

Review

Update on Interfacial Charge Transfer (IFTC) Processes on Films Inactivating Viruses/Bacteria under Visible Light: Mechanistic Considerations and Critical Issues

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Abstract: This review presents an update describing binary and ternary semiconductors involving interfacial charge transfer (IFCT) in composites made up by TiO₂, CuO, Ag₂O and Fe₂O₃ used in microbial disinfection (bacteria and viruses). The disinfection mechanism, kinetics and generation of reactive oxygen species (ROS) in solution under solar/visible light are discussed. The surface properties of the photocatalysts and their active catalytic sites are described in detail. Pathogenic biofilm inactivation by photocatalytic thin films is addressed since biofilms are the most dangerous agents of spreading pathogens into the environment.

Keywords: photocatalysis; bacterial inactivation; viral inactivation; disinfection mechanisms; ROS lifetimes; ROS diffusion lengths



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1. Introduction

This update is designed primarily for students and researchers in the area of photocatalysis, describing the design, synthesis, evaluation and characterization of innovative photocatalysts allowing environmental disinfection. The advantage of supported photoactive materials is that they preclude the use of chlorinated compounds in water bodies to inactivate bacteria/viruses. Chlorination generates dangerous/carcinogenic chloro-species due to the interaction of Cl radical reacting with organic compounds in water under solar irradiation.

Semiconductor and inert oxides/double oxides present a mild alternative approach to disinfect water bodies. No comprehensive mechanism for the degradation of organic matter and pathogens by photocatalysis is available at the present time. There is a need for more work on stable, kinetically fast photocatalysts inactivating virus/bacteria. Recent reviews reporting environmental disinfection methods have been published recently [1–6]. This is a topic of timely interest and will motivate more research on the understanding of viral resistance to catalytic/photocatalytic treatment.

Most studies available on the photocatalytic inactivation of viruses rely on TiO₂, a low-cost, inert, non-toxic semiconductor that is stable within a wide pH range. However, the performance of TiO₂ is limited by its slow inactivation kinetics induced under solar light, resulting in long treatment times. This precludes a wide application of this approach. Doped TiO₂ promoters have been reported to enhance the virus inactivation kinetics by TiO₂ at a modest scale [5]. Highly porous TiO₂ increasing the adsorption of dispersed virus/bacteria is a way to increase the virus disinfection kinetics. Highly porous films such as TiO₂/Ag, TiO₂/Cu and TiO₂/Cu-apatite are not available at the present time [3–5].

Viruses constitute a group of heterogeneous organisms. They range in size from 0.01 to 0.3 microns and are much smaller compared to bacteria. They do not present independent metabolism and are fed by the host. They contain either DNA or RNA as genetic materials, but not both. The nucleic materials are surrounded by a protein and are coated to protect

them from the action of harmful external agents. The virus external envelope contains binding sites made up of protein, lipids and polysaccharides. These functional groups are needed by the virus to attach to the host cells. The binding functional sites fuse with the receptor host cell, allowing viral replication and further infecting the host.

Virus inactivation is due to virus protein/genome damage. The complete set of reactions leading to the photocatalytic oxidation of bacteria/viruses/fungi has not been worked out at the present time [6–9]. Capsid and genetic damage occurs during the TiO₂ photocatalytic inactivation of viruses. The extent of damage required for full virus inactivation is not clear. More work is required to elucidate the nature of the sites inducing viral protein damage. The viruses responsible for influenza, polio and enterococci infections have been extensively reported during recent decades, as well as adenovirus, which is widely found in contaminated food or water effluents. Enterobacteria phage (MS2) until now, is the most widely studied virus and is non-toxic to humans. Several water-borne viruses have been detected in groundwater, treated water in secondary biological municipal stations and natural water bodies. They can be transmitted from person to person, such as the case of the rotavirus, spread by the respiratory channels. Viral infection occurs when the genome/protein contacts the receptors of host cells infecting the patient. Veal, chicken, pig and lamb meat ordinarily transmit coronavirus, but in a form that cannot interact and penetrate human receptors/host cells [10].

Hybrid processes applying microfiltration and ultraviolet light (UV) light have been used for a long time, although at a reduced scale to reduce/inactivate viruses. UV treatment alone is not an effective treatment for highly turbid waters/matrices. This is mainly due to the photon screening in such non-transparent solutions. To disinfect effluents, colloids with small sizes close to the size of bacteria and/or viruses. Semiconductor and metal colloids can be prepared in very small sizes (sub-micron) as colloids presenting a high degree of transparency due to their small size. This allows large light penetration and low scattering with a balanced hydrophobic–hydrophilic behavior and suitable redox properties. This is necessary to disinfect water-borne bacteria/viruses. Coupling microfiltration with the photocatalytic process has been applied when disinfecting turbid wastewaters in a photoreactor. However, this leads to membrane fouling with concomitant costs. During virus removal and inactivation, the reaction media should present the relatively high light penetration required, allowing the disinfection process to take place.

This update presents some selected photocatalysts inactivating bacteria/viruses due to the reactive oxygen species (ROS) generated in the media. These ROS permeate into the host cell, inducing damage to the virus. Photogenerated charges damage bacteria/viruses when they migrate and translocate into the pathogen outer layers.

2. Light-Induced Semiconductor Reactions Inducing Virus/Bacteria Inactivation

TiO₂ activated by light has been used for its stability, low cost and inertness as the model semiconductor to inactivate bacteria and viruses, despite absorbing only 4–5% of the incoming solar light. The absorbed photon by TiO₂ induces an e⁻/h⁺ pair. However, decorating or doping the TiO₂ with oxides of Ag, Au, Cu, Pt, Fe, Ni, Pd and Rh enhances the visible light absorption capability of TiO₂. Interfacial charge transfer (IFCT) processes occur between TiO₂ and the metal oxide, absorbing light in the visible range. In the case of TiO₂, IFCT processes involve electron transition from anatase to rutile in the TiO₂ heterostructure. This is possible due to the higher conduction band position of anatase (cb1) with respect to the rutile conduction band position (cb2).

The incident light induces electron pair separation in the semiconductor followed by the generation of reactive oxygen species (ROS) in solution. These species are short-lived, present high redox potentials and were seen to effectively inactivate *Escherichia coli* (*E. coli*). ROS cause oxidative stress in the bacteria, and the oxygen singlet ¹O₂ generated in solution leads to peroxidation of the bacterial envelope [11,12]. The peroxidation step arises from the H-abstraction from an unsaturated fatty acid in the *E. coli* lipopolysaccharide (LPS) bilayer and leads to lipid radicals, as shown in Figure 1a. Within longer times, the oxidative

radical R leads to the formation of further ROS in solution. This is shown schematically in Figure 1b.

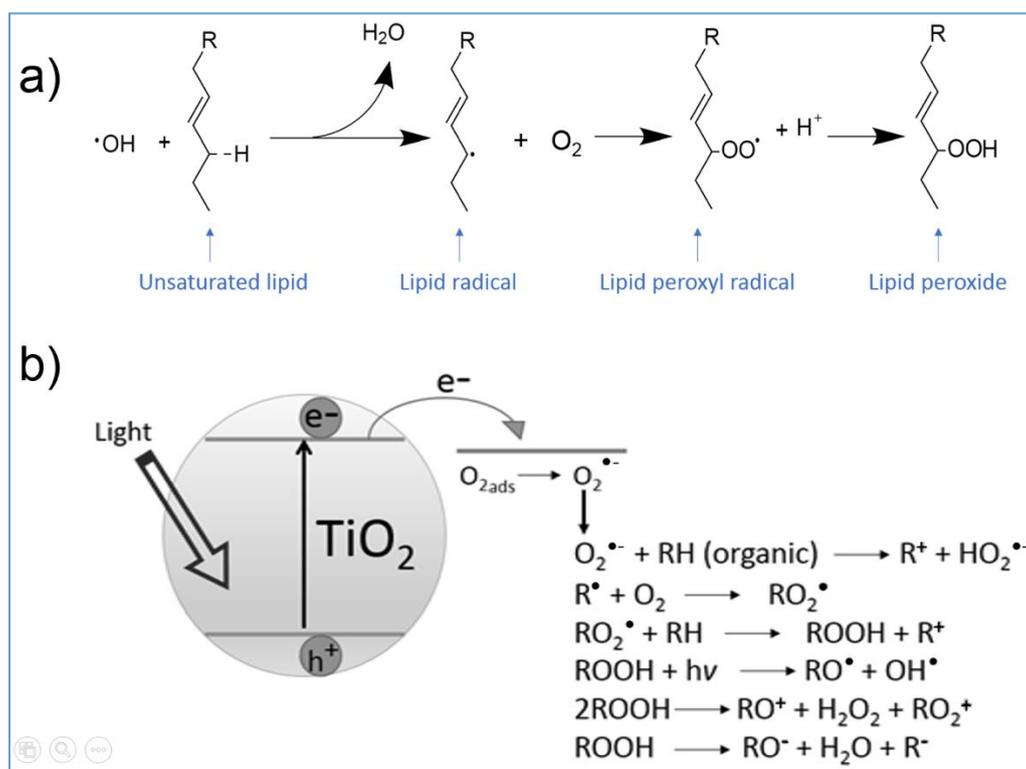


Figure 1. (a) H-abstraction from an unsaturated fatty acid in the lipopolysaccharide (LPS) topmost bilayer of *E. coli*, leading to cell envelope peroxidation. This is the precursor step for the generation of lipid radicals, initiating the mechanism leading to bacteria/virus degradation. (b) Schematic steps suggested for TiO_2 -mediated generation of reactive oxygen species (ROS) in solution under band gap irradiation, leading to bacteria/virus degradation.

The *E. coli* envelope consists of lipopolysaccharides (LPS) and phosphatidyl-ethanolamine (PE) bilayers, and these bilayers photodegrade during TiO_2 -mediated photocatalysis. Within longer times, the most resistant layer of the envelope, the peptidoglycan (PGN), gets damaged, leading to several peroxidation products. Depending on the type of bacteria used, the degradation products consist of aldehydes, ketones, carboxylic acids and residual organic oxidized intermediates. Full mineralization of the bacteria leads to CO_2 , water and inorganic anions such as $\text{NO}_3^-/\text{SO}_4^-$ [13,14].

Light irradiation to inactivate viruses has revealed that adenoviruses are highly resistant to irradiation by UV [15]. The United States Environmental Protection Agency (USEPA) has increased the UV fluence requirements for 4- \log_{10} removal of viruses from 40 to 186 mJ/cm^2 . This latter dose was shown to be necessary to completely deactivate the virus. This implies a higher amount of energy/costs necessary to inactivate this kind of pathogens. More efficient photocatalysts are required at the present time. Adenovirus is resistant to conventional treatment since its genome is mainly DNA and not RNA. This enables the virus to utilize the DNA repair enzymes present in the host cells to repair its genome damage, increasing its resistance to disinfection. Metals can undergo specific redox covalent reactions with the topmost functional groups of the virus or cells' outer layers. This involves redox reactions between the metal/ions and generates organometallic toxic species, leading to cell inactivation in the dark or under light [16–19].

Singlet oxygen has been reported to inactivate MS2 by the oxidation of amino acid residues [3–5,20]. Binary TiO_2 - SiO_2 -mediated photocatalysts have shown improved virus inactivation ability when compared to bare TiO_2 due to the finer dispersion of the TiO_2 nanoparticles (NPs) reached on the SiO_2 . Hydrogen peroxide was found to be an effective

additive in the dark for the reduction of viruses on stainless steel and, when used in conjunction with UV, led to a faster inactivation compared to H₂O₂ applied alone. For this reason, combined hydrogen peroxide–UV treatments to reduce viruses from many foods have been used during recent years.

3. Virus Inactivation by TiO₂ under Light

3.1. Enterococcal Viruses

The enteric group of viruses are not able to replicate in the environment outside of their hosts. Visible light-irradiated TiO₂ was seen to produce ROS in solution, leading to the inactivation of rotavirus and astrovirus. Human noroviruses are the most common cause of viral gastroenteritis. Fluorinated TiO₂ films have been shown to more effectively inactivate human norovirus under visible light compared to TiO₂ by itself. One hour of irradiation with fluorescent light at 10 mW/cm² reduced norovirus approximately 3–4log₁₀. When no light was applied, no detectable reduction was observed. The norovirus destruction was due to ROS and was accelerated by the presence of F-radicals leaching out of the catalyst surface [6]. Different viruses have been found in treated sewage leading to infections, despite the information reported by municipal treating stations that they are complying with the legislation on the standard microbiological water treatment. This revealed insufficient municipal treatment of wastewater viruses and enteric bacteria [15]. A more stringent evaluation of biologically treated water is needed nowadays to ensure the quality of drinking water and ensure that it is free of viruses.

3.2. Poliovirus

Poliovirus was effectively inactivated by titanium dioxide photocatalysis. The rates were more rapid than for the inactivation of coliform bacteria. However, the photocatalytic disinfection of effluents using titanium dioxide under sunlight is severely limited due to the relatively slow inactivation kinetics. Polio virus 1 was 99.9% killed in secondary waste effluent after 30 min of irradiation with 40-W black light in phosphate buffer Dulbecco's Phosphate Buffered Saline (DPBs) TiO₂ P25 dispersions [2].

3.3. Influenza

The influenza virus has been shown to be inactivated by TiO₂ photocatalysis. First, the protein present was deactivated, and subsequently, RNA was attacked. The data suggest that the photocatalytic activity destroys the viral binding proteins. The degradation of the proteins was due to ROS, mainly involving •OH and O₂•⁻ produced by the TiO₂ photocatalysis. The virus inactivation depends on the UV-A intensity and irradiation time. Titanium dioxide photocatalysis under UV light can inactivate 4-log₁₀ influenza virus within a short time at extremely low doses of UV-A (0.01 mW/cm²) [21,22].

3.4. Adenovirus

Adenoviruses are found widely in contaminated food or sewage water. UV-C treatment has been used at a laboratory scale and induced a higher reduction of virus than UV-A and UV-B. Adenoviruses' inactivation in the water supply chain is, in some instances, carried out using monochloramine with the generation of undesirable DBPs. Adenoviruses' contamination of water systems allow adenovirus to cycle through the environment. It is also transmitted from person to person, as in the case of the rotavirus. This virus spreads by respiratory/oral pathways by contact with the host cells of the receptor. After contact with the host cell, adenovirus produces the proteins required for further genetic replication. Adenoviruses proceed when significant damage takes place on their capsid structure. In contrast to MS2, adenovirus proteins were inactivated within acceptable times during the TiO₂-mediated photocatalytic process. The adenovirus undergoes damage to its genomes by UV254 light. However, even when damaged, is still capable of infecting cells. Viruses use the host cell enzymes to repair their damaged DNA and then replicate, allowing for further infection of a variety of host cells.

4. Parameters Controlling the Photocatalytic Virus Inactivation

The sections below briefly describe the solution parameters controlling pathogen inactivation in solution. Most of the available scientific literature reports studies on TiO_2 deactivation of bacteria, but not viruses. The photocatalytic inactivation of viruses follows the methodology and photochemical set-ups used to inactivate bacteria. The experimental use is adjusted for the features of viruses.

4.1. Effect of the Catalyst Concentration

A higher catalyst concentration accelerates the virus inactivation kinetics, but this process is limited by the optical density of the solution. The limit is set by the light penetration into the reactor volume [1–5]. Increasing the concentration of TiO_2 or TiO_2/Ag enhanced ROS production and led to an enhanced virus inactivation. The NP concentration mediating virus inactivation should be much higher than the photons/sec/cm² reaching the photochemical reactor. Saturation with the incoming photons of the photocatalyst NPs by the solution is another limiting parameter to consider during virus inactivation processes. Silver NPs were reported to enhance the TiO_2 photocatalytic inactivation kinetics of viruses. Silver also has a high affinity for sulfur (cysteine) and carboxyl groups present in the topmost virus bilayer. The latter functional groups bind silver [23,24].

4.2. Effect of Light Intensity

A higher light intensity leads to higher bacterial photocatalytic inactivation kinetics, but in certain limits. It increased linearly with the applied light dose when lower light intensities were used. However, the reaction rate only increased with the square root of the intensity at intermediate intensities. At higher light intensities, the rates increase only marginally or not at all. It was observed that the inactivation rates became constant after a certain threshold [25]. The reason for this is that higher light intensities lead to a faster recombination of the photogenerated charges (electrons and holes) in semiconductors such as TiO_2 [1–5]. The range and dose of the applied light have a controlling effect on the bacterial inactivation kinetics. Figure 2 shows the range of the light applied in photocatalytic processes.

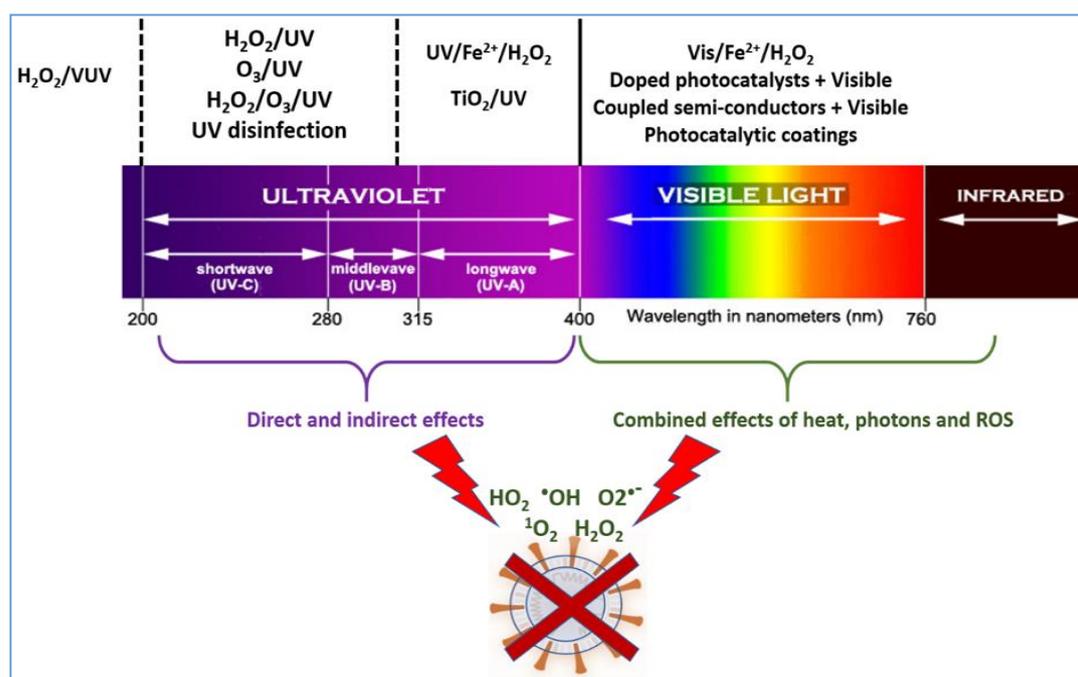


Figure 2. Range of the applied light effective in viral/bacterial inactivation in semiconductors and semiconductors doped by metals or oxides. Decorated, doped semiconductors present interfacial charge transfer (IFCT) processes under solar/visible light.

TiO₂ coatings and metal /oxide composites have been reported to inactivate a wide variety of microorganisms in small- or large-size reactors [1–5]. The variation of the light dose within a day had important effects on the pathogen inactivation kinetics, as the wavelength of the incident light changes during daily solar irradiation as a function of the time. UV-A and UV-B have been widely reported to inhibit cellular activity in the presence of TiO₂ or doped TiO₂, and the observed inactivation kinetics is a function of (a) the type of virus and/or bacteria colonies to be inactivated and (b) the concentration of pathogens. A bovine rotavirus was inactivated within 3 h and a 3log₁₀ concentration reduction was observed under solar light irradiation. Poliovirus was inactivated by applying a solar dose of 85 mW/cm² at 25 °C < 6 h. *E. coli* was disinfected under identical conditions within shorter times.

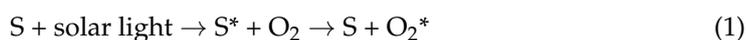
4.3. Contact between the Virus and the TiO₂ Surface

A higher surface area of a TiO₂ NP photocatalyst has been reported to lead to faster bacterial inactivation kinetics compared to samples prepared at higher temperatures and presenting lower specific surface areas (SSAs) [26]. A closer contact of the TiO₂ NPs with bacteria/viruses has been reported to accelerate the pathogen inactivation kinetics [27]. The binding of the TiO₂ NPs to bacterial cells involves electrostatic, van der Waals forces, pH, agglomeration of the catalyst and the NPs' surface potential.

5. Photocatalytic Mechanism Leading to Virus/Bacterial Inactivation

TiO₂ photocatalysis processes damage protein, leading to viral capsids and releasing DNA, involving protein and genome degradation [5,6]. The inactivation sequence of a virus involves the following steps: (a) modification of the protein sequence cross-linking, (b) disruption of the protein conformation, (c) disruption of the protein aggregated size and (d) disruption of the transfer of the viral genome to the host cells. The surface area of the TiO₂ and its size, crystallographic structure and porosity control the amount of ROS produced in the solution leading to pathogen inactivation. The salts, cations and anions found in the solution interfere with the contact between the catalyst and the pathogen, hindering the bacterial/virus inactivation. Impurities and additives can also interfere and partially inhibit the adsorption of the pathogen onto the TiO₂ NPs.

Organic compound sensitizers are found in natural water bodies. Solar sensitizers transmit their energy by contact/diffusion to waterborne viruses [3–8]. These organic sensitizers adsorb solar irradiation and interact with the O₂ dissolved at normal temperature and pressure in water (8 mg/L). Sensitizers are activated by visible light when reacting with the dissolved O₂ present in the ground state as a triplet state. Triplet–triplet energy transfer occurs between S* and ³O₂ or by a second deactivation channel electron transfer from the S* to oxygen. Alternatively, the electron can transfer from the S* to the superoxide radical anion O₂^{•−} in the solution at biological pH values of 5.7–8.0. The sensitizer (S) can either transfer the energy, leading to excited O₂* by reaction (1), or react subsequently by charge transfer with dissolved O₂, as shown in reaction (2):



The excited O* species and the radicals O₂^{•−} in Equations (1) and (2) lead to the oxidation of bacteria/viruses. The O₂^{•−} radical exists in solution at the biological pH (6–8), since the pK_a of the equilibrium HO₂[−] → H⁺ + O₂^{•−} is 4.8. ROS radicals such as the •OH radicals OH/OH[−] presenting a thermodynamic potential Normal Hydrogen Electrode potential (NHE −1.90 eV) and HO₂/O₂[−] (NHE −0.65 eV) present a short diffusion length in solution. The •OH and HO₂^{•−} interact with pathogens, pollutants, salts and dissolved organic matter (DOM) in solution within the range of their respective diffusion spheres. The diffusion of both radicals has been estimated from the simplified Smoluchowski equation. Furthermore, one has to consider the time span in which the sensitizer (S) reacts with the

dissolved O₂ to produce ROS radicals in the solution. The next paragraph describes the estimation of ROS lifetimes and the diffusion length of these ROS radicals.

The concentration of O₂ is 8 mg/L or 0.5×10^{-3} M in water and is used to estimate the lifetime of the reaction pair made up by the sensitizer (S) and O₂. The reaction time (τ) can be estimated from the relation $1/\tau = 0.5 \cdot 10^{-3} \text{ M} \times 6 \cdot 10^9 \text{ M}^{-1} \cdot \text{s}^{-1} = 3 \cdot 10^6 \text{ s}^{-1}$ or 0.33 μs . The diffusion distance (x) of the $\bullet\text{OH}$ radical away from the catalyst surface is estimated by way of the simplified Smoluchowski relation. The reaction rate between $\bullet\text{OH}$ and the organic compound (RH) in aqueous solution is $k_{\text{OH}} = 10^{-9} \text{ M}^{-1} \cdot \text{s}^{-1}$, and if the concentration of (RH) is 10^{-2} M, the lifetime of the reaction pair is noted in Equation (3) below:

$$\frac{1}{\tau} = k_{\bullet\text{OH}}[\text{RH}] = 10^7 \text{ s}^{-1} \text{ and } \tau = 0.1 \text{ } \mu\text{s} \quad (3)$$

The motility/diffusion of the $\bullet\text{OH}$ radical can also be estimated by the Smoluchowski relation $x^2 = D\tau$ inserting $D = 5 \times 10^{-6} \text{ cm}^2 \cdot \text{s}^{-1}$, the average value for the diffusion of small molecules in aqueous solution. The diffusion length for the $\bullet\text{OH}$ radical in solution is $x = 0.7 \times 10^6 \text{ cm}$ or 70 Å. The diffusion length (x) of the HO₂ \bullet radical can also be estimated by the simplified Smoluchowski diffusion equation. The reaction rate constants for the HO₂ \bullet radical have been reported by fast kinetic spectroscopy in the range $k = 10^5\text{--}10^7 \text{ M}^{-1} \cdot \text{s}^{-1}$.

$$\text{Then, } \frac{1}{\tau} = 10^4 \text{ s}^{-1} \quad (4)$$

and the lifetime found is of τ is 0.1 ms. The diffusion of the HO₂ \bullet in solution is estimated as $x = 2000 \text{ \AA}$, taking the average reaction rate k for HO₂ \bullet radicals as $\sim 10^6 \text{ M}^{-1} \cdot \text{s}^{-1}$.

ROS leading to virus inactivation considers (a) that mass transfer does not play a role in the virus ROS transfer process and (b) that no activation energy is required for ROS radical reactions with viruses. The quenching of singlet oxygen ¹O₂ by NaN₃ has been used frequently to identify ¹O₂ and quantify its presence in solution. However, this approach presents a drawback: azides also react with the $\bullet\text{OH}$ radical at a rate close to its reaction with ¹O₂. Singlet oxygen ¹O₂ has been reported with a lifetime of 3.6 ms and a diffusion motility of $\sim 75 \text{ nm}$ in solution. This length is above the observed diffusion length for the $\bullet\text{OH}$ radical motility in solution of $\sim 70 \text{ \AA}$. The reason for this is that the singlet oxygen is a species in the gas phase and the OH radical is a species in solution. ¹O₂-mediated processes lead to *E. coli*, adenovirus and MS2 and many other pathogens' inactivation [27,28]. Singlet oxygen has also been reported to hinder MS2 genome replication and oxidize virus protein.

The OH radical presents an oxidation potential $\bullet\text{OH}/\text{OH}^-$ 1.9 eV NHE [5] and is high enough to lead to inactivation of the MS2 virus and oxidize capsid proteins. However, the quantification of the intermediate ROS leading to virus inactivation has not been fully reported until now.

Bacteria and viruses can sorb on dissolved organic matter (DOM) in solutions and water bodies. This increases the probability of encounters between ROS and the DOM particles adsorbed on the surface of bacteria/viruses. DOM–virus interactions have been reported for capsid proteins due to cation bridges, electrostatic, steric, hydrophobic and carboxylate effects.

6. IFCT Ag/TiO₂-Mediated Bacterial/Virus Inactivation

In recent years, photocatalytic heterostructures have been reported, presenting increased absorption in the visible region. This is attained by decorating/doping TiO₂ and ZnO semiconductors with metals, oxides and double oxides absorbing in the visible region. In the latter case, IFCT processes take place and (a) increase the separation of charge carriers in the semiconductor, (b) suppress the recombination rate of photoinduced electron–hole pairs, thus improving photocatalytic efficiency, (c) induce synergistic effects induced by the components in the heterostructure and, (d) in some cases, increase photostability of the composite [29].

Metallic oxides with oxygen vacancies such as W-oxides are of growing interest in the field of visible light-driven photocatalysis as they exhibit a wide absorption tail in the Near Infrared Region (NIR). This is due to oxygen defects. The combination of TiO₂ with poly-oxometalla (tes leads to a hybrid photocatalyst absorbing in the UV and visible region [30].

Ag is the oldest antimicrobial agent and has been known for 2000 years. Its antibacterial features have been reported in a series of reviews [5–7]. Silver is known to be antimicrobial, biocompatible and non-toxic to humans at levels < 0.1 mg/L. The virus inactivation kinetics mediated by TiO₂/Ag is enhanced by the photogenerated charges in the TiO₂ surface increasing the Ag disinfection activity due to the presence of ROS radicals in solution. Higher Ag contents decrease the amount of light reaching the TiO₂ surface and reduce the generation of electrons, precluding the ability to inactivate pathogens. The interfacial charge transfer (IFCT) induced by TiO₂ nanoparticles (TiO₂/Ag NPs) in solution under visible light is shown below in Figure 3.

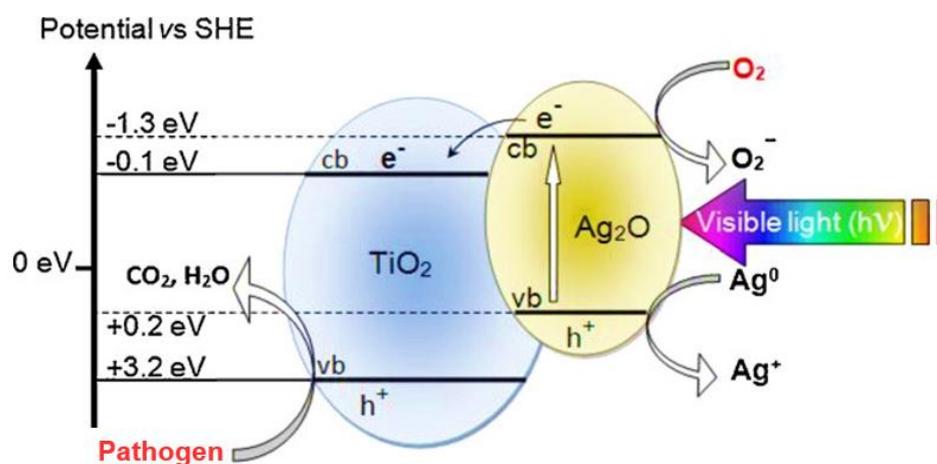
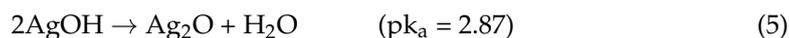


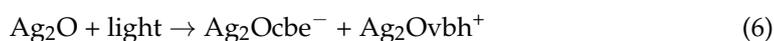
Figure 3. Interfacial charge transfer (IFCT) in TiO₂/Ag NPS in solution under visible light.

Ag surface plasmon resonance (SPR) sites build up in Ag clusters of TiO₂/Ag surfaces and this favors light absorption. TiO₂/Ag increases the generation of ROS radicals such as HO₂[•] and [•]OH radicals and ¹O₂, leading to the degradation of viruses such as adenovirus, rotavirus and astrovirus. TiO₂/Ag has been reported to increase the sorption/adsorption of pathogens compared to bare TiO₂ in the dark and under light. The TiO₂/Ag photocatalytic action is commonly rationalized in terms of an enhanced charge separation at the surface of the TiO₂ by the Ag dopant hindering the e⁻/h⁺ charge recombination.

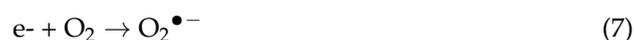
Variation of the of the rutile/anatase ratio in the catalyst make-up led to a variation of the kinetics of MS2 virus inactivation. Ag photocatalysis leads to the formation of ROS species [23,24]. Ag reacts almost instantaneously with air (O₂) and atmospheric water vapor, leading to AgOH. Only 2–3 atomic layers of AgOH (0.4–0.6 nm thick) are found on NPs' surfaces [31]. AgOH decomposes spontaneously to Ag₂O.



Ag₂O is thermodynamically stable in the pH 6–8 range, the physiological pH range of bacteria. Visible light irradiation activates Ag₂O, generating electrons and holes:



and the photogenerated build-up of O₂^{•-}, [•]OH and HO₂[•] radicals as outlined next:





Light excitation on catalysts leads to faster pathogen inactivation compared to runs in the dark. This is due to the additional release of $\bullet OH$ radicals as suggested next in the reaction (11)



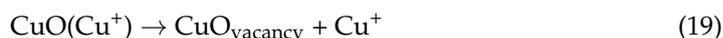
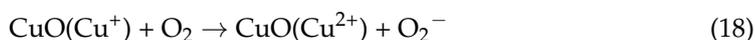
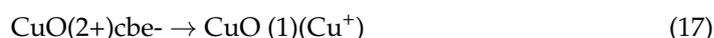
The Langmuir–Hinshelwood (L-H) kinetic model is frequently employed to describe bacterial inactivation kinetics [6,7]. This model is applied to process the data obtained during virus inactivation mediated by TiO_2 and metal- TiO_2 photocatalysts [5–8]. In the L-H model, the reactants are adsorbed and considered in equilibrium with the catalysis surface. This model assumes (a) that the system is in a steady state, (b) that the product formation is the rate limiting step, (c) that the products desorb quickly after the reaction and, finally, (d) that one of the reactants is present in a large excess. Viruses are made up of complex structures, presenting features that differ from chemical compounds, and the L-H model does not take into account the complexities of viruses. Any description of virus inactivation kinetics by this approach is not possible.

7. IFCT-Mediated TiO_2/Cu Bacterial/Virus Inactivation

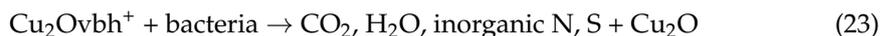
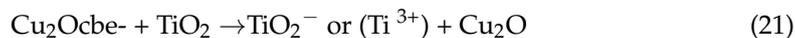
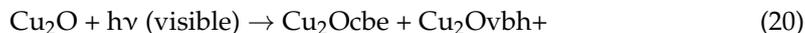
TiO_2/Cu inactivation of pathogens was first reported by a Sunada et al. [32]. Later, several laboratories reported Cu ions/Cu composites leading to *E. coli* inactivation by Fenton-like reactions and ROS radicals and peroxides being generated and partly adsorbed on TiO_2 . Inactivation by Cu_xO and TiO_2/Cu films has been the focus of some recent reports [33–35]. TiO_2/Cu lead to a higher degree of cell inactivation compared to TiO_2 due to the Cu toxicity and the additional ROS generation detected in the solution compared to bare TiO_2 .

Why does doping enhance TiO_2/Cu photocatalytic disinfection? Cu improves the TiO_2 photocatalytic pathogen inactivation compared with TiO_2 alone considering the following parameters: (a) the biocidal nature of Cu and (b) the shift of the NPs' absorption to the visible region up to 600 nm. TiO_2 , absorbing around 4–4.5% of the solar irradiation, has been widely reported to activate pathogen inactivation processes. The bacterial protein for many pathogens absorbs light at up to 320–325 nm [36,37]; (c) the TiO_2 band gap is made narrower by the added Cu/CuO and (d) the Cu clusters on TiO_2 as electron traps hinder, to some extent, e^-/h^+ recombination [38], and this occurs in conjunction with the charge transfer between the TiO_2/Cu surfaces and the bacteria. A scheme for the IFCT processes taking place is shown below in Figure 5.

A simplified mechanism for bacterial/virus inactivation is suggested for CuO NPs in Equations (16)–(23) [33,34]. By the use of photon energies exceeding the CuO band gap, the photogenerated cbe^- can react with O_2 , leading to the ROS $O_2^{\bullet -}$ or alternatively reducing the Cu^{2+} to Cu^+ as noted below:



Sequentially, binary sputtered films of Cu on polyethylene terephthalate CuO_x/TiO₂-PET under visible light promote the interfacial charge transfer (IFCT) of the cbe⁻ from Cu₂Ocb to TiO₂cb, as noted in Equation (21). Then, the TiO₂vbh⁺ holes with an appropriate positive potential oxidize bacteria, as shown below by Equation (23).



The bacteriophage f2 has been photocatalytically inactivated under visible light irradiation. The removal efficiency of the virus increased with catalyst concentration, light intensity and temperature and decreased at a higher initial virus concentration. Zheng et al. reported on TiO₂ doped with non-noble metals able to inactivate virus under light irradiation [39]. Until now the role of metal-ions in photocatalytic viral disinfection have not been explored in a systematic, comprehensive and detailed way. Composite material semiconductors containing Ag and to a lesser extent Cu have shown to form plasmon under UV-visible light irradiation [40–42]. The plasmons play a significant role in bacteria and virus inactivation. This remains a task of the future and seems necessary to increase the virus inactivation kinetics. This is one of the critical issues hindering the commercial scale application of photocatalytic technology for disinfection purposes [43].

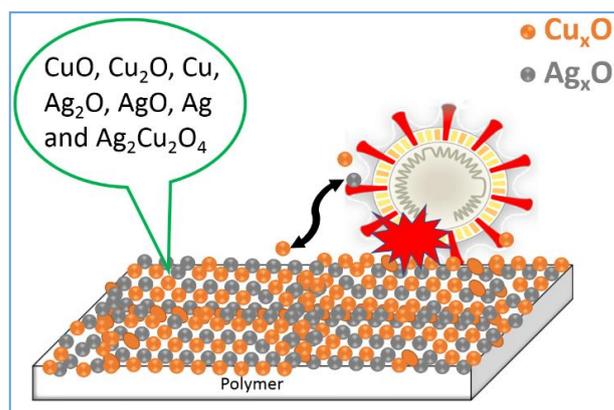


Figure 4. Schematic illustrating the species involved in virus inactivation in the dark or under light when composites made up of Ag-Cu were used as catalysts/photocatalysts.

8. IFCT-Mediated Ag-Cu Bacterial/Virus Inactivation

Studies on Ag-Cu NP films show higher bacterial inactivation kinetics compared to either Ag or Cu alone. Studies on Ag-Cu NP films leading to the destruction of bacteria have been reported recently [44,45]. The composition and species found in these composites are shown in Figure 4.

The faster kinetic inactivation of pathogens by bimetallic/trimetallic oxides is used due to two main reasons: (a) it increases the absorption of semiconductors such as TiO₂ into the visible range and (b) it also increases the number of potential couples available to catalyze/photocatalyze chemical transformations on the catalyst surface. Pathogen inactivation kinetics by bimetals such as Ag-Cu films compared to either Ag or Cu films by themselves. Work on Fe-doped/decorated wide-band semiconductors such as TiO₂, mediating bacterial inactivation, has been reported recently [1]. Using fast kinetics spectroscopy, the short-lived precursors leading to bacterial inactivation have been sorted out [43,46].

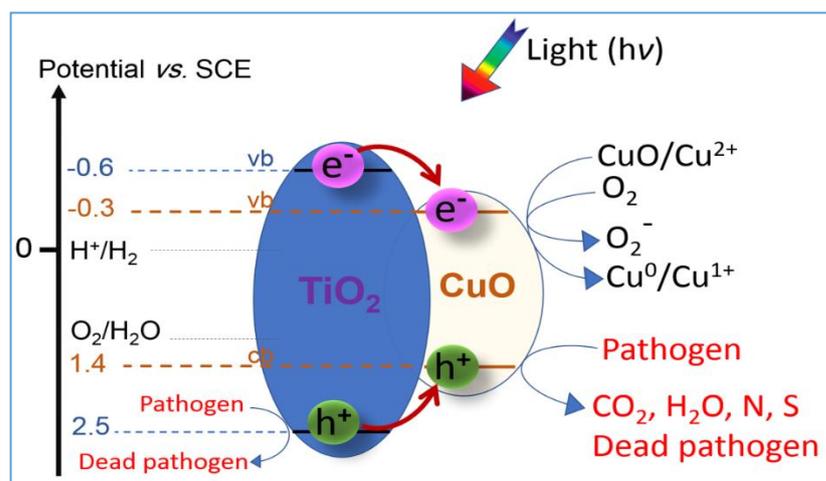


Figure 5. Interfacial charge transfer (IFCT) between CuO and TiO₂ in TiO₂/CuO nanoparticles (NPs), leading to the inactivation of pathogens in solution under visible light.

Killing bacteria on metallic copper surfaces occurs primarily due to the release of copper ions. However, it also involves slower processes such as the contact of the bacteria with the Cu/CuO surface [35,36]. Copper cations participate in Fenton-like reactions, leading to fast cellular damage by ROS species in mammalian cell membranes. This is due to the high Cu cation cytotoxicity. There are no standard tests to evaluate NPs' cytotoxicity which may allow to compare the results found in different laboratories [47–49]. The translocation of Cu NPs/Cu ions and Ag ions into the pathogen cytoplasm is far from being understood at the present time. The relative index of the toxicity on mammalian cells of Ag NPs and CuO NPs is shown in Table 1 below. The parentheses show the number of experimental runs performed to report the median value.

Table 1. Mammalian cells in vitro relative cytotoxicity index.

	Relative Index of Toxicity	Number of Studies Used to Report the Median Value
Ag NPs	11	(25)
Ag ⁺ ions	2.0	(18)
CuO NPs	25	(21)
Cu ²⁺ ions	53	(10)

It is difficult to differentiate the toxicity introduced individually by Ag-Cu NPs' surfaces and the separate effects induced by Ag and Cu NPs by themselves under light in aqueous media [8,16,25]. Recent studies indicated that Cu NPs induced oxidative damage to bacteria with higher kinetics compared to Ag NPs [24]. Figure 6 suggests the IFCT for Ag₂O/CuO composite films under light irradiation and the subsequent generation of radicals, leading to the degradation of organics (RH) or bacteria.

Like many other metal film composites, Ag-Cu films accelerate the inactivation of pathogens [50,51]. This presents an alternative to the use of antibiotics without significantly increasing the risk of resistance when antibiotics are administered for long periods of time. The efficiency of these films is dependent on the coating technique used to prepare them. A more advanced nanotechnological approach to prepare uniform and reproducible NP polymer films would benefit the inactivation of pathogens within acceptable times [52,53]. The quantitative contribution of each NP to the production of reactive oxygen species is still an open issue. Cu(II) ions are reduced to Cu(I) ions by the enzymes regulating the human respiratory cycle [35]. An excess of copper cations cannot be removed by these enzymes [36] and leads to respiratory cycle dysfunction and death within short times.

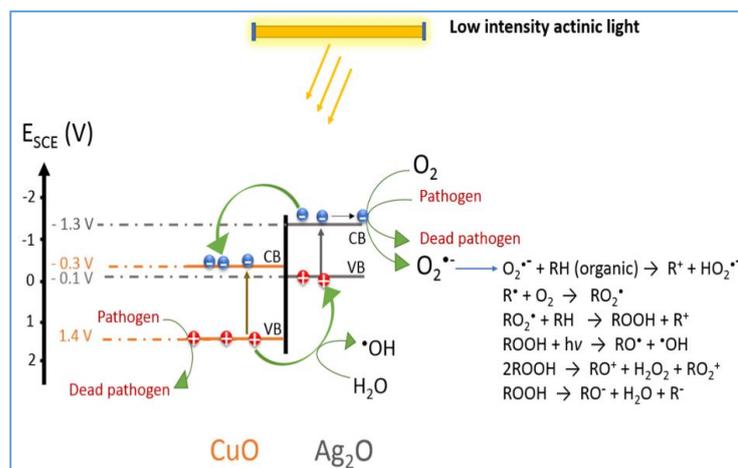


Figure 6. Interfacial charge transfer (IFCT) for $\text{Ag}_2\text{O}/\text{CuO}$ composites under light doses of $<4 \text{ mW/cm}^2$, showing the generation of ROS, leading to the inactivation of pathogens under light irradiation and the potential energy level of both semiconductor components.

Surface plasmon resonance (SPR) bands have been investigated for some years for Au, Ag and Cu nanoparticles. Cu SPR films are also applied in optical catalysis, detection and conductive devices used in printing technologies [52]. Until now, no comprehensive mechanism has been reported for the detailed intervention of Cu plasmons leading to bacterial/virus inactivation, and Au plasmons intervening in bacterial inactivation have recently been reported [43]. Visible light irradiation of SPR Au/ TiO_2 bands activates the electron injection of Au NPs into TiO_2 with a slight dissolution of the Au. This dissolution leads to (a) the injection of hot Au electrons into TiO_2 followed by femtosecond spectroscopy and (b) the generation of Au ions in the solution [43]. This study also reports on Au ions' mechanism of interaction with bacterial cells. The bactericidal inactivation mechanism by Au ions is significantly different to the mechanism reported for Ag SPR and Cu SPR activated by light.

Higher toxicity has been extensively reported for Cu relative to Ag. Extremely low amounts of Cu species (in the nano range) induce pathogen inactivation in the dark due to the high cytotoxicity of Cu per unit weight compared to Ag. Cu is not a noble metal and is readily available compared to Ag. This makes Cu/ CuO/Cu ions a preferred choice for pathogen inactivation studies. Copper is number 29 of the periodic table of elements. The Cu atomic structure is made up of negatively charged electrons in completely filled orbitals close to the positively charged atomic nucleus. The electrons of the unfilled orbitals $4s^1 3d^{10}$ require a small input of energy to activate chemical reactions. Cu(I) ($4s^0 3d^{10}$) and Cu(II) ($4s^0 3d^9$) oxidation states require a much lower one-electron redox potential compared to Ag(I)/Ag(II) and Fe(II)/Fe(III) in electron transfer reactions. Gram-negative bacteria including Cu in the periplasm can handle the toxicity of copper cations up to a certain level in their own metabolism.

9. IFCT in $\text{Fe}_2\text{O}_3/\text{TiO}_2$ Giving Rise to Fenton-Like Reactions Mediating Virus/Bacterial Inactivation

$\text{Fe}_2\text{O}_3\text{-TiO}_2$ double oxides have been reported to induce a more accelerated photocatalysis compared to bare Fe_2O_3 under visible light. This is due to the presence of the $\text{Fe}_2\text{O}_3\text{-TiO}_2$ heterojunction, in which Fe_2O_3 acts as the TiO_2 photosensitizer under visible light. Figure 7 presents a schematic for the $\text{Fe}_2\text{O}_3\text{-TiO}_2$ intervention [54]. The conduction band (cb) of Fe_2O_3 is located at a lower potential energy level with respect to TiO_2 , as shown in Figure 7. However, electron transfer is possible since the TiO_2 lattice presents trapping sites $\sim 0.8 \text{ eV}$ below the position of the TiO_2 cb [4,55,56]. The charge transfer between the two semiconductors led to (a) an increase in the photogenerated charge separation in Fe_2O_3 , (b) an increase in the pollutant oxidation kinetics by the generated holes and (c) improved

reduction kinetics of the $(\text{O}_2)_{\text{air}}$ by the photogenerated (cb) electrons. The TiO_2 vb generates the $\bullet\text{OH}$ radicals leading to bacterial/virus inactivation due to the more favorable potential energy position of its vb. The charge transport in the Fe_2O_3 - TiO_2 controls the photocatalytic-mediated reaction kinetics [57–60]. Under light, the Fe^{2+} in the Fe_2O_3 - TiO_2 undergoes a redox reaction and oxidizes to Fe^{3+} , as noted in Equation (24) below:

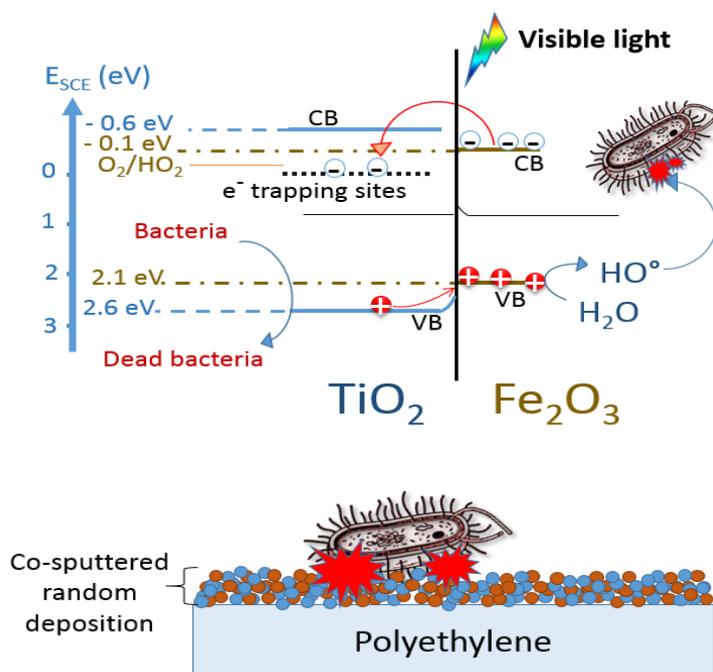


Figure 7. Mechanism suggested for bacterial inactivation by co-sputtered by FeOx-TiO₂-PE films under visible light, showing the IFCT process taking place between Fe₂O₃ induced by visible light.

The schema in Figure 8 suggests that with the FeOx/TiO₂-PE, bacterial inactivation on the film is possible from the Fe oxide into TiO₂ due to low-lying states detected below the conduction band (cb) of TiO₂. This allows to suggest a scheme of reactions leading to a short-lived unstable bacteria cation(+) as noted below in Equation (25) and reported in reference [43].



During the bacterial abatement process shown in Figure 7, some Fe ions leach out of the FeOx-TiO₂-PE films under light irradiation in an aqueous solution. This gives rise to Fenton-like reactions with ions such as Fe, Mn or Cu ions [20]. ROS radicals are generated in solution [20]. The reaction sequence leading to virus/bacterial inactivation proceeds through several steps as detailed next: (a) virus adsorption on the catalyst; (b) ROS generation and surface and bulk reactions and (c) desorption of the inactivated viruses. The diffusion of the virus within the inoculate plays a role in the inactivation process [9]. Mononuclear metalloenzymes include Fe(II) in their structure since Fe ions are important in human/animal metabolism. These mononuclear Fe enzymes are made up of Fe and Zn(II) and Mn(II). Fenton chemistry occurs when FeOx corrodes, leading to Fe ions in solution [61]. Fe enzyme Fe-S dehydratase such as Fe(II)-cysteine is oxidized through Fenton reactions to the sulphinic RSO(=H) or sulphonic RS=O₂-OH species, displacing Fe(II) from the metalloenzyme. The latter reaction involves several steps: (a) oxidant depletion and (b) additional (excess) ROS and thiol depletion from the cell bilayer topmost functional groups [62]. Thiol depletion due to Fenton-like reactions is shown schematically

in Figure 9. The outcome is that normal cell metabolism is inhibited, leading to cell death. Fe-S-type dehydratases are particularly sensitive to the destruction of the topmost bilayer functional binding groups containing S-moieties [63].

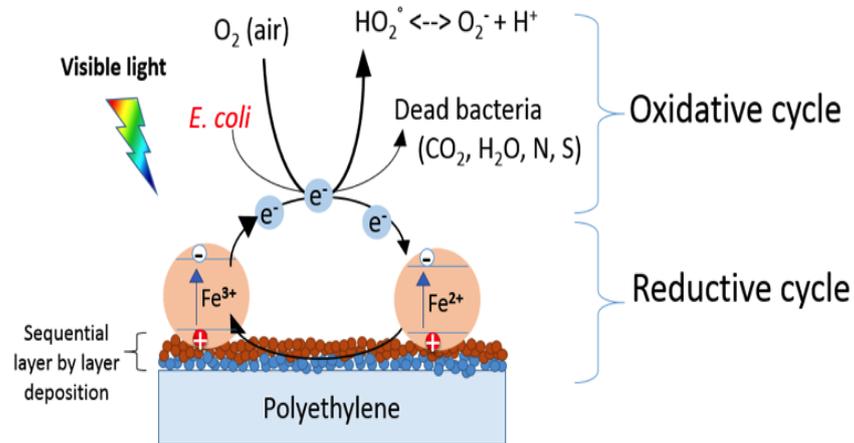


Figure 8. Mechanism suggested for bacterial inactivation mediated by sequentially sputtered polyethylene films (FeOx/TiO₂-PE) films under visible light.

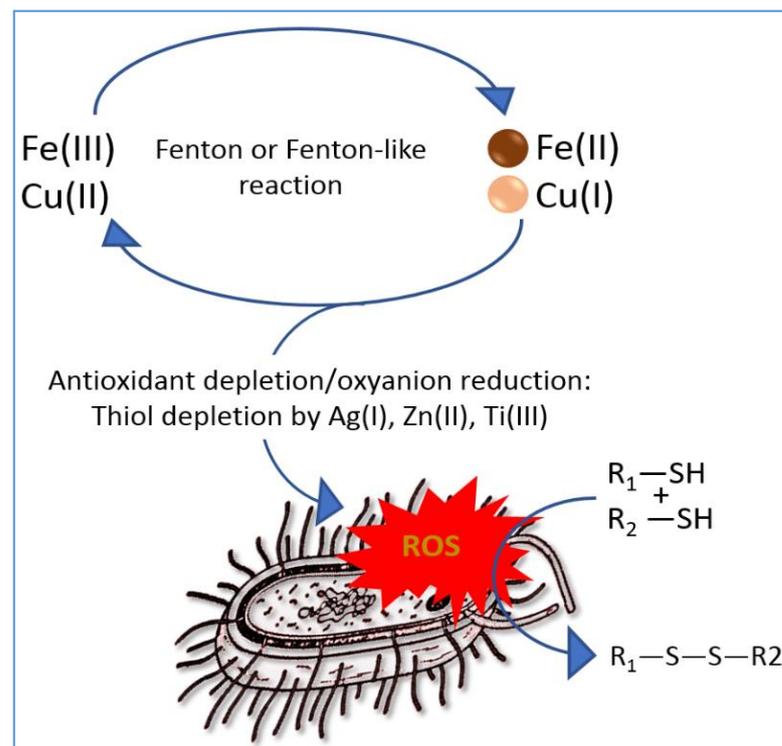


Figure 9. Schematic of Fenton chemistry due to ions of Fe(III) or Cu(II) associated with ROS intermediates leading to thiol depletion in the bacteria and virus topmost layers.

When the Fenton reagent (Fe²⁺ + H₂O₂) is added in solution, an increase in the OH radicals is observed compared to solutions with only added H₂O₂. This is due to the fine dispersion of nanosized Fe NPs leaching out Fe ions during the partial dissolution of the Fe NPs. The catalytic species Fe(H₂O)₆²⁺ in solution with a lifetime < 1 ns induce peroxide decomposition in solution in the dark and more so under visible band-gap light irradiation (Haber–Weiss-related type of reactions). The concentration of OH radicals when an organic

compound (RH) with a concentration of 10^{-3} M is added to a solution made up of $\text{Fe}^{2+}/3+$ 10^{-3} M and H_2O_2 10^{-2} M can be estimated from the reactions:



$d(\text{OH})/dt = k_1\text{Fe}(\text{Fe}^{2+})(\text{H}_2\text{O}_2) - k_2(\bullet\text{OH})(\text{RH})$ is ~ 0 , a quasi-stationary state, since the concentration of OH^\cdot is very small. The OH^\cdot radical is generated in sub-nanomolar concentrations as shown by the equation $(\bullet\text{OH}) = k_1(\text{Fe}^{2+})(\text{H}_2\text{O}_2)/k_2(\text{RH})$ in quantities of about $\sim 10^{-10}$ M. This latter value corresponds to a sub-nanomolar concentration. The concentration estimated for the OH^\cdot radicals is similar to the concentration of OH^\cdot radicals found in many natural water bodies.

10. IFCT in Ternary Semiconductors Leading to Bacterial/Virus Inactivation

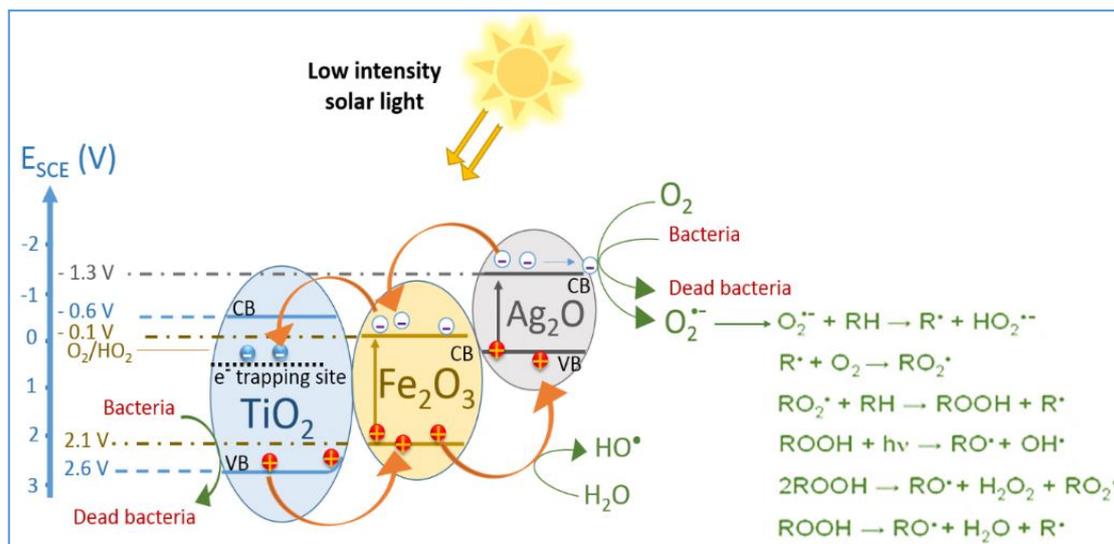
A mechanism is suggested in Figure 10a for the interfacial charge transfer (IFCT) in the $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$ photocatalyst mediating bacterial inactivation [64]. The TiO_2cb position in $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$ is more cathodic compared to FeO_xcb , and the TiO_2vb is more anodic in respect to FeO_x . The potential energy positions of the potential electronic bands of the three oxides determine the IFCT kinetics taking place in Figure 10a. The release of Fe-ions in aqueous solutions by the magnetic $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$ composite has been reported for several years when using Fe-composites. In the case of the $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$, a high calcination temperature was used during the preparation of this composite. The calcination step strongly binds the Fe to the TiO_2 through Ti-OH bonds. This precludes the Fe-ions release into solution within the bacterial inactivation process. The FeO_x (mainly Fe_2O_3) in the $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$ during the calcination step becomes deeply buried in the catalyst bulk. If present in the surface, the FeO_x would photo-corrode in aqueous media under light irradiation [65]. The significant acceleration in the bacterial inactivation kinetics by $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$ is due to the localized electric field leading to more reactions per unit time compared to the non-magnetized catalyst $\text{Ag}(3\%)\text{-TiO}_2$. This leads to the polarization in the $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$ heterojunction enhancing the separation of electrons and holes [66]. Repetitive recycling of the $\text{Ag}(3\%)\text{-TiO}_2\text{-FeO}_x(3\%)$ did not lead to loss of catalytic activity. This provides evidence for the stable nature of the FeO_x present in the composite photocatalyst.

The use of heterostructures made up by wide-band semiconductors, oxide and metals reduces the band gap of TiO_2 (bg 3.2 eV). As a result, development of visible light active titanium dioxide materials was observed. This is one of the main challenges in the field of semiconductor photocatalysis. Heterostructures composed of ternary catalysts, doped or not, enhance light absorption in the visible region. Figure 10b shows an example of IFCT over ternary semiconductor photocatalysts. More work to identify appropriate dopants to improve visible light absorption and electron-hole separation is needed at this time to enhance photocatalytic activity. One serious problem encountered in binary or ternary semiconductors designed to accelerate bacterial/virus inactivation is the electrostatic recombination of the cbe- of one semiconductor with the vbh+ holes of the other semiconductor. This hinders the diffusion of photogenerated charges to the semiconductor surface, lowering the production of readily available photogenerated species [67].

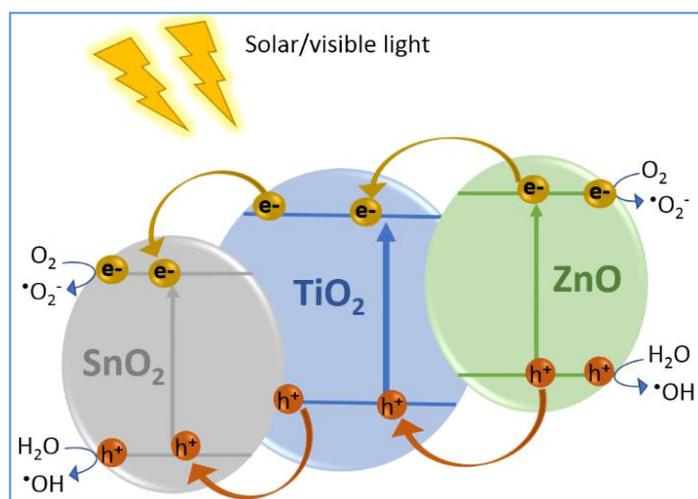
Other compounds presenting heterostructures with a TiO_2 component, leading to bacterial inactivation, have recently been reported such as C70-TiO_2 [68], $\text{TiO}_2/\text{Ag}_3\text{PO}_4$ [69], $\text{TiO}_2\text{-graphene}$ [70], $\text{TiO}_2\text{-MWCNT}$ [71] and $\text{TiO}_2\text{-graphene}$ [72]. The inactivation of bacteria by other composites involving IFCT processes has also been described in the open literature [73–78]. The photocatalytic inactivation of staphylococcus, streptococcus and fungi can be found in references [79–83].

Semiconductors have many advantages over nanoparticle dispersions or colloidal dispersions in photocatalytic reactions activated by solar/visible light irradiation. Long et al. [84] reported on the beneficial effect of intrinsic structural defects in a semicon-

ductor on their chemical reactivity. This affects the stability and electronic properties of semiconductors. In some cases, these defects enhance the potential energy position of the semiconductor electronic bands. Fu et al. recently reported that the photocatalytic efficiency of a semiconductor varied with the type of defects in the composite, whether it was a surface, bulk or adsorbate defect. Defects modify the semiconductor band bending and, therefore, the semiconductor electronic structure. Both factors introduce changes in the semiconductor IFCT process mediating the chemical transformation. The potential to improve semiconductor performance by the introduction of adequate defects in the semiconductor structure is important for any future application of semiconductor devices.



(a)



(b)

Figure 10. (a) Ag(3%)-TiO₂-FeO_x(3%) ternary photocatalyst being irradiated by visible light. The interfacial charge transfer (IFCT) within the composite photocatalyst is shown as a function of their relative potential energy levels leading to the bacterial inactivation process. (b) IFCT and electron transfer mechanism for the stable ternary semiconductor photocatalyst made up of TiO₂/SnO₂/ZnO.

11. Viral Biofilms

Biofilms are aggregates of bacteria/viruses on a surface surrounded by a protective coating that keep for long times on surfaces. They have been recognized as the most effec-

tive way of spreading dangerous pathogens into the environment. Biofilms induce diseases that account for more than 80% of viral/bacterial infections. The biofilms once anchored on supports are resistant to their removal by conventional antiseptics or disinfectants. Viruses are capable of forming complex biofilm-like assemblies, similar to bacterial biofilms, and disseminate viruses dangerous pathogens into the environment for long periods of time, especially within hospitals and healthcare facilities, depending on local conditions. Biofilms are usually resilient to stress, lack of water, pH effects and mechanical stress or impacts. The stages of biofilm formation are schematically suggested in Figure 11 [85].

- (a) Biofilm formation: The first stage in biofilm formation is the adhesion of a virus on a surface by adsorption, involving hydrophobic effects through covalent bonding. Roughness of the surface favors this step and avoids liquids flowing near the biofilms to preclude biofilm formation. The second stage involves bacteria reproduction to form a colony matrix, with this step being concomitant to the growth of an extracellular polymeric shell protecting the colony. In the third/last step, the colony attains its critical mass, ingests nutrients and eliminates metabolic residuals with a kinetics regulated by its enzymes [86].
- (b) The degradation of recalcitrant biofilms occurs in different ways and is suggested next in Figure 12, involving several steps: (a) surface structural modification to preclude biofilm adhesion; (b) the use of bactericidal agents inducing quorum quenching/enzymatic/immunological disruption and (c) the use of catalysts under light or in the dark, leading to bacterial interference in the biofilm [87]. Biofilm destruction is important since many films are resilient to degradation by antibiotic metal/oxide or chloro-compounds. Human coronavirus (HCoV) films lead to respiratory diseases [38]. TiO_2/Ag , TiO_2/Cu and $\text{TiO}_2/\text{Fe}_2\text{O}_3$ composites under light irradiation involving IFCT processes have been discussed above in this study and illustrated with a few relevant examples.

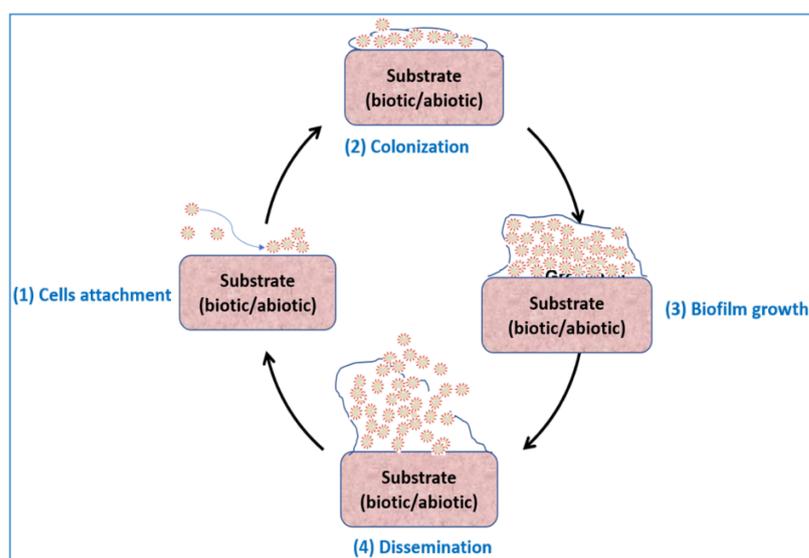


Figure 11. Growth of a virus biofilm. Initially, viruses settle down onto a surface that can either be inanimate or living tissue. Then, secretion of extracellular matrix allows the cells to colonize and anchor on the substrate. This step repeats itself, leading to biofilm growth. Finally, a fully developed biofilm with water channels allows oxygen and nutrients to penetrate to the biofilm and allows processes disseminating viruses into the environment for a long time.

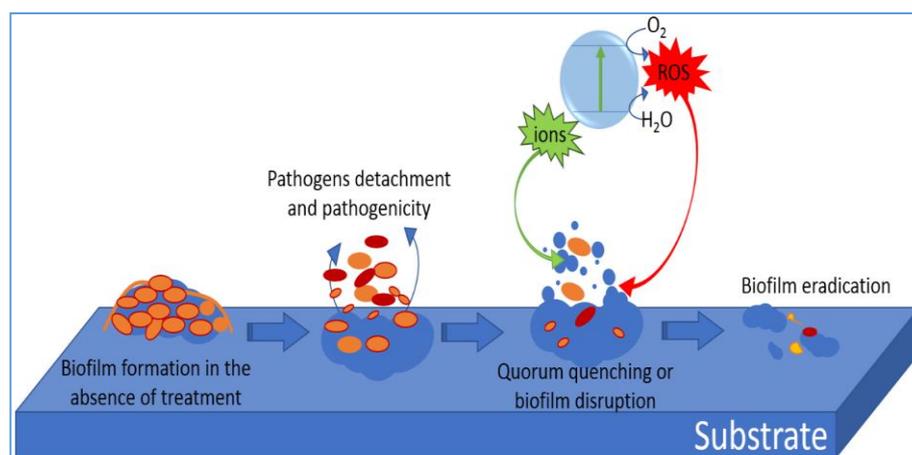


Figure 12. Strategies to neutralize the damage of toxic films spreading pathogens into the environment.

12. Conclusions and Outlook for Future Work

This update briefly describes the scientific bases for the use of semiconductor and metal/semiconductor materials, addressing critical issues in the photocatalytic pathogen inactivation field. Some fundamental issues related to the inactivation of pathogens by materials under visible light irradiation were summarily described. The advantages and limitations of these innovative materials leading to pathogen inactivation are briefly described. The advantages of photochemical processes in bacterial/virus disinfection compared to the use of chlorinated compounds were addressed in detail. The objective of this update was to guide the readership working in microbiology and photocatalysis on disinfection by a mild and environmentally friendly technique.

Future work should address the synthesis of innovative, better performing catalysts. These materials may be applied in conjunction with oxidants commonly present in water reservoirs such as H_2O_2 /organic peroxides. The virus inactivation kinetics of supported films has been reported to be below the one found for clusters/colloids/powders. This is due to the lower surface area available in the film compared to dispersions. However, microparticles dissolved in solution have to be separated after the disinfection process. This is not the case for bactericide films. Films presenting high adsorption for bacteria/viruses could remove this disadvantage and enable their use in photoreactors, useful in continuous processes.

Little is known, to date, on the subject of the viral disinfection kinetics mechanism. Virus disinfection kinetics does not follow the first-order kinetics reported for organic pollutant degradation induced by TiO_2 under light irradiation. This is due to the more complicated viral structure, envelope and repair mechanisms against ROS damage. Much of the present work to inactivate viruses by photocatalytic treatment follows the approach used to inactivate bacteria. However, the structure, surface, size, hydrophobic balance and redox properties of viruses are quite different to those found in bacteria. Innovative materials should be biocompatible, stable and present during long operational times and should present an acceptable corrosion resistance. This enables the release of NPs into solution within the threshold fixed by sanitary regulations. Novel Cu disinfection materials precluding Cu release from the matrix should be designed but should still present contact catalytic sites on their surface that are able to inactivate pathogens. A precedent for this type of material is the case of Fe catalysts/photocatalysts, proceeding without the release of Fe- from TiO_2 matrixes [54]. Future research should address the synthesis of more advanced double metal oxides and other composites made up of biocompatible metal/ions with high redox potentials such as Ag, Mn, Fe and Au compared to traditional semiconductors such as TiO_2 , ZnO and MnO_2 . Controlled release of metals/ions in sub-nanomolar amounts into water bodies has not been addressed, until now, to induce virus inactivation.

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