Surface-Enriched Boron-Doped TiO$_2$ Nanoparticles as Photocatalysts for Propene Oxidation

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ABSTRACT: A series of nanostructured boron-TiO$_2$ photocatalysts (B-X-TiO$_2$-T) were prepared by sol–gel synthesis using titanium tetraisopropoxide and boric acid. The effects of the synthesis variables, boric acid amount (X) and crystallization temperature (T), on structural and electronic properties and on the photocatalytic performance for propene oxidation, are studied. This reaction accounts for the remediation of pollution caused by volatile organic compounds, and it is carried out at low concentrations, a case in which efficient removal techniques are difficult and costly to implement. The presence of boric acid during the TiO$_2$ synthesis hinders the development of rutile without affecting the textural properties. X-ray photoelectron spectroscopy analysis reveals the interstitial incorporation of boron into the surface lattice of the TiO$_2$ nanostructure, while segregation of B$_2$O$_3$ occurs in samples with high boron loading, also confirmed by X-ray diffraction. The best-performing photocatalysts are those with the lowest boron loading. Their high activity, outperforming the equivalent sample without boron, can be attributed to a high anatase and surface hydroxyl group content and efficient photo-charge separation (photoelectrochemical characterization, PEC), which can explain the suppression of visible photoluminescence (PL). Crystallization at 450 °C renders the most active sample, likely due to the development of a pure anatase structure with a large surface boron enrichment. A shift in the wavelength-dependent activity profile (PEC data) and the lowest electron–hole recombination rate (PL data) are also observed for this sample.

KEYWORDS: photocatalysis, titanium dioxide, boron, interstitial doping, propene oxidation, photoelectrochemical characterization

1. INTRODUCTION

Environmental legislation drives increasing emphasis on removing toxic organic pollutants, like volatile organic compounds (VOCs), from air. Among VOCs, propene is one of the main components of tobacco smoke, and it is present in vehicle emissions and exhaust gases from several industries (petrochemicals, foundry processes, etc.). It has important harmful effects on human health, even at low concentrations. Photocatalysis has shown to be one of the most suitable techniques to successfully degrade organic pollutants both in the gas and the liquid phase (leading to its complete mineralization). These properties and issues confer to TiO$_2$ advantages over other semiconductors like ZnO, CeO$_2$, CdS, ZnS, and so forth. However, its photocatalytic efficiency is hindered by the fast recombination of photogenerated holes and electrons and its relatively wide band gap (3.2 eV) that limits its activity to the near-UV region and, thus, only about 5% of solar irradiation can be used. Many research studies have focused on combining TiO$_2$ with other elements or compounds with the purpose of increasing the TiO$_2$ efficiency.

Recently, boron incorporation into nanostructured TiO$_2$ has been proposed as an alternative to overcome some of the drawbacks mentioned above. A potential benefit of the presence of boron is the creation of electronic levels that could hinder the recombination of the electron–hole pairs. Density functional theory calculations suggest that the occupation of interstitial sites or O substitution are more energetically favorable than substitution of Ti by B atoms. It has been reported that the addition of boron to TiO$_2$ sol–gel precursors generally leads to interstitial B-doping, together with the formation of B$_2$O$_3$ on the surface of...
the TiO₂ particles.⁶,¹⁹,²⁷,²⁸ Some studies have also shown that surface boron species can introduce residual charge, which can increase the number of surface OH groups, being them correlated with the incorporated B.¹⁹,²⁹

The positive influence of boron has also been related to changes in the TiO₂ structure as it could inhibit the growth of crystalline TiO₂, leading to materials with increased surface area and anatase content.¹⁹,²⁰

Several studies have reported a photoactivity improvement upon B doping of TiO₂,⁶,¹⁹,²⁰–²⁶ but the interpretation is still controversial, especially regarding the local structure around the boron impurity and the electronic effects of the dopant species.¹⁶,¹⁸,¹⁹ Zhang et al. reported highly efficient photocatalysts consisting of B-TiO₂ (3.5 at. B %) for gas phase degradation of benzene under UV–vis light, which are more active than the commercial TiO₂-P25. This improvement is ascribed to a significant enhancement of the UV–vis light absorption.²² Anson-Casas et al. prepared B-TiO₂ photocatalysts by sol–gel synthesis using boric acid,¹⁹ leading to a slightly improved photocatalytic efficiency in the degradation of diphenhydramine (drug pollutant model) in water. UV–vis diffuse reflectance analysis allowed to discard visible light absorption, and the activity improvement was attributed to the crystalline structure and composition of the photocatalysts. Jeong et al.²⁰ prepared a boron-doped TiO₂ anode (sol–gel, 2, 5, and 10 wt % B) for lithium-ion batteries, and they found a significant improvement with respect to pristine TiO₂, which was attributed to the increased interplanar spacing of the TiO₂ lattice due to interstitial B atoms and/or to a larger amount of surface hydroxyl groups. Doozzi et al. studied B- and F-doped TiO₂ materials, and they found that the photodegradation of formic and acetic acid (in the liquid phase) can be correlated with the photocatalyst’s structural features. The higher activity of the doped TiO₂ with respect to the bare one was attributed to its higher surface area (from 45 to 100 m²/g) and the hindered transformation of anatase into rutile, whereas the surface B₂O₃ formed (observed by XPS) had no effect on the photoactivity.²⁷

Many of the indicated effects can be positive, enhancing the photocatalytic activity of TiO₂ for propene oxidation, which is the target reaction of the present study.

With the aim of shedding light on the role of boron in the physicochemical properties of TiO₂ modified with this element, the present research focuses on the study of nanostructured B-TiO₂ samples, prepared by sol–gel synthesis using titanium tetraisopropoxide and different amounts of boric acid and crystallization temperatures. A detailed characterization has been carried out using a plethora of techniques (gas adsorption, X-ray diffraction (XRD), X-ray photoelectron spectroscopy (XPS), UV–vis spectroscopy, analysis of photoluminescence (PL), photoelectrochemical characterization (PEC), and Raman scattering). The prepared photocatalysts have been tested in the gas phase photocatalytic oxidation of propene (100 ppmv), which accounts for remediation of pollution caused by low-concentration VOCs, a case in which efficient removal techniques are difficult and costly to implement.³,⁴,²³

The general mechanism of propene oxidation involves the following steps (where R represents the organic substrate)

\[
\text{catalyst} + h\nu \rightarrow h^+ + e^- \quad (1)
\]

\[
\left( O_2 \right)_{ads} + e^- \rightarrow O_2^- \quad (2)
\]

\[
H_2O + h^+ \rightarrow ^\circ OH + H^+ \quad (3)
\]

\[
OH_{ads} + h^+ \rightarrow ^\circ OH \quad (4)
\]

\[
R - H + ^\circ OH \rightarrow R^+ + H_2O \quad (5)
\]

\[
R^+ + h^+ \rightarrow R^{*+} \rightarrow \text{mineralization products} \quad (6)
\]

Reactions 5 and 6 are an example of the possible interaction of the substrate with oxidant species. In a general way: \[R - H + ^\circ OH/O_2^-/h^+ \rightarrow \text{mineralization products} \]

In the case of propene oxidation, the global reaction can be written as

\[
2C_3H_6 + 9 O_2 \rightarrow 6CO_2 + 6H_2O \quad (7)
\]

The thorough characterization of the B-TiO₂ samples, using techniques that allow an approach to the photocatalyst properties from very different points of view and that will help to find a suitable correlation between properties and activity of the photocatalysts, is one of the main novelties of this study.

### 2. EXPERIMENTAL SECTION

#### 2.1. Materials. Titanium(IV) isopropoxide (TTIP, C₃H₆O₄Ti, 97%, Sigma-Aldrich), boric acid (H₃BO₃, 99.9%, Gilca), and 2-propanol (C₃H₇O₃, >99.8%, Panreac AppliChem) were used as received. The well-known standard Aeroxide P25 TiO₂ photocatalyst from Degussa was used as a reference.

#### 2.2. Synthesis of B-TiO₂ Photocatalysts. The synthesis procedure is based on a previously optimized sol–gel method.³⁰ In a typical preparation, 5 mL of TTIP were diluted in 50 mL of isopropanol and added dropwise, under constant stirring, to 50 mL of a water/isopropanol (1:1) mixture containing a certain quantity of boric acid. The resulting gel was aged overnight, filtered, and dried in an oven at 90 °C. The as-prepared amorphous TiO₂ was thermally treated in a horizontal quartz reactor under an ambient atmosphere to induce crystallization. The obtained materials are labeled as B-X-TiO₂-T, where X refers to the amount of boric acid (mg) added during the synthesis and T (°C) to the crystallization temperature. Scheme 1 illustrates the preparation and structure of the catalysts.

![Scheme 1. Illustration of the Photocatalysts Preparation and Composition/Structure](https://www.acsanm.org/doi/suppl/10.1021/acsnanm.2b02217/suppl_file/12528ACSappnanomat2022512527-12539.pdf)

The nominal B content in the resulting samples was estimated taking into account the yield of the TiO₂ synthesis and assuming that all the boron from the added boric acid remains in the final product. More specifically, the average hydrolysis yield for 5 mL TTIP was 1.5 ± 0.1 g, and the yield of the thermal crystallization treatment was 81.5 ± 1.5%. Therefore, the boron content ranges from 0.07 to 3.4 wt %.

#### 2.3. Characterization. N₂ adsorption–desorption at −196 °C (QuantaChrome Autosorb-6B Instrument) after sample degasification (250 °C, 4 h) was used to characterize the porous texture (the apparent Brunauer– Emmett–Teller (BET) specific surface area \(S_{BET}\) and the total micropore volume \(V_{DR} N_0 \phi < 2 \text{ nm}\) applying BET and Dubinin Radushkevich equations, respectively).† An estimation of the mesopore volume \(2 \phi < \phi < 50 \text{ nm, } V_{meso}\)
was calculated as \( V_{\text{ads}} (P/P_0 = 0.9) - V_{\text{N_2}} (P/P_0 = 0.2) \), expressed as liquid. \(^{34}\) Total pore volume \( (V_p) \) was determined as the volume of nitrogen adsorbed at \( P/P_0 = 0.99 \) (as liquid).

XRD measurements were carried out in a MiniFlex II Rigaku (30 kV/15 mA) equipment at room temperature, with Cu Kα radiation, 2°/min scanning rate, and 6–80° 2θ range. The average crystallite size \( (B) \) was calculated with the Scherrer equation:

\[
B = \frac{K\lambda}{\beta \cos \theta}
\]

(8)

where \( \lambda \) is 0.1540 nm (Cu Kα radiation wavelength), \( K \) is a constant (taken as 0.93 considering spherical grains), \(^{36}\) and \( \beta \) and \( \theta \) are, respectively, the full width at half-maximum intensity (FWHM) and the diffraction angle of the main peak for each crystalline phase.

Crystalline TiO\(_2\) was characterized and quantified using XRD data of 50/50 (wt/wt) B-TiO\(_2\)/CaF\(_2\) mixtures, as described in ref 37. Further explanations on the calculation of total crystallinity and percentage of crystalline phases are reported in Section 3.1.

High-resolution transmission electron microscopy (HRTEM) in combination with an energy-dispersive X-ray spectroscopy EDX (FEI Tecnai G2 system with a Schottky field emission electron gun operated at 200 kV) were used for characterizing TiO\(_2\) microstructure and chemical composition in some prepared samples. For that purpose, TiO\(_2\) nanopowders were suspended in ethanol and dispersed on TEM grids.

XPS was measured in a VG-Microtech Multilab 3000 spectrometer with an Al anode working at 6 mA and 12 kV and with a pass energy of 50 eV. Binding energies were referenced to the C 1s line at 284.6 eV. \(^{38}\)

PL and Raman scattering spectra of cold-pressed pellets of the powder catalysts were recorded using a laser diode as the excitation source \( (\lambda_{\text{exc}} = 405 \text{ nm}) \) and a \( \times50 \) microscope objective to achieve an irradiance of about 10\(^6\) W/m\(^2\). The laser light was rejected by a longpass edge filter, which cutoff all signals below 200 cm\(^{-1}\). Spectra were recorded with a HORIZA Jobin Yvon iHR320 spectrometer equipped with a Peltier-cooled CCD detector. All measurements were carried out at room temperature and atmospheric pressure with the samples in contact with air.

Diffuse reflectance measurements were carried out on a Shimadzu UV–vis 2501PC spectrophotometer equipped with an integrating sphere. BaSO\(_4\) was used as reference material and for background measurement. The measurement range studied was 220–900 nm, with a 1.0 nm resolution and a spectral bandwidth of 5 nm. The spectra recorded were transformed using the Kubelka–Munk function.

For the PEC characterization, powder photocatalysts were suspended in isopropanol using an ultrasonic bath and immediately deposited on fluorinated tin oxide (Solems SA, 80 \( \Omega/\text{sq} \)) substrates by spray coating. \(^{39}\) The as-prepared films were sintered at 450 °C for 2 h under ambient air. Specimens were connected as working electrodes in a three-electrode configuration, with a graphite bar as the counter electrode, the Ag/AgCl, 3 M NaCl \( (E^\circ = 0.210 \text{ V vs SHE}) \) as the reference electrode, and 0.1 M Na\(_2\)SO\(_4\) as the electrolyte. The electrolyte was deoxygenated by flowing N\(_2\) for longer than 10 min before the measurements. The glass PEC cell was provided with a quartz window and irradiated with a laboratory-scale solar simulator (150 W Xe arc lamp by LOT-Quantum Design GmbH). The simulated solar spectrum was the AM1.5G with a maximum irradiance of approximately 100 mW/cm\(^2\) (see Figure S1A). The wavelength-dependent response was measured using a LOT Quantum Design monochromator (MSH-300).

2.4. Photocatalytic Oxidation of Propene. The photocatalytic tests of propene oxidation were carried out in an experimental setup that consists of a quartz reactor (AFORA) and a UV-A lamp (Philips, TL8 W/05 FAM, \( \lambda_{\text{max}} = 365 \text{ nm} \)) located parallel to the photocatalyst at a distance of about 1 cm. Some experiments have been carried out using a visible lamp (Sylvania, BW8/TS/54–765, Daylight). The spectra of both lamps can be seen in Figure S1, Supporting Information. A scheme of the setup is shown elsewhere,\(^{3,43}\) and a detailed description of the experimental conditions has been previously reported by Cano-Casanova et al.\(^{40,41}\) The UV-A irradiance of the Philips lamp was determined at 1 cm distance with a portable Delta Ohm photo radiometer (model HD 2102.2) equipped with a UV-A probe \( (LP 471 \text{ UVA}) \), being 2.24 mW/cm\(^2\).

In a typical experiment, the photocatalyst (0.11 g) was placed on a quartz wool plug inside the reactor and, after purging with helium \( (60 \text{ mL/min}) \), a stream of propene in air \( (100 \text{ ppmv propene, 30 or 60 mL/min (STP)}) \) was passed through the reactor at 25 °C. The outlet stream was continuously monitored by mass spectrometry (ThermoStar GSD 301 O1, from Balzers), and once the signal was stabilized, the UV-lamp was switched on, and the illumination was maintained for 3 h. The experiments were repeated at least twice to check reproducibility.

Blank experiments (in the absence of the catalyst) and measurements in dark show that propene conversion does not occur in these conditions.

Propene conversion was calculated, as shown in eq 9

\[
\text{propene conversion (\%) = } \frac{C_{\text{initial}} C_2H_6 - C_{\text{stationary}} C_2H_6}{C_{\text{initial}} C_2H_6} \times 100
\]

(9)

where \( C_{\text{initial}} C_2H_6 \) is the initial propene concentration (100 ppmv) and \( C_{\text{stationary}} C_2H_6 \) is the stationary propene concentration under illumination (3 h reaction time). This is illustrated in Scheme 2.

**Scheme 2. Drawing of the Variation of the Propene Concentration under Illumination (in Presence of a Photocatalyst), Parameters as Indicated Above.**

Reaction products, i.e., carbon dioxide and water, were also analyzed by mass spectrometry. A calibration cylinder containing 300 ppmv CO\(_2\) in He was used for CO\(_2\) quantification. Mass scan measurements (up to m/z = 100) allowed to discard any additional oxidation compound, and the total mineralization of propene was confirmed by the carbon balance,\(^{43}\) according to the following reaction

\[
2C_2H_6 + 9O_2 \rightarrow 6CO_2 + 6H_2O
\]

This condition was verified for each experiment.

3. RESULTS AND DISCUSSION

This section includes the results of different characterization techniques with the aim of analyzing: (i) the effect of the amount of boric acid used and (ii) the effect of the crystalization temperature, on the photocatalytic activity.

3.1. Characterization of the Prepared Samples. 3.1.1. XRD Analysis and Textural Properties. XRD patterns of the B-X-TiO\(_2\)-550 samples \( (\text{Figure 1}) \) show the characteristic peaks of anatase \( (2 \theta \text{ values of } 25.3^\circ (101), 37.8^\circ (004), 48.0^\circ (200), 54.5^\circ (105), 55.0^\circ (211), 62.7^\circ (204), 70.4^\circ (116), \text{ and } 74.5^\circ (220)) \) and rutile \( (2 \theta \text{ values of } 27.5^\circ (110), 36.1^\circ (101), \text{ and } 54.4^\circ (211)) \). B-100-TiO\(_2\)-550 and B-230-TiO\(_2\)-550 samples show an additional peak at \( 2 \theta = 28.1^\circ \), which corresponds to boric oxide \( (B_2O_3) \).\(^{46,47}\) (Inset of Figure 1), resulting from the thermal decomposition of boric acid.
The XRD profiles of the B-X-TiO$_2$-550 series (X ≠ 0) show that boron incorporation does not take place in the bulk TiO$_2$ crystal lattice and, thus, it can be considered that boron atoms are not extensively inserted in the anatase or rutile unit cells. The calculated percentage of crystalline and amorphous phases and the average crystallite size for all the prepared photocatalysts and P25 are listed in Table 1. The percentages of crystalline and amorphous TiO$_2$ determined from the XRD pattern.

Thus, the percentages of crystalline and amorphous TiO$_2$ have been calculated using the following equations:

\[ A_{\text{Cryst}} = \frac{A_{\text{Anatase, 101}}}{A_{\text{CaF$_2$, 220}}} \times 100 \times 1.25 \]  
\[ R_{\text{Cryst}} = \frac{A_{\text{Rutile, 110}}}{A_{\text{CaF$_2$, 220}}} \times 100 \times 0.90 \]

where \( A_{\text{Anatase, 101}} \), \( A_{\text{Rutile, 110}} \), and \( A_{\text{CaF$_2$, 220}} \) are the areas determined from the XRD pattern.

Thus, the percentages of crystalline and amorphous TiO$_2$ are, respectively, \( W_{\text{Cryst}} = A_{\text{Cryst}} + R_{\text{Cryst}} \) and \( W_{\text{Am}} = 100 - W_{\text{Cryst}} \).

These data show that, in general, the presence of boron acid during the TiO$_2$ synthesis hinders the formation of rutile, stabilizing up to some extent the anatase metastable phase, in agreement with the literature. Although differences between the B-X-TiO$_2$-550 samples regarding phase composition are small, it can be highlighted that B-5-TiO$_2$-550 is the one with the highest anatase and lowest rutile content, whereas B-50-TiO$_2$-550 has the lowest anatase content and the highest proportion of amorphous TiO$_2$.

The specific surface area \( (S_{\text{BET}}) \) of the B-X-TiO$_2$-550 samples is about half that of P25, ranging from 24 to 33 m$^2$/g (see complete textural properties in Table S1, Supporting Information), and it does not significantly vary with the amount of boron present in the sample.

### 3.1.2. HRTEM–EDX Characterization

The comparative TEM–HRTEM analysis of samples B-0-TiO$_2$-550 and B-5-TiO$_2$-550 (see Supporting Information, Figures S2 and S3) shows that the TiO$_2$ nanoparticles have a similar particle size (50 nm average size), but some interesting differences in the particles structure have been observed. Thus, in sample B-0-TiO$_2$-550 both monocrystal and some polycrystalline nanoparticles are present, and most of them also show amorphous domains, whereas B-5-TiO$_2$-550 contains mainly monocrystals.

EDX analysis (Figure S2) has shown no differences between B-5-TiO$_2$-550 and B-0-TiO$_2$-550 samples. The lack of detectable signal at the K-alpha emission of boron (expected at 0.183 keV) in the spectrum of B-5-TiO$_2$-550 in not surprising due to the low boron content in this sample and the proximity of the K-alpha peak of C (carbon is present in the TEM grids and produces a strong peak).

### 3.1.3. XPS Analysis

The XPS spectra survey of the B-X-TiO$_2$-550 samples series is presented in Figure S4. Figure 2a–c shows the XPS spectra of the B-X-TiO$_2$-550 photocatalysts for B 1s, O 1s, and Ti 2p, respectively (the latter in the 461–456 eV range, see full-range Ti 2p spectra in Figure S5). The corresponding binding energy (BE) values are collected in Table 2. Note that these values correspond to the maxima of the experimental spectra except for O 1s, for which three binding energies have been obtained from the fitted deconvoluted spectra of each oxygen contribution.

As shown in Figure 2a and in Table 2, the B 1s BE is around 192.1 eV in B-5-TiO$_2$-550, B-10-TiO$_2$-550, and B-50-TiO$_2$-550 samples, while it is 192.8 eV in B-100-TiO$_2$-550 and B-230-TiO$_2$-550. As the B 1s BE in B$_2$O$_3$ is 192.9 eV,\(^{46,47}\) it can be assumed that boron is largely present as B$_2$O$_3$ on the surface of the B-100-TiO$_2$-550 and B-230-TiO$_2$-550 samples, which is in agreement with the presence of B$_2$O$_3$ determined by XRD (see Figure 1). In the case of the B-5-TiO$_2$-550, B-10-TiO$_2$-550, and B-50-TiO$_2$-550 samples, boron is present in a less deficient electronic state, which can be related to a lower interaction with electronegative atoms such as O. Quesada-González et al.\(^{48}\) in their study of B-TiO$_2$ samples reported that the B 1s BE of about 192 eV corresponds to interstitial B and that a peak at 192.6 eV can be attributed to Ti$_2$O$_3$, or B$_2$O$_3$. The assignment of a 191.8 eV BE to interstitial B is also reported by Feng et al.\(^{21}\) and Zhang et al.,\(^{22}\) who also found a 191.9 eV BE for B$_2$O$_3$ samples. They proposed that B is incorporated into the TiO$_2$ lattice and that its chemical environment might be Ti--O--B.\(^{22}\)

These results evidence the presence of interstitial B in our photocatalysts, being easily detected by XPS analysis on the surface of the samples with lower B loading, whereas it could probably be covered by B$_2$O$_3$ in the B-100-TiO$_2$-550 and B-230-TiO$_2$-550 samples. In the study of Quesada-González et al.,\(^{48}\) the interstitial boron was detected by in-depth XPS analysis.

### Table 1. Crystalline Properties and Surface Area in the B-X-TiO$_2$-550 Series and P25

| sample       | crystalline TiO$_2$ (%) | amorphous TiO$_2$ (%) | average crystallite size (nm) | A | R | S$_{\text{BET}}$ (m$^2$/g) |
|--------------|------------------------|-----------------------|------------------------------|---|---|-----------------|
| B-0-TiO$_2$-550 | 65 12                  | 23 28                 | 32 28                        | 28 | 28 | 32              |
| B-5-TiO$_2$-550 | 75 2                   | 23 25                 | 19 23                        | 33 | 33 | 33              |
| B-10-TiO$_2$-550 | 70 6                   | 24 25                 | 27 25                        | 30 | 30 | 30              |
| B-50-TiO$_2$-550 | 56 6                   | 38 25                 | 29 24                        | 24 | 24 | 24              |
| B-100-TiO$_2$-550 | 66 5                   | 29 24                 | 23 23                        | 27 | 27 | 27              |
| B-230-TiO$_2$-550 | 70 7                   | 23 27                 | 25 25                        | 25 | 25 | 25              |
| P25          | 73 14                  | 13 22                 | 28 55                        | 55 | 55 | 55              |

*A = anatase, R = rutile.*

Figure 1. XRD spectra for B-X-TiO$_2$-550 samples. Inset image: amplification of Figure 1 in the 24–30° 2θ range.
The O 1s spectra show that oxygen is present in different electronic states. In the case of B-0-TiO$_2$-550, the main peak is due to O bonded to Ti (at 529.6 eV, TiO$_2$ lattice oxygen), and a shoulder (at 531.1 eV) corresponding to O in surface OH groups can be observed [identified as O(I) and O(II), respectively] in Table 2.

For the boron-containing samples, apart from the main O−Ti peak, a two-component band, with its maximum at BE ≥ 532 eV, can be observed corresponding to O in surface OH groups. Such a band has been deconvoluted to distinguish these two contributions (Figure 3), and the one assigned to O−B has been identified as O(III) in Table 2. As in the case of the B1s data, there are differences between B-100-TiO$_2$-550 and B-230-TiO$_2$-550 samples on one side, and B-5-TiO$_2$-550, B-10-TiO$_2$-550, and B-50-TiO$_2$-550 on the other side, both in the intensity and in the BE of these O−B species. In agreement with the higher B content, the proportion of O bonded to B species (O(III)) is noticeably higher in the B-100-TiO$_2$-550 and B-230-TiO$_2$-550 samples and, in line with B 1s BE and XRD data, these species largely correspond to B$_2$O$_3$ on the TiO$_2$ surface. In the case of the B-5-TiO$_2$-550, B-10-TiO$_2$-550, and B-50-TiO$_2$-550 samples, the O−B bonds should correspond to the surface interstitial species present in these samples.

The BE of Ti 2p$_{3/2}$ in sample B-0-TiO$_2$-550 (458 eV) shows that, as expected, titanium is present as Ti$^{4+}$ (Figures 2 and 3).
For the B-X-TiO$_2$-550 series of samples, the BE is slightly shifted to higher values, indicating that the presence of boron reduces the electron density of Ti 2p. This can be a consequence of the proximity of boron atoms (more electronegative than Ti) located in interstitial positions of the TiO$_2$ structure. The displacement seems to increase with the boron amount, being related to more boron atoms located in interstices. It is important to point out that the distortion of the TiO$_2$ lattice produced as a consequence of boron insertion has been detected by means of XPS data and not by XRD.

Figure 3. O 1s XPS deconvoluted spectra for the B-X-TiO$_2$-550 samples.
which indicates that it is not a bulk extensive phenomenon. Then, it could be assumed that it happens mainly in a surface region. The $\text{B}_2\text{O}_3$ identification by XRD analysis allows considering that only a certain amount of boron can be hosted in the TiO$_2$ lattice, and the rest is deposited on the surface of the TiO$_2$ nanoparticles as $\text{B}_2\text{O}_3$.

3.2. Photocatalytic Activity. Figure 4 shows propene conversion data obtained with the two flow rates tested.

![Figure 4](image-url)

Figure 4. Propene conversion (%) at 30 and 60 mL/min for B-X-TiO$_2$-550 photocatalysts and P25.

Photolysis does not occur, and these results indicate that all the prepared photocatalysts are active, being two of them more active than the reference P25. Furthermore, the same trend is found with the two propene flow rates, which proves the reproducible performance of these catalysts. The lower flow rate leads to a higher conversion because of the larger contact time and/or the lower amount of propene molecules to be oxidized per unit of time.

The catalyst stability has been stated after confirming that the MS propene signal remains stable during the whole experiment. As no decay was observed at all, it can be assumed that the catalysts remain stable.

The order in the activity of B-X-TiO$_2$-550 samples is: B-5-TiO$_2$-550 > B-10-TiO$_2$-550 > B-0-TiO$_2$-550 > B-50-TiO$_2$-550 > B-100-TiO$_2$-550 > B-230-TiO$_2$-550.

Samples B-5-TiO$_2$-550 and B-10-TiO$_2$-550 are more active than the prepared bare TiO$_2$ (sample B-0-TiO$_2$-550) and also than P25. However, B-X-TiO$_2$-550, with $X = 50, 100,$ and 230 samples are less active, and the photocatalytic activity decreases as the amount of boric acid used increases, being sample B-230-TiO$_2$-550 practically inactive. These results imply that interstitial boron doping likely leads to electronic modifications that improve the TiO$_2$ photoactivity. However, above a certain amount of boron, there is an excess that segregates as $\text{B}_2\text{O}_3$ and partially covers the interstitially boron-doped TiO$_2$ surface, with the consequent photocatalytic activity decrease. Thus, the very good photocatalytic performance, shown by B-5-TiO$_2$-550 and B-10-TiO$_2$-550, seems to be related to the low, but suitable, quantity of boron present and its location. The B-5-TiO$_2$-550 sample contains the highest anatase and the lowest rutile amount, and it has a certain content of surface interstitial boron. These characteristics likely enhance the photocatalytic propene oxidation.

Some selected samples were also tested under visible light. The results are discussed in Section 3.3, but it can be anticipated that the activity is significantly lower. Propene conversion (using 30 mL/min propene flow) of sample B-5-TiO$_2$-550 decreases from 74% (under UV light) to 18% (under visible light) (compare data in Figures 4 and S10).

In order to better understand and to explain why the photocatalysts that contain the lowest amount of boron show the best photocatalytic activity, they were further characterized by means of PEC and PL measurements.

3.2.1. PEC and PL Characterization of the Most Active Samples. Figure 5 shows the analysis by cyclic voltammetry (CV) in the dark and under irradiation in a solar simulator. In the dark, the shape of the CV scans reflects the accessibility of electronic energy states that are near the conduction band level, which is located at around $-1$ V versus Ag/AgCl. The CV plot of P25 is quite different from that of the B-X-TiO$_2$-550 samples tested. Specifically, the hysteresis of P25 at potentials below $-0.7$ V versus Ag/AgCl is wider, which can be associated to the higher $S_{\text{BET}}$ and, thus, to a larger charge accumulation on the surface. Besides, trap states around $-0.5$ V are more clearly defined for P25, in particular in the...
Raman scattering spectra in detail (the scale has been changed to wavenumbers, and intensity is represented in the logarithmic scale, for clarity).

observed at values and, thus, similar charge accumulation capacities.

backward (negative currents) scan. All the B-X-TiO$_2$ samples tested show a similar scan shape, as all of them have close $S_{\text{BET}}$ values and, thus, similar charge accumulation capacities.

Under full solar spectrum irradiation, a photocurrent is observed at $E > -0.35$ V for the tested B-X-TiO$_2$-550 samples, being the highest one for B-5-TiO$_2$-550 and B-0-TiO$_2$-550 anodes. Besides, both electrodes show a particular distortion in the charge accumulation region, with a clear feature at around $-0.7$ V (positive currents scan). This feature points to the creation under irradiation of certain surface states close to the conduction band, being particularly pronounced for B-5-TiO$_2$-550, which could explain the better photocatalytic performance of this sample.

Figure 6a shows the PL spectra obtained for B-0-TiO$_2$-550, B-5-TiO$_2$-550, and P25, with an excitation source of 405 nm (see Figure S6 for the PL spectra of the complete B-X-TiO$_2$-550 series). This excitation energy is slightly lower than rutile or anatase TiO$_2$ band-gap energies and, therefore, the absorption should correspond to transitions from the valence band to shallow trap states that appear as an exponential continuous tail just below the conduction band, being associated to structural disorder and unbalanced Ti atoms on the surface.\(^{30}\) The PL spectrum of P25 shows a wide band, which can be deconvoluted into three different peaks centered at 442 nm (blue), 530 nm (green), and 600 nm (yellow-red), respectively. The blue peak is attributed to transitions between the shallow continuous states and the valence band, while the green and red PL peaks involve deep intermediate gap states originated by intrinsic point defects, such as oxygen vacancies, and are typical of anatase TiO$_2$.\(^{50,51}\) In nanostructured TiO$_2$, oxygen vacancies are located close to the surface, and their densities can be affected by environmental conditions. It has been widely reported that O$_2$ adsorption results in a significant reduction of the PL intensity of anatase TiO$_2$, and the opposite occurs upon O$_2$ desorption.\(^{19}\)

Sample B-0-TiO$_2$-550 presents a PL spectra similar to that of P25, but with a more intense blue band, indicating a less efficient charge relaxation toward trapped states near the surface. On the other hand, B-5-TiO$_2$-550 exhibits a drastic quenching of all PL bands (red line in Figure 6a). The absorbance spectra of these two samples (see Figure S7) are very similar, indicating that absorption is not affected by B incorporation (absorption in the visible region does not occur), ruling out one of the possible mechanisms by which B can modify the activity of TiO$_2$ (substitution of oxygen atoms would produce the TiO$_2$ band gap narrowing). However, the mentioned PL quenching reveals an overall reduction in the recombination rate of photo-excited electrons and holes, which can be explained by a more efficient charge separation and transfer toward the surfaces.\(^{52-55}\) This supports the interpretation of its highest photocatalytic activity. The suppression of PL allows the observation of several sharp features present in the blue region. These findings correspond to the resonant Raman spectrum of the sample and have been plotted in more detail in Figure 6b. The Raman spectrum is dominated by three very intense peaks (below 1000 cm$^{-1}$) labeled as the principal vibrational modes of anatase TiO$_2$. The low-frequency $E_g$ mode of anatase is not detected as it has been cut off by the Raman edge filter. There are no traces of rutile modes, further confirming that anatase is the most abundant crystallographic phase in these materials. Similar peaks are observed in the spectrum of sample B-0-TiO$_2$-550 but, surprisingly, not in the case of P25, which shows no resonant Raman signal. Finally, sample B-5-TiO$_2$-550 exhibits two additional Raman bands at higher wavenumbers. These bands, which are not seen in the Raman spectrum performed with green laser excitation (not shown), have been linked to vibrational H--O--H bending modes at 1600 cm$^{-1}$ and to the stretching vibrational modes of hydroxyl groups between 3000 and 3500 cm$^{-1}$.\(^{56}\) These observations would indicate, on the one hand, the chemisorption of water molecules and, on the other hand, the production of hydroxyl radicals under UV irradiation.\(^{57}\) The presence of the latter band only in the Raman spectrum of sample B-5-TiO$_2$-550 could be linked to its higher photocatalytic efficiency. Although the presence of boron had no measurable effect in band gap narrowing, the highest photocatalytic performance of this sample could be linked to its best photocurrent response ability, the additional production of hydroxyl radical groups, and the lowest recombination rate of electron/hole pairs.
Table 3. Physicochemical Properties and Propene Conversion for the B-5-TiO$_2$-T Samples and P25

| Sample          | Crystalline TiO$_2$ (%)$^a$ | Amorphous TiO$_2$ (%)$^a$ | Average Crystallite Size (nm)$^b$ | BET (m$^2$/g)$^c$ | B at. (%)$^d$ | Propene Conversion (%)$^e$ |
|-----------------|-----------------------------|-----------------------------|----------------------------------|--------------------|----------------|-------------------------|
| B-5-TiO$_2$-550| A (75)–R (2)                | 23                          | A (25)                           | 33                 | 2.8            | 74                      |
| B-5-TiO$_2$-450| B (76)                      | 24                          | A (18)                           | 29                 | 5.4            | 84                      |
| B-5-TiO$_2$-350| A (70)                      | 30                          | A (11)                           | 104                | 3.6            | 58                      |
| P25             | A (73)–R (14)               | 13                          | 22                               | 55                 | 61             |                         |

$^a$Determined as indicated in the characterization by XRD. $^b$Calculated from Scherrer’s equation. $^c$Calculated from N$_2$ adsorption data. $^d$Estimated from XPS analysis. $^e$Determined by eq 9 at 30 mL/min.

Figure 7. (a) Transient photocurrent measurements at a constant potential of 0 V vs Ag/AgCl and solar irradiation power of 70 mW cm$^{-2}$ for electrodes containing P25, B-5-TiO$_2$-450, B-0-TiO$_2$-550, and B-5-TiO$_2$-550. (b) Normalized photocurrent as a function of the irradiation wavelength at a constant potential of 0.4 V vs Ag/AgCl.

3.3. Effect of the Crystallization Temperature. Sample B-5-TiO$_2$-550, the most active in the B-X-TiO$_2$-550 series, has been selected to study the effect of the crystallization temperature on the properties of these photocatalysts. Thus, after the sol–gel synthesis, B-5-TiO$_2$ sample has been heat-treated at 350 and 450 °C to be compared with B-5-TiO$_2$-550. XRD patterns of the three B-5-TiO$_2$-T photocatalysts (Figure S8 in Supporting Information) show the characteristic peaks of anatase (see Section 3.1.1), and only for B-5-TiO$_2$-550, an incipient peak of rutile can be observed. Inset of Figure S8 shows the enlarged 22–30° 2θ range, revealing changes in the crystallinity of those samples. As expected, the peaks become broader as the crystallization temperature decreases, which means that the sample is less crystalline. Table 3 summarizes the main properties of the B-5-TiO$_2$-T series and the measured propene conversion. Data corresponding to P25 are also included.

Regarding the calculated percentage of crystalline and amorphous phases and the average crystallite size for B-5-TiO$_2$-T samples (calculated following the same procedure as for the B-X-TiO$_2$-550 set of samples, data included in Table 3), the B-5-TiO$_2$-350 catalyst presents the lowest crystallinity (30% of the amorphous phase) and smallest average crystal size (11 nm), whereas the B-5-TiO$_2$-550 has the highest crystallinity, even with a small amount of the rutile phase (2%). In addition, these data agree with specific surface areas ($S_{BET}$) obtained for these materials (Table 3). The material prepared at 350 °C presents the highest surface area, which is consistent with the largest proportion of the amorphous phase and the smallest crystal size. Samples B-5-TiO$_2$-450 and B-5-TiO$_2$-550 show similar surface area and crystallinity, although the effect of the temperature is reflected in the higher average crystal size of the sample treated at 550 °C.

B-5-TiO$_2$-350 and B-5-TiO$_2$-450 were also studied by XPS, and they show features similar to sample B-5-TiO$_2$-550 (see Figure S9 and Table S2 and Table S3 in Supporting Information). However, the sample prepared at 450 °C shows the highest boron atomic percentage, 5.4% (Table 3), meaning that the crystallization temperature most probably has an effect on the distribution of boron atoms on the TiO$_2$ surface. Finally, data in Table 3 show that the catalyst crystallization temperature influences propene conversion, being B-5-TiO$_2$-450 the most active sample (84% propene conversion) and B-5-TiO$_2$-350 the least active one.

The higher activity of sample B-5-TiO$_2$-450 compared to B-5-TiO$_2$-550 (both have similar surface area) can be explained considering that the first contains pure anatase, of lower average crystal size, and larger amount of surface interstitial boron atoms. The lower activity of sample B-5-TiO$_2$-350 can be related to the lower anatase content.

The photocatalytic propene oxidation under visible light using B-5-TiO$_2$-450, B-5-TiO$_2$-550, and P25 samples has also been studied. A previous blank experiment shows that, as expected, propene photolysis does not occur, while the tested photocatalysts led to relatively low propene conversion (around 15%), see Figure S10. The activity measured with visible light is probably mainly related to the contribution of UV radiation in the used visible lamp (see spectra in Figure S1, Supporting Information), although it seems that the three tested photocatalysts have some features by which the relative activity order under visible and UV light do not agree (Figure S10). In any case, the different lamp irradiances should be taken into account: UV irradiance of the visible lamp, 0.049 mW cm$^{-2}$, is much lower than that of the UV-lamp, 22.4 mW cm$^{-2}$, and the activity is not linearly related with irradiance. Nevertheless, the poor response under visible light is in
agreement with the lack of band gap modification of TiO$_2$ upon B incorporation (Figure S7 compares results of B-5-TiO$_2$-550 and B-0-TiO$_2$-550) and supports discarding substitutional B doping.

In order to explain the highest photocatalytic activity of sample B-5-TiO$_2$-450 under UV light, the series of B-5-TiO$_2$-T samples has been characterized by means of PEC and PL measurements (like in the case of the B-X-TiO$_2$-550 set of samples discussed above).

3.3.1. PEC and PL Characterization of B-5-TiO$_2$-T Samples. The PEC characteristics of the B-5-TiO$_2$-450 electrode under CV conditions in the dark (not shown) are analogous to those of B-0-TiO$_2$-550 and B-5-TiO$_2$-550 (Figure 5). Under full solar light irradiation, all the specimens develop an induced photocurrent at positive potentials versus Ag/AgCl. The transient photocurrent response for those selected specimens is shown in Figure 7a. A high photocurrent could be related to an improved ability of this sample to decrease the radiative recombination of photogenerated electron/hole pairs. The intensity of light pulses shows an exponential decay, which relates to the capacitive filling of surface states. The dynamics shown by P25 is quite different from those of the sol–gel prepared photocatalysts in the considered time scale, again connecting with the link of higher surface area and charge accumulation. The B-5-TiO$_2$-550 sample produces the highest intensity response. However, when the photocurrent is analyzed as a function of the light excitation wavelength using a monochromator (Figure 7b), samples B-0-TiO$_2$-550 and B-5-TiO$_2$-550, and P25 as well, show nearly identical wavelength-dependent profiles, while B-5-TiO$_2$-450 has clearly a shift in its wavelength activity profile.

Furthermore, diffuse reflectivity (Figure S7) shows that regardless the crystallization temperature, all samples present a linear uprise of the absorption pointing to the same indirect band gap wavelength (427 nm), but B-5-TiO$_2$-450 shows an Urbach tail (usually due to defects or loss of crystallinity) into the visible which is not present in the other two samples. This analysis reveals subtle differences between samples B-5-TiO$_2$-450 and B-5-TiO$_2$-550 and allows to propose that the wavelength shift of the photocurrent intensity maximum has a larger influence on the photocatalytic activity for propene oxidation under 365 nm UV light than the absolute intensity value.

Finally, Figure 8 contains, for comparison purpose, PL spectra (excitation source of 405 nm) of B-5-TiO$_2$-T.

Figure 8 reveals that the whole B-5-TiO$_2$-T photocatalyst series exhibits a quenching of all PL bands (compared with the B-0-TiO$_2$-550 sample shown in Figure 6). Thus, the reduction in the recombination rate of photo-excited electrons and holes when boron is (mostly interstitially) present is again confirmed. B-5-TiO$_2$-450 and B-5-TiO$_2$-550 samples show the lowest PL intensity within this series (lowest recombination rate of electron and holes), supporting the interpretation of their higher photocatalytic activities.

4. CONCLUSIONS

Nanostructured B-TiO$_2$ photocatalysts (B-X-TiO$_2$-550) with different amounts of boron (X) have been prepared by the sol–gel method and treated at different crystallization temperature (T). All the B-X-TiO$_2$-550 catalysts have similar $S_{BET}$ values (25–30 m$^2$/g). The presence of boric acid during the TiO$_2$ synthesis hampers the formation of rutile and, thus, the B-TiO$_2$ samples mostly consist of anatase with minor quantities of rutile (<12%). XPS analysis reveals that B occupies interstitial positions (close to the surface), and when X is in excess, it is externally deposited as B$_2$O$_3$. The activity of the B-TiO$_2$ samples for propene photo-oxidation is highly influenced by the boron content. Samples with low boron amount (X = 5 and 10) show the highest activity. The optimal behavior of these samples can be explained by the presence of interstitial B, together with a high crystallinity and high proportion of anatase. B$_2$O$_3$ deposits are formed in samples with higher boron loading, which cover the active titania surface with the subsequent activity decrease.

PEC and PL data reveal significant differences between samples. The highest photocatalytic performance achieved is related to the improvement of the photocurrent response ability (PEC), the production of hydroxyl radicals (Raman scattering), and the lower recombination rate of electron/hole pairs (PL), which points to a more efficient charge separation measurements.

The B-5-TiO$_2$-450 sample is about 10% more active than B-5-TiO$_2$-550. This improvement is related to the increased anatase content, lower average crystal size, and larger surface boron enrichment. Also, its intrinsic photo-activity suitably influences the photocatalytic activity of TiO$_2$. Also, it must be pointed out that the thorough characterization with complementary techniques has allowed to understand the effect of the interstitial B atoms in the photocatalytic behavior of the prepared samples.

ASSOCIATED CONTENT

+ Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/DOI/10.1021/acsanm.2c02217.

Spectra of used lamps; TEM and EDX data of samples B-0-TiO$_2$-550 and B-5-TiO$_2$-550; HRTEM images of
samples B-0-TiO$_2$-550 and B-5-TiO$_2$-550; XPS survey spectra of B-X-TiO$_2$-550 samples; Ti 2p XPS spectra of B-X-TiO$_2$-550 samples; PL spectra of B-X-TiO$_2$-550; square root of absorbance data of samples B-5-TiO$_2$-550, B-5-TiO$_2$-450, and B-0-TiO$_2$-550; XRD spectra of the B-S-TiO$_2$-T samples; Ti 2p, O 1s, and B 1s XPS spectra of B-S-TiO$_2$-T samples; propane conversion of B-S-TiO$_2$-550 and B-S-TiO$_2$-450 catalysts; textural properties of B-X-TiO$_2$-550 samples and P2S; binding energies of B 1s, O 1s, and Ti 2p in B-S-TiO$_2$-T samples; and composition of samples B-S-TiO$_2$-T determined by XPS (PDF).

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**Notes**

The authors declare no competing financial interest.

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