Comparison of endothelin receptors in normal versus cirrhotic human liver and in the liver from endothelial cell-specific ETB knockout mice

Citation for published version:

Digital Object Identifier (DOI):
10.1016/j.lfs.2012.02.003

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Publisher's PDF, also known as Version of record

Published In:
Life Sciences

Publisher Rights Statement:
Available under Open Access

General rights
Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.
Comparison of endothelin receptors in normal versus cirrhotic human liver and in the liver from endothelial cell-specific ET\(_B\) knockout mice

Lowell Ling \(^a\), Rhoda E. Kuc \(^a\), Janet J. Maguire \(^a\), Neil J. Davie \(^b\), David J. Webb \(^c\), Paul Gibbs \(^d\), Graeme J.M. Alexander \(^e\), Anthony P. Davenport \(^a,⁎\)

\(^a\) Clinical Pharmacology Unit, Box 110 Addenbrooke's Hospital, Cambridge CB2 0QQ, United Kingdom
\(^b\) Pfizer Ltd., Sandwich, Kent, United Kingdom
\(^c\) University of Edinburgh, Queen's Medical Research Institute, Edinburgh EH16 4TJ, United Kingdom
\(^d\) Department of Surgery, Addenbrooke's NHS Trust, United Kingdom
\(^e\) Department of Medicine, Addenbrooke's NHS Trust, United Kingdom

**ABSTRACT**

**Aims:** Endothelin (ET) antagonists show promise in animal models of cirrhosis and portal hypertension. The aim was to pharmacologically characterise the expression of endothelin receptors in human liver, hepatic artery and portal vein.

**Main methods:** Immunofluorescence staining, receptor autoradiography and competition binding assays were used to localise and quantify ET receptors on hepatic parenchyma, hepatic artery and portal vein in human cirrhotic or normal liver. Additional experiments were performed to determine the affinity and selectivity of ET antagonists for liver ET endothelin receptors. An endothelial cell ET\(_A\) knockout murine model was used to examine the function of sinusoid endothelial ET\(_B\) receptors.

**Key findings:** ET\(_A\) receptors predominated in normal human liver and displayed the highest ratio (ET\(_B\):ET\(_A\) 63:47) compared with other peripheral tissues. In two patients examined, liver ET\(_A\) expression was up-regulated in cirrhosis (ET\(_B\):ET\(_A\) 83:17). Both sub-types localised to the media of normal portal vein but ET\(_B\) receptors were downregulated fivefold in the media of cirrhotic portal vein. Sinusoid diameter was fourfold smaller in endothelial cell ET\(_B\) knockout mice. The liver morphology of ET\(_B\) knockout mice was markedly different to normal murine liver, with loss of the wide spread sinusoidal pattern. In the knockout mice, sinusoids were reduced in both number and absolute diameter, while large intrahepatic veins were congested with red blood cells.

**Significance:** These data support a role for the ET system in cirrhosis of the liver and suggest that endothelial ET\(_B\) blockade may cause sinusoidal constriction which may contribute to hepatotoxicity associated with some endothelin antagonists.

© 2012 Elsevier Inc. All rights reserved.

**Introduction**

Current medical treatment of portal hypertension (PH) is inadequate and patient mortality remains high resulting from complications of chronic liver disease, including variceal haemorrhage. The aim of therapy and prophylaxis is to lower portal pressure to levels that reduce the risk of variceal haemorrhage and death. Meta-analysis has demonstrated that a portal pressure of less than 12 mmHg or a greater than 20% reduction from baseline portal pressure reduces variceal bleeding and mortality (D’Amico et al., 2006). Current pharmacological interventions, including beta-blockade, aim to reduce splanchnic flow and portal pressure, but 30–40% of patients either do not respond or target reduction in portal pressure is not achieved (Garcia-Tsao et al., 1986; Merkel et al., 2000). Liver transplantation is an effective treatment for both cirrhosis and PH, but the demand for donor organs continues to outstrip supply. With an increasing prevalence of chronic liver disease and limited treatment options, there is clinical urgency for new medical therapy.

A role for the endothelin (ET) system in liver cirrhosis has been suggested by clinical studies that demonstrated a correlation between the severity of cirrhosis and elevated plasma endothelin-1 (ET-1) levels (Tsai et al., 1995; Uchihara et al., 1992). ET-1 mediates its actions via two receptors, ET\(_A\) and ET\(_B\) (Davenport, 2002; Davenport and Maguire, 2006) and in healthy rat liver, both sub-types are concentrated along sinusoids, mainly on hepatic stellate cells and sinusoidal endothelial cells (Gondo et al., 1993; Houssset et al., 1993). In rats with cirrhosis, expression of both receptors on hepatic stellate cells was reported to be up-regulated, but receptor...
expression on sinusoidal endothelial cells was unchanged (Yokomori et al., 2001a). Using immunohistochemistry, ETb expression was also reported to be up-regulated in human cirrhotic liver, with low ETb expression detected in both normal and diseased liver (Yokomori et al., 2001b). Endothelin may also be an important regulator of sinusoid vascular resistance mediated by vasoconstriction of stellate cells, the main effector of sinusoid calibre (Rockey and Weissiger, 1996). In animal studies infusion of ET-1 caused a dose-dependent decrease in sinusoid diameter (Okumura et al., 1994; Zhang et al., 1994) and both selective ETb and mixed antagonists were shown to decrease portal pressure (Feng et al., 2009; Sogni et al., 1998). Additionally, ET antagonists may also decrease the fibrotic response in cirrhosis (Khimji and Rockey, 2011).

Our aim was to determine the distribution and density of ET receptors in human normal and cirrhotic liver parenchyma, hepatic artery and portal vein. Immunofluorescence staining and autoradiography were used to localise and quantify ET receptors on normal and diseased hepatic artery and portal vein. Receptor autoradiography and competition binding assays were used to quantify ET receptors and to determine the binding characteristics of ET antagonists in human liver. Lastly, an endothelial cell specific ETb knockout murine model was used to investigate the consequence of loss of ETb receptors to provide evidence that selective ETb antagonism may be the treatment of choice in cirrhosis and PH.

Materials and methods

BQ123, cyclo-[d-Asp-L-Pro-d-Val-L-Leu-d-Trp-], BQ788, (N-cis-2,6-dimethylpiperidinocarbonyl-L-γ-MeLeu-d-Trp(CooMe)-d-Nle-ONa) and BQ3020, [Ala11, 15]Ac-ET-16-21 were synthesised by solid phase t-Boc chemistry. [125I]-ET-1 was from PerkinElmer, while unlabelled ET-1 was from Peptide Institute. Ambisentan and sitaxentan were from Gilead Sciences, Inc and Pfizer, Inc, respectively. ETa and ETb rabbit antiserum were raised to the C-terminus of ETa(413–427) and to ETb(302–313) (Davenport and Kuc, 2005a). Primary goat anti-von Willebrand factor and anti-smooth muscle α-actin were from Dako; secondary fluorescent antibodies were from Invitrogen.

Human tissue samples

Tissue samples collected at the time of operation were obtained with ethical approval (REC 10/H0305) and informed consent. Diseased liver parenchyma, hepatic artery and portal vein were obtained from patients undergoing liver transplantation for non-infectious causes of end-stage liver disease. Normal liver parenchyma, hepatic artery and portal vein were obtained from donor liver and vessels that were not used for transplantation surgery or were normal tissue from liver resections. Unspecified otherwise n-values refer to the number of patients from whom tissue was obtained.

Endothelial cell specific ETb knockout mice

Mice were generated using the Cre-loxP system (Bagnall et al., 2006). After euthanasia, consecutive, cryostat-cut frozen sections (10 μm) of knock-out and control mouse torsos were mounted onto gelatine coated glass microscope slides. Sinusoid diameter in ETb knockout and control mice were measured under light microscopy. A two-sided unpaired t-test was used to compare changes in sinusoid number and between control and knock-out mice, with significance set at P<0.05.

Dual-labelled immunofluorescence microscopy

Methods were as previously described (Davenport and Kuc, 2005a). Briefly, tissue sections (10 μm) were dried overnight at room temperature and fixed in ice-cold acetone for 10 minutes. Slides were incubated with 5% non-immunised donkey serum (DS) in phosphate-buffered saline (PBS) for 1 hour at room temperature to block non-specific protein interactions and then incubated overnight at 4 °C with primary rabbit anti-ETa (1:50) or anti-ETb (1:50) antiserum and either primary goat anti-von Willebrand factor (1:100) or goat anti-smooth muscle α-actin (1:100) antibody diluted in 1% PBS/0.1% Tween-20/3% DS. Slides were then washed (3 × 5 minutes) in cold 1% PBS/0.1% Tween-20 before incubation for 1 hour at room temperature with Alexa Fluor 488 conjugated donkey anti-rabbit (1:200), Alexa Fluor 568 conjugated donkey anti-goat (1:100) secondary antibodies and Hoechst (1:100) diluted in 1% PBS/0.1% Tween-20/3% DS. Tissue sections were washed again (3 × 5 minutes) in cold 1% PBS/0.1% Tween-20 and mounted with ProLong Gold (Invitrogen). Confocal imaging was performed using a Leica TCS-NT-UV confocal laser-scanning microscope (Leica Microsystems, Heidelberg, Germany).

Quantitative autoradiography

Adjacent 10 μm tissue sections were incubated with HEPES buffer (50 mM HEPES, 5 mM MgCl2, 0.3% bovine serum albumin, pH 7.4) for 20 minutes at room temperature. Sections were incubated for 2 hours at room temperature with [125I]-ET-1 (0.1 nM) alone for total binding and either with unlabelled ET-1 (1 μM) to determine non-specific binding, with BQ3020 (100 nM) to label ETa receptors or with BQ123 (100 nM) to label ETb receptors (Molenaar et al., 1993). Sections were washed (3 × 5 minutes) in ice-cold Tris–HCl buffer (50 mM, pH 7.4), air dried and apposed, together with [125I]-ET-1 standards, to Kodak MR-1 autoradiography film for 2 days at room temperature. The resulting autoradiographs were analysed using computer-assisted densitometry (Quantimet 970, Leica, Milton Keynes, UK) (Davenport and Kuc, 2005b) and receptor densities expressed in amol/mm².

Competition binding assays

Adjacent 10 μm tissue sections were incubated with HEPES buffer (50 mM HEPES, 5 mM MgCl2, 0.3% bovine serum albumin, pH 7.4) for 20 minutes at room temperature and then for 2 hours at room temperature with [125I]-ET-1 (0.1 nM) and either sitaxentan (20 pM to 200 nM), BQ788 (20 pM to 200 nM) or ambisentan (20 pM to 200 nM). Non-specific binding was determined by inclusion of 1 μM ET-1. Sections were washed (3 × 5 minutes) in ice-cold Tris–HCl buffer (50 mM, pH 7.4) and the amount of tissue bound [125I]-ET-1 measured by gamma counting. Data were analysed using EBDA software (McPherson, 1983) to provide initial estimates of equilibrium dissociation constant (Kd) and maximum binding density (Bmax) and LIGAND software (Munson and Rodbard, 1980) to determine final estimates based on the F-ratio test (P<0.05) of 1, 2 or 3 site models. The Bmax was normalised to protein concentration (Davenport and Kuc, 2005b). Results were expressed as the mean ± standard error of the mean.

Results

Dual-labelled immunofluorescence microscopy

Confocal photomicrographs of a transverse section through a human portal vein from a patient with cirrhosis are shown in Fig. 1A and B. Antiserum to ETa (Fig. 1A) was visualised as green fluorescence and a second antiserum to the smooth muscle marker α-actin was visualised in red. Co-localization of both antiserum within the medial and intimal smooth muscle layers of portal vein is shown in yellow. In Fig. 1B, antiser to ETb was visualised as green fluorescence and α-actin in red, showing that both sub-types were detectable on smooth muscle. ETa and ETb immunoreactivity was also localised to the adventitial layer of
both vessels. Results at the cellular level were supported by radioligand binding using $^{125}$I-ET-1 where the distribution of ETA (C) and ETB (D) receptors is shown by the blackening of autoradiography film. Adjacent sections stained with haematoxylin and eosin were used to delineate the endothelium (red arrow), media and adventitia.

Confocal photomicrographs (Fig. 2) illustrating the localization of antisera to ETA (A, shown in green) the endothelial cell marker vWF (B), shown in red to visualise the single layer of cells in the vascular endothelium in a transverse section of human portal vein from a patient with cirrhosis. Digitally overlaid confocal photomicrographs (C) illustrating the co-localization are shown in yellow. Corresponding images of ETb immunoreactivity are shown in (Fig. 2D), vWF (Fig. 2E) and the digital overlay demonstrating co-localization in Fig. 2F.

**Autoradiography**

In normal liver parenchyma, there was a homogenous distribution of ET receptors within the tissue, measured by radioligand binding and quantitative autoradiography. Total ET receptor density measured using $^{125}$I-ET-1 which binds with the same affinity to both sub-types was $152 \pm 11$ amol/mm$^2$ with ETA receptors comprising $41 \pm 6$ amol/mm$^2$ (37 ± 5%) and ETB receptors $81 \pm 23$ amol/mm$^2$, (63 ± 5%) (Fig. 3A and B). In preliminary studies using tissue from two patients with cirrhosis the total receptor density was reduced to about 100 amol/mm$^2$ and the ratio of ETA:ETB was 17:83. Therefore,
in liver parenchyma, cirrhosis was associated with a downregulation of ET\textsubscript{A} (19 amol/mm\textsuperscript{−2}) and small increase in ET\textsubscript{B} (96 amol/mm\textsuperscript{−2}) receptor density (Fig. 3A and B). Cirrhotic liver parenchyma had a characteristic nodular pattern with ET\textsubscript{A} localised mainly in the fibrous septa and at lower levels within lobules. Similarly, ET\textsubscript{B} was also concentrated along the fibrous connective tissue between lobules in the diseased liver, but ET\textsubscript{B} binding was also high within the lobules (Fig. 4).

Autoradiographs revealed the expression of ET\textsubscript{A} and ET\textsubscript{B} receptors determined using binding of \textsuperscript{125}I-ET-1 in portal vein (Fig. 1C and D) and hepatic artery were consistent with that obtained by immunohistochemistry. In normal portal vein ET receptor density was 149±25 amol/mm\textsuperscript{2} in the medial smooth muscle layer with an ET\textsubscript{A}:ET\textsubscript{B} ratio of 57:43 (96±15 amol/mm\textsuperscript{2} ET\textsubscript{A} and 70±8 amol/mm\textsuperscript{2} ET\textsubscript{B}). In portal vein from patients with cirrhosis the overall density of ET receptors was lower (131 amol/mm\textsuperscript{2}) but the ratio of ET\textsubscript{A}:ET\textsubscript{B} was 90:10. Therefore, in contrast to parenchyma, in portal vein there was no change in medial ET\textsubscript{A} receptor expression but there was a downregulation of ET\textsubscript{B} receptors (Fig. 3C and D).

**Competition binding**

The results of the ability of the three antagonists, BQ788, sitaxentan and ambrisentan to compete for the binding of \textsuperscript{125}I-ET-1 are shown in Fig. 5. Pooling data from competition binding curves for the three antagonists, BQ788, sitaxentan and ambrisentan (Fig. 5) total ET receptor density were not different in cirrhotic liver (150.0±22.7 fmol/mg) compared to normal liver (125.6±12.2 fmol/mg). The ET\textsubscript{B} receptor comprised the predominant subtype in normal liver and the proportion of ET\textsubscript{B} was increased further in disease with a shift in ET\textsubscript{A}:ET\textsubscript{B} ratio from 20:80 in normal tissue to 5:95 with cirrhosis. As expected, the ET\textsubscript{B} selective antagonist, BQ788 exhibited high affinity for the ET\textsubscript{B} receptor in both normal (K\textsubscript{D} 37±8 nM) and diseased (133±34 nM) liver with lower micromolar affinity for the ET\textsubscript{A} receptor (normal ET\textsubscript{A} K\textsubscript{D} 65±88 μM; cirrhosis ET\textsubscript{A} K\textsubscript{D} 46±108 μM). Since the majority of the ET\textsubscript{B} receptors in liver were of the ET\textsubscript{B} subtype, the density of ET\textsubscript{A} receptors in normal and particularly cirrhotic liver was too low to derive an accurate affinity constant at the ET\textsubscript{A} receptor for sitaxentan and ambrisentan. However, from the competition curves it
was clear that both compounds competed for the ET<sub>B</sub> receptors in the low micromolar range.

**Endothelial cell specific ET<sub>B</sub> knockout mice**

Livers from endothelial cell ET<sub>B</sub> knockout mice were examined and compared to normal murine liver under light microscopy (Fig. 6). The liver morphology of ET<sub>B</sub> knockout mice was markedly different to normal murine liver, with loss of the wide spread sinusoidal pattern seen in normal livers. In the knockout mice, sinusoids were reduced in both number and absolute diameter, while large intrahepatic veins were congested with red blood cells. Sinusoidal diameter was reduced from 22.3±1.5 μm in normal murine liver to 6.0±0.4 μm in knockout mice (P=0.0004).

**Discussion**

Studies on the ET pathway in cirrhosis have thus far focused on the liver parenchyma itself (Ikura et al., 2004). This study characterises ET receptors for the first time in human portal vein. Consistent with studies on other human vessels, we found that smooth muscle cell ET<sub>A</sub> is the main receptor subtype in the media of hepatic artery and portal vein (Bacon and Davenport, 1996; Davenport et al., 1995). As there was a six fold decrease in ET<sub>B</sub> density in portal vein media, the ET<sub>A</sub>: ET<sub>B</sub> ratio in portal vein media changed from 57:43 in health to 90:10 in cirrhosis. It is possible that elevated plasma ET-1 in cirrhosis coupled with relatively high ET<sub>A</sub> expression in portal vein media may increase pre-hepatic portal resistance and cause PH. Therefore, ET<sub>A</sub> antagonism may be beneficial in dilating the portal vein in PH. It should be emphasised that these are preliminary descriptive studies and the results justify further experiments to determine reproducibility in a larger number of individuals.

Immunofluorescence staining for ET<sub>A</sub> and ET<sub>B</sub> was found along the endothelium of normal and cirrhosis hepatic artery and portal vein. While it is well established that ETB is present on the endothelium of human vessels, endothelial ET<sub>A</sub> is less well documented (Nishimura et al., 1995). Endothelial ET<sub>A</sub> mediates NO release from the endothelium, and is an important counter mechanism to ET-1 induced vasoconstriction on smooth muscle ET<sub>A</sub> (Hirata et al., 1993; Honoré et al., 2002). Moreover, endothelial ET<sub>B</sub> functions as clearing receptors that remove excess ET-1 from the circulation (Johnstrom...
et al., 2005; Kelland et al., 2010). While ET\textsubscript{A} antagonism may be beneficial in PH, whether ET\textsubscript{B} antagonism is also helpful is less straightforward, as the opposing functions of endothelial ET\textsubscript{B} (dilatation and reducing plasma ET-1) and smooth muscle ET\textsubscript{B} (constriction) complicate therapeutic choice. Nevertheless, animal studies showed that mixed antagonists such as bosentan are as effective as ET\textsubscript{A} selective antagonists in reducing portal pressure (Feng et al., 2009; Rockey and Weisiger, 1996). The choice between selective ET\textsubscript{A} and mixed antagonism continues to be debated in the treatment of pulmonary arterial hypertension (Vachiery and Davenport, 2009) and renal failure (Davenport and Maguire, 2011).

Interestingly sinusoid diameter was reduced three-fold in the knockout compared to wild type mice and intrahepatic branches of the portal vein were congested with red blood cells (Fig. 6). This suggests that blood was unable to pass through sinusoids effectively owing to reduced sinusoidal diameter and produced congestion in the portal venous system. Hence, sinusoidal endothelial cell ET\textsubscript{B} may be critical in maintaining adequate sinusoidal diameter. Therefore, ET\textsubscript{B} blockade which causes sinusoid constriction and consequent liver ischaemia may contribute to the liver toxicity observed with mixed antagonists. This suggests additional benefit of ET\textsubscript{A} selective compounds.

This study has identified changes in ET receptor ratio and density within human cirrhotic liver and portal vein. ET receptor densities from quantitative autoradiography and competition binding experiments are consistent with those suggested by previous studies on the human liver (Yokomori et al., 2001b). Both techniques demonstrated a change towards upregulation of ET\textsubscript{B} and downregulation of ET\textsubscript{A} in the cirrhotic liver. The significance of these changes is underdetermined, but it may be a physiological response towards normalising elevated portal pressure in cirrhosis. Adding the results from this study to data from animal models, selective ET\textsubscript{A} antagonism may be a beneficial therapy for cirrhosis and PH by four distinct mechanisms. First, it may reduce prehepatic portal resistance by dilatation of the portal vein. Second, it may decrease intrahepatic resistance by reducing hepatic stellate cells contraction and increasing sinusoid diameter. Third, it may also reduce intrahepatic resistance by reversing fibrosis and restoring normal liver architecture. Fourth, it preserves the beneficial sinusoidal endothelial cell ET\textsubscript{B} mediated dilatation and clearance of elevated plasma ET-1. Further in vitro studies using human tissues are required with antagonists of the propionic acid class of compounds (such as ambrisentan) that do not display significant hepatotoxicity but as our results show are ET\textsubscript{A} selective in human liver, for the treatment of cirrhosis.

**Conflict of interest statement**

N.J.D. was an employee of Pfizer. The research was supported in part by an investigator led grant from Pfizer.

**Acknowledgements**

This study was supported by an investigator led grant from Pfizer. J.J.M. was supported by the British Heart Foundation (PC/09/050/27734). L.L. was part of the Wellcome Trust Translational Medicine and Therapeutics Programme. The authors would like to thank the NIHR Cambridge Biomedical Research Centre, Pfizer for providing sitaxentan and Gilead Sciences for ambrisentan.

**References**


Davenport AP, Kuc RE. Radioligand-binding and molecular-imaging techniques for the quantitative analysis of established and emerging orphan receptor systems. Methods Mol Biol 2005b;306:93–120.


