples of technical grade: Quinalphos (>95%) and Monocrotophos (>95%) were obtained from Crops Chemical Limited, Kotkapura, Punjab (India) and used as received for present study. Titania P-25 having 85% anatase and 15% rutile phase (surface area 50 m²/g) was used as received for present study. Titania P-25 having >95% was obtained from Degussa and ZnO (surface area 5 m²/g) from Merck. Potassium dichromate, sulfuric acid, silver sulphate, and mercuric sulphate were purchased from S.D. Fine Chemical, India. HCl and NaOH were obtained from Merck, India. All other chemicals used were of analytical grade. Millipore water was used for preparation of various solutions. The pH of the solutions was adjusted with 1 M HCl or 1 M NaOH.

Instruments
The degradation process was carried out in specially designed reaction vessels (diameter 0.08 m and volume 500 ml) in the photoreactor equipped with 4 UV tubes each of 30W (Philips). The intensity of UV light was 2.4×10⁵ einstein/minute measured by chemical method i.e., potassium ferrioxalate actinometry. Constant stirring of solution was insured by using magnetic stirrers, and aeration was done with the help of aquarium aerator. The experimental set-up used for photodegradation experiment was reported earlier. The spectra were taken with UV–vis spectrophotometer (DR 4000); pH meter (Thermo Orion 920A) was used to adjust the pH of the solution.

PROCEDURE

Preparation and Characterization of Nanosized Photocatalysts
The nanosized heterostructured ZnO/TiO₂ (Z₉T) photocatalyst was prepared using TiO₂ and Zn(NO₃)₂·6H₂O as starting materials, and NaOH was used as the precipitant without further purification. Zn(NO₃)₂·6H₂O was dissolved in a minimum amount of deionized water with constant stirring until clear solution was obtained. 1M NaOH solution was added till white amorphous precipitates of Zn(OH)₂ were formed. Then the required amount of TiO₂ was added with constant stirring for the preparation of the heterostructured ZnO/TiO₂ photocatalyst with the Zn/Ti molar ratios of 9:1, by maintaining pH to about 7. The precipitates were stirred for 20 hrs for homogeneous mixing. The precipitates were filtered and washed with deionized water to remove the soluble ions. The wet powder obtained was dried at 110 °C in air to form the precursors of the heterostructured ZnO/TiO₂ photocatalyst which were further calcined at 250 °C in air for 3 hrs. The nanosized photocatalyst was obtained in powder form labeled as Z₉T.

Characterization was done by XRD, SEM and UV-DR. X-ray powder diffraction (XRD) analysis of the heterostructured ZnO/TiO₂ photocatalyst powder was carried out at room temperature by using a Rigaku D/max-III C diffractometer with Cu-Kα radiation (λ = 0.15418 nm) over the 2θ collection range of 20–100 °C. The accelerating voltage of 40 kV, emission current of 30 mA and the scanning speed of 4.4 counts per second were used. UV-vis transmittance spectra (UV-vis) were recorded in air at room temperature in the wavelength range of 200–700 nm using Perkin Elmer Lambda 750 UV/vis/NIR spectrophotometer. Microscopic feature of catalyst was studied with scanning electron microscopy (SEM, Hitachi-S-3400N) using fine catalyst powder supported on carbon tape and coated with gold.

Photocatalytic Degradation Experiments
For the degradation experiments, 0.1 gm of photocatalyst was added to 100 ml of pesticide solution, and suspension was subjected to irradiation under UV light. The aqueous suspension was magnetically stirred and aerated throughout the experiment. At different time intervals, aliquot was taken out with the help of syringe and then filtered through Millipore syringe filter of 0.45 µm. The absorption spectra were recorded at λmax 320 nm for Quinalphos and 216 nm for Monocrotophos. The rate of degradation was studied in terms of changes in absorption spectra. The degradation efficiency (%) has been calculated as follows:

Efficiency (%) = [(C₀ – C) / C₀] × 100

where C₀ is the initial concentration of pesticide and C is the concentration of pesticide after photoirradiation. The nanophotocatalysts used in the present study are TiO₂, ZnO, TiO₂/ZnO in mixed ratio and synthesized heterostructured nanophotocatalyst.
RESULTS AND DISCUSSION

The molecular structures and properties of Quinalphos and Monocrotophos are illustrated in Table 1. Monocrotophos is highly soluble in water while Quinalphos has less solubility in water but high in ethanol. Quinalphos shows characteristic absorption peaks at 240 nm and 320 nm and Monocrotophos shows at 216 nm in UV region (Table 1). The rate of degradation was recorded with respect to the change in intensity of absorption peaks at 216 nm for Monocrotophos and 320 nm for Quinalphos. The absorption peaks corresponding with pesticides diminished during reaction, which showed that the pesticides had been degraded.

Characterization of Catalyst

The prepared heterostructured TiO$_2$/ZnO photocatalyst (Z$_9$T) was characterized by XRD, SEM and UV-vis transmittance spectroscopy. The X-ray diffraction pattern of prepared sample (Z$_9$T) is shown in Fig. 1. From the XRD pattern and corresponding characteristics 20 values of the diffraction peaks, it can be confirmed that TiO$_2$ is in rutile form having tetragonal structure as the peaks matched with Joint Committee Powder Diffraction Standards (JCPDS) card number 21-1276 while ZnO is in zincite phase (JCPDS card number 36-1451). The h, k, l values are shown in figure at respective peaks. The average crystallite size of Z$_9$T was calculated to be about 21.48 nm using the Scherrer Equation,$^{14,22}$

$$t = \frac{K \lambda}{B \cos \theta},$$

where the number K is a coefficient having value 0.9, $\lambda = 0.1541$ nm is the X-ray wavelength, B is the full width half maximum (FWHM) of the catalyst, $\theta$ is the diffraction angle.

The energy band gap of sample has been calculated according to the equation widely adapted for crystalline semiconductors.$^{24,25}$

$$\alpha h\nu = A (h\nu - E_g)^{n/2}$$

![Figure 1. XRD pattern of Z$_9$T.](image)

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$$\alpha h\nu = A (h\nu - E_g)^{n/2}$$

<table>
<thead>
<tr>
<th>Common name</th>
<th>Monocrotophos</th>
<th>Quinalphos</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pesticide group</td>
<td>Organophosphate</td>
<td>Organophosphate</td>
</tr>
<tr>
<td>IUPAC name</td>
<td>dimethyl [(E)-4-(methylamino)-4-oxobut-2-en-2-yl] phosphate</td>
<td>O,O-diethyl O-quinoxalin-2-yl phosphorothioate</td>
</tr>
<tr>
<td>Molecular formula</td>
<td>C$<em>9$H$</em>{14}$NO$_5$PC</td>
<td>C$<em>{12}$H$</em>{15}$N$_2$O$_3$PS</td>
</tr>
<tr>
<td>Molecular weight</td>
<td>223.2</td>
<td>298.3</td>
</tr>
<tr>
<td>Activity</td>
<td>Aliphatic organophosphate insecticide and Acaricide</td>
<td>Aromatic organophosphate insecticide and Acaricide</td>
</tr>
<tr>
<td>Solubility in water</td>
<td>100% at 20 °C</td>
<td>20 mg/L</td>
</tr>
<tr>
<td>Toxicity Class</td>
<td>I</td>
<td>II</td>
</tr>
<tr>
<td>Acute oral LD$_{50}$ for rats</td>
<td>14 mg/kg</td>
<td>14–37 mg/kg</td>
</tr>
</tbody>
</table>

![Structure](image)

![Absorbance maxima at 216 nm(1.057 nm)](image)

<table>
<thead>
<tr>
<th>$\lambda_{max}$</th>
<th>240 nm and 320 nm</th>
</tr>
</thead>
</table>
where $\alpha$, $\nu$, $A$, $E_g$, and $n$ are the absorption coefficient, incident light frequency, constant, band gap, and an integer (normally equal to 1, 2, 4, or 6), respectively. The $\alpha$ and $\nu$ values at the steep edges of the transmission spectra were used to construct the plots of $(\alpha h \nu)^{1/2}$ vs photon energy. As seen in Fig. 2, the band gap of sample was determined by the intersection of the extrapolated linear portion of the plot with the energy axis. The calculated band gap value of synthesized photocatalyst was 3.60 eV.

The structure and morphology of the $Z_9T$ photocatalyst was studied using Scanning Electron Microscope (SEM). The SEM unit was operated at 15 kV of accelerating voltage. Fig. 3 shows the SEM photograph of the $Z_9T$ under 15000X magnification. The coupled photocatalyst $Z_9T$ consist of nanosized particles and their size seems to be less than 100 nm. TiO$_2$ nanoparticles and ZnO nanorods aggregate to form irregular shaped clusters.

**Control Experiments**

The degradation of quinalphos and monocrotophos was carried out under different experimental conditions viz UV light and TiO$_2$/UV light. The degradation rate was recorded in terms of change in intensity of characteristic peak at 320 nm for quinalphos and 240 nm for monocrotophos. Initially blank experiments were performed under UV irradiation without addition of catalyst (UV alone) and only 4% and 13.29% degradation of quinalphos and monocrotophos was observed respectively. The photocatalytic degradation experiment was performed by taking 0.1 gm photocatalyst in 100 ml solution of the pesticide solution. About 89% degradation of quinalphos and 60% degradation of monocrotophos were achieved in the presence of the semiconductor (TiO$_2$) and UV light in 4 h. The results from the degradation of both the pesticides reveal that TiO$_2$ shows good photocatalytic activity for the degradation of pesticides.

**Effect of Parameters**

To determine the effect of catalyst dose (TiO$_2$), the experiments were performed by varying the catalyst dose from 0.5 to 2.0 g/L for pesticide solutions. The results reveal that efficiency of degradation increases up to concentration 1.5 g/L for quinalphos after that the rate of degradation decreases but for monocrotophos degradation efficiency increases up to 2.0 g/L catalyst dose. This is in accordance with the previous studies and can be explained on the basis of optimum catalyst loading which depends on initial solute concentration. The total active surface area increases with the increase of catalyst dose. At the same time, higher dose of photocatalyst increases turbidity of the suspension which leads to decrease the penetration of UV light. Hence photo activated volume of suspension decreases.

The experiments to check the effect of pH were carried out in the pH range 2–10 for both of the pesticides. The maximum degradation was obtained at pH 8 for the quinalphos and pH 4 for the monocrotophos while sharp decrease occurred at both high and low pH. The results are in accordance with the earlier findings for degradation of trichlorfon in suspension of titanium dioxide. The dependence of degradation on pH depends not only on the pKa of TiO$_2$ but that of the substrate as well. Otherwise, the trend (degradation vs pH) would have been uniform for both the substrates.

**HPLC Studies**

The aqueous solutions of photo degraded pesticides are further analyzed by high performance liquid chromatography (HPLC). The HPLC chromatograms after 2 h of irradiation are shown in Fig. 4. HPLC chromatogram confirms...
the formation and disappearance of intermediates during photodegradation of quinalphos and monocrotophos. Several small organic intermediates were formed, out of which the major products of quinalphos are with retention time 11.988 and 3.788 min. For the monocrotophos the major products with retention time 6.058 and 8.819 min are formed.

**Kinetics**

*Fig. 5* shows the kinetic of disappearance of quinalphos and monocrotophos for an initial concentration 20 ppm respectively. The results show that the photocatalytic degradation of pesticide in aqueous TiO$_2$ can be described by the pseudo first order kinetic model $\ln(C_0/C) = kt$, where $C_0$ is an initial concentration and $C$ is concentration at time $t$. The correlation constant for the fitted line is $R^2 = 0.99664$ for quinalphos and $R^2 = 0.9758$ for monocrotophos. The rate constant has been calculated to be 0.00567 min$^{-1}$ for quinalphos and 0.00199 min$^{-1}$ for monocrotophos.

**Degradation of pesticides with mixed and synthesized catalyst**

Photodegradation experiments were also performed by using fixed amount of pesticide (20 ppm) with catalyst dose, i.e., 1 g/l of ZnO/TiO$_2$ mixed in different ratio (9:1, 8:2, 7:3 and 6:4), synthesized heterostructured nanophotocatalyst (Z$_9^T$). The results are compared with photocatalytic efficiency of TiO$_2$ and ZnO at natural pH in UV light for 4 hours. The percentage degradation of pesticide by using various catalysts is reported in *Fig. 6*.

Experimental studies show that the degradation occurs at higher rate with TiO$_2$ as compared with ZnO. The present result is in confirmation with earlier findings of using TiO$_2$ for the photocatalytic degradation of pesticides and the proposed mechanism of degradation for TiO$_2$ and ZnO.$^{30}$ However, when the photocatalysts were mixed, the degradation was comparatively high as compared to use TiO$_2$ and ZnO alone. The maximum degradation (81%) of Monocrotophos has been observed with ZnO and TiO$_2$.

*Figure 4.* HPLC chromatograms of (a) Quinalphos and (b) Monocrotophos after 2 h of irradiation.

*Figure 5.* Kinetic study of degradation of (a) Quinalphos and (b) Monocrotophos.
mixed in 7:3. However, 93.50% degradation of Quinalphos has been observed with ZnO and TiO\textsubscript{2} in 8:2. In earlier findings higher rate of photocatalytic degradation of Malachite Green\textsuperscript{80} and Methyl Blue\textsuperscript{81} were reported with TiO\textsubscript{2} and ZnO mixed in the ratio 1:9 and 9:1 respectively.

Quinalphos and Monocrotophos were also subjected to photocatalytic degradation of Quinalphos and Monocrotophos in presence of synthesized heterostructured nanophotocatalyst (Z\textsubscript{9}T). Z\textsubscript{9}T showed the best photocatalytic degradation efficiency (96% of Quinalphos and 84.2% of Monocrotophos) in comparison to all other photocatalysts (Fig. 6). The better degradation efficiency of Z\textsubscript{9}T may be explained on the basis of smaller crystallite size (21.48 nm). As the catalyst grain size is reduced to a few nanometers, an elevated density of active sites for substrate adsorption and catalysis can be guaranteed. Thus, higher surface-to-volume ratio as compared to the bulk material results in improved efficiency.

CONCLUSION

The photocatalytic degradation of Quinalphos and Monocrotophos in aqueous solution was carried out using TiO\textsubscript{2}, ZnO, both the catalysts mixed in different proportions and synthesized heterostructured nanophotocatalyst. The rate constant for degradation of Quinalphos and Monocrotophos with TiO\textsubscript{2} has been calculated to be 0.00567 min\textsuperscript{-1} and 0.00199 min\textsuperscript{-1} respectively. XRD studies of synthesized photocatalyst Z\textsubscript{9}T reveal the existence of TiO\textsubscript{2} in rutile form and ZnO in zincite phase. The smaller crystalline size of Z\textsubscript{9}T was found to be responsible for the better photocatalytic efficiency in comparison to others. The studies related to synthesis of new photocatalyst with improved efficiency will helpful in designing photocatalytic degradation based water treatment plants.

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