A Pathogenetic Role for Endothelin-1 in Peritoneal Dialysis-Associated Fibrosis

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ABSTRACT

In patients undergoing peritoneal dialysis (PD), chronic exposure to nonphysiologic PD fluids elicits low-grade peritoneal inflammation, leading to fibrosis and angiogenesis. Phenotype conversion of mesothelial cells into myofibroblasts, the so-called mesothelial-to-mesenchymal transition (MMT), significantly contributes to the peritoneal dysfunction related to PD. A number of factors have been described to induce MMT in vitro and in vivo, of which TGF-β1 is probably the most important. The vasoconstrictor peptide endothelin-1 (ET-1) is a transcriptional target of TGF-β1 and mediates excessive scarring and fibrosis in several tissues. This work studied the contribution of ET-1 to the development of peritoneal damage and failure in a mouse model of PD. ET-1 and its receptors were expressed in the peritoneal membrane and upregulated on PD fluid exposure. Administration of an ET receptor antagonist, either bosentan or macitentan, markedly attenuated PD-induced MMT, fibrosis, angiogenesis, and peritoneal functional decline. Adenovirus-mediated overexpression of ET-1 induced MMT in human mesothelial cells in vitro and promoted the early cellular events associated with peritoneal dysfunction in vivo. Notably, TGF-β1–blocking peptides prevented these actions of ET-1. Furthermore, a positive reciprocal relationship was observed between ET-1 expression and TGF-β1 expression in human mesothelial cells. These results strongly support a role for an ET-1/TGF-β1 axis as an inducer of MMT and subsequent peritoneal damage and fibrosis, and they highlight ET-1 as a potential therapeutic target in the treatment of PD-associated dysfunction.


Continuous ambulatory peritoneal dialysis (PD) is an alternative to hemodialysis for the treatment of ESRD. The process uses the patient’s peritoneum as a semi-permeable membrane to clear wastes and extra fluid and return electrolyte levels to normal.1,2 The primary advantage of PD, greater patient mobility and autonomy, is counterbalanced by serious complications, including a higher risk of infection and the development of an ultrafiltration failure during long-term exposure to PD fluids.3–5 The peritoneal membrane is lined by a monolayer of mesothelial cells, a specialized cell type that has some characteristics of epithelial cells, acts as a permeability barrier, and secretes various substances involved in the regulation of peritoneal permeability and local host defense.6,7 Exposure to the hyperosmotic, hyperglycemic, and acidic solutions used in dialysis often causes low-grade chronic inflammation and subsequent injury to the peritoneum, which progressively becomes

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denuded of mesothelial cells and undergoes fibrosis and angiogenesis.\textsuperscript{8–12} Such structural alterations are considered to be the principle cause of failure of ultrafiltration, which affects up to 20% of patients undergoing continuous ambulatory PD.\textsuperscript{4,5,13,14}

Several pathogenetic factors, such as inflammatory mediators, high glucose content, the presence of glucose degradation products, and low pH, can induce peritoneal mesothelial cells to lose certain epithelial characteristics and progressively acquire a fibroblast-like phenotype soon after the initiation of PD.\textsuperscript{14–17} This so-called mesothelial-to-mesenchymal transition (MMT) serves as a trigger for peritoneal fibrosis, for which over-activation of the TGF-\(\beta\)1 signaling has been proposed as a pathogenetic mechanism.\textsuperscript{18–22} As such, blocking the action of TGF-\(\beta\)1 by using several strategies has been shown to significantly ameliorate fibrosis and angiogenesis, leading to improved peritoneal function.\textsuperscript{23–27} However, because TGF-\(\beta\)1 is physiologically pleiotropic, nondiscriminate targeting of TGF-\(\beta\)1 signaling may result in undesirable side effects. Thus, the identification of potential TGF-\(\beta\)1 downstream targets involved in the action of this cytokine will provide more specific strategies for the preservation of the peritoneal membrane with limited secondary consequences.

Several lines of evidence indicate that endothelin-1 (ET-1) may be a mediator in the development of excessive scarring and fibrosis in several organs and tissues.\textsuperscript{28,29} ET-1 is normally produced by endothelial cells, but it has been shown also to be overexpressed by fibroblasts in certain fibrotic conditions, including scleroderma and idiopathic pulmonary fibrosis.\textsuperscript{30} Recent reports, including our own work, have shown that ET-1 expression is highly induced by TGF-\(\beta\)1 stimulation, and like TGF-\(\beta\)1, ET-1 has been described to promote the epithelial-to-mesenchymal transition.\textsuperscript{31–33} Some reports have shown that ET-1 is also able to induce the expression of TGF-\(\beta\)1; these observations indicate the existence of reciprocal regulation of these factors in some cellular contexts.\textsuperscript{34–36} Concerning the role of ET-1 in peritoneal fibrosis, early studies have shown that volume stress stimulates peritoneal ET-1 release in PD and that increased osmolarity induces collagen type I RNA synthesis in human peritoneal mesothelial cells through a mechanism inhibited by ET receptor blockade.\textsuperscript{37,38} Nevertheless, despite the pre-eminent role of ET-1 in fibrogenesis in different pathologic contexts, the contribution of this factor to the development of PD-induced peritoneal impairment remains largely unknown.

In this work, we provide evidence arising from both \textit{in vitro} and \textit{in vivo} models supporting a pathogenetic role for ET-1 in the PD-associated peritoneal damage. We show that the main components of the ET system, namely the ligand ET-1 and ET receptors type A (ET\textsubscript{A}) and B (ET\textsubscript{B}), were upregulated in the peritoneum on exposure to PD fluid. We also show that the antagonism of ET receptors by two different dual ET\textsubscript{A}/ET\textsubscript{B} blockers, bosentan and macitentan, with efficacy that has been proved in preclinical and clinical studies\textsuperscript{39,40} resulted in a marked attenuation of PD fluid-induced peritoneal membrane structural and functional alterations. Using human mesothelial cells \textit{in vitro}, we found that ET-1 promoted the expression of the mesenchymal differentiation marker \(\alpha\)-smooth muscle actin (\(\alpha\)-SMA). This action was abrogated by TGF-\(\beta\)1 blockade, therefore suggesting ET-1–induced TGF-\(\beta\)1 expression. Reciprocally, TGF-\(\beta\)1 was also found to increase the expression of ET-1 in this cell model, therefore indicating the existence of a positive feedback loop between these factors to induce the differentiation process. In parallel to the \textit{in vitro} model, adenoviral-mediated overexpression of ET-1 in the peritoneum triggered MMT and fibrocyte recruitment, early events of PD-induced peritoneal damage, and this action was prevented by the blockade of TGF-\(\beta\)1 signaling. Our observations strongly support a role for ET-1 as a contributor of PD-induced peritoneal fibrosis through a TGF-\(\beta\)1–dependent mechanism in mesothelial cells.

RESULTS

ET-1 and Its Receptors Are Expressed in the Peritoneal Membrane and Upregulated in a Mouse Model of PD Fluid Exposure

We initially studied the expression of the main components of the ET system, namely the ligand ET-1 and the receptors ET\textsubscript{A} and ET\textsubscript{B}, in a mouse model of PD fluid exposure.\textsuperscript{41} Immunohistochemical analysis of the parietal peritoneum of animals exposed to saline solution (control group) displayed a significant enrichment of ET-1, ET\textsubscript{A}, and ET\textsubscript{B} immunoreactivity in the outermost layer of the peritoneal membrane (Figure 1A). This staining was associated with the mesothelial cell layer, which was assessed by staining of consecutive sections with an anti-cytokeratin (mesothelial marker) antibody (data not shown). Exposure to PD fluid induced a loss of the mesothelial monolayer and the thickening of the peritoneal membrane. Interestingly, the expression of ET-1 and ET\textsubscript{A}/ET\textsubscript{B} receptors was strongly increased in the submesothelial thickened zone on exposure to PD fluid, suggesting that cells contributing to peritoneal damage and fibrosis might display enhanced ET-mediated signaling (Figure 1A). We have also analyzed the accumulation of ET-1 in the peritoneal cavity. As shown in Figure 1B, ET-1 is significantly accumulated in the effluents of PD fluid-treated mice compared with control animals, an observation that confirms the increased expression found by immunohistochemistry.

ET Receptor Blockers Abrogate PD-Induced Peritoneal Damage and Dysfunction

To analyze the contribution of ET-1–mediated signaling to peritoneal damage, we tested the effect of two different dual ET\textsubscript{A}/ET\textsubscript{B} receptor blockers, bosentan and macitentan, because both receptor types were observed to be upregulated on PD fluid exposure. Administration of either ET receptor blocker reduced (in a dose-dependent manner) PD fluid-induced peritoneal thickening and preserved the mesothelium (Figure 1B).
ET-1 Promotes a Mesenchymal Phenotype in a TGF-β1-Dependent Manner in Human Mesothelial Cells

To gain insight into the contribution of ET-1 to peritoneal fibrosis and damage, we investigated whether ET-1 induces MMT in vitro. For that purpose, mesothelial transdifferentiation as monitored by the presence of the mesenchymal differentiation marker α-SMA was analyzed using immunofluorescence in human omentum mesothelial cells incubated with increasing doses of ET-1 ranging from 1 to 100 nM. As shown in Figure 4, both ET-1 and TGF-β1 induced dose-dependent increases in the expression of α-SMA. Additionally, we infected mesothelial cells with adenoviruses for the overexpression of ET-1 and TGF-β1. Figure 4 also shows that, similar to the ligands, both adenoviruses upregulated the expression of α-SMA.

We then examined whether there exists a reciprocal regulation in the actions of ET-1 and TGF-β1 on the induction of α-SMA expression in mesothelial cells by using the ET receptor antagonist bosentan and the TGF-β1–blocking peptide P17. As shown in Figure 5, whereas bosentan was able to inhibit the action of ET-1 but not that of TGF-β1, P17 peptide blocked the responses induced by both stimuli. These results indicate that TGF-β1 is involved in the action of ET-1 on α-SMA expression.

We then investigated whether ET-1 is able to promote enhanced expression of TGF-β1. For that purpose, human mesothelial Met5A cells were infected with adenoviruses overexpressing ET-1. As shown in Figure 6A, ET-1 overexpression induced significant increases in TGF-β1 mRNA levels, which...
were associated with augmented accumulation of active TGF-β1 in the cell supernatants as assessed with the mink lung epithelial TGF-β reporter cell line.42 We have also studied whether ET-1 expression is regulated by TGF-β1 in Met5A cells. Figure 6B shows that treatment with the growth factor induced the expression of ET-1 mRNA and the accumulation of ET-1 peptide in the extracellular medium, thus indicating that a reciprocal relationship indeed exists for the control of the expression of ET-1.

**DISCUSSION**

Peritoneal fibrosis is invariably observed in patients undergoing long-term PD. Exposure to PD fluids, which contain high concentrations of glucose and glucose degradation products, damages the peritoneum, inducing loss of mesothelial cells and enlargement of the submesothelial compact zone caused by interstitial fibrosis accompanied by changes in the structure and number of blood vessels.3,8,9 These pathologic changes are closely associated with an increased peritoneal transport rate and loss of ultrafiltration capacity. In addition to a significant improvement in the biocompatibility of PD fluids, specific aims of this research on PD include the identification and characterization of factors contributing to peritoneal fibrosis. Among a number of different factors, TGF-β1 is considered the master molecule in the development of peritoneal dysfunction because of its capacity to induce MMT both *in vitro* and *in vivo*.16,18–20,23 Nevertheless, from a clinical perspective, the blockade of TGF-β1 signaling is a double-edged sword, because it plays important roles in the immune and inflammatory responses. TGF-β1–regulated genes, such as ET-1, represent, therefore, potential targets for the study of their involvement in the pathologic process. In this work, we describe that the expression of the main components of the ET system, ET-1 and its receptors ET<sub>α</sub> and ET<sub>β</sub>, which are restricted to the mesothelial monolayer of the peritoneal membrane under control conditions, was significantly upregulated in the submesothelial thickened zone on exposure to PD fluid. These results suggest that mesothelial cells undergoing MMT might constitute an active source of cells contributing to enhanced ET signaling. One limitation of our study is that the antibodies did not allow for performing of double immunofluorescence, therefore precluding the precise identification of cells expressing ET-1/ET receptors. We also show here that the blockade of ET signaling by dual ET<sub>α</sub>/ET<sub>β</sub> receptor antagonists significantly ameliorated fibrosis and angiogenesis, reduced the number of cells undergoing MMT, and as a result, improved peritoneal function in the mouse model of exposure to PD fluids. Finally, we show that increased ET-1 expression by adenoviral infection induced MMT in human mesothelial cells *in vitro* and when applied directly into the...
peritoneal cavity of mice, promoted the early cellular events associated with peritoneal damage on exposure to PD fluid, namely MMT and fibrocyte recruitment. In contrast to the chronic exposure to PD fluid or overexpression of TGF-β1, ET-1 did not promote increased angiogenesis, likely indicating a late-onset, low-magnitude response of the peritoneal membrane to this factor. Interestingly, these actions of ET-1 did not promote increased angiogenesis, likely because of its important modulating functions of the immune and inflammatory responses. Gene-tailored approaches proven to have an effect on TGF-β1 expression and/or activity directly in the peritoneum might represent options for the clinical intervention. On the basis of the results of this work, the blockade of the ET-1 signaling by the use of specific antagonists may fulfill this requirement. The addition of ET receptor antagonists to the PD fluid may, therefore, provide protection from the deterioration of the mesothelial membrane. To this respect, bosentan (Tracleer) is currently indicated for the treatment of pulmonary arterial hypertension and has shown its anti-inflammatory properties in reducing the number of new digital ulcers in patients with systemic sclerosis, a prototypic fibrotic disorder.37–49 However, macitentan (Opsumit) has been developed as an improved dual ET receptor antagonist that has been recently approved for the treatment of pulmonary arterial hypertension and is also under study for systemic sclerosis-associated digital ulcers.39,50 In vitro, macitentan has been recently described to reduce the profibrotic response of dermal fibroblasts from systemic sclerosis patients.51,52 It should be, however, noted that these drugs have shown significant side effects that have so far hampered their use as a general anti-fibrotic therapy.29 This fact, together with the observation that PD fluid-associated complications develop in the long term, raises concerns about the widespread use of these drugs against functional deterioration of the peritoneal membrane. It is, therefore, mandatory to carefully design and perform a corresponding clinical trial, which as a first approach, may be focused on a subset of renal patients already displaying signs of membrane deterioration. Additional studies are, thus, needed to unequivocally show whether these drugs may be safely translated to the clinical practice of patients receiving PD.

Figure 3. Treatment with ET receptor antagonists decreases the number of activated fibroblasts derived from mesothelial cells and bone marrow-recruited cells and reduced angiogenesis. Mice received through a peritoneal access port a daily 2-ml instillation of either saline (control) or standard PD fluid (PDF) containing bosentan (Bos; 50 mg/kg per day) or macitentan (Mac; 10 mg/kg per day) for 5 weeks. Peritoneal samples were prepared and processed for immunofluorescence studies. (A) Two-color immunofluorescence analyses with anti-FSP1 (fibroblast marker) antibody together with anti-cytokeratin (mesothelial marker) and CD45 (leukocyte marker) to estimate cells undergoing MMT (Cyto+/FSP1+; quantified in B) and recruited fibrocytes (CD45+/FSP1+; quantified in C) show that ET receptor antagonists significantly reduced the number of activated fibroblasts. In addition, the blockade of ET signaling decreased PD-induced angiogenesis as assessed by CD31 (endothelial marker) immunofluorescence (quantified in D). Cell counting was restricted to the submesothelial thickened zone. Box plots represent the median, minimum, and maximum values as well as the 25th and 75th percentiles (n=10). *P<0.005 versus PDF; †P<0.01 versus PDF; ‡P<0.05 versus PDF.

From a clinical perspective, together with the development of more bio compatible fluids that better preserve the mesothelial cell monolayer, the specific blockade of the TGF-β1 signal transduction pathway may provide an interesting therapeutic approach. Nevertheless, the use of TGF-β1–blocking small molecules is unacceptable because of its important modulating functions of the immune and inflammatory responses. Gene-tailored approaches proven to have an effect on TGF-β1 expression and/or activity directly in the peritoneum might represent options for the clinical intervention. On the basis of the results of this work, the blockade of the ET-1 signaling by the use of specific antagonists may fulfill this requirement. The addition of ET receptor antagonists to the PD fluid may, therefore, provide protection from the deterioration of the mesothelial membrane. To this respect, bosentan (Tracleer) is currently indicated for the treatment of pulmonary arterial hypertension and has shown its anti-inflammatory properties in reducing the number of new digital ulcers in patients with systemic sclerosis, a prototypic fibrotic disorder.37–49 However, macitentan (Opsumit) has been developed as an improved dual ET receptor antagonist that has been recently approved for the treatment of pulmonary arterial hypertension and is also under study for systemic sclerosis-associated digital ulcers.39,50
In conclusion, we report here that ET-1 plays an important role as an inducer of the process of MMT and subsequent peritoneal damage and fibrosis and show a causal role for TGF-β1 in the actions of ET-1 on the peritoneum. Our observations that the administration of ET receptor blockers significantly reduced the peritoneal thickness and preserved the mesothelium and the filtration function in the mouse model of PD fluid exposure also support the notion that ET signaling antagonism may represent a useful strategy for the treatment of PD complications.

**COMPLETE METHODS**

**Reagents**

Dual-type A/B ET receptor blockers bosentan (Tracleer) and macitentan (Opsumit) were provided by Martine Clozel (Actelion Pharmaceuticals Ltd., Allschwil, Switzerland). Control and ET-1-overexpressing adenoviruses were from Vector Biolabs (Philadelphia, PA). The adenoviral vector expressing active TGF-β1 was provided by David Dichek (University of Washington, Seattle, WA) and has been previously described. TGF-β1–blocking peptides P17, a soluble hydrophilic peptide derived from a phage display peptide library (KRIWFIPRSSWYERA), and P144, a hydrophobic peptide derived from the sequence of the extracellular region of type III receptor for TGF-β (amino acids 730–743 from β-glycan; TSLDASHWMMQN), as well as corresponding control peptides were from Digna Biotech (Madrid, Spain). These peptides have shown an antagonist effect over TGF-β–dependent processes in cellular cultures and a strong TGF-β inhibitory effect in different animal models.23,55–57

**PD Fluid Exposure Model in Mice**

Female C57BL/6 mice between 12 and 16 weeks of age were used in this study (Harlan Interfauna Iberica, Barcelona, Spain). The experimental protocol used was in accordance with the National Institutes of Health Guide for Care and Use of Laboratory Animals and approved by the Animal Ethics Committee of the Centro de Biología Molecular "Severo Ochoa."
cell line Met5A was obtained from the American Type Culture Collection (Manassas, VA).

Omentum-derived and Met5A cells were treated for 24 hours with ET-1 (1–100 nM), TGF-β1 (0.25–10 ng/ml), or ET-1 and active TGF-β1—overexpressing adenoviruses in the presence or absence of the ET receptor antagonist bosentan (10 μM), TGF-β1—blocking peptide P17 (150 μg/ml), or its corresponding control peptide. Omentum-derived mesothelial cells were used for immunofluorescence analysis of the expression of the mesenchymal marker α-SMA. Cells were fixed in 4% formaldehyde in PBS for 10 minutes and then permeabilized with 0.2% Triton X-100 in PBS for 5 minutes at room temperature. Cells were washed with PBS, blocked with 1% BSA in PBS for 1 hour, and then incubated overnight at 4°C with an anti–α-SMA antibody (Master Diagnostica, Granada, Spain) followed by the corresponding fluorescent secondary antibody. Cell fluorescence was visualized by microscopy with a Nikon Eclipse T2000U (Nikon, Amstelveen, The Netherlands). Images were analyzed by computerized digital image analysis (NIS-Elements Imaging Software; Nikon).

Experiments requiring a higher number of cells, such as ET-1 and TGF-β determinations and RT-PCR analysis, were done with Met5A cells. For RT-PCR studies, total RNA was isolated by guanidium thiocyanate/phenol/chloroform extraction, cDNA was synthesized using an iScript cDNA synthesis kit, and quantitative PCR for the detection of TGF-β1 mRNA expression was performed using iQ SYBR green Supermix and primers as previously described (Bio-Rad, Hercules, CA). Sequences of primers were as follows: human TGF-β1 forward: ACCTGAACCCGTGTTGCTCT; human TGF-β1 reverse:
Overexpression of TGF-β1 and ET-1 in the Peritoneum by Adenovirus-Mediated Gene Transfer

For *in vivo* experiments involving the overexpression of ET-1 and TGF-β1, mice were intraperitoneally injected with 100 μl suspensions containing 1×10⁶ plaque-forming units of the corresponding adenoviruses as previously described. Animals treated with P144 peptide or its corresponding control peptide (4 mg/kg per day diluted in 1 ml saline solution) were injected intraperitoneally every day (with the exception of the days that adenovirus was applied). Four or twelve days after the injection, animals were euthanized, and the parietal peritoneum was collected for immunofluorescence analysis.

Statistical Analyses

Results are presented as 25th and 75th percentiles; median, minimum, and maximum values in box plot graphs; or means±SEMs in bar graphs. The data groups were compared with the nonparametric Mann–Whitney rank sum *U* test and Wilcoxon signed rank test for *in vivo and in vitro* studies, respectively (Prism version 4.0; GraphPad, La Jolla, CA). *P*<0.05 was considered statistically significant.

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DISCLOSURES

Actelion Pharmaceuticals Ltd. supplied the endothelin receptor antagonists bosentan and macitentan. Actelion Pharmaceuticals Ltd. had no role in the study design, collection, analysis, or interpretation of the data, or writing the manuscript. Actelion Pharmaceuticals Ltd. has not revised the content of the submitted manuscript or made decisions on the approval of the publication of the article. J.D. is an employee of Digna Biotech.
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