THE REMOVAL OF NEONICOTINOID INSECTICIDE IMIDACLOPRID IN AN ANNULAR PHOTOREACTOR

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Abstract: Heterogeneous photocatalysis has recently attracted an increasing interest of scientists and experts who deal with the waste water and air treatment. An important area of application is removal of persistent organic pollutants, which can not be easily destroyed by conventional methods. Among these pollutants are neonicotinoid insecticides, which are widely used all over the world and currently are included in the watch list of substances of the European Commission within the Water Framework Directive. Therefore, there is a need to study their influence on the environment and to develop appropriate technologies for their removal. In this work, the photolytic and photocatalytic degradation of neonicotinoid insecticide imidacloprid in an annular photoreactor with recirculation under different working conditions (irradiation source, pH, type and concentration of the catalyst) was studied. The photolytic degradation was examined using lamps that emit UVA, UVC and simulated sun light. The photocatalytic experiments in the suspension involved the use of commercial titanium dioxide (TiO2 P25, Degussa/Evonik), commercial TiO2 P25 pre-treated with UVC irradiation prior to use in the catalytic system and nitrogen-doped TiO2 (CCR 200 N produced by Cinkarna Celje). The catalysts were characterized using XRD, UV/Vis-DRS and BET analysis. The most efficient photocatalyst was then immobilized on the glass waving fibre, using peroxytitanic acid (produced by Cinkarna Celje) as a binder. The degree of degradation of imidacloprid was determined using high performance liquid chromatography (HPLC). According to the obtained results, UVC-treated TiO2 showed the best efficiency among the examined catalysts in the slurry reactor while using lamp that simulates the sun irradiation. The imidacloprid degradation rate increases with the increase in the catalyst concentration. The immobilized UVC-treated catalyst gave satisfying results in terms of stability, activity and reuse.

Keywords: heterogeneous photocatalysis, neonicotinoid insecticides, imidacloprid, annular reactor

1. INTRODUCTION

Neonicotinoid insecticides are relatively new group of insecticides, widely applied for the seed treatment. Because of their high solubility in water and low adsorption to the soil, they can reach underground and surface water during rainfall rinsing. In the last ten years several investigations connected some neonicotinoids with bees poisoning all over the world (Žabar et al. 2012). At present, their use is restricted, but still not completely banned. Some insecticides from the group of neonicotinoids, e.g. imidacloprid, are included in the watch list of substances for Union-wide monitoring in the field of water policy, established by the European Commission Implementing Decision 2018/840 pursuant to Directive 2008/105/EC (Commission implementing decision (EU) 2018/840).

The existing water treatment processes cannot remove persistent organic compounds sufficiently, which is why the advanced oxidation processes (AOPs) are increasingly explored as an alternative to conventional methods. Among the most investigated AOPs is heterogeneous semiconductor photocatalysis based on the excitation of semiconductor particles in the aqueous media using corresponding source of irradiation (UV or visible irradiation) (Ibhadon & Fitzpatrick 2013). First photocatalytic studies began with titanium dioxide (TiO2), which has been the most studied photocatalyst till now. TiO2 has many advantages, for instance high chemical stability, non-toxicity, availability, environmental acceptability and low prices compared to other semiconductor oxides. The most commonly studied TiO2 catalyst is commercial P25 (80% anatase structure, 20% rutile structure) from Evonik/Degussa. The bandgap energy of the TiO2 P25 is about 3.2 eV, corresponding to the wavelength of activation ≤ 390 nm. High bandgap energy and the ability to absorb only UV radiation, which consists only 5% of the solar spectrum, limits the use of TiO2 for commercial purposes (Yang et al. 2018). Therefore, there is a large number of photocatalytic investigations directed to the modification of TiO2 in order to achieve photocatalytic activity in the visible area of the solar spectrum (λ> 390 nm). These procedures include the modification of surface TiO2 by sensitization to organic dyes, polymers or surface-bound organic complexes.
and creating composites with other semiconductor materials (Rehman et al. 2009). In addition to surface modification, the activity of TiO₂ in the visible part can be achieved by narrowing the bandgap by doping/codoping TiO₂ with nonmetals and transition metals (Zaleska-Medynska 2008). Also, some studies have shown that reduction of Ti⁴⁺ to Ti³⁺ ions results in creating oxygen vacancies due to the release of oxygen ions from the band structure. These vacancies are formed between the valence and conduction bands in the TiO₂ band structure and they act as an electron captures from the valence band, which consequently enhances photocatalytic activity of TiO₂ in the visible region (Zuo et al. 2010). Reduction of TiO₂ in this way was carried out in several studies using different strategies, for instance plasma, vacuum TiO₂ heating, steam chemical vapor deposition or bombarding TiO₂ with high energy particles.

In this paper we studied pretreating of TiO₂ with UVC light in order to obtain reduced TiO₂ with enhanced photoactivity in the visible part of the sun spectrum. The obtained catalyst, the so called UVC-TiO₂ was compared to two commercial catalysts in the suspended form. Also, some experiments were performed using UVC-TiO₂ in the immobilized form.

2. MATERIALS AND METHODS

2.1. Chemicals

An analytical standard imidacloprid, used for HPLC analysis was purchased from Sigma Aldrich Company Ltd. The experiments were performed using commercial plant protection product Boxer 200 SL supplied by Sharda Ltd. Peroxotitanic acid (PTA) and nitrogen-doped TiO₂ (N-TiO₂ CCR 200 N) were donated by Cinkarna Celje. TiO₂ P25 was purchased from Evonik. Formic acid 98 % p.a. and acetonitrile HPLC grade were purchased from VWR. All aqueous solutions were prepared with ultra-pure water.

2.2. Characterization of the catalysts

The specific surface area, total pore volume and average pore diameter of the powder samples were determined with nitrogen adsorption-desorption isotherms, using the Brunauer–Emmet–Teller (BET) method on the Micrometrics ASAP 2000. The crystalline phase composition of the prepared samples was determined by X-ray diffraction measurements (Shimadzu 6000) with CuKα radiation. The UV–vis spectra of the prepared powder samples were obtained using DRS (Perkin-Elmer Lambda 35) equipped with an integrating sphere. The spectra were recorded at room temperature in the wavelength range of 200–800 nm. BaSO₄ was used as a reference.

2.3. Apparatus

The photocatalytic experiments were performed in an annular batch reactor (including recirculation in a closed system) with the source of irradiation placed in the central part of the reactor. The same reactor was used for 2-hour pretreatment of commercial TiO₂ using UVC lamp before photocatalytic reaction. The reaction mixture was recirculated using recirculating pump and the temperature was kept constant using a thermostat (Julabo–Model ED, Heating Immersion Circulator). The reaction conditions were given in the Table 1.

| Table 1. Reaction conditions |
|-----------------------------|
| **INITIAL CONCENTRATION OF IMIDACLOPRID, ppm** | 10 |
| **TEMPERATURE, °C** | 30 |
| **REACTOR VOLUME, mL** | 450 |
| **INITIAL pH OF THE SOLUTION** | 6.5-7.0 |
| **RECIRCULATION FLOW RATE, (L/min)** | 1.4 |
| **FORM OF THE CATALYST** | suspended | immobilized | without catalyst-photolysis |
| **TYPE OF THE CATALYST** | TiO₂ P25, N-TiO₂ (CCR 200 N), UVC-TiO₂ | UVC-TiO₂ | - |
| **CATALYST CONCENTRATION, (g/L)** | 0.05-0.5 | 0.036-0.110 | - |
| **IRRADIATION SOURCE** | Arcadia Natural Sunlight 8 W | Arcadia Natural Sunlight 8 W | UVC 8 W, UVA 8 W, Arcadia Natural Sunlight 8 W |
The immobilized type of the catalyst was prepared using peroxotitanic acid (PTA) as a binder. The suspension of catalyst and binder was prepared by mixing 1 g of catalyst in 100 ml of PTA. The prepared mixed solution was applied in thin layers on the glass wooling fibre. Different amount of catalyst on the support was achieved by applying several layers and drying between each layer. The immobilized TiO$_2$ layer was placed close to the reactor walls using specially designed inert photocatalyst holder. After “dark” period (establishing of the adsorption equilibrium), the reaction mixture was illuminated with the corresponding lamp. The concentration of model compound imidacloprid was analyzed at different reaction times on a 5 μm, 250 x 4.6 mm Zorbax SB-C18 column by high performance liquid chromatography (Shimadzu LC 20AT Prominence) equipped with an autosampler and diode array detector (DAD).

3. RESULTS AND DISCUSSION

3.1. Characterization of the powder catalysts

Figure 1 presents UV-Vis spectra of the commercial nitrogen doped TiO$_2$, N-TiO$_2$ (produced by Cinkarna Celje) and UVC-treated TiO$_2$ P25 (UVC-TiO$_2$). The band gap energy, $E_g$, was estimated by plotting modified Kubelka-Munk function for indirect electron transition, $(F(R)\psi)^{1/2}$ vs. photon energy ($h\nu$), the so-called Tauc’s plot, followed by extrapolation of the linear region onto the energy axis (Valencia et al. 2010). The estimated band gap energies, together with wavelength of activation, are shown in Table 2. Pretreatment of TiO$_2$ P25 with UVC radiation during 2 hours led to decreasing of $E_g$ to 2.6 eV compared to 3.2 and 3 eV as characteristic values for the anatase and rutile phase of TiO$_2$ P25, respectively. The decrease in value of $E_g$ resulted in a change of wavelength activation from 390 nm for TiO$_2$ P25 to 476 nm for UVC-TiO$_2$ and, consequently, enabled photocatalytic experiments in the visible part of solar spectrum ($\lambda > 390$ nm).

![Figure 1. Diffuse reflectance spectra of N-TiO$_2$ (black line) and UVC-TiO$_2$ (red line)](image)

The XRD patterns of the N-TiO$_2$ and UVC-TiO$_2$ are shown in Figure 2. The XRD spectra of the UVC-TiO$_2$ correspond to the spectra of TiO$_2$ P25 before UVC irradiation that was found in literature (Wang et al. 2011). It can be concluded that irradiation of TiO$_2$ P25 with UVC light did not affect the crystalline structure of the photocatalyst, but probably led to formation of oxygen vacancy states between the valence and conduction bands in the TiO$_2$ band structure, that take part in the photoexcitation process. That is, the electron may be excited to the oxygen vacancy states from the valence band even with the energy of visible light (Nakamura et al. 2000). DRS spectroscopy of N-TiO$_2$ (Fig. 1) showed that this photocatalyst is also active in the visible part of spectrum, but from the XRD spectra it is visible that this photocatalyst consists only of rutile crystal phase, which is less active than anatase (Figure 2). The lower activity of N-TiO$_2$ was confirmed while performing photocatalytic experiments, as mentioned later in the text. The results of the N$_2$ adsorption/desorption analysis showed that treating TiO$_2$ P25 with UVC light did not affect the textural properties of the catalyst (Table 3) As expected, addition of PTA as a binder for immobilization of TiO$_2$ on a glass wooling fibre lead to decreasing of specific surface area of prepared UVC-TiO$_2$/PTA sample in comparison to the pure UVC-TiO$_2$. This decrease can be explained in partial blocking of pores of TiO$_2$. The specific BET surface area for N-TiO$_2$ was higher than for TiO$_2$ P25.
Table 2. Values of indirect band gap energy for the examined TiO$_2$ photocatalysts

| MATERIAL      | $E_g$, eV | $\lambda$, nm |
|---------------|-----------|---------------|
| TiO$_2$ P25   | A: 3.2; R: 3.0 | 390           |
| UVC-TiO$_2$   | 2.6       | 476           |
| N-TiO$_2$     | 2.8       | 443           |

Figure 2. XRD spectra of N-TiO$_2$ (a) and UVC-TiO$_2$ (b)

Table 3. Textural properties of powder catalysts determined using the nitrogen adsorption/desorption analysis

| MATERIAL            | $S_{BET}$, (m$^2$/g) | $V_p$, (cm$^3$/g) | $d_p$, nm |
|---------------------|----------------------|-------------------|-----------|
| TiO$_2$ P25         | 56                   | 0.25              | 17.50     |
| UVC-TiO$_2$         | 54.28                | 0.27              | 20.22     |
| UVC-TiO$_2$/PTA (50 g/L TiO$_2$) | 17.93 | 0.05 | 11.44 |
| N-TiO$_2$           | 87.92                | 0.47              | 21.40     |
3.2. Testing of the photocatalytic properties

The photolytic and photocatalytic activity of the suspended and immobilized TiO$_2$ photocatalysts is evaluated by photodegradation of imidacloprid in the simulated sunlight conditions. In order to study the contribution of photolysis on the degradation of imidacloprid, three different irradiation sources were examined. The results showed that the lamp of interest (Arcadia natural sunlight), that closely simulates sun spectrum, did not degrade imidacloprid even after 3.5 hours of irradiation in an annular photoreactor. On the other side, the strong UVC light completely degraded imidacloprid after 1 hour of irradiation (Figure 4).

Further experiments were performed using simulated sunlight under the presence of photocatalysts. Three TiO$_2$ catalysts in the suspended form were compared in the same reaction conditions and it was shown, according to our expectations, that the best efficiency was obtained using UVC-TiO$_2$ (Figure 5). N-TiO$_2$ was not efficient in our reaction system, although the same photocatalyst exhibited excellent photocatalytic activity in the presence of natural or artificial UV and visible light for the decomposition of nitrogen monoxide (NO).

Furthermore, the influence of different concentration of UVC-TiO$_2$ photocatalyst on the degradation efficiency was examined. The higher catalyst concentration led to greater decrease in the imidacloprid concentration (Figure 6). This is obvious since the higher catalyst concentration leads to more photocatalytically active sites available for the photoexcitation process. Some authors have reported on the optimal catalyst concentration in the suspended form that gives the best degradation efficiency. In these studies higher concentration than optimal led to decrease in the degradation rate of model pollutant (Kitsiou et al. 2009). This was explained with possible agglomeration of TiO$_2$ particles and decreased system opacity, which consequently blocked reaching of light to the surface of catalyst particles (Affam & Chaudhuri 2013). This phenomenon was not noticed in the concentration range of TiO$_2$, studied in this paper (0.05 – 0.5 g/L).
Afterwards, the immobilized UVC-TiO$_2$ was evaluated in the photocatalytic degradation of imidacloprid in the same reaction conditions as in the suspended form (concentration of imidacloprid, irradiation source, recirculation flow rate and reaction temperature). „Dark“ experiment during 1 hour showed that the adsorption process did not contribute to decrease of imidacloprid concentration (Figure 7).

The influence of different number of layers of immobilized UVC-TiO$_2$ that consequently led to different amount of catalyst on the support, was examined (Table 1). It was shown that there was no significant change in the degradation rate with catalyst concentration in the prepared layer (Figure 8). The degradation rate obtained with the immobilized TiO$_2$ was much lower than with suspended TiO$_2$ as expected. Regardless of the lower efficiency, the prepared immobilized layer of TiO$_2$ showed good stability and no Ti leaching. Peroxotitanic acid as a binder ensured good adherence strength of the immobilized photocatalyst layer to the substrate surface.
4. CONCLUSION

The UVC-treated TiO$_2$ photocatalyst showed the higher visible light activity for imidacloprid removal among the examined TiO$_2$ catalysts under simulated sunlight conditions. Neither the crystal structure nor the specific surface area was observed to change between the commercial TiO$_2$ P25 and UVC-treated TiO$_2$ P25 photocatalyst. The appearance of the visible light photocatalytic activity in UVC-treated TiO$_2$ is due to a new photoexcitation process by the formation of the oxygen vacancy state located between the valence and the conduction bands. This phenomenon will be confirmed performing ESR measurements in the future work. The imidacloprid degradation rate increased with the increase in the catalyst concentration in the studied concentration range. The immobilized UVC-treated TiO$_2$ gave satisfying results in terms of stability, activity and reuse.

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