Scleral ossicles: angiogenic scaffolds, a novel biomaterial for regenerative medicine applications†

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Given the current prolonged life expectancy, various pathologies affect increasingly the aging subjects. Regarding the musculoskeletal apparatus, bone fragility induces more susceptibility to fractures, often not accompanied by good ability of self-repairing, in particular when critical-size defects (CSD) occur. Currently orthopedic surgery makes use of allografting and autografting which, however, have limitations due to the scarce amount of tissue that can be taken from the donor, the possibility of disease transmission and donor site morbidity. The need to develop new solutions has pushed the field of tissue engineering (TE) research to study new scaffolds to be functionalized in order to obtain constructs capable of promoting tissue regeneration and achieve stable bone recovery over time. This investigation focuses on the most important aspect related to bone tissue regeneration: the angiogenic properties of the scaffold to be used. As an innovative solution, scleral ossicles (SOs), previously characterized as natural, biocompatible and spontaneously decellularized scaffolds used for bone repair, were tested for angiogenic potential and biocompatibility. To reach this purpose, in ovo Chorioallantoic Membrane Assay (CAM) was firstly used to test the angiogenic potential; secondly, in vivo subcutaneous implantation of SOs (in a rat model) was performed in order to assess the biocompatibility and the inflammatory response. Finally, thanks to the analysis of mass spectrometry (LCMSQE), the putative proteins responsible for the SO angiogenic properties were identified. Thus, a novel natural biomaterial is proposed, which is (i) able to induce an angiogenic response in vivo by subcutaneous implantation in a non-immunodefl exhibits good ability of self-repairing, in particular when critical-size defects (CSD) since the bone lesion is so severe that it prevents the skeletal segment from self-repairing. Recovery of significant skeletal defects could be partially abortive due to the perturbations that affect the regenerative process. Nowadays, surgical techniques include: bone grafting and joint arthroplasties, autografts (which are considered the gold standard in the field), and allografts (that are often used to treat such bone defects). However, although all these strategies are commonly used in orthopedic surgery, these treatments have some limitations concerning their costs and their side effects such as potential infections, non-unions, donor site morbidity,

1. Introduction

Given the current prolonged life expectancy and the evermore aging world population, there is a rapid increase in musculoskeletal pathologies such as bone fragility inducing more susceptibility to fractures, often not accompanied by good ability of self-repairing and hence subjected to some complications like non-unions, as well as vertebral collapses, osteomyelitis, etc. As a result, costs and bone-related medical treatments increase constantly. Also, as a consequence of injuries, it is possible to incur large-sized fractures called critical-size defects (CSD) since the bone lesion is so severe that it prevents the skeletal segment from self-repairing. Recovery of significant skeletal defects could be partially abortive due to the perturbations that affect the regenerative process. Nowadays, surgical techniques include: bone grafting and joint arthroplasties, autografts (which are considered the gold standard in the field) and allografts (that are often used to treat such bone defects). However, although all these strategies are commonly used in orthopedic surgery, these treatments have some limitations concerning their costs and their side effects such as potential infections, non-unions, donor site morbidity,
difficulties in finding adequate donors and a limited amount of tissue to harvest. Moreover, severe fractures and replacements of injured skeletal segments are currently treated by means of metal implants that, instead, show some limitations such as post-transplant rejection and reduced bioactivity.\textsuperscript{10–12} Furthermore, implant mobility, inflammation and bone resorption are associated with failure of some metal implants. In many cases the biological system is not able to replace the bone loss that may be also compensated by the surrounding soft-tissue environment. Furthermore, the destruction of the local vascular network often occurs at the fracture site and for the bone healing process to be assured it is required to restore the vascularization previously. It is to be noted, in fact, that an essential prerequisite of fracture healing is the proper development of blood vessels and the complex interplay among blood vessel formation, oxygen supply, growth factors and cell proliferation influences the final healing outcome in large bone defects.\textsuperscript{13–17}

One of the most important aspects to take in consideration during the setting up of a new material for bone regeneration is its angiogenic potential.\textsuperscript{18} Angiogenesis is a complex process in which, during development, endothelial cells (ECs) switch from the “quiescent” to the “angiogenic phenotype” in response to the balance between anti- and pro-angiogenic factors.\textsuperscript{19} To do this, EC growth, migration, and tube formation are processes that occur under precise stimuli which regulate the cell–extracellular matrix (ECM) interactions in order to form new blood capillaries from the pre-existing ones.\textsuperscript{20,21}

Angiogenesis and osteogenesis are tightly coupled: the angiogenesis process is essential and crucial for osteogenesis and bone regeneration, and osteogenesis is critical for the maintenance of a healthy and fully functional skeletal system.\textsuperscript{22} The blood vessels mediate the transport of circulating cells, oxygen, nutrients and waste products and provide angiogenic and angiocrine signals. Therefore, normal vascular formation plays a key role in both physiological and pathological processes of the skeletal system. The crucial role of the vessels has been already established during the various phases of bone osteogenesis, where two types of bone formation (\textit{i.e.} static and dynamic) have been demonstrated to occur in close relationships with the vascular frameworks.\textsuperscript{23–26} In this sophisticated and finely modulated process, a variety of angiogenic factors are all widely expressed as primary inducers of vascular development.

In this regard in the last decade, regenerative medicine and TE have played a central role, proposing strategies for bone reconstitution with the ambition to overcome the complications associated with traditional techniques. In particular, many different scaffolds are developing with different properties, proposing new materials (\textit{i.e.} 3D printable matrices/scaffolds) to be used as new strategies in order to trigger the onset of bone regeneration and optimize the cell growth.\textsuperscript{27–32} A variety of different exogenous chemical or physical stimuli were tested, such as soluble growth/differentiation factors, as well as the application of mechanical forces (\textit{i.e.}, loading).\textsuperscript{33–40} Finally, many types of cells have been used alone or in co-culture.\textsuperscript{41–48} An engineered scaffold-based strategy is the starting point of the recent innovative therapies, for which the choice of material properties, manufacturing methods and component treatments, is crucial. They are followed by preclinical \textit{in vitro} studies and \textit{in vivo} experimentation, which can be a prelude to a production of devices/constructs that meet the need/expectations of patients. Many of these materials display good mechanical properties but to the detriment of retaining a high porosity, so, scaffolds for TE which have demonstrated good outcomes \textit{in vitro} have failed when implanted \textit{in vivo} because of their insufficient capability of vascularization not permitting the cell infiltration and hence a proper vascularization.\textsuperscript{49}

In this work it is proposed that the use of scleral ossicles (SOs) is an innovative solution.\textsuperscript{1} SOs are peculiar bony plates forming a ring at the scleral–corneal border of the eyeball of lower vertebrates. This proposed model is interesting because once SOs reach the definitive size, in the adult animal, they are subjected to mechanical stereotyped stress for their lifetime, solely with a protective function. Therefore, the bone remodeling processes (in response to metabolic need) must be avoided and, to do this, the osteocytes undergo massive apoptosis, making the ossicles like naturally decellularized bones.\textsuperscript{50–53} SOs have been suggested as natural biocompatible scaffolds for bone repair, inasmuch it has been reported that SOs are able to sustain cell adhesion, proliferation and differentiation and, moreover, they seem to have angiogenic potential \textit{in vitro}.\textsuperscript{1} Definitely, the SOs act as a natural ECM-like platform to support angiogenesis and vascularization.

The aims of this study are: (I) to evaluate the angiogenic potential of SOs \textit{in ovo} and \textit{in vivo}, (II) to verify the absence of inflammatory potential upon \textit{in vivo} SO subcutaneous implantation and finally (III) to investigate the factors responsible for the angiogenesis.

2. Experimental section

2.1. Extraction and preparation of the scleral ossicles (SOs)

Poultry processing waste, donated by a local butcher, was used as the source of SOs. Each eyeball of adult chickens (50–70 days old) was incised with a scalpel in order to extract the scleral ring. Under a stereomicroscope, the tissue membranes that cover the scleral ring were removed and each of the 13 SOs were disjoined. All bony plates were cleaned with PBS pH 7.4 (phosphate buffered saline) to eliminate the residues of eye liquid humor, dried and finally sterilized by exposure for 30 min under UV radiation.

2.2. Chorioallantoic membrane (CAM) assay

Fertilized White Leghorn chicken eggs (Istituto Zooprofilattico di Puglia e Basilicata, Foggia, Italy) staged according to Hamburger and Hamilton (HH)\textsuperscript{56} were placed, at the onset, into an incubator and kept under humidity at 37 °C (day 0). At stage HH3 (day 3), a square window was opened into the egg-
were anesthetized with 4% isoflurane and maintained at 2% 
twelve rats, sterile SO was placed subcutaneously. The rats 
after, one control rat was sacrificed (day 0). In the remaining 
cycle and the temperature was controlled.

All animals were kept under a 12 h light/dark
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2016030408537165 v8). Animal housing and caretaking were
by the French Ministry for Education, Research, and
Animals, and approved by the Bordeaux Ethical Committee
the European Guidelines for Care and Use of Laboratory
Facilities of the University of Bordeaux, France.

The animal experiments were conducted at the Animal 
subcutaneous implantation of scleral ossicles

Thirteen 12-week-old male Wistar rats weighing 350–400 g 
were purchased from Janvier Labs (Saint Berthevin, France).
The animal experiments were conducted at the Animal 
Facilities of the University of Bordeaux, France.

All animal procedures were performed in accordance with 
the European Guidelines for Care and Use of Laboratory 
Animals, and approved by the Bordeaux Ethical Committee 
and by the French Ministry for Education, Research, and 
Innovation (agreement number APAFIS #4375-
2016030408537165 v8). Animal housing and caretaking were 
provided by the animal facility in accordance with the national 
guidelines. All animals were kept under a 12 h–12 h light/dark 
cycle and the temperature was controlled.

After one week of adaptation, rats were shaved and the day 
after, one control rat was sacrificed (day 0). In the remaining 
twelve rats, sterile SO was placed subcutaneously. The rats 
were anesthetized with 4% isoflurane and maintained at 2%

using a mask during the surgery. After scrubbing the skin with 
0.2% sterile chlorhexidine, 2 cm dorsal incisions were made 
bilaterally on the sacral–lumbar area. Subcutaneous pockets 
(4–5 cm in length and 1.5 cm in width) were made bilaterally, 
parallel to the rostro-caudal axis in the direction of the head, 
and one SO per side was inserted (Fig. 2A). All the incisions 
were sutured with 3–4 sterile staples and the animals were fol-
lowed throughout the post-implantation period up to euthana-
sia. After 1, 2, 4, and 12 weeks post-implantation, animals 
were euthanized using CO₂ (TEM SEGA rodent system, 
Automate d’euthanasie v 4.5). The skin areas containing the 
SO were excised from each rat (implant areas) as well as skin 
control samples (control areas) from the dorsal thoracic region 
(Fig. 2B).

2.4. Histological staining and immunohistochemistry

Samples with/without SOs were decalcified for 4 days in 10% 
EDTA and then processed for paraffin embedding (Leica 
EG1150 H) following the procedure here described: 70% 
ethanol for 30 min, three times in 95% ethanol, three times in 
100% ethanol, three times in toluene and two times in 
paraffin, 1 h for each. After paraffin embedding, twelve sec-
tions (5 µm) were obtained from each sample.

Slides were deparaffinized by means of xylene (two times 
for 5 min each), hydrated through a graded series of alcohol 
(100%, 95%, 80% and 70%) for 5 min each and rinsed first 
with tap water and then with deionized water for 1 min each. 
After that, four slides (with two sections each) of both control 
(CTR) and implanted (IMPLANTED) samples, for each time 
point (weeks 1, 2, 4 and 12), were stained both with 
Hematoxylin & Eosin (H&E) or Masson’s Trichrome (MT).

Briefly, for H&E staining, after the dehydration procedure, 
the slides were stained with Carazzi hematoxylin solution 
(DiaPath C0203), rinsed with tap water, stained with alcoholic 
0.5% eosin Y (DiaPath C0353) and then rinsed with deionized 
water for 1 min each.

For Masson’s trichrome staining, dehydrated slides were 
stained with Mayer’s hemalum solution for 10 min, rinsed 
first with tap water for 10 min and then with deionized water 
for 1 min, before being exposed to 1% Fuchsin-Ponceau (1v/2v) 
for 1.5 min. After 1 min in 3% (w/v) phosphomolybdic acid, 
slides were run through acidified water (0.1% (v/v) glacial
acetic acid) for 1 min, stained with 1% (w/v) light green for 1 min and rinsed with acidified water again for 1 min.

Immunohistochemistry (IHC) was performed using the automatic system Ventana Bench Mark XT according to the following protocol: samples were incubated for 15 min with buffer EZ prep (Ventana 950-102) at 72 °C; the antigen retrieval was performed by arranging slides in a Cell Conditioning 1 Buffer (Ventana 950-124) for 36 min; slides were incubated for 24 min with macrophage specific anti-CD68 clone PG-M1 (DakoCytomation M0876) at a concentration of 1:300 in an antibody diluent (Ventana 251-018); samples were then treated with an ultraView Universal DAB Detection Kit (Ventana 760-500) and finally, slides were counterstained by immersion in hematoxylin II (Ventana 790-2208) for 12 min and mounted on slides. Stained sections were evaluated under a Nikon Eclipse microscope (Nikon) equipped with a DS-Fi2 camera (Nikon) and processed by means of NIS-Elements D 5.11.00 software (Nikon).

2.5. Mass spectrometry, protein identification and annotation

Three replicates were prepared for mass spectrometry analysis according to the following protocol: four SOs were finely chopped and incubated with 50 µl of 50 ng µl⁻¹ trypsin (PIERCE 900578) at 37 °C for 16 h. Next, the trypsin was inactivated with 5% formic acid (with a volume equal to 10% of the final digestion volume) and subsequently dried in the Speed-vac for 1 h. Before performing the LCMSQE analysis, the samples were re-suspended in 40 µL of a mixture of water : acetonitrile : formic acid (97 : 3 : 2), sonicated for 20 min and centrifuged at 12 100 r.c.f. for 10 min. Analyses were performed on an ESI Q Exactive Mass spectrometer (Thermo Scientific), controlled by an Xcalibur (v. 29 build 2926) and interfaced with an Ultimate 3000 UHPLC pump. The column (Zorbax SB-C18 RRHT, 2.1 × 50 mm, 1.8 µ particle size, Agilent Technologies), thermostated at 30 °C, was equilibrated with 0.3 ml min⁻¹ of water and 0.1% formic acid (A) with 2% acetonitrile (B). Two different methods were used: 80 min for samples 1 and 2, and 3 h for sample 3. Briefly, in the first method after sample injection (20 µl), B% was kept at 2% for 2 min, then it was increased from 2 to 3% in 5 min, then linearly increased from 3 to 28% in 59 min; B% was then brought to 90% in 4 min and kept at 90% for 3 min, before the reconditioning step. The total runtime was 80 min. For the second method after sample injection (30 µl), B% was kept at 2% for 2 min, then it was linearly increased from 2 to 28% in 153 min; B% was then brought to 90% in 10 min and kept at 90% for 5 min, before the reconditioning step. The total run time was 3 h.

ESI (electrospray ionization) was performed in positive mode; the probe was heated at 290 °C, the capillary temperature was set at 270 °C; the following nitrogen flows (arbitrary units) were used to assist the ionization: Sheath Gas 40, Aux Gas 30, and Sweep Gas 3; capillary voltage was set to 3.5 kV and the S-Lens RF level was set at 55 (arbitrary units).

Centroided MS and MS² spectra were recorded from 200 to 2000 m/z in FULL MS/dd-MS² (TOPS) mode, at a resolution of 70 000 and 17 500, respectively. The five most intense multi-charged ions were selected for MS² nitrogen-promoted collision-induced dissociation (NCE = 28). A precursor active exclusion of 10 seconds was set; peptide-like isotope pattern ions were preferred. The mass spectrometer was calibrated before the start of the analyses.

MS raw data files were converted with MSConvert (version 3.0.19135-311725741 ProteoWizard) into mascot generic format and the peak lists were searched against Trembl (21902 sequences for Gallus gallus) for peptide sequences and C-Rap for contaminants with MASCOT (version 2.4, Matrix Science, London, UK).

Trypsin was set as a proteolytic enzyme, and deamidated-NQ and oxidized-M were set as variable modifications. One missed cleavage was allowed. Mass tolerances were set at 10 ppm for the precursor ions and 0.01 Da for the product ions. Automatic decoy database search was used to estimate the false discovery rate, which was adjusted to <1%.

All proteins identified from each experimental approach are listed in Table 1, ESI† GO annotation was used to identify protein clusters, this was obtained using Protein Analysis Through Evolutionary Relationships (PANTHER version 14.1) (released 2019-03-12) contains 15 524 protein families, divided into 107 627 functionally distinct protein subfamilies), STRING database (version 11.0 since January 19, 2019 currently covers 9 643 763 proteins from 2031 organisms) and the DAVID Bioinformatics Resources database (version 6.8).

2.6. Statistical analysis

Experiments carried out in triplicate (as indicated in the Experimental section and in the respective paragraphs in the “Results and discussion” section) are defined respectively as two or three independent samples that were treated, isolated and analysed separately. All data are expressed as mean ± Standard Deviation (SD). Differences between the experimental conditions were evaluated by one-way analysis of variance (ANOVA) with Dunnett’s corrections for multiple comparison or two-way analysis of variance (ANOVA) with Bonferroni corrections for multiple comparison; the type of analysis and significant p values are indicated in the figure legends. Statistical analyses and relative graphs were performed using GraphPad prism 6 (version 6.01).

Differences are classified as significant with p < 0.05.

3. Results and discussion

3.1. In ovo angiogenic potential of scleral ossicles

In order to evaluate the angiogenic potential of the SOs in ovo, a peculiar analysis was conducted on the chick embryo chorioallantoic membrane (CAM). The CAM is an extra embryonic membrane which serves as a gas exchange surface and its function is supported by a dense capillary network. Some side considerations concern the ethical choice of the CAM
model: the chick embryo that has not reached the 14th day of its gestation period does not feel pain and can therefore be used for experimentation without any ethical restrictions or prior protocol approval.\textsuperscript{62,63} Additionally, according to the “three Rs” rule, we have used a reduced number of animals for the successive \textit{in vivo} experimentation ("reduction"), we minimized the pain and suffering from animals ("refinement") and we have used an alternative (\textit{in ovo} instead of \textit{in vivo}) for the first experimental phase of the study ("replacement").\textsuperscript{64,65} Because of its easy accessibility, the chicken CAM (frequently used as an experimental model, also as an alternative to the animal model, as just mentioned) has been used, in the present study, for the investigation of the angiogenic potential of SO.\textsuperscript{66–68} Moreover, in the development stages used in this experimentation (from day 8 to day 12) the immunocompetent chicken system is not fully developed and for this reason it is not responsive to an exogenous material.\textsuperscript{69}

The angiogenic potential of the SO was evaluated by the CAM assay, incubating firstly SO and subsequently media conditioned by SOs. Experiments were started at the stage HH8 of embryo development (when crucial steps as the development of the circulatory system occur) and carried out for four days, as described in the Experimental section.

In a preliminary experiment, SO was brought directly into contact with the growing CAM resulting in a strong vasculo-proliferative activity. Fig. 3 shows the appearance of CAM’s vessels at stage HH12; the magnification of the dashed squared area shows a linear and orderly course of the vessels (Fig. 3A). The SO implant (removed because of its opacity and outlined with a dashed yellow line in Fig. 3B) induces a strong vasculo-proliferative reaction resulting in the formation of neo-formed vessels (yellow arrows) on the pre-existing ones. As an interesting aspect, it should be noted that the vascular course is extremely tortuous and irregular, representing the formation of new capillaries during tumour onset.

![Fig. 3 Appearance of CAM at stage HH12 with (B) or without SO (A). The yellow dashed line indicates the site of implantation of SO on the growing CAM (scale bar = 2 mm).](Image)

### Table 1 Composition of the conditioned media used in the angiogenic CAM assay.

| Media composition | CTR | SO | GF |
|-------------------|-----|----|----|
| αMEM + gluta + P/S | +   | –  | –  |
| #1                | +   | +  | –  |
| #2                | +   | +  | +  |
| #3                | +   | +  | +  |
| #4                | +   | –  | +  |
| #5                | +   | –  | +  |

To better understand whether this vasculo-proliferative reaction is induced by factors released by the SO, six different conditioned media, obtained by culturing for 48 h a combination of serum free medium, SO and growth factors, were tested through the CAM assay (medium compositions are specified in Table 1).

Both the medium conditioned by the presence of the SO-released-factors alone (#1) and the medium containing the combination of the two (#2 and #3) or the growth factors alone (#4 and #5) induce the formation of numerous allantoic vessels that develop radially with respect to the implant in a “spoked-wheel” pattern (white arrows in Fig. 4A). These observations are supported by the quantification of the number of vessels that moved away from the areas of medium inoculation under all conditions. The histogram in Fig. 4B is related to the quantification performed on the images of the CAM vessels at stage HH12 under all the conditions analyzed.

In particular, the number of vessels is significantly higher under conditions #1 to #5 compared to the CTR (****p < 0.0001). It has been highlighted that the SO alone is able to induce an angiogenic response not so different (*p < 0.5) from conditions #4 and #5 (representing our positive controls). It is of note that the VEGF (#2) is able to potentiate the SO angiogenic response (1#) (**p < 0.01) suggesting an additive effect that is not observable with the FGF (#3). These findings suggest that the SO induces an important angiogenic reaction triggered by angiogenic factors released in the media where it is located. Furthermore, the VEGF acts as an enhancer of the SO angiogenic response while the FGF in combination with the SO effect does not affect the angiogenic potential of the SOs.\textsuperscript{70–72} The last evidence probably means that the FGF angiogenic response is not potentiated because of a “saturation” effect (\textit{viz.} every CAM, despite the sum of favorable conditions, is not able to develop vessels beyond a certain number\textsuperscript{73}). As the final consideration, concerning the data so far obtained on this aspect, it is correct to assert that the SOs have a good angiogenic potential \textit{in vivo}.

#### 3.2. \textit{In vivo} subcutaneous implantation of scleral ossicles

Based on the previous CAM results, indicating a strong angiogenic potential of SOs, it was decided to test the biocompatibility and inflammation response in a rat model through the
subcutaneous implantation in the dorsal region for 1, 2, 4 and 12 weeks.

No loss of animals was observed throughout the procedure, not even after surgery. All rats had good recovery and no animals showed pain after surgery. Following euthanasia, a longitudinal incision was made in the dorsal region above the spine; then, both sides were opened to (i) retrieve the samples of skin (sample area of $1 \times 1$ cm) including the SO (implant area in Fig. 2B) and (ii) take the CTR samples (control areas in Fig. 2B). All implanted SOs were visible by gross inspections at all the time points and no sepsis or evidence of infection was seen (Fig. 5).

Furthermore, histological analysis showed interesting aspects. The SOs, under both the macroscopic (Fig. 5) and histological views (Fig. 6), neither show induction of any inflammatory/edema signs nor macrophage migration and infiltration of new fibroblasts or necrosis at all the time points (1, 2, 4 and 12 weeks). This is also in line with the IHC analysis: as expected, the presence of CD68 positive cells was observed both in the control rat tissue (Fig. 7-CTR) and in the implanted animals, in proximity to blood vessels at a distance from the implant site (right panels in Fig. 7-IMPLANTED). Interestingly, in the area close to the SO, the connective tissue displays a well-organized...
and compact collagen texture (asterisks in Fig. 8) compared to the adjacent collagen that has a looser organization and presents a greater quantity of vessels (circles in Fig. 8). This different organization of the connective tissue is visible immediately adjacent to the SO at all the time points tested.

Quantification of blood vessels is showed in Fig. 9: the histogram clearly shows that the amount of blood vessels found in the IMPLANTED samples is statistically significant compared to the CTR. Furthermore, significant differences in the quantity of vessels in the IMPLANTED samples are visible especially between weeks 1–2, 1–12 and 4–12, but decrease with time. The authors speculate that these data suggest the hypothesis that at the onset, the early vascularization process depends on factors issued by SOs, and overtime the consolidation of the vascular environment is strongly aimed at the osteogenesis induction. As an element of discussion, it is worth emphasizing that neovascularization is a positive response to the introduction of a foreign body in an organism, that ensures a proper supply of biochemical signals, oxygen and nutrient to the cells of the tissue surrounding the implant.82

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**Fig. 6** Representative overview of the rat skin control (CTR) and skin grafted with SO (IMPLANTED). H&E = Hematoxylin and Eosin staining shows connective tissue in pink with nuclei stained in dark violet; MT = Masson’s Trichrome staining shows collagen fibers in green, muscle fibers and glands in violet, and nuclei in blue. Images were taken in the 2nd week but the panel is representative of all the other time points.

**Fig. 7** A representative overview of the rat skin control (CTR) and skin grafted with SO (IMPLANTED). Immunohistochemistry analysis for CD68 positive cells. On the left, positive cells in the subcutaneous area of rat skin in the CTR sample; on the right, two different fields of the IMPLANTED samples in which the absence of positive cells close to the implanted SO is clearly visible along with some positive cells in the area far from the implant. The yellow area indicates the SO, detached during the immunohistochemistry procedures.
The extensive blood vessel formation suggests the onset of a series of events able to speed up the vascularization, thus enhancing the efficacy of tissue formation. These features are very important for a biomaterial able to stimulate angiogenesis without amplifying inflammation, underlining that, in this case, angiogenesis is an essential prelude to osteogenesis processes.83,84

3.3. Mass spectrometry analysis

Following in ovo and in vivo observations, demonstrating that SO has a very strong ability to induce angiogenesis, we conducted further investigation by mass spectrometry in order to determine which factors were responsible for these effects.

Since both the SOs and the conditioned media were inducers of angiogenesis, the protein content of SOs was analyzed by mass spectrometry analysis (LCMSQE). All proteins were categorized in GO annotations analyzing molecular functions, biological processes and protein classes.

In Fig. 10, it is clearly visible that the majority of proteins found inside the SOs (panel A) are involved in binding activity, catalytic activity and structural molecular activity (such as extracellular matrix structural constituent, etc.).

They are also mainly involved in cellular processes, localization and metabolic processes (Fig. 10B). Finally, the most abundant protein classes are cytoskeletal proteins (mainly actin and microtubule families), receptors and signaling molecules (Fig. 10C).

These pieces of evidence could suggest that several proteins belonging to these categories are also involved in the angiogenesis process.85–87

Next, in order to validate the previous observations concerning the SO angiogenic effects, we analyzed all the hits found by LCMSQE analysis by means of proteomic and interatomic data-bases. A series of proteins were identified as belonging to the Gene Ontology (GO) annotations: vasculature development (GO0001944), cardiovascular system development (GO0072338), regulation of vasculature development (GO1901342) and negative regulation of vasculature development (GO1901343).

All the proteins defined in Table 2 are involved in the pathways of angiogenesis or vascular development and act as posi-
In the literature, it is reported that the angiogenic processes of endothelial cell growth, migration and tube formation are finely regulated by the balance between pro- and anti-angiogenic factors, cell–extracellular matrix interactions and matrix-degrading proteases. Therefore, to complete the previous investigations on the proteins found in SOs, the LCMSQE analysis confirmed the presence of proteins with angiogenic properties capable of inducing angiogenic reactions in the surrounding tissue.

In particular, the first protein with the highest MASCOT protein score is decorin (DCN), a chondroitin/dermatan sulphate proteoglycan, belonging to the small leucine-rich proteoglycan family, which is involved in the collagen fibril assembly to control fibril formation and regulating their mechanical properties. It also plays an important role in the induction of a specific set of metalloproteinases and the concomitant formation of cord-like structures and cell-lined cavities. Furthermore, decorin has also been found to be expressed in a significant amount around neovessels after varicose vein surgery in patients, and it has been shown in animal models that the injured cornea deficient for decorin has compromised angiogenesis. Moreover, it is worth noting that decorin activates angiogenesis through the IGF1R/AKT/VEGF pathway, one of the most important signal transduction mechanisms involved in speeding the cell proliferation in normal and cancer cells. In parallel decorin seems to be involved in anti-angiogenic processes; fragments of decorin seem to depress VEGF-induced focal adhesion kinase phosphorylation and assembly of focal adhesions. A retardation in cornea neovascularization has also been observed via a downregulation of proangiogenic molecules including VEGF, linked with an overexpression of decorin.

Therefore, decorin’s role in angiogenesis (pro- or anti-angiogenic) seems to be dependent on the molecular microenvironment (where angiogenesis is induced). Decorin can either promote or inhibit angiogenesis and thus, impacts the fate (life and death) of endothelial cells.

Other relatively abundant proteins found on SOs are myosin heavy chain 9 and 10. There are two non-muscle myosin II also namely NMHC IIA and NMHC IIB, respectively. In particular myosin-9, identified as a nucleolin-binding protein, is the physical linker between nucleolin and...
the cytoskeleton. This actin-based motor protein provides an essential function, together with those of the VEGF and ECM, in linking nucleolin and the cytoskeleton in order to mediate nucleolin’s function in angiogenic processes.97 Nucleolin has also been described as a nucleolar protein involved in cell proliferation, cytokinesis, replication, embryogenesis and nucleogenesis. Moreover, cell surface nucleolin is essential for migration and tube formation of endothelial cells. During angiogenesis, the upregulation of cell-surface nucleolin is attributed to the shuttle of nucleolin from the nucleus to the cell membrane, mediated by myosin-9.95 In summary, the LCMSQE analysis supports the previous in vitro and in vivo analysis showing that the proteins present in SOs could induce angiogenic effects.

4. Conclusions

In this research, we demonstrated the capability of the previously characterized SOs to release bioactive molecules able to induce a clear angiogenic response. A variety of techniques were used to investigate, firstly, the angiogenic response in ovo and in vivo and, secondly, the biocompatibility in vivo. After observing both the angiogenic potential and the biocompatibility, the putative actors of these effects were also sought by highlighting which angiogenic factors could be released from this peculiar material. In conclusion, the scaffold investigated in the present study might be considered a good candidate as a bone substitute or a TE scaffold for future clinical applications.

Conflicts of interest

There are no conflicts to declare.

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