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Local administration of stem cell-derived extracellular vesicles in a thermoresponsive hydrogel promotes a pro-healing effect in a rat model of colo-cutaneous post-surgical fistula†

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Extracellular vesicles (EVs), especially from stem/stromal cells (SCs), represent a cell-free alternative in regenerative medicine holding promises to promote tissue healing while providing safety and logistic advantages in comparison to cellular counterparts. Herein, we hypothesize that SC EVs, administered locally in a thermoresponsive gel, is a therapeutic strategy for managing post-surgical colo-cutaneous fistulas. This disease is a neglected and challenging condition associated to low remission rates and high refractoriness. Herein, EVs from a murine SC line were produced by a high-yield scalable method in bio-reactors. The post-surgical intestinal fistula model was induced via a surgical cecostomy communicating the cecum and the skin in Wistar rats. Animals were treated just after cecostomy with PBS, thermoresponsive Pluronic F-127 hydrogel alone or containing SC EVs. A PET-monitored biodistribution investigation of SC EVs labelled with ⁸⁹Zr was performed. Fistula external orifice and output assessment, probe-based confocal laser endomicroscopy, MRI and histology were carried out for therapy follow-up. The relevance of percutaneous EV administration embedded in the hydrogel vehicle was indicated by the PET-biodistribution study. Local administration of SC EVs in the hydrogel reduced colo-cutaneous fistula diameter, output, fibrosis and inflammation while increasing the density of neo-vessels when compared to the PBS and gel groups. This multi-modal investigation pointed-out the therapeutic potential of SC EVs administered locally and in a thermoresponsive hydrogel for the management of challenging post-surgical colon fistulas in a minimally-invasive cell-free strategy.

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Introduction

Extracellular vesicles (EVs) from stem/stromal cells (SCs) are recent players in the evolving landscape of regenerative medi-

cine bringing along new promises and challenges. In the front-line of cell therapy, SCs are known to provide the microenvironment with trophic and survival signals including cytokines released by means of EVs, pointing out EV role in paracrine

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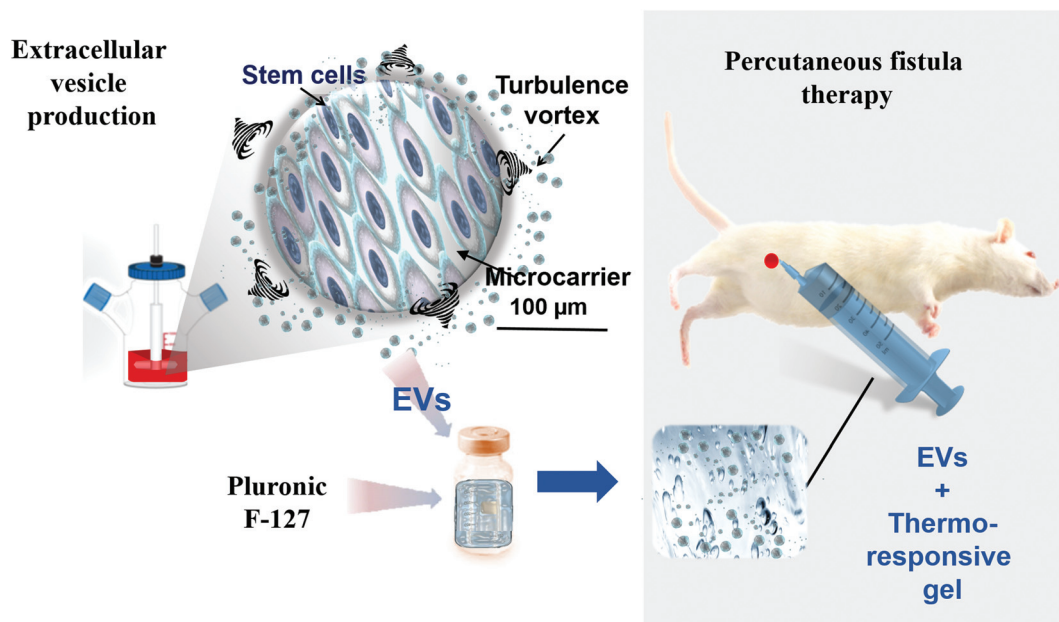


Fig. 1 Schematic representation of the experimental approach based on the production of EVs at high yield by turbulence stimulation from murine stem cells cultured on microcarriers in 1 L bioreactors. EVs were combined to Pluronic F-127 thermoresponsive hydrogel and administered percutaneously at $<19\text{ }^{\circ}\text{C}$ (below gel transition temperature) in a colo-cutaneous fistula model in rats. The formulation gelling *in situ* at body temperature is intended to increase EV residence time at the site of interest.

Experimental section

EVs and gel production and characterization

Cell culture in 2D flasks. Murine C3H/10T1/2, Clone 8 (ATCC®) cells were cultured at $37\text{ }^{\circ}\text{C}$ and 5% CO_2 . Dulbecco's Modified Eagle Medium (DMEM) supplemented with 10% fetal bovine serum (FBS) and 100 U mL^{-1} penicillin and 100 U mL^{-1} streptomycin was used. Cells were cultured in 150 cm^2 flasks until confluence.

3D cell culture in spinner flask bioreactors. Commercial dextran microcarriers of about $200\text{ }\mu\text{m}$ (Cytodex 1, GE Healthcare) were dispersed in PBS, and autoclaved for sterility. PBS was changed to DMEM without phenol red, and stored at $4\text{ }^{\circ}\text{C}$. Previous to cell seeding, microcarriers were incubated in $37\text{ }^{\circ}\text{C}$ complete medium at a 6 g L^{-1} concentration during 3 h. Cells were seeded with a cell to microcarrier ratio of 5/1 and submitted to 24 cycles of 45 min of rest followed by 3 min of a gentle mixing at 60 rotations per minute (rpm) in 1 L bioreactor to enable homogeneous adhesion of cells on microcarriers. After cell adhesion, microcarriers were diluted to 3 g L^{-1} with complete medium at $37\text{ }^{\circ}\text{C}$ at 60 RPM until reaching confluence on microcarriers (about 7 days). Every 2–3 days, 30 to 70% of the medium was changed according to the cell confluence.

Turbulence-triggered EV production in spinner flask bioreactors. Once cells were confluent on microcarriers, turbulence-triggered EV production was launched. The protocol consisted in changing the complete medium to heated ($37\text{ }^{\circ}\text{C}$) serum-free DMEM media without phenol red with 100 U mL^{-1} penicillin and 100 U mL^{-1} streptomycin after 3 rinsing steps.

By controlling the spinner flask stirring, cells on microcarriers were submitted to a turbulent flow featuring a Kolmogorov length of $35\text{ }\mu\text{m}$ during 4 hours to stimulate EV release.⁵² After that, cellular debris of the supernatant were removed by centrifugation at $2000g$ for 10 min. The supernatant was ultracentrifuged at $110\text{ }000g$ for 70 min to obtain an EV pellet.

Nanoparticle tracking analysis. EV size distribution and concentration were determined by nanoparticle tracking analysis (NTA) using a Nanosight LM10-HS (NanoSight, UK) with a 405 nm laser. Before measurements, EVs were diluted to an appropriate concentration (between 3×10^8 and 2×10^9 particles per mL) with sterile PBS (confirmed to be particle-free by NTA measurement). For each sample, 5 movies of 30 s were recorded using a camera level of 16. Data were analyzed with NTA Analytical Software.

Cryo-transmission electron microscopy. A total of $2\text{--}7\text{ }\mu\text{L}$ of SC EVs was deposited onto an electron microscopy grid coated with a perforated carbon film (Ted Pella, Redding, CA, USA), the excess liquid was blotted off with a filter paper, and the grid was then quickly plunged into liquid ethane. Analysis was carried out in MET Jeol 2100 (LaB6) at 80 kV.

EV marker analysis by ExoView device. A murine SC EV batch produced by the turbulence method was analysed by ExoView (NanoView Biosciences, USA). All samples were diluted in PBS with 0.05% Tween-20 (PBS-Tw). The samples were incubated on the ExoView Tetraspanin Chip for mouse EVs placed in a 24-well plate for 16 h at room temperature. The chips were washed 3 times with PBS-Tw. Chips were incubated with ExoView Tetraspanin Labelling antibodies that consist of anti-CD81 Alexa-555, anti-CD63 Alexa-488, anti-CD9



Histological analysis. At D60, rats were sacrificed and the fistula site as well as its periphery (1 cm³ specimen) were collected and transferred to a formaline solution. Specimens were embedded in paraffin and sectioned perpendicular to the center of the fistula to obtain thin tissue sections of 7 μm, which were stained with hematoxylin and eosin (HE) and Sirius Red (fibrosis assessment). Slices were analyzed with an optical microscope (Leica DMIL). Two investigators, blinded to treatment allocation, performed histological analyses. All slides were digitally scanned (Digitiser Hamamatsu Photonics®, Massy, France) and analyzed with dedicated software (NDP.view software®, Massy, France). The density of cell type, fibrosis and inflammation were determined by semi-quantitative analysis using a score from 0 to 5. Fibrosis score (from 0 to 25) was calculated by semi-quantitative density of fibrosis (score from 0 to 5) multiplied by semi-quantitative fibroblast score (from 0 to 5). A histological scoring was used for evaluation inflammation.

Statistics

The results are presented as means ± standard deviation for continuous variables, and as percentages for categorical variables. Fischer's exact test was carried out for comparisons between categorical variables and the nonparametric Mann-Whitney test was used for non-paired continuous variables. Comparisons between more than two groups were performed with the nonparametric Kruskal-Wallis test. An estimation of the p value by the Chi-square test was carried out for the comparison concerning the number of cases per group. A P-value of <0.05 was considered indicative of significance. Statistical analysis was conducted using SPSS software version 24.0 (IBM Corp., Armonk, NY). GraphPadPrism (Graphpad Software, La Jolla, CA, USA) software was used for statistical analysis information displayed in figures.

Results

High-yield production of EVs in bioreactors triggered by turbulent flow

Murine SCs were cultured in 3D on the surface of micro-support beads in spinner flasks. Stirring speed in these bioreactors was tuned in order to generate a turbulent flow featuring a Kolmogorov length of 35 μm. Murine SCs were stimulated during 4 hours. Typically, this turbulence set-up may enable the production of about 500 000 EVs per cell in 4 hours while about 10 times less is produced in 48 h of starvation.⁵² Herein, a total of 10¹³ murine SC EVs were obtained in 1 L bioreactors and were then isolated by ultracentrifugation and characterized by cryo-TEM, NTA and Exoview. Cryo-microscopy analysis evidenced membrane-enclosed round-shaped structures featuring high size polydispersity between 70 and 500 nm (Fig. 2A). Size distribution characteristic to EVs mainly spanning from 100 to 250 nm was observed. The values for the mean size and mode obtained by NTA were 136.2 nm and 94.8 nm, respectively, with a size distribution standard deviation

of 70.4 nm (Fig. 2B and C). In more details, 90% of the particles were smaller than 228.6 nm, 50% of the particles were smaller than 112.7 nm, while 10% of the particles were smaller than 81.6 nm.

EV markers were characterized for a murine SC EV batch produced by the turbulence method *via* Exoview analysis. The obtained data indicated that the particles could be immune-captured by anti CD63 and CD81 antibodies. Minimal capture was observed by anti CD9 antibody and the same applied to negative control mouse IgG antibody (Fig. 2D). Fluorescence images enabled to visualize of single immune-captured EVs further labelled by fluorescence antibodies, especially anti CD81 (Fig. 2E). Minimal fluorescence was detected for calnexin negative cytosolic marker when using an anti-calnexin fluorescent antibody in permeabilized immune-captured EVs.

As a final formulation, EVs were combined to PF-127 hydrogel at 20% just before use. The rheological properties of PF-127 hydrogel were assessed by the evolution of dynamic storage modulus (*G'*) and loss modulus (*G''*) as a function of the temperature, confirming the thermoresponsive properties of the obtained hydrogels (Fig. 2F). A transition temperature near 20 °C was determined at *G'* and *G''* cross over.

Cecostomy enabled the induction of a post-surgical colocolocutaneous fistula model in rats

A surgical colocolocutaneous fistula model was performed *via* a cecostomy at day 0 (Fig. 3A and B). After the surgical procedure, all fistulas featured an external orifice diameter ≥4.5 mm. Besides, all fistulas were permeable, featuring faeces secretion output, in 100% of the cases. The model allowed the formation of a communication resembling a digestive fistula-like disease with skin and intestinal lesions. The presence of a fistula tract was observed by MRI (Fig. 3C). There was no complication related to fistula induction surgery. The mean of weight before to surgery was 265.2 ± 8.3 g. Therapy follow-up after surgery was 60 days. At the end of the follow-up mean weight was 274.4 g ± 20.8 g.

Percutaneous EV delivery in the thermoresponsive gel allowed EV retention in the fistula tract

We investigated the interest of the percutaneous EV administration embedded in the thermoresponsive gel (Fig. 3D) compared to other administration strategies. A thorough biodistribution study was out of the scope of the present paper. Our aim was to appraise the potential relevance of the proposed administration strategy providing a general picture of biodistribution patterns obtained by other delivery approaches. For that, an EV radiolabeling procedure with [⁸⁹Zr]zirconium was conceived. The direct labelling of DFO-EVs with [⁸⁹Zr][Zr(ox)₄]⁴⁻ was unsuccessful. Therefore, a [⁸⁹Zr][Zr(8-HQ)₄] labelling procedure was used and this complex was obtained in 52% yield (67.6 MBq), with radiochemical purity of 97.5%. [⁸⁹Zr]Zr-EVs were obtained in a 32.5% yield (22 MBq) with a radiochemical purity of 89.5%. Animals were treated with [⁸⁹Zr]Zr-EVs *via* percutaneous intra-fistula administration in saline, percutaneous intra-fistula administration embedded in



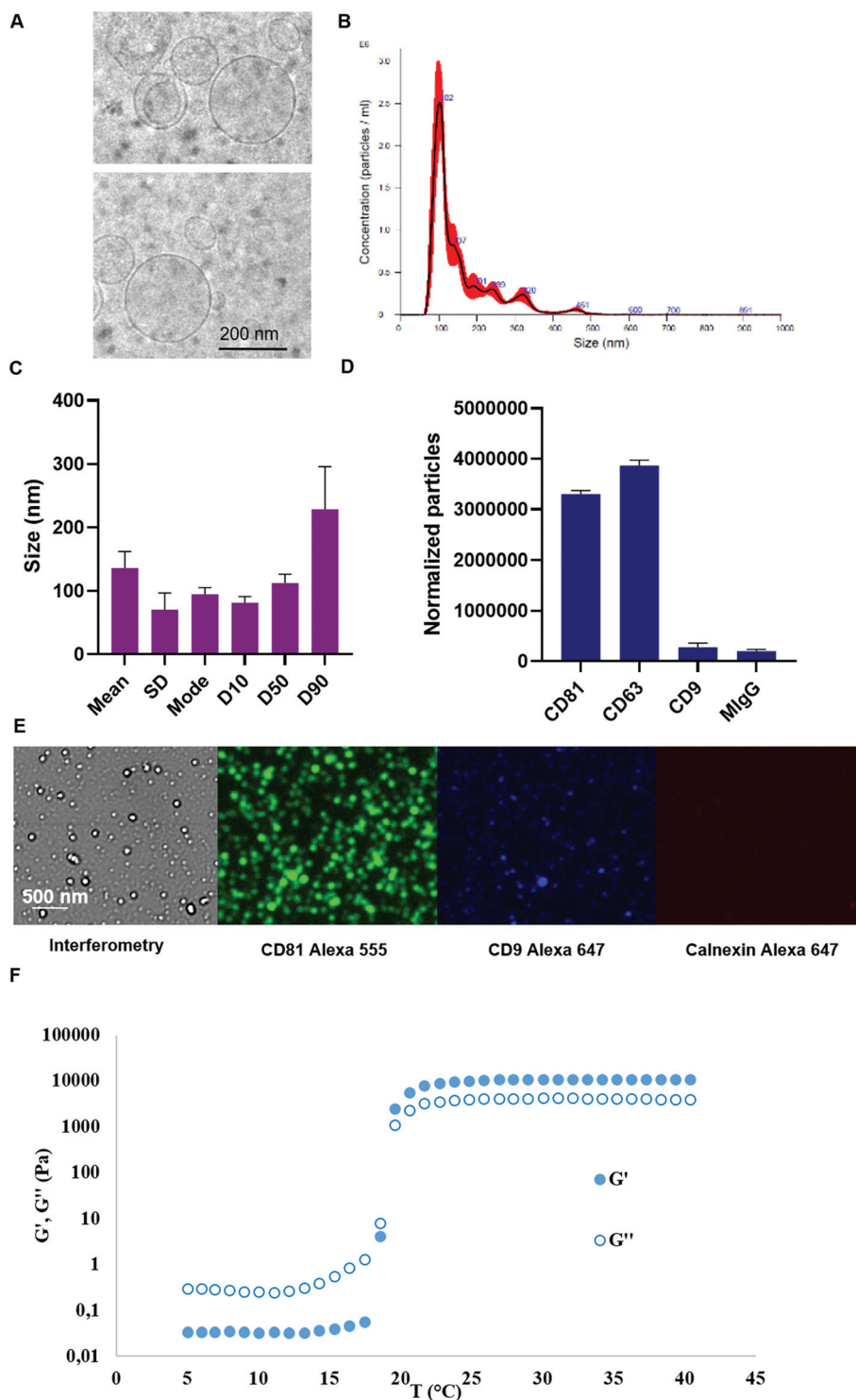


Fig. 2 Characterization of murine stem cell (SC) EVs and Pluronic F-127 hydrogel. Cryogenic transmission electron microscopy images of SC EVs (A). Size distribution of SC EVs, averaged on 5 recorded videos (B) and the mean, mode, standard deviation and percentiles of the size of EV samples, represented as the mean \pm SEM of three independent batches by nanoparticle tracking analysis (C). EV marker analysis by ExoView for SC EVs using CD9, CD63 and CD81 capture antibodies with a mouse IgG2a (MlgG2a) used as negative control capture antibody (D). Representative fluorescence images of the multiplexed EV marker analysis by ExoView for SC EVs using CD81 capture antibodies. Refractometry images are shown as well as fluorescence images using CD81 and CD9 detection antibodies (positive markers) and calnexin fluorescence detection antibody (negative marker) (E). Evolution of dynamic storage modulus (G') and loss modulus (G'') of a Pluronic F-127 at 20% at a heating rate of $5\text{ }^{\circ}\text{C min}^{-1}$ from $5\text{ }^{\circ}\text{C}$ to $40\text{ }^{\circ}\text{C}$ for the determination of the sol-gel transition temperature (G' and G'' cross over) (F).



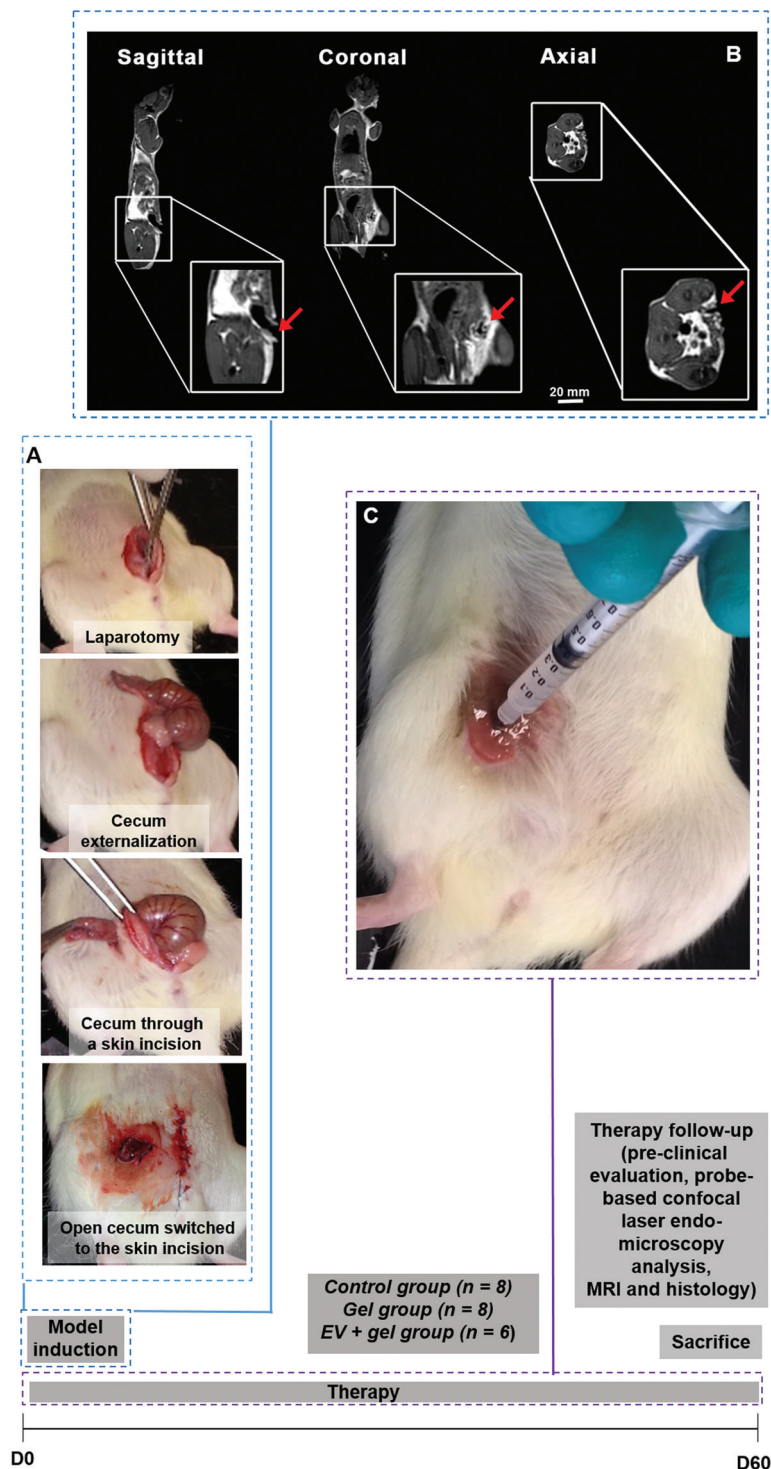


Fig. 3 Design of the *in vivo* experiments and their main steps. Study timeline and sample size per group are displayed. Surgical procedure for the colo-cutaneous fistula model induction from laparotomy to cecum communication to the skin in rats (A). T1 weighted MRI illustrating the colo-cutaneous fistula model. The red arrows indicate the fistula in different anatomical planes: sagittal, coronal, and axial (B). Percutaneous administration of the EVs into the Pluronic F-127 gel *via* the fistula external orifice (C).

the thermoresponsive gel or *via* intravenous injection in saline (Fig. 4). We observed that the percutaneous [^{89}Zr]Zr-EVs treatment in saline induced a disseminated tracer uptake in the digestive tract at 1 h. A fast decrease in the SUV of 75% in the

fistula site was evidenced between 1 h and 24 h (56 and 14, respectively). Concerning the [^{89}Zr]Zr-EVs delivery percutaneously in the thermoresponsive gel, PET signal mainly remained limited to the site of interest and the SUV decrease



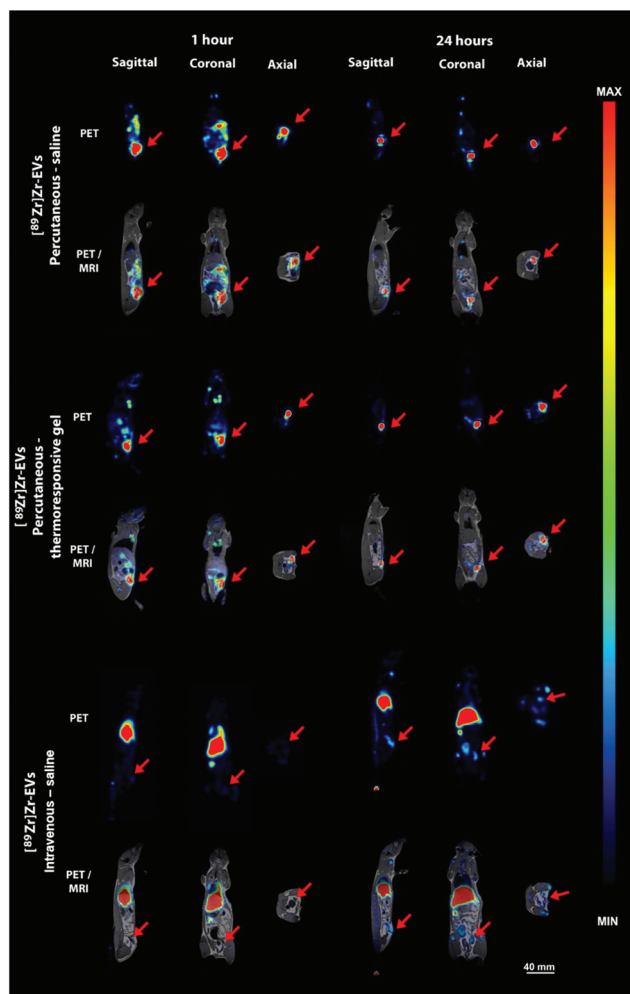


Fig. 4 PET and PET/MRI images illustrate the biodistribution and local uptake of radiotracer labeled EVs ($[^{89}\text{Zr}]\text{Zr-EVs}$), 1 hour and 24 hours after local percutaneous administration, in suspension or embedded in the thermoresponsive gel, or intravenous injection (3 anatomical planes are illustrated). Red arrows indicate the radiotracer uptake in the colcutaneous fistula. Note that the percutaneous treatment without gel (first line) revealed faster standardized uptake value (SUV) decrease in the fistula site, SUV mean between 1 hour and 24 hours decreased around 75%. For the percutaneous treatment with gel (middle line), the SUV at the fistula site decreased around 30% between 1 hour and 24 hours. For the intravenous injection (bottom line), there was a $[^{89}\text{Zr}]\text{Zr-EVs}$ uptake increase of 400% between 1 hour and 24 hours. However, SUV levels were 3 times lower when compared to local percutaneous administration.

was of 30% between 1 hour and 24 hours (30 and 21, respectively). After intravenous injection, $[^{89}\text{Zr}]\text{Zr-EVs}$ in saline mainly displayed a liver accumulation pattern along with minor uptake in the fistula site and joints. An increase of 400% was observed for the SUV in the fistula site between 1 hour and 24 hours (1.5 and 7.5, respectively) indicating EV homing. However, this SUV level were 3 times inferior when compared to local percutaneous administration. At day 7th post-administration, imaging was performed for animals treated with EVs embedded in the gel administered percutaneously or following

EV intravenous injection in saline. The tracer signal was still mainly limited to the fistula site featuring a SUV decrease of just 15% compared to SUV at 24 hours for the percutaneous EV administration into the thermoresponsive gel condition (Fig. S1†). Comparatively, an almost 4-fold decrease in the SUV at the fistula site was observed for the intravenous injection condition when comparing time points 7 days and 24 hours. As control conditions, the PET biodistribution of the of the $[^{89}\text{Zr}][\text{Zr}(\text{ox})_4]^{4-}$ tracer alone or linked to the conjugation $[^{89}\text{Zr}][\text{Zr}(\text{DFO})]$ molecule was recorded at 24 hours and 7 days after intravenous administration in saline or percutaneous administration in the thermoresponsive gel. Data showed a biodistribution pattern different from the ones observed for the $[^{89}\text{Zr}]\text{Zr-EVs}$ (Fig. S2†). For instance, the intravenous administration of $[^{89}\text{Zr}][\text{Zr}(\text{ox})_4]^{4-}$ indicated a bone biodistribution (spine) quite different for the liver biodistribution observed for the intravenous $[^{89}\text{Zr}]\text{Zr-EVs}$ injection. This seems to indicate the effective tracer conjugation to EVs.

EV therapy in the thermoresponsive gel decreased fistula output and external orifice diameter

Repeated weekly treatment during 60 days with the gel or the gel with EVs was well tolerated by animals. No evident signs of local irritation, allergy or toxicity were observed. No statistically significant difference in mean body weight was evidenced for animals in the control, gel and EVs + gel group at day 60th after the begin of the therapy (Fig. S3†). Fistula output was evaluated at day 30th. One case of no fistula output (absence of faeces secretion) from 8 animals was observed for the control and also for the gel group, corresponding to 12.5%, for both. Two cases of no fistula output from 6 animals were observed for the group treated with EVs and gel, corresponding to 33% of the animals (Fig. S4†). Fistula closure, output and external orifice diameter were evaluated at day 60th. One case of fistula closure per group was observed for each group, corresponding to 12.5%; 12.5% and 16.7% of the cases for the control, gel and EVs + gel groups. The frequency of no fistula output was significantly different when comparing the control group (1 case from 8: 12.5% of the animals) to the EVs + gel group (4 cases from 6: 66.7% of the animals) (Fig. 5A). No significant difference was observed when comparing the frequency of no output for the control group to the gel one (2 from 8 cases: 25% of the animals) nor the gel group to the EVs + gel one. The external orifice diameter distribution showed a clear shift towards smaller size ranges. The size of external orifice diameter ranges increased in the following sequence EVs + gel < gel < control animals (Fig. S5†). The frequency of fistulas featuring an external orifice diameter <3 mm was significantly different when comparing the control group (1 case from 8: 12.5% of the animals) to the EVs + gel one (4 cases from 6: 66.7% of the animals). No significant difference in the frequency of fistulas featuring an external orifice diameter <3 mm was observed when comparing the control group and the gel one (3 cases from 8: 37.5% of the animals) nor the gel group to the EVs + gel one (Fig. 5B). The macroscopic view of the fistulas revealed a higher and





Fig. 5 Potency evaluation *in vivo* by a multi-modal approach in rats treated with the gel or EVs + gel in comparison to an untreated negative control. Preclinical evaluation of fistulas at day 60 indicating the percentage of animals per group featuring the absence of faeces (no output) at the external fistula orifice (A) as well as percentage of animals per group featuring an external orifice diameter inferior to 3 mm (B). For both A and B, *p* values were estimated by Chi-square test based on the number of cases per group featuring the presence (output+) or absence of feces (output-) at the external fistula orifice (A) and number of cases featuring an external orifice diameter inferior or superior to 3 mm (B). Macroscopic aspect of external fistula orifices at day 60 for the three groups (C). Therapy follow-up by probe-based confocal laser *endo*-microscopy analysis at day 60 showing vessel network (D) and the quantification of total vessel area (E) for each group. Therapy follow-up by MRI analysis at day 60 showing fistula area quantification (F) and representative related images (G) for each group (fistula tracts were indicated by yellow dotted circles). For both E and F, *p* values were computed using Mann Whitney test.

extended zone featuring an inflammatory aspect surrounding the external orifice for control and gel groups in comparison to fistulas from the EVs + gel group (Fig. 5C).

EV therapy in the thermoresponsive gel seemed to improve vascularization while reducing fistula area: an image-based therapy follow-up

Vascularization parameters were analysed by probe-based confocal laser endomicroscopy. No significant difference was observed for group comparison in terms of vessel diameter, total vessel length, total vessel area, mean vessel diameter nor functional capillary density length, for instance (Table S1†). There was a trend towards a higher functional capillary density

area and total vessel area for animals from the group gel or EVs + gel one in comparison to control (Fig. 5D, E and Table S1†). This trend did not reach statistical significance probably due to the reduced sample size.

Fistula metrics was assessed by MRI analysis right after the percutaneous injection of a gadolinium contrast agent *via* the external fistula orifice at day D60. There was a trend towards a reduced fistula length when comparing EVs + gel group to control (Table S2†). MRI data evidenced a significant reduction of the fistula area when comparing EVs + gel group to control (Fig. 5F and G). No significant reduction of the fistula area was observed when comparing the gel group to control nor the gel group to the EVs + gel one.



Histological analysis indicated a decrease in inflammation and fibrosis and an increase in neo-vessel density induced by EV therapy in the thermo-responsive gel

Histological analysis was carried out to assess inflammation, fibrosis and vascularization. A histology score (integrating epithelial damage, mucosal inflammation and submucosal inflammation parameters as indicated in the Table S3†) evidenced a significant difference when comparing the control group to the EVs + gel one (Fig. 6A). A significant difference was also observed when comparing the gel group and EVs + gel one. Accordingly, the density of polynuclear neutrophils (PNN) was significantly lower for the EVs + gel group than observed for the control one (Fig. 6B). The histological evaluation validated the trend observed by probe-based confocal laser endomicroscopy as the density of neo-vessel score was significantly higher for the EVs + gel group in comparison to control (Fig. 5C). This is also in agreement with *in vitro* pro-angiogenic (pro-migratory) properties of the EVs (Fig. S6†). A significant reduction in the fibrosis score was also evidenced when comparing the EVs + gel group to the control one (Fig. 6D). Besides, a significant difference in the fibrosis score was also observed when comparing the gel group and the EVs + gel one. Extensive fibrosis and mucosa damage was observed for the control group (Fig. 6E). Fibrosis and mucosa damage seemed more limited in the gel group (Fig. 6F). Fibrosis was limited in the gel and EVs + gel group, which featured minimal mucosa damage (Fig. 6G). Table S4† provides additional histology data.

Discussion

Herein we investigated a biomaterial-based cell-free local therapy for post-surgical fistulas. This combined therapy relied on turbulence SC EVs embedded in a PF-127 thermo-responsive hydrogel administered percutaneously in colo-cutaneous fistulas.

As far as we know, this is the first time that SC-EVs incorporated in a thermo-responsive hydrogel are investigated for the therapy of post-surgical colo-cutaneous fistulas as an alternative to cell therapy. In a previous report from our team, SC organized in double cell sheets showed a therapeutic effect in an equivalent fistula model in mice.⁵⁹ In the attempt to shed light on the mechanisms of cell therapy for fistula treatment, our group performed *in vivo* experiments with doubled labelled SCs for fluorescence and MRI tracking. SC detection at fistula site was showed just following transplantation, but it could not be clearly evidenced 7 and 14 days post-treatment. We showed that SC cell sheet therapeutic benefits on fistula healing were accompanied by a paracrine effect with an increase in the expression of anti-inflammatory cytokines and intestinal growth factors involved in tissue repair.⁵⁹

As EVs are known to participate in SC paracrine role,^{1,2} we decided to test herein an EV-based cell free therapy. SC EVs were produced *via* a high-yield scalable strategy in bioreactors. The turbulence method in 4 h may enable up to a 10-fold

increase in EV production yield compared the state-of-the-art starvation approach in 48 h.⁵² Our turbulence strategy is fully integrated to the cell culture in stirred tank in cGMP bioreactors. We evidenced that the turbulence approach was scalable based on turbulence physical laws *via* the Kolmogorov equation.⁵² The choice of the turbulence approach enabling high-yield, scalability and cGMP compliance is one of the translational strengths of our investigation. The comparison between turbulence SC EVs and state-of-the-art ones in terms of properties and potency *in vitro* is provided elsewhere.⁵²

Turbulence SC EVs were combined to PF-127 thermo-responsive hydrogel for local colo-cutaneous fistula therapy. The aim of the hydrogel was two-fold: (i) to act as a sealant promoting fistula mechanical occlusion and (ii) to constitute a delivery system for EVs modulating their release and biodistribution. Concerning fistula sealants, other biomaterials such as porcine plug or fibrin glue⁶⁰ have been investigated in a clinical setting for post-surgical fistula therapy. However, the use of these biomaterials may be limited by abscess formation⁶¹ and spontaneous expulsion.⁶² We have recently evidenced in a case report that PF-127 gel may be a thermo-actuated fistula sealant of interest. It was well-tolerated by the patient while inducing a reduction in the external fistula orifice diameter and secretion output volume in combination to a stent placement.⁶³ Due to its thermo-responsive properties, this biomaterial can be administered as a liquid *via* a catheter. Its gelation *in situ* at body temperature is expected to enable its retention at the fistula tract, limiting the circulation of secretions and thus favoring the healing process.

As we have mentioned, the use of the PF-127 gel herein goes beyond a sealant agent also providing a delivery system for EV administration locally in the rat model of colo-cutaneous post-surgical fistula. In this regard, we developed an EV labelling strategy with a radiotracer in order to identify the biodistribution pattern of EVs following their administration either percutaneously in the gel or in saline in comparison to an intravenous administration in saline. We selected [⁸⁹Zr]zirconium as the radionuclide as its half-life of 3.3 days enables the *in vivo* tracking from hours to days post administration. EVs were tagged with [⁸⁹Zr]zirconium *via* a double labelling strategy. In a first attempt, we investigated EV labelling by the direct reaction between DFO-EVs⁵⁴ with [⁸⁹Zr][Zr(ox)₄]⁴⁻. This strategy was not successful. Therefore, we decided to use [⁸⁹Zr][Zr(8-HQ)₄] to react with DFO-EVs which could concurrently (i) transfer part of the ⁸⁹Zr⁴⁺ to the DFO bounded to EVs or (ii) cross the lipid membrane delivering ⁸⁹Zr⁴⁺ into the EV inner compartment. Control experiments showed that the biodistribution of [⁸⁹Zr][Zr(ox)₄]⁴⁻ or [⁸⁹Zr][Zr(DFO)] was quite distinct from the one observed for [⁸⁹Zr]Zr-EVs. This supports the assumption of a successful EV radiolabelling by our proposed double strategy. As far as we know, this is the first time that EVs are tagged with [⁸⁹Zr]zirconium *via* this double labelling strategy, which allowed tracking up to 7 days post-administration. For instance, different EV radiolabelling strategies with [^{99m}Tc]Tc⁶⁴ or [¹²⁴I]I⁶⁵ were reported in the literature with a tracking time up to and 5 h and 72 h, respectively.





Fig. 6 Potency evaluation *in vivo* by histology in rats treated with the gel or EVs + gel in comparison to an untreated negative control. Therapy follow-up for the three groups by histological analysis at day 60 assessing a histology score based on epithelial damage, mucosal inflammation and submucosal inflammation parameters (A). Therapy follow-up for the three groups by histological analysis at day 60 assessing the polynuclear neutrophils (PNN) density (B), neo-vessel density (C) and fibrosis (D) scores. Mann Whitney test was used for computing p values in group comparison for all scores. Histology analysis (Sirius red staining) of the colo-cutaneous fistulas from control (E), gel (F) and EVs + gel (G) groups. Note that the fistula orifice was delineated in blue color and main fibrosis regions, in red staining, were delimited by *. The epithelium was identified by an orange staining. Important epithelial damage was observed for the control group and its extent was indicated by black arrows.

By using the [⁸⁹Zr]Zr-EVs, we observed that the highest SUV at the fistula region were obtained following percutaneous administration. Besides, signal detection was mainly

restrained to the site of interest when administration was performed percutaneously *via* the thermoresponsive hydrogel. Signal persisted with a 30% reduction in 7 days. In a related



literature approach, a pH-responsive hydrogel featuring immediate gelation upon administration into physiological systems was investigated for local EV delivery. A sustained EV delivery until day 4 post-administration was observed with EV localization in the tissue surrounding the injection site.⁶⁶ In a myocardial infarction model disease, it was reported that SC EV local delivery in the heart of rats *via* an alginate hydrogel improved EV cardiac retention.⁶⁷ *Ex vivo* 2D optical microscopy coupled to 3D optical tomography revealed a stronger fluorescent signal from DiR-labeled EVs for animals treated with in EVs incorporated in the gel compared to those treated with EVs alone.

Considering the potential interest of local percutaneous EV administration in the PF-127 hydrogel indicated by our biodistribution data, we next evaluated the therapeutic outcome of this strategy in colo-cutaneous fistulas.

The combination of EVs and PF-127 gel biomaterial was previously tested in the therapy of post-surgical oesophageal fistulas in pigs by our team.⁶⁸ Of note, a different producer cell type (primary AdSCs) was used in the precedent study. In this preceding investigation, we showed by rheological analysis that the dispersion of EVs into PF-127 gel did not impact its thermoresponsive properties. EV integration into the gel also seemed to have minor impact on EV integrity. We evidenced that EV release was expected to be driven by the saliva flow exerting a dilution effect on the gel. Indeed, we showed that the aqueous dilution of the gel to a concentration inferior to 13% notably induced a viscosity decrease and the release of EVs.⁶⁸ Based on these data, we chose the PF-127 concentration of 20%. In principle, we could choose any concentration $\geq 14\%$. However, at 14% there would be a transition to a liquid state releasing the EVs even at a minimal dilution, resulting in a burst EV release. We chose a concentration superior to this in order to enable the gel to withstand to a dilution factor of about 1.5 before transition to a liquid state to allow EV release in a more sustained way.

In our previous study, an anti-inflammatory, anti-fibrotic and pro-angiogenic effect was observed for animals treated with EVs embedded in the PF-127 hydrogel with total fistula healing being observed in this pig esophagocutaneous disease model.⁶⁸ Considering the differences in fistula tract size and secretion constitution/volume, the EV release and fate as well as the therapeutic outcome following the administration of EVs and PF-127 gel biomaterial in a colo-cutaneous fistula in rats may be indeed distinct from the one observed in the esophagocutaneous fistulas in pigs. Importantly, gel dilution factor is expected to be reduced in the colo-cutaneous model as the faeces secretion features inferior aqueous constitution in comparison to the saliva secretion in the oesophagus. The smaller fistula tract size in the colo-cutaneous rat model in comparison to the esophagocutaneous fistula model in pigs is expected to influence the gel residence time as the “reservoir” volume of these tracts are different. In brief, the combined EV/biomaterial fate and the obtained therapeutic outcome may be influenced by fistula characteristics, which depend on the

involved organs. Despite these differences, we observed a therapeutic effect following the percutaneous EV administration in the gel for both the colo-cutaneous model and the esophagocutaneous one. The investigation of our approach in different fistula settings is of interest in order to extend the clinical relevance of the local delivery of EVs in the PF-127 thermoresponsive hydrogel.

Conclusions

Intestinal fistulas secondary to surgery are a major health problem. Current unmet needs require innovative strategies. We evidenced here the therapeutic potential of a biomaterial-based cell-free local EV therapy for the management of colo-cutaneous post-surgical fistulas. Thanks to a turbulence flow stimulation, EVs were produced in high-yield in an approach that combines scalability and cGMP capabilities, which is a translational asset of our study. EVs were tagged with [⁸⁹Zr]zirconium *via* a double labelling strategy, which allowed tracking up to 7 days post-administration. This radiolabeling approach enabled us to identify the pattern of EV biodistribution following intravenous injection in saline in comparison to percutaneous administration in saline or PF-127 gel in a colo-cutaneous fistula model in rats. We observed that the highest EV retention at the fistula site was observed following percutaneous administration. Besides, fistula retention of EVs was more prominent when their percutaneous administration was performed embedded in the thermoresponsive hydrogel. We assessed therapy outcome following the latter administration strategy by evaluating fistula output and external orifice diameter by pre-clinical macroscopic analysis; vascularization by probe-based confocal laser endomicroscopy and fistula metrics by MRI. Histology was equally assessed. Based on this multi-modal approach, we evidenced that the local administration of SC EVs in the thermoresponsive hydrogel reduced colo-cutaneous fistula external diameter, area, output, fibrosis and inflammation while increasing the density of neo-vessels in comparison to the negative PBS control. Altogether, our study brings prospects for a local minimally-invasive EV delivery relying on a thermo-actuated administration strategy in PF-127 gel in colo-cutaneous fistulas. Importantly, PF-127 gel has been previously approved for clinical use (Legoo, Genzyme), which may facilitate the clinical translation of our approach. This study extends insights from a previous investigation of our team that showed successful post-surgical esophagocutaneous fistula therapy by the equivalent strategy. As a future perspective, additional Crohn's disease and post-surgical fistula models involving different digestive organs should be also investigated to appraise post-surgical fistula disease heterogeneity. Hopefully, these further studies may confirm the versatile potential of the proposed approach as a next-generation therapeutic option for digestive fistula management in a “one-fits-all” basis. The translational assets of the proposed strategy may accelerate and render economically feasible pharmaceutical development steps toward clinical investi-



gation in the perspective to tackle unmet needs and benefit patient care.

Author contributions

Conception, design, and study supervision: Amanda Karine Andriola Silva, Gabriel Rahmi, Florence Gazeau. Acquisition of data: Max Piffoux, Alba Nicolás-Boluda, Arthur Berger, Alice Grangier, Imane Boucenna, Caroline Cristiano Real, Fabio Luiz Navarro Marques, Daniele de Paula Faria, Amália Cinthia Meneses do Rego, Chloe Broudin, Analysis and interpretation of data: Arthur Berger, Olivier Clément, Christophe Cellier, Carlos Alberto Buchpiguel; Writing, review, and/or revision of the manuscript: Amanda Karine Andriola Silva, Gabriel Rahmi, Claire Wilhelm, Florence Gazeau, Arthur Berger

Conflicts of interest

Florence Gazeau, Amanda Karine Andriola Silva, Claire Wilhelm and Gabriel Rahmi are co-founders of the spin-off Evora Biosciences. Amanda Karine Andriola Silva, Claire Wilhelm are co-founders of the spin-off EverZom. The other authors are no conflicts to declare.

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