Shiga toxins, glycosphingolipid diversity, and endothelial cell injury

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Summary
Shiga toxin (Stx)-producing Escherichia coli (STEC) cause an enteric illness that results in a spectrum of outcomes ranging from asymptomatic carriage to uncomplicated diarrhea, bloody diarrhea, and the postdiarrheal haemolytic uremic syndrome (HUS), which leads to renal and other organ microvascular thrombosis. Binding of Stx to the glycosphingolipid (GSL) globotriaosylceramide (Gb3Cer/CD77) on endothelial cells followed by receptor-mediated endocytosis is the linchpin in STEC-mediated disease. Only GSLs that associate strongly with lipid rafts appear to carry Stxs retrogradely from the plasma membrane through the Golgi apparatus to the endoplasmic reticulum where they are translocated to the cytosol and exert their toxic function. Thus, the biophysical features of the lipid moiety of GSL receptors may influence its incorporation into certain membrane domains and thereby affect toxin destination. Consequently, a detailed structural analysis of Stx-binding GSLs is required to illuminate the molecular causes that may underlie the different Stx susceptibilities of endothelial cells derived from various vascular beds. Solid phase overlay binding assays of thin-layer chromatography (TLC)-separated GSL preparations employing specific antibodies and/or Stxs in conjunction with anti-Stx-antibodies are commonly used for the identification of Stx-binding GSLs. Such GSL-profiling combined with matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF-MS) represents a convenient strategy to structurally characterize Stx-receptors from any biological sources such as primary cells, cell lines, or organs. This approach may be helpful to gain insights into Stx-induced impairment of target cells that is suggested to originate at least partly from the structural heterogeneity of the cellular ligands of Stxs.

Keywords
STEC, Gb3Cer, endothelial cells, TLC overlay assay, MALDI-TOF-MS

Shiga toxins and their association with clinical outcome of human infections

Shiga toxin (Stx)-producing Escherichia coli (STEC), especially of serotype O157:H7, cause a food- or waterborne enteric illness that results in a spectrum of outcomes ranging from asymptomatic carriage to uncomplicated diarrhea, bloody diarrhea, and the haemolytic uremic syndrome (HUS), a leading cause of acute renal failure in children (1–3). After ingestion, Stxs are released by STEC in the intestine, translocated across the gut epithelium into the circulation, and transported to capillary endothelial cells in renal glomeruli and other organs. Host cells are then injured by inhibiting the protein synthesis, stimulating prothrombotic messages, or inducing apoptosis. Vascular injury during HUS results from the action of Stxs on vascular endothelial cells (3, 4). The net result is a multi-organ thrombotic process (4).

Stxs are presently the most important and best characterized STEC virulence factors that can cause microvascular endothelial injury. They are AB₅ ribosome-inactivating toxins with rRNA-N-glycosidase activity (5, 6). Based on cell-culture toxin-neutralization assays and nucleotide sequence analysis, Stxs can be divided into two major groups, Stx1 and Stx2 (7). The prototype toxins of these families display 57% and 60% nucleotide sequence identity in their A and B subunits, respectively (8). Each of the Stx groups contains an increasing number of variants (Table 1). A single strain can possess one or more different stx alleles (9–14) and the different stx subtypes are associated with different clinical outcomes of infections (9–11, 14–17) (Table 1). The Stx1 group includes, in addition to the prototype Stx1, two...
Figure 1: Peptide maps of tryptic in-gel-digested A and B subunits of Stx1 and structural assignment.

Three µg of purified Stx1 together with 20 µg of reference proteins were applied to SDS-PAGE under reducing conditions and stained with Coo massie blue. Separated Stx1A and Stx1B were digested in gel with trypsin and the resulting peptide mixture subjected to MALDI-TOF-MS according to Shevchenko et al. (62) and a protocol previously published (171). All experimentally derived m/z-values of peaks 1 to 21 of Stx1A and peaks 1 to 5 of Stx1B were submitted to ProteinProspector (UCSF Proteomic tools, v4.27.2) and ALDENTE (Peptide Mass Fingerprinting tool available at the ExPASy proteomics server from the SWISS Institute of Bioinformatics SIB). Stx1A and Stx1B unrelated fragment ions caused by autoproteolytic digestion of trypsin are labeled with “Try” in the spectra. The ions with low mass (m/z between 1000 and 1250) in the Stx1B peptide map represent detergent-derived impurities of the peptide preparation. The peptide mass search resulted in the identification of Stx1A (Shiga-like toxin I subunit A precursor, EC 3.2.2.22, rRNA N-glycosidase) encoded by the bacteriophage H19 composed of 315 amino acids with a molecular mass of 34,814 Da (accession number: P08026; excluding the signaling peptide: 293 amino acids, 32,398 Da) and Stx1B (Shiga-like toxin I subunit B) composed of 89 amino acids with a molecular mass of 9,743 Da (accession number: P69179, excluding the signaling peptide: 69 amino acids, 7,691 Da), both from *E. coli*. Those data fit well to the apparent molecular masses of ~32 kDa and ~7.7 kDa determined by SDS-PAGE for the A and B subunit, respectively. The database comparison of the submitted list of tryptic peptides resulted in a sequence coverage of 72.2% for Stx1A (highlighted in red) and 88.4% for Stx1B (highlighted in blue).
variants designated Stx1c (10, 18) and Stx1d (19, 20), which are frequently found in STEC isolated from patients with uncomplicated diarrhea (9, 10, 13, 14, 18). The Stx2 family is comparably more heterogeneous. Besides the classical Stx2, it comprises several variants of which Stx2c (21), Stx2c2 (22), Stx2d (23), the mucus-activatable Stx2d (Stx2dactivatable) (24), Stx2e (25), and Stx2f (26–28) were identified in STEC isolated from patients (Table 1). Stx2, Stx2c, and Stx2dactivatable are associated with severe human disease such as HUS and bloody diarrhea (9, 11, 13–15, 17, 29). The remaining Stx2 variants have been mostly identified in STEC isolated from patients with uncomplicated diarrhea or from asymptomatic shedders (9, 10, 14, 18, 23, 25), but are rarely found in HUS patients (14, 30). Stx2e, the major Stx type associated with the edema disease in swine, is rare in human STEC (9, 13, 14, 25). Stx2f is a common Stx variant expressed by STEC isolated from pigeons (31). Its presence in human STEC isolates is the exception (9, 13, 14, 26–28, 32). However, from recent studies focusing on the stx genes it becomes obvious that several E. coli strains represent a highly dynamic course during the course of human infection that can modulate the pathogens in both directions due to the loss or gain of stx-converting bacteriophages (33–37).

Biochemical features and mode of action of AB5 Stxs

All members of the Stx family are composed of a single enzymic ~32 kDa A subunit (38) non-covalently associated with five ~7.7 kDa B subunits. The pentameric B subunit, which may contain as many as 15 binding sites for its preferential glycosphingolipid (GSL) receptor globotriaosylceramide (Gb3Cer) (39), mediates attachment and internalization of the AB5-Gb3Cer complex by receptor-mediated endocytosis via clathrin-coated vesicles (7, 40) or by endocytic routes that do not involve clathrin-coated pits (41–43 and references therein).

In Stx-sensitive cells the toxin-receptor complex undergoes retrograde transport from early endosomes through the cisternae of the trans-Golgi network to the endoplasmic reticulum (ER) and even to the nuclear membrane (44–46). Once Stx has reached the ER, the ~32 kDa A subunit is cleaved by the membrane-anchored protease furin into a catalytically active ~27.5 kDa A1 and a ~4.5 kDa A2 fragment. This processing is required for retro-translocation into the cytosol, where the A1 subunit exerts its rRNA N-glycosidase activity that removes a specific adenine residue from 28S rRNA (47). The proteolytic cleavage is essential for maximal cytotoxicity of Stx1 from E. coli O157:H7 (48), and about one molecule of translocated A1 fragment of Stx1 per cell is sufficient to inhibit protein synthesis and kill a cell (49).

It has been known for some time that following the intracellular routing of Stxs to the nuclear membrane Stxs trigger programmed cell death signalling cascades in intoxicated cells (6, 50, 51). This is important because for unknown reasons a variety of cells succumb to apoptotic cell death rather than to necrosis through inhibition of cytoplasmic protein synthesis. The mechanisms of apoptosis induction are newly emerging and suggest that Stx-induced apoptosis may contribute to the pathogenesis of HUS caused by STEC (4, 52, 53). Furthermore, Stx causes sublethal cell injury by altering cell adhesive properties and by increasing endothelial susceptibility to leukocyte-mediated injury (54). The resulting injured endothelium changes its normal thromboresistant phenotype and becomes thrombogenic, initiating microvascular thrombus formation (55, 56).

Stx purification and structural assignments to A and B subunits

Facile methods for affinity chromatography purifying Stxs are based on their binding specificity using immobilized truncated synthetic analogs of the receptor (57), Gb3Cer-conjugated octyl Sepharose CL-4B (58) or Gb3Cer non-covalently immobilized onto silica gel (59). However, conventional purification procedures of Stxs employing ammonium sulfate precipitation, followed by Sephacryl S200 and Affi-Gel Blue column chromatography, chromatofocusing and high-performance liquid chromatography are still as relevant and useful (60, 61). In addition to a preliminary proof of the apparent purity and structural integrity of a purified toxin by SDS-PAGE, peptide-mass mapping performed by matrix-assisted laser desorption ionization time-of-flight mass spectrometry (MALDI-TOF-MS) of tryptic peptides obtained from in-gel digests represents one of the most commonly employed approaches to rapidly identify proteins from a gel band or gel spot (62, 63). The structural verification of SDS-PAGE-separated Stxs following their purification according to standard procedures (60, 61) is fast and convenient, and it requires only a few micrograms of toxins. The resulting peptide maps of the A and B subunits of Stx1 and Stx2 are shown in Figures 1 and 2 together with the corresponding Coomassie blue-stained bands and the amino acid sequence coverages. The peptide mass search in sequence databases allows the identification of the A and B subunits of both toxins thereby confirming Stx1 and Stx2 as the “unknown” proteins in accordance with high sequence coverages of the subunits of the toxins (for details see figure legends). In addition, the amino acid sequence alignments of the matched peptide fragments illustrate the rather low degree of homology of the A and B subunits of the two closely related members of the Stx family (8, 64). Such control of purity and precise biochemical structure is highly recommended as a first step for the characterization of any Stx-preparation for physiological or receptor binding assays.

Cellular ligands of Stxs

GSLs are components of all vertebrate cells and distribute with high specificity between mammalian species, organs (65, 66), and cell types (67, 68). GSLs are known to play a fundamental role during development and cell differentiation (69, 70) and to mediate a wide variety of cellular processes, including cellular growth (71), signal transduction (72), and cell-cell interaction (73).

GSLs are amphipathic molecules composed of a hydrophilic carbohydrate chain and a hydrophobic ceramide anchor (74, 75). In mammalian cells, the ceramide moiety is typically generated by the long-chain aminoalcohol sphingosine (d18:1), which is linked with a fatty acid varying in chain length from C16 to C24.
Figure 2: Peptide maps of tryptic in-gel-digested A and B subunits of Stx2 and structural assignment. SDS-PAGE and MALDI-TOF-MS of Stx2 as well as the database search were performed exactly as described in Figure 1. All experimentally derived m/z-values of peaks 1 to 33 of Stx2A and peaks 1 to 9 of Stx2B were submitted to the data bases. The peptide mass search resulted in the identification of Stx2A (Shiga-like toxin II subunit A precursor, EC 3.2.2.22, rRNA N-glycosidase) encoded by the bacteriophage 933W composed of 319 amino acids with a molecular mass of 35,714 Da (accession number: P09385; excluding the signaling peptide: 297 amino acids, 33,195 Da) and Stx2B (Shiga-like toxin II subunit B) composed of 89 amino acids with a molecular mass of 9,874 Da (accession number: P09386, excluding the signaling peptide: 70 amino acids, 7,818 Da), both from E. coli. Those data fit well to the apparent molecular masses of ~32 kDa and ~7.7 kDa determined by SDS-PAGE for the A and B subunit, respectively. The database comparison of the submitted list of tryptic peptides resulted in a sequence coverage of 80.8% for Stx2A (highlighted in red) and 92.9% for Stx2B (highlighted in blue).
GSLs are located primarily in microdomains of the plasma membrane of animal cells (76–78) also named lipid rafts (79). Their oligosaccharide chains spread in the aqueous environment at the cell surface and this makes them excellent candidates for cell surface recognition molecules (80, 81). Consequently, GSLs are involved in the pathophysiology of many infections and serve as receptors for viruses (82, 83), bacteria (84–86) and bacterial toxins including Stxs (87–90).

Stx1 and Stx2 both preferentially bind to the GSL Gb3Cer (91, 92) which alone is a functional receptor for Stx1 when incorporated in cells that lack Gb3Cer (93). Prototypic Stx1 and Stx2 bind with high affinity to Gb3Cer, where the Galα1–Gal terminus provides the specific recognition sequence for toxins’ B pentamers. Both Stxs interact weakly with the tetrahexosylceramide globotetraosylceramide (Gb4Cer). Stx1 and Stx2 bind to Gb3Cer with similar specificity, but the oligosaccharide binding affinity of Stx1 is increased 10-fold compared to Stx2 (94). In addition, Stx1 and Stx2 exhibit differential association and dissociation rate constants: Stx2 binds to Gb3Cer more slowly than Stx1. However, once bound, Stx2 is difficult to dissociate as shown in vitro by surface plasmon resonance-based real-time receptor-binding analysis (95) (for biosynthesis and structures of globo-series neutral GSLs see Fig. 3). Differential binding properties may affect toxicities of Stx1 and Stx2 in vivo (96). However, not only the oligosaccharide part is important for toxin binding, but also the ceramide lipid anchor of the receptor (97, 98). The chain length and the degree of unsaturation and hydroxylation of the sphingosine base can vary, but the primary source of heterogeneity of the ceramide lies in the fatty acid composition. It has been shown that the length of the fatty acyl chain of Gb3Cer influences receptor function (99), intracellular sorting and retro-translocation of Stx to the cytosol (100, 101).
and nuclear targeting (46). Those data provide a molecular basis for the different pathology in vivo and suggest a previously unrecognized mechanism for the modulation of membrane GSL receptor function (7).

Stx1, Stx2, and their variants so far investigated bind preferentially to Gb3Cer with the exception of Stx2e. The latter toxin displays greater affinity to Gb4Cer compared to Gb3Cer (102–104). This difference in receptor binding is consistent with the difference in susceptibility of cell lines to the cytotoxicity of Stx2e and Gb3Cer-specific Stx1 and Stx2 (102). Thus, those toxins that are associated with human disease bind to Gb3Cer while Stx2e, which is associated with edema disease of swine, binds preferentially to Gb4Cer. In an elegant study performed by Lingwood et al., Gb3Cer binding specificity of Stx1 and Stx2 was converted to Gb4Cer and vice versa that for Stx2e from Gb4Cer to Gb3Cer by site-directed mutagenesis of the B subunits (105). The altered carbohydrate recognition of modified Stx2e from Gb4Cer to Gb3Cer has biological significance, resulting in Stx1-like disease after intravascular injection into pigs as compared with classical Stx2e-induced edema disease (106). These studies suggest the primary role of the carbohydrate binding specificity of Stxs in determining systemic pathology of STEC.

Stx receptors of endothelial cells

Binding of Stx to its receptor Gb3Cer on endothelial cells is postulated to be the critical event triggering the vascular injury caused by STEC (1, 4). At the target organs, Stxs bind through their B subunits to Gb3Cer, which is expressed in microvascular endothelial cells of the kidney (107), intestine (108), and brain (109). Variability in receptor expression has been ascribed to a number of factors including the degree of confluence of the cell culture (110, 111), cell-cycle-dependent regulation of GSL receptors (112, 113) and their tissue of origin, which is thought to reflect the different sensitivities observed in vivo (114). Though the importance of GSLs as receptors for Stxs is well documented, the exact GSL composition of endothelial cells has generally received low attention, and only limited data are available about the GSL composition of the various types of human endothelial cells. Most studies lack a detailed structural characterization of the relevant receptor GSLs and are limited to the immunological

Figure 4: Scheme of the Stx TLC overlay assay and its application for detection of Stx1 and Stx2 receptors in human endothelial cells (139). A) GSLs are separated by TLC on silica gel precoated glass plates. After plastic fixation of the silica gel, the immobilized GSLs are overlaid with Stx solution, followed by incubation with primary anti-Stx antibody. Secondary enzyme-linked antibody (e.g. alkaline phosphatase) is required to visualize Stx-positive reaction by generating a coloured precipitate from a suitable substrate (e.g. 5-bromo-4-chloro-3-indolylphosphate). B) GSL extracts of HBMECs and EA.hy 926 cells were separated together with 20 µg of neutral GSL references from human erythrocytes (Ery) as a positive control and stained with orcinol or subjected to Stx1 or Stx2 overlay assay. Applied GSLs correspond to 2 x 10⁶ cells in the orcinol stain and to 5 x 10⁵ cells in the Stx1 and Stx2 overlay assays. Vertical white lines indicate assembled non-contiguous lanes.
detection of globo-series Gb3Cer and Gb4Cer using antibodies or Stxs, in combination with anti-Stx antibodies. The majority of investigations focused on human umbilical vein endothelial cells (HUVECs) (115, 116), where the neutral GSLs Gb3Cer and Gb4Cer were found to represent the dominant GSLs (117). We performed a thorough characterization of HUVEC GSLs using specific antibodies combined with methylation analysis and mass spectrometry (118). Gb4Cer and Gb3Cer, both carrying mainly C24 or C16 fatty acid beside sphingosine, were detected as the major neutral GSLs in HUVECs. The comparison of neutral GSLs from HUVECs and bovine aorta endothelial cells revealed striking differences in their GSL profiles. The predominance of neolacto-series and the absence of globo-series neutral GSLs are hallmarks of bovine aorta endothelial cells (119), which explains their resistance towards Stxs (120). As in HUVECs (110, 121), globo-series neutral GSLs were the major GSLs in primary and immortalized human brain microvascular endothelial cells (HBMECs) (122, 123). Upon stimulation with inflammatory mediators, an enhanced expression of Gb3Cer has been reported for HUVECs (114, 124, 125) and primary HBMECs from various sources (126–129) that correlated with increased Stx sensitivity. Thus, as with other cells, endothelial cell susceptibility to Stxs depends largely on Gb3Cer receptor expression. Differing Stx1 susceptibilities have been reported for endothelial cells from different vascular beds and species. The association between the degree of Stx sensitivity and Gb3Cer content is generally recognized (130).

Identification of Stx-receptors by overlay assay detection

Because of its resolving power and easy handling, high-performance thin-layer chromatography (TLC) has become the standard tool for separation and partial characterization of GSLs in mixtures (131–133). It is routinely used for analytical and preparative applications (134, 135). The oligosaccharide portions of GSLs can be visualized by conventional staining with orcinol after GSL-separation on silica gel precoated TLC plates. In conjunction with carbohydrate-binding proteins, such as antibodies or bacterial toxins, GSL species can be further differentiated in complex GSL mixtures by chromatographic resolution and subsequent immunodetection on the TLC plate (overlay assay) (136). The general procedure includes silica gel fixation with plastic to prevent flaking of the silica layer during incubation and washing steps, overlay of the plate with the primary agent (e.g. antibody or toxin), followed by incubation with secondary detection agents (e.g. alkaline phosphatase labeled antibody) and color development with a suitable substrate (e.g. 5-bromo-4-chloro-3-indolylphosphate). The schematic representation of the TLC overlay assay employing Stx is shown in Figure 4A. Figure 4B demonstrates its application to detect the receptors for Stx1 and Stx2 in two types of endothelial cells, human brain microvascular endothelial cells (HBMECs) and EA.hy 926 cells. EA.hy 926 cells represent “immortalized” macrovascular endothelial cells, which have been obtained by fusion of HUVECs with a human epithelial cell line (137). Both HBMECs (138) and EA.hy 926 cells have been widely used for endothelial cell research ([139] and references therein). The orcinol stain of TLC separated GSL mixtures provides rather vague information about their individual GSLs, whereas specific cellular ligands of Stx1 and Stx2 can be easily identified in the GSL extracts of both cell lines by using the Stx overlay assay (Fig. 4B). Results from these experiments indicate that the two cell lines express different amounts of Gb3Cer, and that the major Gb3Cer species are distinguishable due to their chromatographic appearance as double bands. At this stage of investigation the upper and lower bands are supposed to represent Gb3Cer species with long- and short-chain fatty acids, respectively, whereas the sphingosine (d18:1) moiety of the ceramide is most likely constant and does not contribute to the structural heterogeneity of the lipid anchor (132, 133). We can further conclude from the TLC overlay assays of Figure 4B that both Stx1 and Stx2 exhibit identical binding specificity and preferentially bind to Gb3Cer, whereas Gb4Cer is only weakly stained in the reference mixture of neutral GSLs from human erythrocytes.

Structural characterization of Stx-receptors by mass spectrometry

MALDI-TOF-MS represents an excellent technology for sensitive mapping of GSL mixtures and structural characterization of their individual constituents (140, 141). GSL differences observed by TLC overlay assay with antibody or toxin binding (see Fig. 4) can be verified and structurally characterized by conventional ultraviolet (UV)-MALDI-TOF-MS. For this purpose the plastic fixative is removed from the silica gel layer by chloroform extraction, and immunopositive GSL bands of interest are scraped from the support and extracted as described in detail in previous publications (142, 143). This combined preparative TLC-MS technology requires only microgram quantities of GSL mixtures. Our group applied this strategy successfully to the structural characterization of Gb3Cer and Gb4Cer species from human erythrocytes with high- and low-affinity binding to Stx1, respectively (143). For many applications, these newer, joint TLC-MS strategies can replace the laborious and time-consuming HPLC purification of single GSLs (144). Following the outlined strategy, we analyzed Stx-stained GSLs from HBMECs and EA.hy 926 cells by MALDI-TOF-MS. Crude GSL extracts from both cell lines (Fig. 4) are sufficient to generate MALDI-TOF mass spectra as shown in Figure 5. The main species detected in both spectra correspond to the monosodiated [M+Na]+ molecular ions of Gb3Cer (d18:1, C16:0) and Gb3Cer (d18:1, C24:1/C24:0) accompanied by low abundance ions indicative of Gb3Cer (d18:1, C22:0). In contrast to HBMECs, minor Gb3Cer species containing C26:1- and C26:0-fatty acids were noted in EA.hy 926 cells. The data demonstrate that HBMECs contain almost equal amounts of Gb3Cer species with long- and short-chain fatty acids. In contrast, Gb3Cer spectra of EA.hy 926 cells demonstrated a relative increase of long-chain fatty acids. Gb3Cer species with saturated C24:0 fatty acid predominated over those with unsaturated C24:1 fatty acids in EA.hy 926 cells, while HBMECs showed a balanced ratio of these fatty acids. It has been suggested that Gb3Cer species with long-chain fatty acids display greater sensitivity to Stx-mediated cytotoxicity,
likely because they may more effectively mediate the localization of internalized toxin to the ER (145). Differences in the overall GSL composition of Stx-sensitive and resistant cells that express Gb3Cer species of apparently identical composition may also affect Gb3Cer recognition by and sorting of incorporated Stx1 (146).

It should be noted that GSLs can be directly analysed from silica gel plates by TLC-IR (infrared)-MALDI-TOF-MS of fluorochrome- or antibody-stained GSLs (147, 148). Thus, the TLC-overlay assay combined with mass spectrometry represents the state of the art to structurally characterize Stx-receptors from any biological sources, including primary cells, cell lines, or organ tissue. Figure 6 shows an example for the identification of Stx-receptors in human kidney and colon based on Stx1 TLC overlay assays and the corresponding MALDI mass spectra obtained from extracts of both tissues. As expected (149), human kidney shows a high concentration of Stx1-binding Gb3Cer receptors, while there is only scant expression of Gb3Cer in human colon (Fig. 6A). This difference is reflected by the corresponding MS signal intensities of Gb3Cer ions in kidney and colon (Fig. 6B and C, respectively). The different Gb3Cer species varying in their fatty acid substitution from C16 up to C24 acyl chain length are speculated to derive mostly from endothelial and epithelial cells of those organs.

The role of lipid rafts and caveolae in Stx binding, internalization, and retrograde transport

Many investigators have provided evidence for the existence of cholesterol-sphingolipid-enriched microenvironments on the cell surface – known as lipid rafts – which renders the membrane more ordered and less fluid than the bulk plasma membrane. Rafts are rich in GSLs, cholesterol, lipid-modified proteins (e.g. GPI-anchored proteins, doubly acylated Src-type kinases) and transmembrane proteins (150). The raft concept has long been controversial, but studies with improved methodologies have dispelled most doubts (79, 151, 152). Furthermore, evidence has accumulated that GSLs, specifically enriched in the exoplasmic

![Figure 5: Structural characterization of Stx1-detected Gb3Cer species in HBMECs (A) and EA.hy 926 cells (B) by MALDI-TOF-MS.](image-url) The Stx1 TLC overlay assays were performed as described in Figure 4 and the Stx1 positive bands, from which the GSLs were extracted for subsequent MALDI-TOF-MS, are shown in the dotted rectangles of the insets. Mass spectra from extracts equivalent to 7 x 10⁴ cells were acquired in the positive ion mode and all Gb3Cer species were detected as singly charged mono-sodiated ions. The minor signals marked with asterisks represent Gb3Cer species carrying C18 to C26 fatty acids assigned by shifts in 28 atomic mass units in comparison to major molecular ions.
Figure 6: Structural characterization of Stx1-detected receptors in human kidney and colon. A) The orcinol stains and Stx1 overlay assays of TLC separated GSL extracts from representative kidney and colon tissue samples were performed as described in Figure 4. Applied GSLs correspond to 2 mg of wet weight tissue. B, C) MALDI-TOF-MS was performed with extracts from Stx1 positive bands (see dotted rectangles in the inserts) that correspond to 30 µg of kidney (B) and 3 mg of colon wet weight tissue (C). Mass spectra were acquired in the positive ion mode and all Gb3Cer species were detected as singly charged mono-sodiated ions. The signals marked with asterisks in the kidney spectrum, which show a remarkable shift in 16 atomic mass units in comparison to major molecular ions, are assumed to represent related hydroxylated Gb3Cer species. The positions of postulated hydroxylation in the ceramide portions are yet unknown, and the exact localization requires further investigation of each single GSL by tandem mass spectrometry. The Stx1 overlay assay of the kidney extract suggests functioning of the hydroxylated Gb3Cer species as Stx1 receptors. The question whether this modification is specific for the kidney cannot be answered without further structural characterization of a representative number of kidney samples.
leaflet of the plasma membrane, exert an important function as platform for the initiation of signaling cascades (153, 154).

Caveolae comprise one subset of lipid rafts in cell surface (150, 152, 155, 156). These flask-shaped membrane invaginations are formed from lipid rafts by polymerization of caveolins, which are hairpin-like palmitoylated integral membrane proteins that tightly bind cholesterol and sphingolipids (157). Endothelial cells are among the richest in caveolae. Caveolae have been found to be partake in many physiological and pathological processes involving endothelial cells, such as atherosclerosis, hemostasis, and thrombosis (158, 159). Caveolae and rafts are internalized via a common pathway, caveolae/raft-dependent endocytosis. This pathway is clathrin-independent and sensitive to cholesterol depletion (160, 161). Studies that investigated the role of lipid rafts in Stx1-interaction with various cell types demonstrated the clustering of Gb3Cer in lipid rafts (162) and, moreover, binding of Stx B-subunit with raft-localized receptors as a requirement for the retrograde transport (163). Binding of Stx to Gb3Cer induces activation of the Src family kinase Yes and intracellular signals that mediate cytoskeleton remodeling (164, 165). Furthermore, the association of Stx1-Gb3Cer-interaction with a precursor of Gb3Cer, glucosylceramide, is an essential requirement for a cytotoxic effect in the ER (166). Moreover, different pools of Gb3Cer have been described that are thought to influence not only Stx cell binding, but also to affect intracellular trafficking of the toxin (167). These data suggest that only GSLs that associate strongly with lipid rafts carry AB toxins retrogradely from the plasma membrane and sort the toxins to the ER (168).

Despite vast data concerning protein assembly in lipid rafts and caveolae of various cell types, little is known about their GSL composition. As outlined above, the oligosaccharide portion of Gb3Cer determines its binding specificity towards Stxs, but the ceramide contributes to Stx binding properties through its fatty acid heterogeneity. Available evidence also suggests that the structure of the ceramide moiety defines the preferential occurrence of GSLs in the liquid-ordered or liquid-disordered phase of the plasma membrane and the strength of the association of GSLs and toxins with lipid rafts (151). Such biophysical features of the lipid moiety of GSL receptors may influence its incorporation into certain membrane domains, the retrograde intracellular route and thus may affect toxin destination.

**Perspectives**

HUS is a thrombotic disorder. Insights into toxin-receptor interactions and, specifically, the mechanism(s) of Stx-mediated cell injury that induces a prothrombotic response, are important goals. Endothelial cells, particularly those with high globo-series GSL content are plausibly the major target in STEC-related disease. Future investigations aimed at advancing our understanding of the subcellular lipid raft distribution in endothelial cell membranes, will be aided by novel biochemical and biophysical strategies (169, 170). This research will allow us to better understand the complex mechanisms of Stx binding and internalization, and may help to develop new strategies directed at the interruption of the Stx-induced pathological intracellular cascades.

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