GENTAMICIN-MONTMORILLONITE INTERCALATION COMPOUNDS AS AN ACTIVE COMPONENT OF HYDROXYPROPYLMETHYLCELLULOSE BIONANOCOMPOSITE FILMS WITH ANTIMICROBIAL PROPERTIES

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Abstract—The present study introduces an overview of gentamicin-clay mineral systems for applications in biomedicine and then focuses on the development of a series of gentamicin/clay hybrid materials to be used as the bioactive phase of hydroxypropylmethylcellulose (HPMC) to produce bionanocomposite membranes possessing antimicrobial activity of interest in wound-dressing applications. Gentamicin (Gt) was adsorbed from aqueous solutions into a montmorillonite (Cloisite®-Na+) to produce intercalation compounds with tunable content of the antibiotic. The hybrids were characterized by CHN chemical analysis, energy-dispersive X-ray analysis, X-ray diffraction, Fourier-transform infrared spectroscopy, and thermogravimetric analysis, confirming the intercalation of Gt by an ion-exchange mechanism. The release of Gt from the hybrids was tested in water and in buffer solution to check their stability. Hybrids with various amounts of Gt were incorporated into a HPMC matrix at various loadings and processed as films by the casting method. The resulting Gt-clay/HPMC bionanocomposites were characterized by means of field-emission scanning electron microscopy, and were also evaluated for their water-adsorption and mechanical properties to confirm their suitability for wound-dressing applications. The antimicrobial activity of the bionanocomposite films was tested in vitro toward various microorganisms (Escherichia coli, Pseudomonas aeruginosa, Staphylococcus aureus, methicillin-resistant Staphylococcus aureus (MRSA), vancomycin-resistant Enterococcus faecium, Acinetobacter baumannii, and Klebsiella pneumonia), showing a complete bacterial reduction even in films with small Gt contents.

Keywords—Antibacterial activity, Bionanocomposite films, Gentamicin, Hydroxypropylmethyl cellulose, Montmorillonite, Wound dressing

INTRODUCTION

The use of clays in health care has been known by humans since ancient times, probably because they are abundant in nature and have been used for a long time for various purposes, from pottery to early writing, including hybrid and composite materials (Faustini et al., 2018). In this way, raw clays found in soils and muds have been useful in the treatment of infections, as de-toxin agents, antidotes of poisons, or topical dermatologic treatments (Rautureau et al., 2017; Finkelman, 2019; Williams, 2019). Clays can be used directly, e.g. to remove toxins and poisons, in geophagy (Allègre, 2012), or as treatment of specific skin diseases, such as Buruli ulcers (Williams & Haydel, 2010; Williams, 2017) or, more recently, employed as carriers of specific agents, typically of drugs (Viseras et al., 2010; Oh et al., 2012; Yang et al., 2016; Ruiz-Hitzky et al., 2019; Stavitskaya et al., 2019; Viseras et al., 2019) but also of other agents such as DNA (Choy et al., 2000; Castro-Smirnov et al., 2016, 2017) or viruses (Ruiz-Hitzky et al., 2009; Wicklein et al., 2012, 2016), for treatment of diverse diseases.

Clays involved in biomedical and pharmaceutical applications include layered 1:1 and 2:1 phyllosilicates, typically kaolinite and smectites, fibrous and tubular clays, such as sepiolite and halloysite nanotubes, as well as certain layered double hydroxides (LDH). Commonly, the final therapeutic system is associated with other components to reach an improved efficiency and a controlled release of the active principle, though its design is determined by the final application. The route of administration can be oral, transdermal, or other to achieve a local release, targeting organs or cells, such as the eyes, the cheek, the vagina, bones, or blood vessels (Kim et al., 2016). The components involved were selected to introduce biocompatibility, to improve solubility and/or adsorption of the drug or muco-adhesive, and/or to target certain properties, including sustained and/or controlled release. Very often, these components are biocompatible and biodegradable polymers and biopolymers, the role of which often is to protect the drug-clay system providing a sustained release (Park et al., 2008), but, also, they are used increasingly to produce a controlled or even targeted release of a drug (Ribeiro et al., 2014; Rebetski et al., 2020).

Antibiotics are among the diverse types of drugs the release of which has been explored in combination with clays. In general, the association with clay improves the antibiotic stability in the medium and affords a sustained release, which is crucial for certain types of applications, e.g. in wound dressing or tissue engineering. One of the antibiotic species that has attracted most interest is gentamicin (Gt). This antibiotic is an...
aminoglycoside species produced by fermentation of *Micromonospora purpurea* that consists of a mixture of basic, water-soluble compounds containing the aminocyclitol 2-deoxystreptamine and two additional amino sugars (MacNeil & Cuerpo, 1995). The Gt formula (Fig. 1a) shares many structural and functional features with other antibiotics containing streptamine or its derivatives, such as streptomycin, neomycin, and kanamycin (Butko et al., 1990), and has been used to treat many types of bacterial infections, particularly those caused by Gram-negative organisms (Moulds & Jeyasingham, 2010). Although application of Gt in human medicine has declined, its use in veterinary medicine and agriculture is still intense, and it could be relevant in the treatment of certain topical injuries or for avoiding internal infection caused by diverse types of implants.

Gentamicin has been associated with several layered silicates from kaolinite (Hui et al., 2019) to smectites (Rapacz-Kmita et al., 2015, 2017a, 2017b; Iannuccelli et al., 2018; Jeong et al., 2018), and has also been incorporated in halloysite nanotubes (Rapacz-Kmita et al., 2019) and layered double hydroxides (Carja et al., 2007). Most of these studies analyzed the release of the antibiotic from the support and provided results on their antibacterial properties against various pathogens such as *E. coli*, *S. aureus*, or *S. epidermidis*. Very few of the studies, however, carried out a proper and accurate characterization of the Gt-clay compounds. The most systematic and complete study was probably that by Iannuccelli et al. (2018) which reported the adsorption of Gt in a natural bentonite from Iglesias (Sardinia, Italy) before and after purification and activation with Na⁺ ions. The raw mineral adsorbed more Gt and so was the first to be used in the present study. Those authors claimed the possible intercalation of Gt and carried out simulations on the possible arrangement of the molecule as a monolayer in the interlayer region although XRD patterns were not shown. The adsorption was driven by an ion-exchange reaction and the presence of S suggested the incorporation of a certain amount as Gt sulfate. Interestingly, the study showed the incorporation of the system into petroleum jelly and application on the skin (in vivo experiment with three volunteers) revealed that the association of the antibiotic with the clay allowed a sustained release and so a more effective anti-infective therapy.

The combination of Gt-clay systems and biocompatible polymers, in particular biopolymers, has been explored with various biomedical purposes mainly addressing the preparation of bioactive coatings for application in implants and scaffolds with the intention of reducing infections. Thus, for instance, Gt was included in the lumen of halloysite nanotubes that were then combined with methyl methacrylate to produce efficient halloysite/poly(methyl methacrylate) bone-cement composites provided with prophylactic activity (Wei et al., 2012). Typical approaches applied to produce coatings of potential relevance in implant applications included Layer-by-Layer (LbL) and spray-drying techniques. Thus, the production of multilayer films by LbL may involve the assembly of a layer of the clay, e.g. montmorillonite, and then one of an organic component, e.g. hyaluronic acid (Wang et al., 2018) or poly-L-lysine (Xu et al., 2017), in which Gt is embedded on the top of substrates such as glass discs, polydimethylsiloxane (PDMS), or silicon wafers. The presence of the antibiotic in the grown coating acted as a biofilm inhibitor as it was released slowly. Gentamicin could also be loaded once the LbL structure is produced as reported for montmorillonite/polyacrylic acid (PAA) systems in which the swelling properties afforded by PAA and the number of grown multilayers are used to control the amount of Gt entrapped and released (Pavlukhina et al., 2014). The creation of multilayered coatings proved especially useful as the presence of the clay layers provided an effective barrier, making it possible to tune and stage release of the antibiotic (Min et al., 2014). Electro-spraying is an alternative methodology applied to produce coatings of montmorillonite/chitosan composite nanospheres in which an antibiotic (e.g. Gt, vancomycin) is included to provide sustained, in situ delivery at implanted bone sites (Kimna et al., 2019). The related electrospinning technique has been explored recently to produce nanoarchitectured composite membranes based on the formation of poly(ε-caprolactone)/montmorillonite fibers in which Gt was also incorporated in order to produce scaffolds provided with bactericidal properties for wound healing, tissue-engineering, and other applications (Reshmi et al., 2018). Other types of processing included the direct solvent-casting preparation of films from dispersions of the polymer incorporating the clay-antibiotic component as reported by Rapacz-Kmita et al. (2020) for developing biodegradable poly(lactic acid)-montmorillonite nanocomposites with antibacterial properties. Electrospinning can be also applied to produce multilayered membranes of poly(lactic acid) reinforced with halloysite, which are later loaded with Gt to produce bioactive films for bone-regeneration applications (Pierchala et al., 2018). The association of Gt with the clay and further combination with water-soluble polymers allowed the production of hydrogels, e.g. poly(vinyl alcohol)/montmorillonite-Gt (Sirousazar, 2013) or chitosan/halloysite-Gt (Luo & Mills, 2019), that can be further conformed, e.g. as films or beads, and submitted to freeze-drying to produce drug-delivery systems for diverse biomedical applications, including wound dressings.

Burn-wound infections are among the most important and potentially serious complications that occur during the acute period following injury. Systemic treatment against infection is limited by inadequate wound perfusion, which restricts migration of host immune cells and the delivery of antimicrobial agents to the wound. In this case, the local concentration of the antibiotics may be insufficient and may lead to bacterial resistance (Elser et al., 2011). The widespread application of a topical antimicrobial agent on the open burn-wound surface can reduce substantially the microbial load and risk of infection (Murphy et al., 2003). The typical burn wound is colonized initially predominantly by Gram-positive organisms, which are replaced by antibiotic-susceptible Gram-negative organisms within 1 week of the burn injury. Gentamicin is used to treat many types of bacterial infections, particularly those caused by Gram-negative organisms and it can protect the recovering tissue from potential infection or re-infection and therefore...
has been used to cure burn wounds (Moulds & Jeyasingham, 2010). Gentamicin-eluting collagen sponges have been found useful in both partial-thickness and full-thickness burn wounds (Elsner et al., 2011). A new concept of wound dressing, which is based on a polyglyconate mesh coated with a porous poly-(DL-lactic-co-glycolic acid) matrix loaded with Gt, is applied to burn wounds (Aviv et al., 2007; Elsner et al., 2011).

Within this general context, the present contribution addresses a systematic study on the adsorption of Gt to montmorillonite and the further incorporation of some of the intercalation compounds produced into hydroxypropylmethylcellulose (HPMC) to produce bionanocomposite films for wound-dressing applications. HPMC (Fig. 1b) has been selected because of its properties as a hydrophilic carrier material including applications in oral drug-delivery systems (Colombo, 1993). The incorporation of the Gt-clay mineral component as a reinforcing and active nanofiller is expected to lead to HPMC bionanocomposite films of the necessary mechanical strength, while the combined binding matrix is aimed at providing adequate moisture control and release of antibiotics to protect the wound bed from infection and promote healing. Mechanical properties, moisture-adsorption behavior and the antimicrobial activity of bionanocomposite films of various compositions were also examined with a view to establishing their appropriateness for potential application as wound-dressing tissues.

EXPERIMENTAL

Materials and Reagents

Gentamicin sulfate from Micromonospora purpurea was supplied by Sigma-Aldrich (Tres Cantos, Spain). The montmorillonite commercialized as Cloisite® Na⁺ Nanoclay, with a cationic exchange capacity of 92.6 meq/100 g, was obtained from Southern Clay Products (Rockwood Additives, Gonzales, Texas, USA) and used in this study without further purification. The polysaccharide hydroxypropylmethylcellulose (HPMC; -22 kDa; viscosity 40–60 cP, 2% in H₂O, 20°C, methoxyl content: 28–20%, hydroxypropoxyl content: 7–12%), sodium phosphate monobasic monohydrate (ACS reagent, ≥98%), and phthalaldehyde solution reagent (P0532), which contains phthalaldehyde (1 mg/mL), Brij® 35, methanol, 2-mercaptoethanol, potassium hydroxide, and boric acid, pH 10.4, were obtained from Sigma-Aldrich (Tres Cantos, Spain). Deionized water (resistance of 18.2 MΩ cm) was obtained with a Maxima Ultra Pure Water system from Elga (Celle, Germany).

Preparation of Gentamicin-montmorillonite Hybrids

A series of Gt sulfate solutions was prepared by dissolving certain amounts of Gt sulfate into deionized water in order to obtain concentrations ranging from 25 to 7300 mg/L, as detailed in Table 1. 50 mg of montmorillonite was added to 5 mL of each Gt solution, reaching a final clay content of 1 wt.%.

Preparation of HPMC/gentamicin-montmorillonite Bionanocomposite Films

The HPMC bionanocomposite films were prepared by incorporation of the freshly prepared hybrid labeled as Gt-Mnt-500, obtained from a starting amount of 500 mg of Gt per g of montmorillonite, in the HPMC solution obtained by dissolving 2 g of HPMC in 100 mL of distilled water. To study the effect of the Gt-Mnt hybrid loading in the HPMC film

![Fig. 1. Chemical structures of a the major components of Gt sulfate and b the polysaccharide hydroxypropylmethylcellulose (HPMC)](https://doi.org/10.1007/s42860-021-00156-3)
matrix, various amounts of nanohybrid were added to 10 mL of the 2% w/v HPMC solution to obtain various film compositions as indicated in Table 2. The mixtures of HPMC solution and different amounts of Gt-Mnt-500 hybrid were homogenized by stirring at 400 rpm at 40°C for 4 h, and then dropped onto plastic plates (100 mm×100 mm) and placed in a laminar flow hood provided with a UV lamp. The samples were sterilized by UV irradiation for 15 min, and then the films were formed by solvent evaporation for 4 days.

**Characterization**

The amount of organic matter in the Gt-Mnt samples was determined by CHN elemental chemical microanalysis using a Perkin-Elmer 2400 analyzer (Waltham, Massachusetts, USA). Powder X-ray diffraction (XRD) patterns of pristine montmorillonite and the antibiotic-loaded hybrids were obtained using a Bruker D8 Advance diffractometer (Billerica, Massachusetts, USA) with CuKα radiation and a scan speed of 0.02°/s. The thermal behaviors of the various materials were analyzed from the simultaneously recorded thermogravimetric (TG) and differential thermal analysis (DTA) curves (using a Seiko (Chiba, Japan) SSC/5200 instrument in experiments carried out under air atmosphere (flux of 100 mL min⁻¹) from room temperature to 1000°C at a heating rate of 10°C min⁻¹). Fourier-transform infrared (FTIR) spectra were measured in transmission absorption mode on a Bruker (Billerica, Massachusetts, USA) IFS 66v/S spectrophotometer, with samples prepared in KBr pellets, throughout the 4000–250 cm⁻¹ region. The surface and cross-sections of the films were observed directly using a Field Emission-Scanning Electron Microscope FEI NOVA NanoSEM 230 (Eindhoven, Netherlands) without conducting coating required. Moisture sorption was investigated using an Aquadyne DVS dynamic water vapor sorption analyzer from Quantachrome Instruments (Boynton Beach, Florida, USA). Moisture-sorption isotherms were recorded at 25 ± 0.3°C in the range of relative humidity from 0 to 95% by using ~7 mg of sample.

**Mechanical Properties**

Films used for tensile tests were conditioned at ~65% RH at 15°C prior to the measurements. A Model 3345 Instron Universal Testing Machine (Instron Engineering Corporation, Canton, Massachusetts, USA) was used to determine the maximum percentage elongation at break (%), and elastic modulus E (or Young’s modulus) according to ASTM standard method D 882-88. The sample films with a rectangular shape (~60 mm×15 mm) were mounted between the grips with an initial separation of ~25 mm. The cross-head speed was set at 5 mm min⁻¹. Three replicates were run for each film sample.

**In vitro Gentamicin Release from the Gentamicin-montmorillonite Hybrids**

The amount of Gt released was detected in the supernatant by UV-Vis spectrophotometry after reaction with phthaldialdehyde. For this purpose, a weighed amount of Gt-Mnt-600 (~60 mg) was added to 5 mL of bidistilled water, kept under shaking at room temperature for various time intervals, and then the samples were centrifuged (15 min, 10,000 rpm, 14,087×g) in order to obtain the supernatants. The determination procedure consisted of mixing 1 mL of the supernatant

| Sample code | Gentamicin concentration (mg/L) | Mass of gentamicin (mg) | Starting amount of gentamicin (mg) per g of clay |
|-------------|---------------------------------|-------------------------|-----------------------------------------------|
| Gt-Mnt-25   | 250                             | 1.25                    | 25                                            |
| Gt-Mnt-50   | 500                             | 2.5                     | 50                                            |
| Gt-Mnt-200  | 2000                            | 10                      | 200                                           |
| Gt-Mnt-500  | 5000                            | 25                      | 500                                           |
| Gt-Mnt-600  | 6000                            | 30                      | 600                                           |
| Gt-Mnt-730  | 7300                            | 36.5                    | 730                                           |

| Sample label | Hybrid content (%) | Clay content (%) | Gentamicin content (%) |
|--------------|--------------------|------------------|------------------------|
| 6.0%Gt-Mnt-HPMC | 62.4              | 56.4             | 6.0                    |
| 3.4%Gt-Mnt-HPMC | 35.6              | 32.2             | 3.4                    |
| 2.1%Gt-Mnt-HPMC | 21.7              | 19.6             | 2.1                    |
| 1.4%Gt-Mnt-HPMC | 14.2              | 12.9             | 1.4                    |
| 0.5%Gt-Mnt-HPMC | 5.2               | 4.7              | 0.50                   |
| 0.26%Gt-Mnt-HPMC | 2.7               | 2.4              | 0.26                   |
| HPMC         | 0                  | 0                | 0                      |
| Mnt-HPMC     | 0                  | 60               | 0                      |
with 1 mL of phthalaldehyde solution reagent and 2 μL of 2-
mercaptoethanol. The mixtures were allowed to stand for
30 min at room temperature, and then the absorbance was
measured between 450 and 320 nm using quartz cuvettes
(10 mm path length) in a Shimadzu (Kyoto, Japan) UV-
2401PC spectrophotometer. The Gt concentrations were ob-
tained for each sample using a calibration curve obtained with
Gt solutions between 2 μg/mL and 15 μg/mL.

Alternatively, 20 mg of Gt-Mnt-200 hybrid and 2 mL of
PBS (phosphate buffer solution, pH 7) were added to plastic
tubes wrapped in aluminum film, and kept in a shaking ther-
mostatic bath at 37°C and 75 rpm. The tubes were removed
from the bath at different time intervals and the suspension was
filtered through a 0.45 μm pore size membrane (MF-Millipore
Membrane Filter, 13 mm diameter) placed on a 13 mm diam-
eter Swinnex® filter holder from Millipore (Burlington, Mas-
achusetts, USA). The solid was air-dried at 40°C under dark
conditions for 24 h, and then the remaining amount of antibi-
otic was determined by chemical analysis.

Antimicrobial Activity
of the HPMC/gentamicin-montmorillonite Films

Study of the antimicrobial properties of the prepared films
was done at the NanoBioMatters S.L., Paterna, Valencia
(Spain) laboratories. Some pathogenic microorganisms present
in hospital environments: *S. aureus* (CECT240, ATCC 6538P),
*E. coli* (CECT 516, ATCC 8739), methicillin-resistant *S. aureus* (CECT 5190, ATCC 43300),
*P. aeruginosa* (CECT 110, ATCC 10145), vancomycin-resistant *E. faecium* (CECT 5253), *A. baumannii* (CECT452,
ACTT 15308), and *K. pneumoniae* (CECT 8453, ATCC 4352),
were purchased from the Spanish Type Culture Collection
(Valencia, Spain) and used to determine the antimicrobial
activity of HPMC films with Gt-based clays. The strains were
stored in Tryptone Soy Broth (TSB) with 20% glycerol at
–80°C until needed. For experimental use, the stock cultures
were maintained by regular subculture on Tryptic Soy Agar
(TSA) slants at 4°C and transferred monthly. A loopful of each
bacterium was placed in 10 mL of TSB and incubated at 37°C
overnight. A 100 μL aliquot from each stock culture was
again transferred to TSB and grown at 37°C to the mida-
exponential phase of growth. These cultures served as the
inoculum in the susceptibility studies according to ASTM
E2149, starting with ~10^5 CFU/mL in the test tubes. These
CFU counts were obtained accurately and reproducibly by
inoculation of 0.1 mL of the culture having an absorbance
value of 0.2, as determined by optical density at 600 nm by
means of UV-Vis spectroscopy.

The antimicrobial effectiveness of the films obtained was
evaluated according to the standard ASTM E2149-01 (Stan-
dard test method for determining the antimicrobial activity of
immobilized antimicrobial agents under dynamic contact con-
ditions, 2001). For this method, 0.05 g of each film was
introduced into test tubes each containing 5 mL of phosphate
buffer solution. A sample with a film without clay was used as
a control. Subsequently, each tube was inoculated with 10^5
cells/mL of *S. aureus* in mid-exponential phase and incubated
in a wrist action shaker (160 rpm) at 25°C for 24 h. Bacterial
counts were then enumerated by sub-cultivation on TSA
plates. All samples were run in duplicate.

Furthermore, the modified disc-diffusion method (Bauer
et al., 1966) was used to evaluate by visual inspection the
antimicrobial activity of the films obtained against all the
bacterial isolates. For this method, tryptic soy agar plates
were seeded with 200 μL of bacterial inoculum (concentration 2.8–
5.3×10^5 cells/mL). 6 mm diameter samples were placed on
inoculated plates and then incubated at 37°C for 24 h. All
samples were run in duplicate. An inhibition zone was moni-
tored after incubation, and the presence of a bacterial growth
inhibition halo around the samples was studied.

RESULTS AND DISCUSSION

Preparation of Gentamicin-montmorillonite Hybrids
and Study of the Adsorption Isotherm

The amino groups of the sugar rings of Gt exhibit variable
pKa values which range from 5.7 to 9.9 (Lesniak et al., 2003).
Thus, Gt carries a net positive charge up to pH values of ~10
which makes possible its intercalation in Na+-montmorillonite
by means of cationic exchange processes. The adsorption
isotherm at 294 K from Gt sulfate solutions on Na+-montmo-
rillonite (Fig. 2) shows the amount of Gt adsorbed, deduced
from the CHN chemical analyses of the Gt-Mnt hybrids. In
order to investigate the adsorption process, three models of
equilibrium isotherms were applied, the Langmuir (Eq. 1), the
Freundlich (Eq. 2), and the Redlich-Peterson (Eq. 3)
isotherms:

\[
q_e = \frac{q_m K_a C_e}{1 + K_a C_e}
\]

where \(q_e\) is the equilibrium adsorption capacity (mg/g), \(C_e\) is
the equilibrium liquid phase concentration (mg/L), \(q_m\) is the
maximum adsorption capacity (mg/g), and \(K_a\) is adsorption
equilibrium constant (L/mg).

\[
q_e = K_F C_e^{1/n}
\]

where \(C_e\) is the equilibrium concentration in the solution (mg/
L), \(q_e\) is the equilibrium adsorption capacity (mg/g), and \(K_F\)
and 1/n are empirical constants.

\[
q_e = \frac{A C_e^g}{1 + B C_e^g}
\]

where \(A, B, \) and \(g\) (0 < g < 1) are three constants of this model
which combines both the Freundlich and Langmuir isotherms.
As shown in Table 3, the Langmuir and the Redlich-
Peterson isotherms provided the best fit to the adsorption data
of Gt on Na+-montmorillonite at 294 K, indicating the homo-
genicity in the adsorption sites.

The Gibbs energy of adsorption, \(\Delta G^o\), calculated for these
adsorption data was ~18.6 kJ mol\(^{-1}\), which is of the order of
values obtained for adsorption of other antibiotics in clay
minerals (Song et al., 2019). The negative value of Gibbs
energy confirms the feasibility of the process and the
spontaneous nature of adsorption of Gt on Na+-montmorillonite. The cationic exchange sites in the clay, with a CEC of 92.6 mEq/100 g, are assumed to be able to adsorb a certain amount of charged Gt due to protonation of $–\text{NH}_3^+$ and $–\text{NH}_2^2–$. The total charge depends on the protonation state of hydroxyl and amine groups at a given pH. In the present work, the pH value in the isotherm experiment was ~5.5. According to the study by Lesniak et al. (2003), the ionic forms of Gt at the experimental pH should be mainly Gt$^{5+}$, with a small amount of Gt$^{4+}$. As a result, the total amount of Gt that could be retained by the clay used in this study should be between 0.185 mmol/g and 0.232 mmol/g. The maximum amounts of adsorbed Gt according to the adsorption isotherm were ~0.22–0.24 mmol/g. Note, however, that these values were obtained for unwashed samples, where the amount adsorbed can exceed the CEC of the clay due to molecules adsorbed on the external surface. The chemical analysis of the Gt-Mnt-600 hybrid after washing gave a value of 0.183 mmol/g, which would correspond mainly to intercalated Gt and suggests that Gt$^{5+}$ is the Gt species intercalated in the clay interlayer space.

Characterization of Gentamicin-montmorillonite Hybrids

The XRD patterns of the pristine clay and the Gt-montmorillonite nanohybrids (Fig. 3) confirm the intercalation of Gt in the clay interlayer space by the shift of the 001 reflection toward lower 2θ values as the Gt:clay ratio increased. As shown in XRD results, the $d_{001}$ spacing reached 1.43 nm when the adsorption reached equilibrium, corresponding to an interlayer distance of 0.47 nm. The lower-energy configuration of Gt, with a two-dimensional size of a Gt molecule, was reported by Doadrio et al. (2004) to be 0.52 nm × 1.53 nm. For Gt-Mnt-500, Gt-Mnt-600, and Gt-Mnt-700 hybrid samples, the distance between two consecutive smectite layers was ~0.47 nm, similar to the size of a Gt molecule in one dimension. The result suggests that the Gt molecule is intercalated as a monolayer with the three rings of the Gt molecule positioned roughly parallel to the smectite layers. This finding is also in agreement with results and modeling of Gt adsorbed on bentonites reported by Iannuccelli et al. (2018).

In order to provide experimental confirmation of the optimal layer spacing, the FWHM profile as a function of $d_{001}$ of Gt-Mnt samples was examined. Changes in FWHM have been attributed previously to the evolution of different configurations of the intercalated molecules, whereby, in a profile of FWHM values as a function of $d_{001}$, an increase in FWHM typically indicates interstratifications of different layer types. As reported previously (Aristide et al., 2013), a minimum FWHM would be reached when the clay interlayers are saturated with adsorbed Gt, and the corresponding layer spacing would be considered optimal for the binding of Gt in the interlayer. With the increasing amount of Gt intercalated in the clay, the FWHM decreased from values of ~0.9°2θ for Mnt and Gt-Mnt-25 to values of ~0.5°2θ for the other hybrids with greater Gt loadings. Such a decrease in FWHM was proposed to be due to a gradual change in the interlayer configuration occurring homogeneously within all layers, wherein the molecules would change position or orientation, e.g. from being stretched to a relatively flat chain molecule with high energy to being a stable molecule chain which is less distorted.

Energy-dispersive X-ray analysis (EDX) measurements were performed on the starting montmorillonite, and on the nanohybrid samples labeled as Gt-Mnt-50, Gt-Mnt-200, and Gt-Mnt-730. The EDX results indicate a decreasing Na+ ratio with the increasing amount of Gt adsorbed on the clay Wt(Na+) (Fig. 3B). In this way, the predominance of ion-exchange mechanisms driving the intercalation of the Gt into the phyllosilicate substrate is confirmed experimentally.

The FTIR spectra were registered for Na+-montmorillonite, the Gt-montmorillonite nanohybrid labeled as Gt-Mnt-730, and pristine Gt sulfate (Fig. 4). Beside the vibrational bands characteristic of the silicate, $\nu_{\text{HOH}}$ of Al,Mg(OH) 3633 cm$^{-1}$; $\nu_{\text{HOH}}$ of H$_2$O 3451 and 3249 cm$^{-1}$; $\delta_{\text{HOH}}$ 1637 cm$^{-1}$; and $\nu_{\text{SiO}}$ of Si–O–Si 1045 cm$^{-1}$ (Fig. 4a), the bands attributed to the intercalated Gt are also observed in the spectrum of Gt-clay hybrid (Fig. 4b). The characteristic vibration bands of Gt in the hybrid material are observed at practically the same frequencies as the Gt monolayer. The FTIR data of Gt on Na+-montmorillonite at 294 K were fitted to the non-linear Langmuir, Freundlich, and Redlich-Peterson equations for the adsorption data of Gt on Na+-montmorillonite at 294 K.

Table 3. Isotherm parameters obtained using the non-linear Langmuir, Freundlich, and Redlich-Peterson equations for the adsorption data of Gt on Na+-montmorillonite at 294 K

| Isotherm model    | Parameters                  | $r^2$ |
|-------------------|-----------------------------|-------|
| Langmuir          | $q_m$ (mg/g)                | 117 ± 5 | 0.983 |
|                   | $K_m$ (L/mg)                | 0.0039 ± 0.0009 |
|                   | $n$                         | 3.1 ± 0.7 | 0.907 |
| Freundlich        | $K_n$ (mg/g)(L/mg)$^{1/n}$   | 8.0 ± 4.8 |
|                   | $g$                         | 1.2 ± 0.1 | 0.987 |
| Redlich-Peterson  | $B$ (L/mg)                  | 0.0006 ± 0.0008 |
|                   | $A$ (L/g)                   | 0.33 ± 0.08 |

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wavenumbers as in the pristine Gt sulfate (Fig. 4c). This confirms that Gt is present as a protonated species in interaction with the silicate layers, which now act as counterions. Also, the N–H in-plane deformation vibration bands of amine groups of Gt appeared at a value very close to that of δ\(\text{NH}_3^+\) of the silicate, and a band due to the overlapping of those bands was observed at 1632 cm\(^{-1}\) in the spectrum of the Gt-Mnt hybrid, which suggests a possible intermolecular hydrogen bonding between water and Gt molecules in the interlayer region (Rapacz-Kmita et al., 2015). At ~1048 cm\(^{-1}\) in the spectrum of Gt-clay sample, the vibration band for C–O–C of Gt overlaps the band of Si–O–Si of the clay. The vibration band at 1531 cm\(^{-1}\) in the spectrum of Gt (Fig. 4c), which corresponds to the symmetric deformation vibration (\(\delta_{\text{NH}_3^+}\)) of the protonated primary amine group, shifted to 1529 cm\(^{-1}\) after Gt was intercalated in the clay.

According to the TG curves (Fig. 5), the mass loss due to water-molecule desorption takes place from 30–87°C and at 160°C for starting silicate and Gt-clay, respectively. This may be due to the interaction of the water molecules with intercalated Gt in the interlayer region, as suggested by FTIR data. Such mass loss was ~4.6% in the starting silicate, while the Gt-clay hybrid showed a slightly larger value of 5.4%, which may be due to the high water-retention capacity of Gt. The thermal decomposition process of Gt was similar in the hybrid material to what it was in the pristine compound, although the different decomposition steps are not as well defined in the intercalated material. The last mass loss ascribed to dehydroxylation of the clay started at a slightly lower temperature, ~500°C in the hybrid instead of 580°C in the pristine clay, partially overlapping the last decomposition phase of Gt. This effect is probably due to the different interlayer cation present in the hybrid.

The stability of Gt in the montmorillonite-based hybrids was evaluated using the Gt-Mnt-200 and Gt-Mnt-600 hybrid samples. The Gt-Mnt-600 sample contains 108.6 mg/g clay (0.234 mmol/g clay), i.e. with excess Gt probably adsorbed on the external surface of the clay. This sample was immersed in deionized water and, after given intervals, the values of Gt released were determined in the supernatant by UV spectro-photometry following reaction with phthaldialdehyde. The remaining amounts of Gt in the hybrid after various time intervals (Fig. 6) show that, after 24 h, ~15% of Gt was released, with ~0.198 mmol/g remaining in the hybrid. If the same study had been carried out with a sample previously washed to remove the excess adsorbed antibiotic, no Gt was detected in the supernatant by the phthaldialdehyde procedure after 24 h.

The stability of the Gt-Mnt hybrids was also evaluated in a buffered solution at pH 7 using the Gt-Mnt-200 hybrid sample, which contains 0.172 mmol of antibiotic per g of clay. As this value is less than the CEC of the clay, the antibiotic molecules are probably intercalated in the clay interlayer space with no excess Gt adsorbed on the external surface. The results of chemical analyses of the solids recovered at each studied time showed that large amounts of Gt were not released from the Gt-Mnt-200 hybrid under these experimental conditions (Fig. 6). The quantity of Gt adsorbed on the clay decreased slightly from the initial 0.172 mmol/g to 0.167 mmol/g after 24 h. At pH 7, Gt was protonated as a mixture of Gt\(^{3+}\), Gt\(^{4+}\), and Gt\(^{5+}\), which causes Gt compounds to be retained in the interlayer of montmorillonite because of high electrostatic attraction; they cannot be replaced easily by sodium or any other ions present in the medium.

**HPMC/gentamicin-montmorillonite Bionanocomposite Films**

A set of HPMC/Gt-montmorillonite bionanocomposite films was prepared by dispersing appropriate amounts of the Gt-Mnt-500 hybrid sample, containing 106.1 mg/g of clay, in aliquots of HPMC aqueous solution, as detailed in the Experimental Section. The compositions of the bionanocomposite films prepared are summarized in Table 2. After drying, the films were separated from the molds as self-standing films showing a uniform aspect. The morphology of the Gt-Mnt-HPMC bionanocomposite films was analyzed by FE-SEM (Fig. 7). The surface aspect of the 0.5%Gt-Mnt-HPMC membrane (Fig. 7a) with a relatively small amount of hybrid (5.2%) shows a smooth aspect where some small dots, probably related to hybrid...
particles which were embedded in the polymer located close to the surface. The presence of such spots was discerned more clearly when the hybrid content increased, as observed in the images of the 1.4%Gt-Mnt-HPMC membrane (Fig. 7b, c). In this case, the image at higher magnification (Fig. 7c) allowed observation of particles of the clay inserted in the polymer bulk. The cross-section image of this membrane (Fig. 7d) confirms the homogeneous distribution of the nanofiller on the polymer matrix which appeared as a continuous phase with a consistent texture.

The success of antibiotic-loaded films depends heavily on favorable mechanical properties. The pure HPMC film has an elastic modulus of 2.7 GPa and a percentage of elongation at break of ~17%, as determined from the corresponding stress-strain curves (Fig. S1, Supplementary Material). The effects of

![IR spectra](https://example.com/ir spectra.png)

**Fig. 4.** IR spectra (4000–250 cm\(^{-1}\) region) of (a) the starting Na\(^{+}\)-montmorillonite, (b) the Gt-clay nanohybrid labeled as Gt-Mnt-730, and (c) starting Gt sulfate

![TG and DTG curves](https://example.com/tg dtg curves.png)

**Fig. 5.** A TG and B DTG curves in the 25–1000°C temperature range, obtained under air flow, of (a) Na\(^{+}\)-montmorillonite, (b) Gt-Mnt-730 hybrid sample, and (c) starting Gt sulfate

![Release profiles](https://example.com/release profiles.png)

**Fig. 6.** Release profiles from Gt-Mnt-600 hybrid sample in deionized water and from Gt-Mnt-200 in 50 mM PBS (pH=7) at 37°C, showing the amount of Gt remaining adsorbed in each hybrid at different time intervals
increasing Gt-Mnt content on the elastic or Young’s modulus and elongation at break values of the Gt-Mnt-HPMC films obtained are listed in Table 4. The incorporation of hybrid in the HPMC matrix produced an increase in the elastic modulus values with respect to that of pure HPMC film, indicating an increase in the films’ stiffness. The increase in hybrid loading also gave rise to a decrease in the value of elongation at break, which is observed commonly in reinforced nanocomposites and is probably attributed to the reduction in mobility of the polysaccharide chains due to their interaction with the reinforcing agent.

The water-absorption tendency of the films containing different loadings of Gt-Mnt hybrid was studied in comparison to the pristine HPMC film. The water sorption isotherms (Fig. 8) show the weight increase due to water uptake expressed as grams of water incorporated per 100 g of dry sample as a function of the relative humidity (%). The adsorption-desorption isotherms of the bionanocomposite films show a very similar behavior, also close to that of the pristine HPMC, with just minor differences at relative humidity values of >80% where a slight reduction in water uptake was observed as the hybrid amount was increased. The

### Table 4. Elastic modulus and percentage of elongation at break of HPMC bionanocomposite films at different loadings of the Gt-Mnt hybrid

| Sample           | Gt-Mnt content (%) | Elastic modulus (GPa) | Elongation at break (%) |
|------------------|--------------------|-----------------------|-------------------------|
| HPMC             | 0                  | 2.7 ± 0.1             | 17 ± 6                  |
| 0.26%Gt-Mnt-HPMC | 2.7                | 3.08 ± 0.05           | 9 ± 2                   |
| 0.5%Gt-Mnt-HPMC  | 5.2                | 3.26 ± 0.30           | 4.8 ± 0.9               |
| 1.4%Gt-Mnt-HPMC  | 14.2               | 4.10 ± 0.32           | 3.1 ± 0.4               |
results confirmed that incorporation of Gt-Mnt hybrids does not reduce the hydrophilicity of HPMC, which is beneficial for wound-dressing applications (Jin et al., 2016).

The antimicrobial activity of the Gt-Mnt-HPMC bionanocomposite films was studied according to ASTM E2149. The HPMC-based films showed antimicrobial properties compared with the control tube with peptone water. The HPMC or/and the montmorillonite samples used in the present study, evaluated as controls, i.e. free of Gt, showed no detectable antimicrobial effect. The antimicrobial properties against *S. aureus* (CECT 240, ATCC 6538P) of the samples obtained are listed in Table 5. These samples presented maximum antimicrobial properties regardless of the amount of Gt within the film.

The antimicrobial properties of HPMC films labeled as 0.5%Gt-Mnt-HPMC, containing 4.7% of clay and 0.5% of Gt, were evaluated by visual inspection according to the modified disc diffusion method (Fig. 9). The HPMC film with modified clay was able to reduce completely microbial development around the sample for all of the microorganisms studied except for methicillin-resistant *S. aureus* (MRSA), where only a slight colony reduction was observed. Experiments carried out with HPMC films or Whatman filters loaded directly with Gt showed a clear inhibition zone in agar plates seeded with *E. coli* and *S. aureus* (Fig. S2, Supplementary Material). The control samples, without clay, were dissolved completely during the incubation process due to their low water resistance, but also because they could serve as nutrient media for the microbial growth. On the other hand, the film with Gt-modified clay does not allow microbial growth on itself and, consequently, it cannot be degraded by microorganisms. Moreover, the incorporation of clays within the polymer matrix could enhance the physical properties of the film, increasing their water resistance.

The antimicrobial properties of Gt incorporated in a biopolymer matrix for medical purposes have already been studied by several authors as described in the introduction section. The effectiveness of PHBV/Gt formulations for therapeutic modalities in implant-related staphylococcal infections was demonstrated by Rossi et al. (2004). A biodegradable PLA/PEG Gt delivery system that showed great potential for prophylaxis of post-operative infection was described by Huang and Chung (2001). Great antimicrobial properties in PLA composites with silver-organomodified clays were demonstrated by Busolo et al. (2010). The combination of antibiotic-modified clays with inorganic antimicrobials such as silver-modified clays could lead to additive or even synergistic biocide effects against a broad range of microorganisms. Consequently, the amount of antibiotic or antimicrobial agents required to achieve a satisfactory effect could decrease significantly. In fact, some recent works have addressed the use of clay-based systems in combination with biopolymers to produce effective bionanocomposites with a biocide effect (Aguzzi et al., 2014; Ambrogi et al., 2017; Rebitski et al., 2018; Lisuzzo et al., 2020). In particular, examples of Gt-clay bionanocomposites for uses in implants and scaffolds have been described (e.g. Kimna et al., 2019), though few approaches have been made in terms of systems for applications in wound- and burn-dressings tissues, where the bionanocomposite system reported here is very promising.

**Table 5.** Antimicrobial properties of the obtained films against *S. aureus* according to ASTM E2149

| Sample description | CFU/mL | CFU/mL | Mean | % Bacterial reduction |
|--------------------|--------|--------|------|-----------------------|
| Control without film | 2.02E+07 | 2.02E+07 | 2.02E+07 | - |
| Control with HPMC film | 1.92E+07 | 1.56E+07 | 1.74E+07 | - |
| 0.26%Gt-Mnt-HPMC | 1.00E+00 | 1.00E+00 | 1.00E+00 | 100.00 |
| 0.5%Gt-Mnt-HPMC | 1.00E+00 | 1.00E+00 | 1.00E+00 | 100.00 |
| 1.4%Gt-Mnt-HPMC | 1.00E+00 | 1.00E+00 | 1.00E+00 | 100.00 |
| 2.1%Gt-Mnt-HPMC | 1.00E+00 | 1.00E+00 | 1.00E+00 | 100.00 |
| 3.4%Gt-Mnt-HPMC | 1.00E+00 | 1.00E+00 | 1.00E+00 | 100.00 |
| 6.0%Gt-Mnt-HPMC | 1.00E+00 | 1.00E+00 | 1.00E+00 | 100.00 |

CONCLUSIONS

Amongst other antibiotics, Gt has been associated with clay minerals of diverse characteristics (e.g. smectites,
kaolinite, halloysite) to produce, alone or in combination with polymers, bioactive systems designed mainly for use in implants and scaffolds in bone-regeneration and tissue-engineering applications. In the present study, Gt was adsorbed to montmorillonite (Cloisite®-Na+) by means of an ion-exchange mechanism, being arranged as a monolayer of Gt⁵⁺ cations compensating the negative charge of the silicate layers. The interaction of the Gt ions with the silicate is strong enough to provide a stable hybrid material, where the release in water and buffer media is slowed. This is promising in terms of the development of a sustained-delivery system. Intercalation compounds with a tuned amount of the drug were later assembled to HPMC at different loadings. These bionanocomposites can be processed easily as films offering better mechanical resistance than the pristine biopolymer, although the elasticity decreases with the hybrid loading. The presence of the hybrids does not reduce the hydrophilic character of HPMC showing similar water sorption isotherms. Besides, the presence of the clay reduces the swelling ability of HPMC films, avoiding their disintegration. Antimicrobial, in vitro tests for Staphylococcus aureus (ASTM E2149) show a maximum efficacy even for small Gt contents, proving interest in applications as a healing-wound tissue. Such significant antibacterial activity with small Gt contents is advantageous because low hybrid loadings in the HPMC films contribute to the preservation of the elasticity of the films. The bacterial reduction was also confirmed in agar plates with other microorganisms (E. coli, P. aeruginosa, S. aureus, methicillin-resistant S. aureus, vancomycin-resistant E. faecium, A. baumannii, and K. pneumoniae), showing, in all cases, an inhibition halo except for the case of MRSA, where the inhibitory effect is only partial. These bionanocomposite films show promising properties for application as wound dressings.

SUPPLEMENTARY INFORMATION
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Compliance with Ethical Statements
Conflict of Interest
The authors declare that they have no conflict of interest.

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