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Expression of receptors for verotoxin 1 from Escherichia coli O157 on bovine intestinal epithelium

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Human enterohaemorrhagic Escherichia coli (EHEC) infection most commonly arises, either directly or indirectly, from cattle, which act as a reservoir host for these bacteria. In man, EHEC disease can be severe, whereas EHEC do not normally cause disease in cattle. Verotoxins (VTs) are the main virulence factors in human disease but no role for VT has been ascribed in cattle; however, this study shows for the first time that VT receptor is expressed by the bovine intestinal tract. VT bound to crypt epithelial cells of the small (ileum and jejunum) and large (caecum and colon) intestine independently of the animals’ age. VT also bound to discrete cell subsets in the bovine kidney and to submucosal lymphoid cells but not to vasculature. Analysis of tissues for isoforms of the VT receptor, Gb3, confirmed the presence of the receptor in the bovine intestinal epithelium and kidney. A distinct pattern of Gb3 receptor isoform mixtures was observed in the bovine kidney. This, together with the general absence of receptors on vasculature, could contribute to the apparent resistance of cattle to systemic effects of VT. Expression of Gb3 on the bovine intestinal epithelium, together with previously described effects, may affect EHEC colonisation in its reservoir hosts and hence the potential for distribution to man.

Introduction

Enterohaemorrhagic Escherichia coli (EHEC) are a subclass of verotoxigenic E. coli (VTEC) [1], among which those belonging to serogroup O157 have been successful in the widespread colonisation of cattle [2, 3]. E. coli O157 has emerged as a significant cause of severe food-borne disease in man [4] with outbreaks occurring regularly [5–8]. Although E. coli O157 can cause severe diseases in man (e.g., haemorrhagic colitis and haemolytic uraemic syndrome), infection in cattle is normally asymptomatic [2, 3, 9]. It is possible that the lack of disease in cattle may have aided its spread; however, the basis for the success of EHEC in colonising cattle and their asymptomatic carriage remains undefined.

Multiple factors are involved in EHEC virulence in man. These include Esp proteins (encoded by locus of enterocyte effacement; LEE) [10], intimin and Tir (also encoded by LEE) [11], plasmid-encoded determinants [12] and verotoxins ( VTs), which are considered to be responsible for the severe damage in EHEC infection in man. VTs represent a family of bacterial compound toxins, the prototype of which is present in Shigella dysenteriae type 1 [13]. The holotoxin (c. 70 kDa) comprises a single 32-kDa A subunit and five B subunits arranged as a pentamer [14]. In VTEC there are two main classes of verotoxins (VT1 and VT2) which are encoded on lysogenic bacteriophages that can be carried by strains in various combinations [15]. Targeting of VT and susceptibility is dictated primarily by expression of the globotriaosyl-ceramide (Gb3) family of glycolipids [16] which have been shown to be necessary in mediating cytotoxicity [17, 18]. VT binds specifically to the galabiose terminal disaccharide moiety of globotriaosyl ceramide-Gb3 (Galβ1-4-Galβ1-4 glucosyl ceramide) [19, 20], also known as CD77 [21]. Gb3 may be distributed in different cell and tissue

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types in different hosts, thus defining the pathology of VTI-induced diseases. Expression of Gb3 in the cell membrane generally reflects sensitivity to the cytotoxic activities of VTs ultimately leading to cell death, for which the following mechanisms have been proposed. The A subunit of VTs catalytically inactivate the 28S RNA of 60S ribosomal subunits leading to an inhibition of cellular protein synthesis [22, 23]. Furthermore, the holotoxin and the B subunit pentamer are capable of inducing programmed cell death (apoptosis) through a pathway independent of the suppression of protein synthesis [24]. VTs can induce necrosis [25], apoptosis [26, 27] or both in several cell types depending on the circumstances. Therefore, current evidence indicates that VT is a potent cytolethal toxin.

*E. coli* O157 has become widely distributed in cattle populations and factors that increase the success of EHEC in colonising cattle consequently increase the threat to human health. The successful colonisation by EHEC of cattle and their apparent resistance to disease caused by EHEC are the subject of much speculation. VT genes are retained in at least 95% of cattle isolates [3, 28, 29], hence VT may confer some selective advantage in this host. VT is produced in the intestine of cattle [30, 31] and, although it does not appear to cause overt damage (as it does in man), may possess other important function(s).

The distribution of VT receptor along with other receptor characteristics is important in determining whether VT may be of significance in the interaction between EHEC and cattle. The present study examined tissues from cattle of two different age groups for VT receptor with a larger sample size than the study reported previously [32]. Bovine tissues were examined in detail for Gb3 by immunohistochemical and biochemical approaches. VT receptors have a novel distribution in cattle, an important finding that is in contrast to the conclusions of the previous study [32]. This suggests that VT may have a role in colonisation of cattle and these possible roles are discussed.

**Materials and methods**

**Preparation of verotoxin**

VT1 was prepared as described previously [20]. After removal of endotoxin by passage through a polymyxin B column, the concentration of VT1 was adjusted to 0.165 mg/ml in PBS and stored in small volumes at −70°C.

**Animals and tissue sampling**

Tissues were obtained from 1- and 10-month-old cattle (n = 3 for both age groups) submitted for routine post-mortem examination (UK) or presented at abattoir (Canada). Necropsy commenced immediately following euthanasia and within 15 min of death duplicate tissue samples (rumen, jejunum, ileum, caecum, colon and kidney) were frozen in liquid nitrogen (for glycolipid extraction) or mounted in OCT (Merck) and frozen in a slurry of dry ice in isopentane (for immunohistochemical detection of VT receptor). Histopathological assessment of haematoxylin and eosin (H&E)-stained sections indicated no pathological alterations in intestinal or renal samples; thus these samples were further processed for Gb3 analysis or immunohistochemical analysis of VT1 binding.

**Immunoperoxidase detection of VT1 binding to bovine tissue**

Sections (6 μm) of frozen autopsy material derived from 1- and 10-month-old cattle (n = 3 for both age groups) were cut with a Leica RM2155 microtome, mounted on to Biobond-coated slides (Biologicals Biocell International) and frozen at −70°C until used. Air-dried frozen sections [33] of rumen, jejunum, ileum, caecum, colon and kidney were blocked sequentially with Peroxidase Block (DAKO), avidin, biotin (15 min each) and finally normal goat serum (NGS) 1% in phosphate-buffered saline (PBS) for 20 min, with extensive washings with PBS between each step. Sections were overlaid with VT1 (50 or 200 ng/ml), mouse anti-VT1 (1 in 250) and biotinylated anti-mouse IgG (1 in 500) (Jackson Laboratories) for 30 min each. VT binding was detected with a Vectastain Elite Universal ABC elite kit (Vector Laboratories) and visualised with 3,3'-diaminobenzidine tetrachloride (DAB; Vector Laboratories); the sections were counter-stained with haematoxylin. Sections were observed with a Leica DMLB epifluorescence microscope under incident illumination.

**Glycolipid extraction from bovine tissues**

Tissues of both 1- and 10-month-old cattle were homogenised (1 g/ml) in PBS and the glycolipids were extracted with 20 volumes of chloroform:methanol (C:M, 2:1 v/v) overnight. A Folch partition against water was performed with each of the extracts [34]. The lower phase was dried and saponified with 1 M NaOH in methanol for 2 h at 37°C to remove phospholipids from the mixture. The dried lower phase was dissolved in C:M 98:2 v/v and separated by silica chromatography according to the method described by Boyd and Lingwood [35]. The Gb3 present was detected by thin-layer chromatography (TLC) overlay binding with VT1.

**Assay of Gb3 content by VT1 TLC overlay**

Samples (20–50 μg/ml) of the glycolipid extracts of bovine tissue were separated by TLC (C:M-water 65:25:4 v/v/v) in parallel with appropriate standards (0.5 μg/ml), i.e., glucosylceramide (GlcCer; Sigma); lactosylceramide (LacCer); Gb3 and Gb4. The plates were dried and blocked with gelatin (1% in water) at
37°C overnight. They were then washed three times with 50 mM Tris-buffered saline (TBS), pH 7.4, for 5 min and incubated with VT1 100 ng/ml for 1 h at room temperature. After further washing with TBS, the plates were incubated with mouse anti-VT1 antibody (2 μg/ml) [36] followed by horseradish peroxidase-conjugated goat anti-mouse IgG (1 in 2000) (DAKO) for 1 h at room temperature. Finally, after washing the plates with TBS, 4-chloro-1-naphthol peroxidase substrate was added to localise toxin binding. A similar plate was prepared and sprayed with orcinol to determine total glycolipid content.

Results

Localisation of VT1 binding in bovine tissues

Frozen tissue sections from the rumen, jejunum, ileum, caecum, colon and kidney from 1- and 10-month old cattle (three of each age) were incubated with purified VT1, either 50 ng/ml or 200 ng/ml. Immunohistological examination of the jejunum, ileum, caecum and colon from both age groups identified VT1 bound to a subpopulation of epithelial cells (Figs. 1 and 2 show representative examples). Specifically, toxin binding localised to the crypts adjacent to the submucosa, a binding pattern that was observed with both concentrations of VT1. Non-specific binding of secondary antibody (with or without binding of primary antibody) to the sections as assessed by incubation without VT1, was not observed in any of the tissues examined (negative control, data not shown).

During histological examination of intestinal tissues from animals of both age groups, binding to subepithelial cells was also observed. VT1 attached to cells present in lymphoid aggregates within the lamina propria, a feature that was particularly evident in the ileum from both age groups (Fig. 2).

The results with kidney sections from 1- and 10-month-old cattle, obtained with a well-established method for detection of the Gb3 receptor [33], show a similar binding pattern to that recently reported with acetone-fixed sections [32]. Although Gb3 could be solubilised from tissue by acetone treatment, as kidney retained receptor in the previous study [32], this treatment is presumably insufficient to remove all the glycolipids from this tissue (in contrast to intestine). Predominant staining was apparent in the convoluted tubules and collecting ducts within the renal cortex, with a complete absence of binding to the renal glomeruli and blood vessels (Fig. 3). VT1 binding to vasculature was not observed in the bovine jejunum, ileum, caecum, colon and kidney (n = 3 for both age groups).

Expression of VT1 receptor

Bovine intestinal samples were further examined for VT receptors by extraction and isolation of glycolipids from mucosa. Glycolipids (20–50 μg/ml) from each tissue extract were analysed by a thin-layer chromatography (TLC) VT overlay assay. Two bands (which are mixtures of isoforms) of Gb3 to which VT bound were identified in enteric (jejunum, ileum, caecum and colon) mucosa. Fig. 4a shows results from caecum and colon which are representative of all intestinal sites. VT bound to glycolipid bands from bovine intestinal mucosa which migrated at the same positions as those in the control Gb3 sample (human adult kidney). The extraction of the Gb3 receptor from these sites correlated with the VT1 binding observed in these tissues by immunohistochemistry.
Kidney from these animals was also examined for the expression of Gb3 receptor. A triplet of bands was observed indicating three isoform mixtures of Gb3 receptor (Fig. 4b). One band of Gb3 in the kidney of both age groups of cattle was absent in the corresponding human tissue. In contrast to man, there were no tissue- and age-related differences in toxin binding in cattle. VT1-binding glycolipids were extracted and quantified in three independent experiments and the above findings are representative of the results obtained.

Discussion

Cattle are the main reservoir of E. coli O157 and other EHEC [37, 38]. In this host, infection is asymptomatic, although in man infection is serious and potentially fatal. Among the many virulence factors of EHEC, verotoxins (VTs) are the main factors responsible for the severe features of disease in man. Cattle are frequently colonised by VT-positive E. coli and as VTs can be detected in cattle faeces [30, 31] these animals are presumably less sensitive to VT toxicity. It has been suggested recently that this refractory nature of cattle may correspond to absence of VT receptor (Gb3) in most cattle tissues [32]; however, findings presented here contradict this suggestion and demonstrate that cattle do express VT receptor in a novel tissue and cellular distribution.

Immunohistochemical analysis of VT binding to tissue sections demonstrated localisation to epithelial cells in the crypt regions adjacent to the submucosa of jejunum, ileum, caecum and colon from both 1- and 10-month old cattle. The presence of receptor in these sites was confirmed by lipid extraction and TLC overlay assay. These findings contrast with a previously published report that bovine intestinal epithelium lacks Gb3 [32]. However, the two studies differ in details of experimental procedures: the age of animals selected for investigation, tissue quantities and glycolipid isolation protocols and the preparation of tissue for immunohistochemistry. Notably, the present study used animals aged 1 and 10 months from both Scotland and Canada, whereas the study of Pruimboom-Brees et al. [32] used only a single colostrum-deprived neonatal (<24 h) calf to isolate VT receptors, and this animal may not be representative. Also, in the present study, snap-frozen tissue was used in a well-established procedure for assessing VT binding to tissue [33], whereas in the other study [32], tissue was acetone-fixed. This process, in addition to permeabilising cells, may solubilise Gb3 and could account for the apparent absence of receptor suggested by that study. In the present study, this receptor distribution was reproducible on triplicate samples from six animals of different geographic origins. It must be concluded that VT receptor is present on bovine intestinal epithelium, localising specifically to a subpopulation of crypt cells.

Crypts are composed of mixed populations of cells including immature proliferating cells and differentiated cell types [39]. A subset of proliferating cells migrate towards the luminal surface and differentiate [39], acquiring the enzymic and morphological characteristics of the mature apical epithelium. Results

Fig. 3. Immunohistochemical localisation of VT1 binding to bovine kidney. This section overlaid with VT1 200 ng/ml was derived from a 1-month-old calf and is representative of results obtained with kidney from cattle aged both 1 month (n = 3) and 10 months (n = 3). A similar binding pattern was observed with the lower concentration of VT1. Binding of VT1 was detected mainly in the renal cortex, with predominant staining apparent in the convoluted tubules and collecting ducts. The absence of binding of VT1 to glomeruli (arrow) and blood vessels is evident.

Fig. 4. VT receptor isolation from intestinal mucosa and bovine kidney. Glycolipids from 1- and 10-month-old cattle (three animals of each age) were analysed by TLC and the VT receptors were identified by toxin-binding overlay assay. (a) Lane 1, human adult kidney (2 μg/ml); 2, 3, colon of 1- and 10-month old cattle (20–50 μg/ml); 4, 5, caecum from 1- and 10-month-old cattle (20–50 μg/ml). (b) Lane 1, human adult kidney (2 μg/ml); 2, kidney from 1-month-old cow (20–50 μg/ml); 3, kidney from 10-month-old cow (20–50 μg/ml).
showed that apical cells do not bind VT1, which presumably corresponds with loss of this receptor as these cells migrate and differentiate. Correlation between cellular proliferation/differentiation and VT binding has been demonstrated previously [40–42]. The binding of VT to bovine intestinal epithelium contrasts with the absence of receptor in human intestinal epithelium [43, 44] and, because of the noted activities of VT, this may be of significance during colonisation of cattle.

In addition to localisation at intestinal epithelium, VT1 also bound to submucosal lymphoid aggregates. There is growing evidence that VTs may target the immune system of the host. For instance, cattle lymphocytes showed a lower lymphocyte proliferation response after infection of calves with VTEC strains [45] and, more recently, Ferens and Howde [46] reported a variety of immunomodulatory activities of VT1 through suppression of mononuclear cell activation and lymphocyte proliferation. Similar activities of VT are seen against lymphocytes from other sources, e.g., man [21, 47]. Therefore, VT may modulate immune responses and hence affect pathogen–host interaction. To act as a modulator of immune responses VT would need to gain access to submucosal sites. The presence of VT receptors on a subpopulation of epithelial cells may prevent toxin translocation and it remains to be determined whether VT crosses the bovine intestinal epithelium and gains access to submucosal tissues.

VT bound to bovine kidney in a pattern that resembled that described in human adult kidney [33], i.e., tubules and collecting ducts. This distribution corresponds to reduced susceptibility to HUS as a consequence of toxicity of systemic VT. Receptor localisation was similar to that observed in a previous study [32], in which acetone fixation was applied. This procedure may solubilise Gb3 and other glycolipids; however, as kidney retained receptor in that study, this treatment is presumably insufficient to remove all glycolipids from this tissue. Intestine, in contrast to kidney, does not apparently retain Gb3 after acetone treatment. The differences resulting from this treatment may be due to a variety of structural and compositional differences between intestinal mucosa and kidney, which may account for the tissue-specific variation. Important among these may be receptor isoform(s); e.g., bovine kidney expressed a third form of Gb3 (Fig. 4), which was not detected in musosal samples.

In man, in addition to targeting kidney, VTs also affect intestinal vascular cells as human endothelial cells express Gb3 [48, 49] leading to haemorrhagic colitis [50]. VT1 binding to capillaries was not observed in the bovine jejunum, ileum, caecum, colon or kidney, in corroboration of previously published observations [32]. This was not unexpected as FACS and immunofluorescence microscopy (unpublished results) with CPAE cells, a bovine pulmonary artery endothelial cell line, indicated that these cells lack Gb3. The general scarcity of VT binding to blood vessels and kidney glomeruli may limit systemic effects of VT and further contribute to the absence of overt pathogenicity of E. coli O157 in cattle.

Binding of toxin correlates with the expression of the various forms of Gb3, the VT receptor. To assess receptor forms in bovine tissues, glycosphingolipid species were isolated and analysed by a TLC VT overlay assay. Two VT-binding Gb3 bands were identified in enteric samples and three in kidney, with neither tissue showing age-related alterations. Thus, specific receptors in bovine intestine are responsible for the binding of VT1 to cells in tissue sections.

Intestinal epithelium in animals which do express epithelial Gb3 varies in susceptibility to VT toxicity [51–54]. VTs possess functions in addition to cytotoxicity, which could influence bacterium–host interaction. Firstly, by targeting crypt cells VTs may affect epithelial development. Infection of ruminants with E. coli O157:H7 has been shown by others [55] to reduce epithelial shedding which correlated with prolonged colonisation; the noted role of VT in suppressing cellular metabolism could contribute to this reduced cell turnover. Also, it has been demonstrated that VTs can elicit cytokines as a consequence of interaction with intestinal epithelial cells [56, 57] as well as directly affecting immune function. Thus, VTs may act as immunomodulators by influencing expression of mediators of inflammation, innate and adaptive immune responses.

There are several possible roles for VTs, which may affect both cellular physiology and host defences and thus influence colonisation and other outcomes of infection. All VTEC, and not just EHEC, contain VT-encoding bacteriophage(s). Strains belonging to these types possess a range of other factors associated with virulence [1] and heterogeneity among these, particularly serotype-dependent variation, could account for the differences in pathogenicity for cattle shown by various EHEC and VTEC serotypes. The effects of VT on bovine epithelial cells are of great significance to the success of colonisation, dissemination and persistence of these organisms in cattle reservoirs.

This study and others indicate possible roles for VT; however, the precise function(s) of VT in cattle remain to be fully defined and will require further evaluation of effects of VT on bovine intestinal mucosa. Nevertheless, it can be concluded that the different consequences of E. coli O157 infection in man and cattle involve a multiplicity of host and bacterial factors. Among those, the disparity between cattle and man in the distribution of VT receptor on intestinal epithelium is a fundamental difference which could contribute to the differing consequences of E. coli O157 infection in these two hosts.
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References
VEROTOXIN RECEPTORS ON BOVINE EPITHELIUM


