Article

Benefits of Polydopamine as Particle/Matrix Interface in Polylactide/PD-BaSO₄ Scaffolds

Naroa Sadaba¹, Aitor Larrañaga¹,², Gemma Orpella-Aceret², Ana F. Bettencourt³, Victor Martin⁴,⁵, Manus Biggs², Isabel A. C. Ribeiro³, Jone M. Ugartemendia¹*, Jose-Ramon Sarasua¹, and Ester Zuza¹,

¹ Department of Mining-Metallurgy Engineering and Materials Science, School of Engineering EIB 1, University of the Basque Country (UPV/EHU) and Polymat, 48013 Bilbao, Spain; naroasadaba89@gmail.com (N.S.); aitor.larranagae@ehu.eus (A.L.); jone.munoz@ehu.eus (J.M.U.); jr.sarasua@ehu.eus (J.-R.S.)
² Center for Research in Medical Devices (CÚRAM), National University of Ireland (NUIG), Newcastle Road, H91 W2TY Galway, Ireland; gemma.orpella@gmail.com (G.O.-A.); manus.biggs@nuigalway.ie (M.B.)
³ Research Institute for Medicines (iMed.ULisboa), Faculdade de Farmácia, Universidade de Lisboa, Avenida Prof. Gama Pinto, 1649-003 Lisboa, Portugal; asimao@ff.ul.isboa.pt (A.F.B.); ribeiroi@campus.ul.pt (I.A.C.R.)
⁴ Laboratory for Bone Metabolism and Regeneration, Faculty of Dental Medicine, U. Porto Rua Dr. Manuel Pereira da Silva, 4200-393 Porto, Portugal; victorzmartin@gmail.com
⁵ Portugal/LAQV/REQUIMTE, U. Porto, 4160-007 Porto, Portugal
* Correspondence: ester.zuza@ehu.eus

Received: 30 June 2020; Accepted: 28 July 2020; Published: 31 July 2020

Abstract: This work reports the versatility of polydopamine (PD) when applied as a particle coating in a composite of polylactide (PLA). Polydopamine was observed to increase the particle–matrix interface strength and facilitate the adsorption of drugs to the material surface. Here, barium sulfate radiopaque particles were functionalized with polydopamine and integrated into a polylactide matrix, leading to the formulation of a biodegradable and X-ray opaque material with enhanced mechanical properties. Polydopamine functionalized barium sulfate particles also facilitated the adsorption and release of the antibiotic levofloxacin. Analysis of the antibacterial capacity of these composites and the metabolic activity and proliferation of human dermal fibroblasts in vitro demonstrated that these materials are non-cytotoxic and can be 3D printed to formulate complex biocompatible materials for bone fixation devices.

Keywords: biodegradable; composite; polylactide; barium sulfate; polydopamine; melt processing; template; 3D-printing; scaffolds; antibiotic; levofloxacin

1. Introduction

Polylactides are biodegradable polymers with great potential for the reconstruction of damaged tissues [1]. As is well known polylactide can be either semicrystalline (poly-L-lactide and poly-D-lactide (PLLA/PDLA)) or fully amorphous (poly-D,L-lactide (PDLLA)). For reconstruction of hard tissue stiff and strong PLLA or PDLA is preferred. Having a glass transition temperature around 60 °C, at body temperature semicrystalline PLA will be in glassy state (vs. elastomeric) and being crystalline at the same it will fulfill the requirements of a high stiffness and strength that a polymeric biomaterial requires for bone reconstruction in the form of fixation devices of small size [2]. Shortcomings to the use of PLA in bone reconstruction as opposed to e.g., metals (e.g., stainless steels or titanium alloys), include transparency to X-ray and mechanical brittleness [3,4]. However, high mass element containing fillers may be added to PLA to obtain radiopaque composites and improve its mechanical properties. Barium
sulfate (BaSO$_4$) [5–7], ferrous oxide (Fe$_3$O$_4$) [8] and bismuth oxide [9,10] have been previously reported as radiopaque composite additives.

Inorganic particulate reinforcements can enhance the mechanical properties of polymers and confer additional filler-specific properties to the matrix [11]. It is also well known that the enhancement of strength related mechanical properties (elastic modulus, resistance to yield and ultimate stress) is much more noticeable with continuous fiber composites [12], however continuous fiber formulations do not lend themselves well to free-form manufacturing techniques such as extrusion, injection molding or, as is the case, 3D printing [13–15]. Particularly, 3D printing is progressing very rapidly as an advanced manufacturing technique also for particulate composites. In this domain, there are also considerable achievements in the development of drug delivery systems. Therefore, in this work we selected a 3D printable system based on a BaSO$_4$ particulate and PLA composites with an antimicrobial drug incorporation for the potential use in fixation devices for bone tissue regeneration.

Polydopamine is a newfangled nature inspired adhesive normally used for immobilization on materials surfaces of small molecules as drugs and proteins for biomedical use [16]. In addition, it has been used for improving the thermal stability of composites [17]. In most cases promoted by its biomimetic adhesivity [18], polydopamine has been used to confer anchoring sites for biologically active molecules as drugs and proteins, as nanocapsules or nanocarriers [19,20]. Nevertheless, the main use of polydopamine is as substrate coating due to its ability and efficiency for conjugation with bioactive molecules as growth factors [21] and drugs [22]. Zhang et al. immobilize bone morphogenic protein 2 (BMP-2) and insulin-like growth factor 1 (IGF-1) on polydopamine coated scaffold, reducing the burst release of the factors and endowing long-term osteoconductivity [21]. Furthermore, titanium nanotubes have been coated with polydopamine to adsorb dexamethasone sodium phosphate, an anti-inflammatory and modulator of osteogenic differentiation, leading to slow, sustained and controlled drug release [22]. However, few publications have reported the use of polydopamine in polymer matrix composites reinforced with inorganic and radiopaque compounds. Almost all the publications referred to composites focus their research on dispersion and biocompatibility, for example, on boron nitride nanotubes [23], bioglass in polylactides [17] and multi-walled carbon nanotubes (MWCNT) in polyurethanes (PU) [24]. Few works have reported changes in mechanical properties of polymer composites due to polydopamine coating of the inorganic phase, one of them is that on polylactide reinforced by polydopamine functionalized halloysite nanotubes [25].

The aim of this work is to study a novel polymeric biodegradable composite system having potential for use as a biodegradable polymeric fixation template applicable to bone reconstruction. In this sense a novel composite system is studied and developed including features of radiopacity (RO) [26], mechanical toughness [3,5,11,12] and antimicrobial properties for prevention of biofilm formation [27]. In this aim, barium sulfate particles were functionalized with polydopamine and incorporated into a PLA matrix to obtain scaffolds by 3D-printing. It will be proved that this composites presents X-ray visibility and enhanced mechanical properties (stiffness, strength, ductility and toughness) through the promotion of new deformation mechanisms caused by proper particle size and specific interactions at the fiber/matrix interfaces leading to good adhesion and improved mechanical performance. The incorporation of polydopamine to the systems presents additional benefits since provides specific sites for biologically active molecules that can be incorporated to the polymeric scaffolds leading to improvements in biocompatibility and other requirements for specific tissue engineering in situ.

2. Results and Discussion

2.1. Improvements in the Mechanical Properties by Incorporation of Polydopamine Coated BaSO$_4$ Particles into PLA

The tensile stress–strain curves and mechanical properties of neat polylactide and its composites (PLA/PD-BaSO$_4$) were determined by tensile testing. Neat PLA, not reaching a yield point, showed typical brittle behavior, exhibiting around 6% of elongation before failure and 60 MPa of tensile strength.
In contrast, PLA/PD-BaSO\textsubscript{4} composites showed, for all PD-BaSO\textsubscript{4} amounts studied, a clear yield point and an extended ductile behavior with a dramatic increase in elongation at break (supporting information S1). This is accompanied with a moderate increase in the elastic modulus. As identified in our previous works with PLA/BaSO\textsubscript{4} composites [3,5], the plastic deformation in composites occurs because the rigid particles act as stress concentrators and, after particles debonding from the matrix and being of a 0.7–1.9 \(\mu\)m size [28], activate, at a point of applied stress, crazing and shear yielding deformation mechanisms. Since polydopamine covered particles (PD-BaSO\textsubscript{4}) present 1.25 \(\mu\)m size and interactions with the matrix through the polydopamine interface, in these composites, increases in all mechanical properties (elastic modulus, strength, ductility and toughness) are achieved [29].

Table 1 shows the tensile mechanical properties of PLA/PD-BaSO\textsubscript{4} composites. As can be observed, up to a 2 wt.% BaSO\textsubscript{4} content in composites, Young’s modulus, ductility and toughness improvements can be observed. Further, if these results are to be compared with those of PLA/BaSO\textsubscript{4}, the non PD functionalized composite counterparts, enhancements in stiffness and strength together with ductility and toughness can be noticed in the former. Please also note that a huge >2300% increase in elongation at break are determined in these novel composite formulations in regard to neat PLA, which brings about a dramatic improvement in toughness. This is attributable to specific interactions stablished between the ester groups of polylactide and the alcohol groups of polydopamine coating of the particles surface [18,30] bringing about a stronger fiber/matrix interface. It is also noticeable that a 30% increase in the elongation at break is obtained in PD coated BaSO\textsubscript{4} PLA composites in regard to the composites without polydopamine coating [3]. Finally, beyond the 2 wt.% particle composition, a slight general decrease in mechanical properties are noticeable, suggesting the existence of particle aggregates and bundles beyond this point.

Table 1. Mechanical properties of neat polylactide (PLA) and PLA/polydopamine-barium sulfate (PD-BaSO\textsubscript{4}) composites with respect to wt.% of PD-BaSO\textsubscript{4}. \(\sigma_y\): yield stress, E: Young’s modulus, \(\sigma_t\): stress at break, \(\varepsilon_b\): elongation at break and TT: tensile toughness.

| PD-BaSO\textsubscript{4} (wt.%) | E (MPa) | \(\sigma_y\) (MPa) | \(\sigma_t\) (MPa) | \(\varepsilon_b\) (%) | TT (J/m\textsuperscript{2}) |
|--------------------------|--------|-----------------|-----------------|------------------|------------------|
| 0                        | 1243 ± 96 | -               | 67.1 ± 0.5      | 9.2 ± 0.5        | 3.8 ± 0.6        |
| 0.5                      | 1309 ± 51 | 78.7 ± 0.4      | 56.3 ± 5.4      | 132.2 ± 13.1     | 67.5 ± 6.1       |
| 1                        | 1396 ± 109 | 77.7 ± 0.8      | 56.0 ± 4.8      | 146.3 ± 9.8      | 73.6 ± 4.6       |
| 2                        | 1417 ± 86  | 78.4 ± 1.7      | 62.5 ± 5.2      | 182.2 ± 5.9      | 95.1 ± 1.7       |
| 5                        | 1415 ± 49  | 75.7 ± 0.6      | 57.6± 5.8       | 171.9 ± 2.9      | 85.9 ± 0.8       |
| 10                       | 1350 ± 124 | 74.1 ± 0.8      | 55.3 ± 4.3      | 154.6 ± 8.6      | 76.2 ± 2.8       |

Figure 1 represents the break surface of the PLA/PD-BaSO\textsubscript{4} 2 wt.% composite. The SEM image corroborates the ductile and tough behavior of the PLA containing 2 wt.% barium sulfate PD-coated particles with values provided in Table 1. A high level of dispersion of PD coated particles (indicated with white and straight arrows) within the PLA matrix can be observed. In the pictures, ductile-deformed threads of PLA matrix appear too, thinner than a micrometer in diameter. This means that the matrix has been stretched during the tensile test beyond the yield point leaving long fibers (red dot-line arrows). Therefore, radiopaque PD-BaSO\textsubscript{4} particles were able to act as fixed points that under the external applied stress assisting the PLA matrix to develop specific ductile deformation mechanisms.
It is also remarkable that the values obtained for the 2 wt.% and 10 wt.% composites do not differ much from each other. However, radiopacity values, as expected, show a significant increase with the increasing amount of PD-BaSO₄ in composites (Figures S3) and particularly in the 10 wt.% composite. Consequently, it was concluded that among all compositions studied the 10 wt.% PLA/PD-BaSO₄ composite presents the optimal properties (Figures S1 and S3) and hence an additional analysis is conducted for this composite.

The mechanical properties of the 3D printed scaffolds are also analyzed. Details of the scaffolds geometry design can be seen in Figure S4. In this case, mechanical properties were measured in compression to mimic the working conditions of the device. Non-reinforced polylactide (as reference) and the 10 wt.% PLA/PD-BaSO₄ scaffolds with 55% of porosity have been tested.

Figure 2 shows the stress–strain curve’s behavior under compression of neat PLA scaffolds and those of its 10 wt.% PLA/PD-BaSO₄ composite counterpart. As can be observed the 3D printed PLA and PLA/PD-BaSO₄ scaffolds do show different regimes and each regime corresponds to a specific mechanism of a porous structured material, in agreement with bibliography. In the first stage (region I), the walls contribute to the resistance of the scaffolds under compressive load, which results in an elastic response region at initial loads and corresponding strains. In the second stage (II), the pores collapse by buckling of the walls (barreling effect). The third stage (III) is featured by a large increase in stress over strain which may be explained by the fact that scaffolds are now compressed to a size that the scaffold becomes denser and furthermore more strain resistant to the applied load [31–33]. Despite the fact that both scaffolds have the same porosity (55%), in Figure 2 the neat PLA behaved like a more rigid structure [34], while PLA/PD-BaSO₄ shows again a flexible and softer behavior. Therefore, it can be concluded that the particles confer flexibility to the scaffolds in compression tests.
2.2. Biocompatibility Assessment

In vitro compatibility studies were performed to determine the possible toxicity of BaSO$_4$ and PD-BaSO$_4$ particles and their resulting PLA composites using Human dermal fibroblasts (HDFs) as a basic toxicity test. Figure 3 shows the metabolic activity of Human dermal fibroblasts (HDFs) in the presence of 10, 50, 100 or 500 µg/mL of BaSO$_4$ or PD-BaSO$_4$ particles. The metabolic activity displayed in this figure was normalized at each time-point with respect to the metabolic activity of HDFs seeded in the absence of particles, which was used as a control. The presence of BaSO$_4$ and PD-BaSO$_4$ particles slightly reduced the metabolic activity of HDFs with respect to the control. However, in all the cases, the metabolic activity was higher than 80%, demonstrating that HDFs were able to maintain a normal metabolic activity in the presence of particles (see also Supplementary Materials Figure S5). At day 1, the metabolic activity of HDFs seeded with 10, 50, 100 and 500 µg/mL of BaSO$_4$ particles was, 95, 94, 90 and 90%, respectively, relative to that of the control. In the case of HDFs seeded with 10, 50, 100 and 500 µg/mL of PD-BaSO$_4$ particles, the metabolic activity was 89, 86, 89 and 83% that of the control, respectively. At day 3, the metabolic activity of those cells seeded with 10 or 50 µg/mL of BaSO$_4$ particles, as well as 10 µg/mL of PD-BaSO$_4$ particles was not significantly different from the metabolic activity of the control. At this day, the metabolic activity in all the cases was higher than 90%, suggesting a negligible effect of the particles in the metabolic activity of HDFs. Additionally, cells observed under an inverted microscope showed normal morphology and the BaSO$_4$ and PD-BaSO$_4$ particles seemed to be internalized by cells and located around the nuclei (Supplementary Materials Figure S6).

**Figure 2.** Compression stress ($\sigma$)-strain ($\varepsilon$) curves of scaffolds for neat polylactide (PLA) and scaffolds for composite of polylactide and coated with polydopamine barium sulfate particles (PLA/PD-BaSO$_4$) 10 wt.%.
As can be seen in Figure 5, DNA content increased over culture time in all experimental and control conditions. Accordingly, significant differences were observed between experimental and control conditions. Moreover, significant differences were observed in the metabolic activity of HDFs with respect to the control, indicating normal metabolic activity of cells seeded on the composites developed in this work.

Figure 4 shows metabolic activity of cells seeded in PLA, PLA/BaSO_4, PLA/PD-BaSO_4 with 10 wt.% of filler composites and control tissue culture plastic (TCP). The metabolic activity was normalized at each time-point with the metabolic activity of HDFs seeded onto the TCP, which was used as a control. Except for cells seeded on PLA after 3 days of culture (see Table S1), no significant differences were observed in the metabolic activity of HDFs with respect to the control, indicating normal metabolic activity of cells seeded on the composites developed in this work.

Figure 3. Metabolic activity of Human dermal fibroblasts (HDFs) seeded in the presence of 10, 50, 100 or 500 μg/mL of barium sulfate particles (BaSO_4) or barium sulfate particles coated with polydopamine (PD-BaSO_4). Asterisks indicate significant differences (p < 0.05) with respect to the control.

Figure 4. Metabolic activity of Human dermal fibroblasts HDFs seeded on polylactide (PLA), composite of polylactide and barium sulfate particles (PLA/BaSO_4) and composite of polylactide and coated with polydopamine barium sulfate particles (PLA/PD-BaSO_4) with respect to the control at day 1, 3 and 7. Asterisks indicate significant differences (p < 0.05) with respect to the cells seeded on tissue culture plastic (TCP).

The proliferation of HDFs on PLA, PLA/BaSO_4 and PLA/PD-BaSO_4 composites was evaluated via DNA quantification. As can be seen in Figure 5, DNA content increased over culture time in all
experimental and control conditions. Accordingly, significant differences were observed between DNA content observed at day 7 and that observed at day 1 for all the samples studied. For example, the calculated proliferation rates between day 1 and day 7 were 1.6, 3.2 and 2.4 for PLA, PLA/BaSO$_4$ and PLA/PD-BaSO$_4$, respectively. The metabolic activity and proliferation results demonstrate that the materials employed in this work are not cytotoxic and can provide a cytocompatibility substrate for cells to attach and proliferate. A higher proliferation of HDFs was observed in PLA/BaSO$_4$ or PLA/PD-BaSO$_4$ composites with respect to pristine PLA samples. Is hypothesized that this higher proliferation rate may be associated to surface characteristic of the samples, such as roughness or hydrophilicity [35,36].

Concurrently, PD coating of BaSO$_4$ particles introduced binding sites for biologically active molecules such as proteins or drugs, increasing the use of this composite in second-generation devices for biomedical applications.

2.3. Adsorption/Release Test in 3D Printed Scaffolds

Following confirmation of cytocompatibility, the potential for antibiotic delivery with 3D printed scaffolds of PLA/PD-BaSO$_4$ was evaluated in vitro, with the aim of preventing an infection due to the insertion surgery (open wound) and the consequent rejection of the device [37]. To this end, levofloxacin was incorporated into the material via PD-BaSO$_4$ particle functionalization as a local drug delivery system for avoiding the oral administration common in this kind of surgeries. Levofloxacin is used for fighting and preventing osteomyelitis, since it is a fluoroquinolone with anti-staphylococcal activity in osteoarticular tissues [38].

Here, in order to compare the levofloxacin loading efficiency of the developed composites a polydopamine coated neat PLA sample (termed PD-PLA) is introduced. The release was performed at pH 5 to simulate the state of infection and at body temperature of 37 °C.

Figure 6 shows a non-detectable release of PLA, this is because the scaffolds have been previously washed, therefore there was a very low level of drug in PLA scaffolds, reflecting also that PLA does not adsorb the antibiotic and consequently does not release it either. Furthermore, in the case of PD-PLA a burst release was observed during the first 24 h where almost 80% of levofloxacin eluted. The remaining amount of drug (20%) was eluted in 4 days. This type of release profile is adequate in the context of an infection, since at the beginning a high rate of drug release is desirable followed by...
a slower drug release. When the PLA/PD-BaSO₄ was analyzed, a more moderate burst release was observed where almost 60% of levofloxacin eluted and later, during the next 3 days the 30% of drug was eluted while the last two days shows a very slow release with an 8% release (see Figure S6). In the end the PLA only releases 0.43 µg/mL due to the drug that is trapped in the holes. While PD-PLA was the one with the highest amount of drug released, 3.12 µg/mL, the PLA/PD-BaSO₄ composite drug released a 2.4 µg/mL.

![Figure 6](image)

**Figure 6.** Release profiles over time of levofloxacin in neat PLA, polydopamine coated neat PLA (PD-PLA) and PLA/PD-BaSO₄.

These release results in global show that PD-PLA and PLA/PD-BaSO₄ display potential antimicrobial properties. Comparing these two materials, the release profile of PLA/PD-BaSO₄ composites was more interesting as it generated a second elution stage with greater release, 40% versus 20% in PD-PLA. It is remarked finally that these scaffolds were immersed in trizma buffer (pH 10) prior to analysis so the results of Figure 6 correspond to bound levofloxacin after adsorption.

### 2.4. Antimicrobial Activity of 3D PLA/PD-BaSO₄ Scaffolds with Levofloxacin

To analyze the antibiotic efficacy of the 3D printed scaffolds the Agar Disk Diffusion tests against *Staphylococcus aureus* (*S. aureus*) were carried out. *S. aureus* was chosen because in bone infections is one of the most important pathogens due to its ability to adhere and form biofilms when in contact with tissue [38]. Figure 7 shows the Agar disk diffusion test corresponding to 3D printed scaffolds of PLA/PD-BaSO₄ with levofloxacin, PLA/PD-BaSO₄ scaffold as a negative control without levofloxacin and the disk of levofloxacin (5 µg) as a positive control. Cicuéndez et al. calculated that the minimum inhibitory concentration (MIC) of levofloxacin has a value of 0.06 µg/mL [39]. As observed in the release assay (Figure 6), these scaffolds release larger amounts of the drug.

Analyzing the Agar Disk Diffusion tests is observed that PLA/PD-BaSO₄ scaffolds with levofloxacin effectively inhibit bacterial growth, being the inhibition zone diameters of 38 ± 4 mm (Figure 7a), while the positive control (5 µg of Levofloxacin) exhibited a diameter average 28 ± 0 mm (Figure 7c). No inhibition zone could be observed for the sample without levofloxacin, PLA/PD-BaSO₄ (negative control, Figure 7b). It should be noted here that the scaffolds were washed after the drug adsorption, therefore the effective inhibition of bacterial growth is attributed to the amount of levofloxacin bound to polydopamine coating of the particles and not to an unspecified content of drug that could remain in the scaffold holds. This would still lead to a higher inhibition zone.
From Agar diffusion tests can be concluded that the polydopamine coating method of BaSO₄ particles used for drug tethering within the PLA composites is effective and facilitates the release of the drug inhibiting the *S. aureus* growth. The described method is simple and allows the addition of the antibiotic after a 3D printing process of scaffolds. This fact has two advantages since the drug is not processed with the scaffold avoiding either the contact with organic solvents nor high processing temperatures that could degrade the molecules, and, on the other hand, the adsorption of the medicament by a scaffold or implant can be done when clinically necessary, in situ.

3. Materials and Methods

3.1. Materials

Poly (D-lactide) homopolymer (100,000 g mol⁻¹) (PLA) was supplied by Purac-Corbion (Barcelona, Spain). Dopamine chloride, Barium sulfate (BaSO₄), phosphate buffer saline (PBS) and levofloxacin were purchased from Sigma-Aldrich (Madrid, Spain). Dulbecco’s modified Eagle’s medium (DMEM), fetal bovine serum (FBS), Hank’s balanced salt solution (HBSS) and penicillin-streptomycin (PS) solution Human dermal fibroblasts (HDFs) were purchased from Sigma-Aldrich (Madrid, Spain). Quant-IT® PicoGreen® dsDNA kit was from Invitrogen (Dublin, Ireland) and AlamarBlue® is from ThermoFisher Scientific (Dublin, Ireland).

3.2. Coating with Polydopamine and Blending

Coating of particles covered with polydopamine (PD) to obtain PD-BaSO₄ were obtained as in our previous article using basic pH 8.5 for 24 h followed by filtration and drying in a vacuum oven overnight [17]. The same coating method was used for the PD-PLA scaffolds. The size of the particles was measured by HORIBA Laser Scattering Particle Size Distribution Analyzer LA-350 (Horiba, Kyoto, Japan).

Polylactide samples were filled with 0.5, 1, 2, 5 and 10 wt.% of PD-BaSO₄. PD-BaSO₄ particles and neat polylactide (PLA) were melt mixed in a DSM Xplore micro-compounder (Xplore Instruments, Sittard, The Netherlands) at 200 °C and 150 rpm during 2 min and then conformed by injection molding. The mold was pre-heated at 45 °C and the injection temperature was 200 °C. Gel Permeation Chromatography (GPC) tests were carried out before and after blending process to check that the matrix is not degraded (see Figure S7). The weight average molecular mass (Mw) and dispersity (D) of PLA pellets before processing were respectively Mw = 183758 g mol⁻¹ and D = 1.8 whereas, after processing in a PLA/BaSO₄ composite, the values obtained for PLA were Mw = 174980 g mol⁻¹ and I = 1.8.
Scanning Electron Microscopy (SEM) and Transmission Electron Microscopy (TEM) were used to observe both the degree of dispersion of the particles inside of the polymer matrix and the break surfaces.

3.3. 3D Printing of Radiopaque Scaffolds

Scaffold fabrication was carried out in a 3D-Bioplotter from Envision TEC. Scaffolding 3D model was designed with Computer Aided Drawing (CAD, Autodesk, San Rafael, CA, USA). The CAD model was uploaded to the Bioplotter software (version 3.0.713.1406, Envision TEC, Gladbeck, Germany), which enables slicing of the model, before 3D-printing.

Two types of scaffolds were designed: one for mechanical properties and the other for the antibiotic release tests. Cylindrical geometries having 10 mm diameter and 10 mm height were used for measuring mechanical properties in compression mode, whereas square geometries (10 × 10 × 2 mm) (length × width × height) were used for adsorption/release and Agar Diffusion tests. In both cases the pore size of the scaffolds was 500 µm and it was used a plastic conic needle of 0.25 mm inner diameter for the printing layer dimension (Figure S8). The resulting file was uploaded to the software Visual Machines (version 2.8.126) that allows the user to input the various parameters that control the Bioprinter (Envision TEC).

Due to the specific dimensions of the feeding cartridge and the degradable nature of the polymer, the scaffolds are printed by solution. The materials (PLA and PLA/PD-BaSO₄) were dissolved with chloroform during 48 h, printed at 20 °C. The printing conditions are shown in Table 2. Reproducibility of the PLA and PLA/PD-BaSO₄ support frames was ensured by using the same CAD model for each frame, and by the high XYZ axis resolution of the Bioplotter (0.001 mm).

| Material          | Temperature (°C) | Pressure (Bar) | Speed (mm/s) | Post-Flow(s) | Pre-Flow(s) |
|-------------------|------------------|---------------|--------------|--------------|-------------|
| PLA               | 25               | 5.0           | 3.5          | 0.11         | 0.04        |
| PLA/PD-BaSO₄      | 25               | 4.4           | 4.1          | 0.11         | 0.01        |

3.4. Mechanical Properties

Dumbbell-shaped samples for tensile tests were punched out from sheets following ISO 527-2 (ISO 527-2/5A/5). In the case of 3D-printed scaffolds, the mechanical properties under compression were obtained following ISO 604. In both testing modes the tests were performed with an Instron 5565 testing machine at 23 °C and 50% of relative humidity (RH).

3.5. Adsorption/Release Test

Before starting the release test, the adsorption of levofloxacin in the scaffolds has been carried out. For that, the scaffolds were submerged in a buffer at pH 10 with levofloxacin (2 mg/mL) [40] for 24 h for facilitate the adsorption. Then, the scaffolds were washed with buffer at pH 10 and dried. The washing was done to eliminate the drug from the holes of the scaffolds, and in this way obtain the amount of drug that is absorbed by the material itself.

Levofloxacin release was evaluated using a UV-vis spectrophotometer (BMG Labtech FLUOstar Omega, Ortenberg, Germany). The band in λ = 287 nm for levofloxacin was employed to build a standard curve with known concentrations of the drug in PBS and the measurements were carried out at concentrations lower than 25 ppm. Release of levofloxacin from the scaffolds was tested in 3 mL of PBS at 37 °C, under mild agitation (220 rpm) at pH 5. Samples were taken at the following times: 30 min, 1 h, 2 h, 4 h, 6 h, 8 h, 12 h, 24 h, 48 h, 72 h, 96 h and 144 h. In all materials, three independent tests were carried out.
3.6. Agar Disk Diffusion

The diffusion disk agar tests for *Staphylococcus aureus* inoculum were performed under the Clinical and Laboratory Standards (CLSI) [40]. Briefly, few colonies of *S. aureus* were resuspended in Mueller Hinton Broth (Biokar Diagnostics, Pantin, France) and further diluted in order to achieve 0.5 McFarland units (1 × 10^8 CFU mL\(^{-1}\)) at 600 nm of wavelength, using a spectrophotometer (U-2000, Hitachi, Tokyo, Japan). The inoculum was swabbed on Mueller Hinton Agar (Biokar Diagnostics) plates and the scaffolds (one without drug as a negative control) were tested, as well as 5 µg of levofloxacin Agar (Biokar Diagnostics) plates and the scaffolds (one without drug as a negative control) were tested, as well as 5 µg of levofloxacin (positive control). Petri dishes were further incubated (Ultima, Revco, Thermo Scientific) at 37 °C for 24 h. Assays were performed in three independent experiments.

3.7. Cell Culture

3.7.1. Human Dermal Fibroblasts

HDFs were grown in T75 flasks using DMEM with 10% FBS and 1% PS. The cells were incubated at 37 °C in an atmosphere of 5% CO\(_2\). The culture medium was changed every 3 days. The cells were then harvested and sub-cultured when >90% confluence was observed.

3.7.2. Cell Seeding

To study the metabolic activity of HDFs seeded in the presence of BaSO\(_4\) or PD-BaSO\(_4\), HDFs were seeded at a density of 25,000 cells/cm\(^2\) in a 96 well tissue culture plate and incubated in 0.5 mL of DMEM with 10% FBS and 1% PS (37 °C, 5% CO\(_2\)). After 1 day in culture, the media was replaced by complete media containing 0, 10, 50, 100 or 500 µg/mL of BaSO\(_4\) or PD-BaSO\(_4\) particles that were previously autoclaved.

HDFs were also seeded on sterilized PLA, PLA/BaSO\(_4\) and PLA/PD-BaSO\(_4\) samples (7 mm in diameter) at a density of 25,000 cells/cm\(^2\) in a 48 well tissue culture plate and incubated in DMEM with 10% FBS and 1% PS. First, HDFs were suspended in 40 µL of culture medium, seeded onto each sample and incubated for 2 h to allow cell attachment (37 °C, 5% CO\(_2\), 95% relative humidity). When cells were attached, and additional 0.5 mL of culture medium was added into each well. The culture medium was replaced at day 3 after seeding.

3.7.3. Cell Viability Studies

AlamarBlue® assay was performed to quantify the metabolic activity of HDFs in the presence of BaSO\(_4\) or PD-BaSO\(_4\) particles or seeded on PLA, PLA/BaSO\(_4\) and PLA/PD-BaSO\(_4\). At the selected time points (1, 2 and 3 days for HDFs in the presence of BaSO\(_4\) or PD-BaSO\(_4\) particles and 1, 3 and 7 days for HDFs seeded on PLA, PLA/BaSO\(_4\) and PLA/PD-BaSO\(_4\)), the cells were washed with HBSS and subsequently incubated (6 h, 37 °C, sheltered from light) in 0.5 mL of fresh culture media with AlamarBlue® (10% v/v). Then, 100 µL of assay media was transferred to a 96 well plate, the absorbance at 550 and 595 nm was read on a microplate reader (Varioskan Flash, Thermo Fisher Scientific) and the percentage reduction of the dye was calculated.

To quantify the DNA amount of cells seeded on PLA, PLA/BaSO\(_4\) and PLA/PD-BaSO\(_4\) a PicoGreen® assay was then performed on the same samples used for AlamarBlue®. Cells were repeatedly frozen at ~80 °C and thawed to lyse the cells and release the entire DNA content. Finally, fluorescence was measured at 480 nm.

3.7.4. Statistics

Statistical differences were analyzed using one-way analysis of variance (ANOVA) and p-values of <0.05 were considered significant. Experiments were performed in triplicate and each assay was repeated three times.
4. Conclusions

In this work the mechanical brittleness associated to polylactide homopolymers is overcome via the incorporation of PD-BaSO$_4$ particles into a PLA matrix. The composite materials showed enhanced stiffness, strength, ductility and toughness. This is indeed relevant since in classical polymer/inorganic composites increases in stiffness and strength usually lead to dramatic decreases of ductility and toughness.

The particulate composites studied in this work are proved to be a valid substrate for cells to attach and proliferate. In addition, coating of barium sulfate particles with polydopamine provides functional groups that can act as anchorage points for incorporation of molecules with biological activity i.e., levofloxacin which is an antibacterial drug. The benefits of PD-BaSO$_4$ prove the potential of use of these PLA composites in bone reconstruction applications.

Supplementary Materials: Supplementary Materials can be found at http://www.mdpi.com/1422-0067/21/15/5480/s1. Figure S1. Tensile stress–strain curves of bulk materials. Figure S2. Particle size of BaSO$_4$ (grey) and PD-BaSO$_4$ (black). Figure S3. X-radiographies for neat PLA, PLA/BaSO$_4$ and PLA/PD-BaSO$_4$ composites with 2, 5 and 10 wt.% of BaSO$_4$ and PD-BaSO$_4$. Figure S4. PLA/PD-BaSO$_4$ (left) and neat PLA (right) 3D-printed cylindric scaffolds for compression tests. Figure S5. HDFs seeded in the presence of 500 µg/mL of BaSO$_4$. Figure S6. Release profiles over time of levofloxacin in neat PLA, PD-PLA and PLA/PD-BaSO$_4$. Figure S7.GPC refractive-index signals for PLA/PD-BaSO$_4$ before and after blending process at 200 °C. Figure S8. The microscopy image of PLA scaffold. (a) x10 and (b) x5. Table S1. Summary of the metabolic activity of the BaSO$_4$ and PD-BaSO$_4$ particles as well as the materials PLA, PLA/ BaSO$_4$ and PLA/PD-BaSO$_4$.

Author Contributions: E.Z., A.L. and J.-R.S. developed the idea and established the conceptual framework of the research. E.Z. guided and coordinated the research. N.S. and V.M. carried out the drug delivery and antimicrobial experiments designed by A.F.B. and I.A.C.R. A.L. and G.O.-A. designed and carried out the cell experiments. J.M.U. participated also in the discussion of results. N.S., A.L., E.Z., M.B. and J.-R.S. wrote and edited the manuscript. All authors have read and agreed to the published version of the manuscript.

Funding: The work has been financially supported by the Basque Government Department of Education, Linguistic Politics and Culture for a consolidated research group project IT-927-16, the Spanish Government’s MINECO MAT2016-78527-P (AEI/FEDER, UE) and EU’s COST TD1305 project. N.S. received a short-term scientific mission grant (ECOST-STSM-TD1305-38384). A.B. acknowledges the Science Foundation Ireland (16/BBSRC/3317). A.B. and I.R. acknowledge the Portuguese government, Fundação para a Ciência e Tecnologia (FCT) for project UID/DTP/04138/2019 (iMed.ULisboa). N.S. thanks POLYMAT for her PhD predoctoral grant, and A.L. the Basque Government for a postdoctoral fellowship.

Acknowledgments: The authors express thanks for technical and human support provided by SGIker of UPV/EHU and European funding: European Regional Development Fund (ERDF) and European Social Fund (ESF).

Conflicts of Interest: The authors declare no conflict of interest.

Abbreviations

| Acronym   | Abbreviation                        | Description                                                                 |
|-----------|-------------------------------------|-----------------------------------------------------------------------------|
| BMP-2     | Bone morphogenic protein 2          |                                                                              |
| PLA/BaSO$_4$ | Composite of Polylactide and barium sulfate particles |                                                                              |
| PLA/PD-BaSO$_4$ | Composite of polylactide and coated with polydopamine barium sulfate particles |                                                                              |
| CAD       | Computer Aided Drawing              |                                                                              |
| DNEM      | Dulbecco’s modified Eagle’s medium  |                                                                              |
| ε$_r$     | Elongation at break                 |                                                                              |
| Fe$_3$O$_4$ | Ferrous Oxide                       |                                                                              |
| PBS       | Fetal bovine serum                  |                                                                              |
| GPC       | Gel Permeation Chromatography       |                                                                              |
| HBSS      | Hank’s balanced salt solution       |                                                                              |
| HDFs      | Human dermal fibroblasts            |                                                                              |
| IGF-1     | Insulin growth factor 1             |                                                                              |
| MIC       | Minimum inhibitory concentration    |                                                                              |
| MWCNT     | Multi-walled carbon nanotubes       |                                                                              |
| PS        | Penicillin-streptomycin             |                                                                              |
| PBS       | Phosphate buffer saline             |                                                                              |
| PD-PLA    | Polylactide cover by polydopamine   |                                                                              |
PD Polydopamine
PU Polyurethanes
RO Radiopacity
RH Relative Humidity
SEM Scanning Electron Microscopy
S. aureus Staphylococcus aureus
E Tensile Modulus
σr Tensile strength
TT Tensile toughness
σy Tensile Yield
TCP Tissue culture plastic
TEM Transmission Electron Microscopy
wt.% Weight percent

References
1. Hamad, K.; Kaseem, M.; Yang, H.W.; Deri, F.; Ko, Y.G. Properties and medical applications of polylactic acid: A review. Express Polym. Lett. 2015, 9, 435–455. [CrossRef]
2. Tajbaksh, S.; Hajiali, F. A comprehensive study on the fabrication and properties of biocomposites of polylactic acid/ceramics for bone tissue engineering. Mater. Sci. Eng. C 2017, 70, 897–912. [CrossRef]
3. Martínez De Arenaza, I.; Sadaba, N.; Larrañaga, A.; Zuza, E.; Sarasua, J.R. High toughness biodegradable radiopaque composites based on polylactide and barium sulphate. Eur. Polym. J. 2015, 73, 88–93. [CrossRef]
4. Sarasua, J.R.; Arraiza, A.L.; Balderi, P.; Maiza, I. Crystallinity and mechanical properties of optically pure polylactides and their blends. Polym. Eng. Sci. 2005, 45, 745–753. [CrossRef]
5. Sadaba, N.; Martini, R.; Barthelat, F.; Martínez de Arenaza, I.; Larrañaga, A.; Sarasua, J.R.; Zuza, E. Understanding the toughness mechanism prompted by submicron rigid particles in polylactide/barium sulfate composites. Polym. Test. 2018, 69, 340–349. [CrossRef]
6. Grémare, A.; Guduric, V.; Bareille, R.; Heroguez, V.; Latour, S.; L’heureux, N.; Fricain, J.C.; Catros, S.; Le Nihouannen, D. Characterization of printed PLA scaffolds for bone tissue engineering. J. Biomed. Mater. Res. Part A 2018, 106, 887–894. [CrossRef] [PubMed]
7. Nuutinen, J.-P.; Clerc, C.; Törmälä, P. Mechanical properties and in vitro degradation of self-reinforced radiopaque bioreinforced poly lactide fibres. J. Biomater. Sci. Polym. Ed. 2003, 14, 665–676. [CrossRef] [PubMed]
8. Chang, W.-J.; Pan, Y.-H.; Tzeng, J.-J.; Wu, T.-L.; Fong, T.-H.; Feng, S.-W.; Huang, H.-M. Development and Testing of X-ray Imaging-Enhanced Poly-L-Lactide Bone Screws. PLoS ONE 2015, 10, e0140354. [CrossRef]
9. Noor Azman, N.Z.; Musa, N.F.L.; Nik Ab Razak, N.N.A.; Ramli, R.M.; Mustafa, I.S.; Abdul Rahman, A.; Yahaya, N.Z. Effect of Bi2O3 particle sizes and addition of starch into Bi2O3–PVA composites for X-ray shielding. Appl. Phys. A 2016, 122, 818. [CrossRef]
10. Abnahel, B.M.; Mustafa, I.S.; Noor Azman, N.Z. Characteristics of X-ray attenuation in nano-sized bismuth oxide/epoxy-polyvinyl alcohol (PVA) matrix composites. Appl. Phys. A Mater. Sci. Process. 2018, 124, 1–7. [CrossRef]
11. Zuza, E.; Meaurio, E.; Sarasua, J.-R. Biodegradable Polylactide-Based Composites. In Composites from Renewable and Sustainable Materials; InTech: Rijeka, Croatia, 2016.
12. Xiu, H.; Qi, X.; Liu, Z.; Zhou, Y.; Bai, H.; Zhang, Q.; Fu, Q. Simultaneously reinforcing and toughening of polylactide/carbon fiber composites via adding small amount of soft poly(ether)urethane. Compos. Sci. Technol. 2016, 127, 54–61. [CrossRef]
13. Wang, X.; Jiang, M.; Zhou, Z.; Gou, J.; Hui, D. 3D printing of polymer matrix composites: A review and prospective. Compos. Part B Eng. 2017, 110, 442–458. [CrossRef]
14. Parandoush, P.; Lin, D. A review on additive manufacturing of polymer-fiber composites. Compos. Struct. 2017, 182, 36–53. [CrossRef]
15. Poh, P.S.P.; Chhaya, M.P.; Wunner, F.M.; De-Juan-Pardo, E.M.; Schilling, A.F.; Schantz, J.-T.; van Grienven, M.; Hutmacher, D.W. Polylactides in additive biomanufacturing. Adv. Drug Deliv. Rev. 2016, 107, 228–246. [CrossRef] [PubMed]
16. Liu, Y.; Ai, K.; Lu, L. Polydopamine and Its Derivative Materials: Synthesis and Promising Applications in Energy, Environmental, and Biomedical Fields. *Chem. Rev.* 2014, 114, 5057–5115. [CrossRef]

17. Larrañaga, A.; Ramos, D.; Amestoy, H.; Zuza, E.; Sarasua, J.-R. Coating of bioactive glass particles with mussel-inspired polydopamine as a strategy to improve the thermal stability of poly(l-lactide)/bioactive glass composites. *RSC Adv.* 2015, 5, 65618–65626. [CrossRef]

18. Ahn, B.K. Perspectives on Mussel-Inspired Wet Adhesion. *J. Am. Chem. Soc.* 2017, 139, 10166–10171. [CrossRef]

19. Zhuang, H.; Su, H.; Bi, X.; Bai, Y.; Chen, L.; Ge, D.; Shi, W.; Sun, Y. Polydopamine Nanocapsule: A Theranostic Agent for Photoacoustic Imaging and Chemo-Photothermal Synergistic Therapy. *ACS Biomater. Sci. Eng.* 2017, 3, 1799–1808. [CrossRef]

20. Yu, X.; Tang, X.; He, J.; Yi, X.; Xu, G.; Tian, L.; Zhou, R.; Zhang, C.; Yang, K. Polydopamine Nanoparticle as a Multifunctional Nanocarrier for Combined Radiophotodynamic Therapy of Cancer. *Part. Part. Syst. Charact.* 2017, 34, 1600296. [CrossRef]

21. Zhang, J.; Li, J.; Jia, G.; Jiang, Y.; Liu, Q.; Yang, X.; Pan, S. Improving osteogenesis of PLGA/HA porous scaffolds based on dual delivery of BMP-2 and IGF-1 via a polydopamine coating. *RSC Adv.* 2017, 7, 56732–56742. [CrossRef]

22. Khoshnood, N.; Zamanian, A.; Massoudi, A. Mussel-inspired surface modification of titania nanotubes as a novel drug delivery system. *Mater. Sci. Eng. C* 2017, 77, 748–754. [CrossRef] [PubMed]

23. Fernandez-Yague, M.A.; Larrañaga, A.; Gladkovskaya, O.; Stanley, A.; Tadayyon, G.; Guo, Y.; Sarasua, J.R.; Tofail, S.A.M.; Zeugolis, D.I.; Pandit, A.; et al. Effects of Polydopamine Functionalization on Boron Nitride Nanotube Dispersion and Cytocompatibility. *Biomacromolecules* 2015, 26, 2025–2037. [CrossRef] [PubMed]

24. Mu, C.; Zhang, L.; Song, Y.; Chen, X.; Liu, M.; Wang, F.; Hu, X. Modification of carbon nanotubes by a novel biomimetic approach towards the enhancement of the mechanical properties of polyurethane. *Polymer* 2016, 92, 231–238. [CrossRef]

25. Pekkan, G.; Aktas, A.; Pekkan, K. Comparative radiopacity of bone graft materials. *J. Cranio-Maxillofac. Surg.* 2012, 40, e1–e4. [CrossRef]

26. Clauss, M.; Trampuz, A.; Borens, O.; Bohner, M.; Ichmann, T. Biofilm formation on bone grafts and bone graft substitutes: Comparison of different materials by a standard in vitro test and microcalorimetry. *Acta Biomater.* 2010, 6, 3791–3797. [CrossRef]

27. Zuiderduin, W.C.J.; Westzaan, C.; Huëtink, J.; Gaymans, R.J. Toughening of polypropylene with calcium carbonate particles. *Polymer* 2002, 44, 261–275. [CrossRef]

28. Kim, G.M.; Michler, G.H. Micromechanical deformation processes in toughened and particle filled semicrystalline polymers: Part 2. model representation for micromechanical deformation processes. *Polymer* 1998, 39, 5699–5703. [CrossRef]

29. Subramanian, A.S.; Tey, J.N.; Zhang, L.; Ng, B.H.; Roy, S.; Wei, J.; Hu, X. ‘Matthew’ Synergistic bond strengthening in epoxy adhesives using polydopamine/MWCNT hybrids. *Polymer* 2016, 82, 285–294. [CrossRef]

30. Rodrigues, N.; Benning, M.; Ferreira, A.M.; Dixon, L.; Dalgarno, K. Manufacture and Characterisation of Porous PLA Scaffolds. *Procedia CIRP* 2016, 49, 33–38. [CrossRef]

31. Ghassemieh, E. Morphology and compression behaviour of biodegradable scaffolds produced by the sintering process. *Proc. Inst. Mech. Eng. Part H J. Eng. Med.* 2008, 222, 1247–1262. [CrossRef]

32. Deplaine, H.; Acosta-Santamaria, V.A.; Vidaurre, A.; Gómez Ribelles, J.L.; Doblaré, M.; Ochoa, I.; Gallego Ferrer, G. Evolution of the properties of a poly(l-lactic acid) scaffold with double porosity during in vitro degradation in a phosphate-buffered saline solution. *J. Appl. Polym. Sci.* 2014, 131, 1–10. [CrossRef]

33. Khodaei, M.; Amini, K.; Valanezhad, A. Fabrication and Characterization of Poly Lactic Acid Scaffolds by Fused Deposition Modeling for Bone Tissue Engineering. *J. Wuhan Univ. Technol. Mater. Sci. Ed.* 2020, 35, 248–251. [CrossRef]

34. Chung, T.-W.; Wang, Y.-Z.; Huang, Y.-Y.; Pan, C.-I.; Wang, S.-S. Poly (ε-caprolactone) Grafted With Nano-structured Chitosan Enhances Growth of Human Dermal Fibroblasts. *Artif. Organs* 2006, 30, 35–41. [CrossRef]

35. Navarro, M.; Engel, E.; Planell, J.A.; Amaral, I.; Barbosa, M.; Ginebra, M.P. Surface characterization and cell response of a PLA/CoP glass biodegradable composite material. *J. Biomed. Mater. Res. Part A* 2008, 85, 477–486. [CrossRef]
36. Meaurio, E.; Sanchez-Rexach, E.; Butron, A.; Sarasua, J.R. The conformation of chloramphenicol in the ordered and disordered phases. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc.* 2019, 211, 383–392. [CrossRef]

37. Ferreira, M.; Rzhepishevska, O.; Grenho, L.; Malheiros, D.; Gonçalves, L.; Almeida, A.J.; Jordão, L.; Ribeiro, I.A.; Ramstedt, M.; Gomes, P.; et al. Levofloxacin-loaded bone cement delivery system: Highly effective against intracellular bacteria and *Staphylococcus aureus* biofilms. *Int. J. Pharm.* 2017, 532, 241–248. [CrossRef]

38. Cicuendez, M.; Doadrio, J.C.; Hernández, A.; Portolés, M.T.; Izquierdo-Barba, I.; Vallet-Regí, M. Multifunctional pH sensitive 3D scaffolds for treatment and prevention of bone infection. *Acta Biomater.* 2018, 65, 450–461. [CrossRef]

39. Martin, V.; Ribeiro, I.A.C.; Alves, M.M.; Gonçalves, L.; Almeida, A.J.; Grenho, L.; Fernandes, M.H.; Santos, C.F.; Gomes, P.S.; Bettencourt, A.F. Understanding intracellular trafficking and anti-inflammatory effects of minocycline chitosan-nanoparticles in human gingival fibroblasts for periodontal disease treatment. *Int. J. Pharm.* 2019, 572, 118821. [CrossRef]

40. Clinical and Laboratory Standards Institute. *M02-A12 Performance Standards for Antimicrobial Disk Susceptibility Tests; Approved Standard-Twelfth Edition*; Clinical and Laboratory Standards Institute: Wayne, PA, USA, 2015.

© 2020 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).