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Prakash Radhakrishnan
University of Nebraska Medical Center, pradhakr@unmc.edu

Ming-Fong Lin
University of Nebraska Medical Center, mlin@unmc.edu

Pi-Wan Cheng
University of Nebraska Medical Center, pcheng@unmc.edu

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Elevated expression of L-selectin ligand in lymph node-derived human prostate cancer cells correlates with increased tumorigenicity

Prakash Radhakrishnan,

Department of Biochemistry and Molecular Biology, College of Medicine, 985870, Nebraska Medical Center, University of Nebraska Medical Center, Omaha, NE 68198, USA

Ming-Fong Lin, and

Department of Biochemistry and Molecular Biology, College of Medicine, 985870, Nebraska Medical Center, University of Nebraska Medical Center, Omaha, NE 68198, USA, Eppley Institute for Research in Cancer and Allied Diseases, University of Nebraska Medical Center, Omaha, NE, USA

Pi-Wan Cheng

Department of Biochemistry and Molecular Biology, College of Medicine, 985870, Nebraska Medical Center, University of Nebraska Medical Center, Omaha, NE 68198, USA, pcheng@unmc.edu, Department of Pharmaceutical Sciences, College of Pharmacy, University of Nebraska Medical Center, Omaha, NE, USA, Eppley Institute for Research in Cancer and Allied Diseases, University of Nebraska Medical Center, Omaha, NE, USA

Abstract

Human prostate cancer LNCaP cells including C-33 and C-81 cells were originally derived from the lymph nodes of a patient with metastatic prostate cancer. These two cells were employed for characterization of L-selectin ligand and *in vitro* tumorigenicity, because they mimic the clinical conditions of early and late-stage human prostate cancer. C-81 cells exhibit higher *in vitro* migratory and invasive properties as compared with C-33 cells. We find that the L-selectin ligand and mucin glycan-associated MECA-79 epitope were elevated in C-81 cells. An increase of these glycotopes positively correlates with elevated tumorigenicity and expression of key glycosyl- and sulfotransferase genes. These results suggest that modulated expression of selective glycotopes correlates with altered tumorigenicity of cancer cells.

Keywords

LNCaP cells; MECA-79; L-selectin ligand; Glycosyltransferases; Sulfotransferase

Introduction

Prostate cancer is the most common solid tumor in men in the US and Western Europe, and the second most common cause of cancer-related death in men over age 55 [1]. Metastasis to the brain, bone and lymph nodes is the main cause of prostate cancer death [2]. However, the molecular mechanism by which the late-stage prostate cancer cells metastasize to the distant sites is not completely understood. One possible mechanism involves interactions of P-selectin on platelets and L-selectin on leukocytes with their respective ligands on hematogeneously

disseminating tumor cells to form microemboli to facilitate their arrest in the vasculature of their target sites [3]. Such interactions are similar to the mechanisms employed by leukocytes for migration to the sites of tissue injury and lymphoid tissues [4]. These processes rely on interactions between P-selectin on the endothelial cells and sialyl Lewis x (sLe^x) on mucin glycan and sulfated tyrosine located at the N-terminal region of P-selectin glycoprotein ligand-1 on the leukocytes, and between L-selectin on the surface of leukocytes and the glycotopes that contain MECA-79 antigen in the high endothelial venues [5–7], respectively. Altered glycosylation, which leads to increased production of sLe^x, has been observed in advanced cancers [8], including prostate cancer [9]. Since sLe^x is an essential component of the selectin ligands, elevation of sLe^x coupled with expression of GlcNAc 6-sulfotransferase could enable the cancer cells to utilize the normal leukocyte migration mechanisms to metastasize [10–13].

In this communication, we provide evidence that supports the hypothesis that expression of L-selectin ligand on LNCaP cells plays a key role in leukocyte-mediated metastasis to the lymph nodes. To achieve this goal, we employed a LNCaP cell progression model, including C-33 and C-81 LNCaP cells [14,15]. The properties exhibited by these two cell clones mimic the early and the late stages of human prostate cancer in the clinic. Thus, this is a useful *in vitro* cell model for studying the mechanisms of clinical progression of prostate cancer. As compared with C-33 cells, C-81 cells have higher proliferation rate, tumorigenicity, and androgen-independent secretion of prostate-specific antigen. In this study, we show that as compared to C-33 cells, C-81 cells exhibit a greater degree of *in vitro* migratory and invasive properties, elevated expression of L-selectin ligand, and increased expression of key glycogenes involved in the synthesis of this ligand. Thus, modulation of selective glycogenes in cancer cells could modify the selectin ligands to alter the metastatic potential of these cells. This property resembles the known tumorigenic trend of these two cell clones.

Materials and methods

Cell lines

The human prostate cancer LNCaP cell line was originally purchased from the American Type Culture Collection ATCC (Rockville, MD). The C-33 and C-81 LNCaP cell model used in the present study was previously developed [15] and characterized [14]. C-33 cells include cells between passages 25 and 35, and C-81 cells include cells between passages 81 and 125. These cells were maintained at 37°C under a 5% CO₂ and water saturated environment in RPMI 1640 supplemented with 5% fetal bovine serum, 2 mM L-glutamine, 100 units/ml Penicillin and 100 µg/ml Streptomycin.

Motility and invasion assays

Motility assay of C-33 and C-81 cells was performed using 24-well plate insert chambers. Briefly, 5×10^4 cells suspended in 1 ml medium were plated on the top chamber of noncoated polyethylene terephthalate (8 µm pore size, Becton Dickinson, Bedford, MA). After incubation at 37°C for 24 h in a 5% CO₂ and water saturated environment, the cells that did not migrate through the pores were removed by scraping with a cotton swab. Cells that transversed the membranes were stained with a Diff-Quick cell staining kit (Dade Behring, Inc., Newark, DE). Cells at three random fields per insert were counted with a phase contrast microscope at 40× magnification and expressed as an average number of cells per field of view. Three independent inserts were analyzed in each case. The data were expressed as mean±SD. Invasion experiment of C-33 and C-81 cells was carried out using 24-well BD BioCoat™ Matrigel™ invasion chambers (8 µm pore size; Becton Dickinson). Briefly, 5×10^4 cells suspended in 1 ml medium were seeded on top of the rehydrated Matrigel insert membrane. The cells were incubated for 22 h as described above and then the cells that did not migrate through the pores were removed

by scraping with a cotton swab. Cells that transversed the membranes were stained with a Diff-Quick cell staining kit (Dade Behring, Inc., Newark, DE) and counted as described above.

Western blot analysis of MECA-79 epitope and C1β3GalT-1

Whole cell lysates of C-33 and C-81 cells were used for detection of MECA-79 epitope and C1β3GalT-1 enzyme. Aliquots of the whole cell lysates were boiled in a SDS buffer and resolved by SDS-Polyacrylamide gel electrophoresis (PAGE) on 8–10% gels, and blotted onto Immobilon-P membranes (0.45 μm pore size, Millipore, Bedford, MA). Membranes were blocked with 5% bovine serum albumin (BSA) in TBS/Tween for 1 h at RT, and treated at 4°C overnight with 3% BSA in TBS/Tween containing biotinylated rat anti-mouse MECA-79 epitope monoclonal antibody (Becton Dickinson, Biosciences; 2.5 μg/ml) or mouse polyclonal C1β3GalT-1 antibody (Abnova Corporation, Taipei, Taiwan; 0.01 μg/ml). After washing with TBS/Tween three times at 5 min each, these membranes were treated with HRP conjugated streptavidin (Zymed laboratories, San Francisco, CA; 1:1,000) and HRP conjugated goat anti-mouse antibodies (Santa Cruz Biotechnology, Inc, San Diego, CA; 1:2,000 dilution), respectively. The bound HRP was treated with the enhanced Chemiluminescent substrate, SuperSignal West Pico Chemiluminescent Substrate (Pierce, Rockford, IL), and then detected according to the manufacturer's instructions.

Enzymatic characterization of the properties of MECA-79 epitope

Aliquots of cell lysates were digested with *O*-sialoglycoprotein endopeptidase (OSGP) (Cedarlane, Ontario, Canada) according to the manufacturer's instructions. Briefly, whole cell lysates were incubated with OSGP (50 μl/50 μg protein) at 37°C for 18 h. The cell lysates so treated were resolved on 8–10% SDS-PAGE and blotted onto Immobilon-P membranes. These membranes were treated with MECA-79 antibody and then analyzed as described above.

Flow cytometry analysis of L-selectin interactions with LNCaP cells

Flow cytometry analysis of L-selectin interactions with LNCaP cells was carried out as described previously [16]. Briefly, LNCaP Cells were detached from plates by incubation with PBS containing 2 mM EDTA for 5 min at 37°C and washed three times with Hanks' balanced salt solution (HBSS). The cells were treated with 0.5% BSA in HBSS for 30 min to block nonspecific sites before exposure to L-selectin as described next. The L-selectin used was a chimera of L-selectin fused with Fc region of human IgG purified from LSRg293 Cytel cells [16] (A kind gift from Dr. Ajit Varki, University of California at San Diego, La Jolla, CA). The L-selectin (50 μg/ml) was preincubated with a goat-anti human IgG conjugated with FITC for 1 h at room temperature before incubation with tumor cells at 4°C for 2 h. After washing with HBSS/BSA and HBSS, the cells were fixed with 2% (wt./vol.) paraformaldehyde in HBSS for 5 min. Cells were washed again with HBSS and resuspended in 1 ml of HBSS/BSA for flow cytometry analysis. Controls were stained in the presence of 5 mM EDTA (calcium chelation). Cells incubated with secondary antibody alone served as antibody control.

RT-PCR analysis of the expression of glycosyltransferase and sulfotransferase genes

Total cellular RNAs were isolated from C-33 and C-81 cells using the TRI-REAGENT (Molecular Research Center, Inc, Cincinnati, OH) according to the manufacturer's protocol. For cDNA synthesis, 2 μg of total RNA was used as the template in a 20 μl RT reaction mixture by using Reverse-iT™ 1st strand synthesis kit (ABgene, UK) according to the manufacturer's instructions. The RT-PCR reaction was performed in an Eppendorf Mastercycler Personal using 1 μl of cDNA template in a total volume of 25 μl. RT-PCR of key Glyco- and Sulfotransferase genes involved in the synthesis of 6-sulfo-sLe^x and β-actin were carried out by using sense and anti-sense primers as shown below: C2GnT-1, 5'-CTCGAAA CACCTCTCTTTTCTGGC-3' and 5'-GGTCAGTGTTT TAATGTCTCCAAAG-3'

(NM_001097633); β 3GnT-3, 5'-TTCTTCA ACCTCACGCTCAAGCAG-3' and 5'-AGCA TCTCATAAGGTAGGAAGCGG-3' (NM_014256); β 4GalT-1, 5'-AGTGACGTGGACCTCATTCC and 5'-CCGA TGTCCACTGTGATTTG-3' (NM_001497); β 4GalT-4, 5'-GCTGTTGACTTTGTGCCTGA-3' and 5'-GCCT GGTGGATG ACGTAGAT-3' (NM_003778); FUT7, 5'-CACCTCCGAGGCATCTTCAACTG-3' and 5'-CGTT GGTATCGGCTCTCATTTCATG-3' (NM_004479); GlcNAc6ST-1, 5'-CGTGAGAGCC TACAGGTGGT-3' and 5'-CATGGGCTGGTAGCAAAACT-3' (NM_004267.3); ST3Gal-III, 5'-ACTCCAGTGGGAGGAGACT-3' and 5'-CGTGACCCAGAGACTTGTT-3' (NM_174963); ST3Gal-IV, 5'-CTAGCCATCACCAGCTCCTC-3' and 5'-CCAT GAAGAAGGGG TTGAGA-3' (NM_006278) and β -actin, 5'-GTGGGGCGCCCCAGGCACCA-3' and 5'-CTCC TTA ATGTCACGGACGATTC-3' (NM_001101). The PCR reaction conditions for all the genes were as follows: 94°C for 2 min (1 cycle); 94°C for 30 s, 50°C for 30 s, and 72°C for 1 min (30 cycles); and then 72°C for 5 min. The expression of glycosyltransferase and sulfotransferase genes was normalized with the expression of β -actin gene, which remained constant in all samples. The PCR products were subject to electrophoresis (130 V, constant-voltage field) on 1% agarose gel equilibrated in Tris-borate electrophoresis buffer containing ethidium bromide (50 μ g/ml). Gels were photographed under UV light.

Results

Elevated *in vitro* migration and invasion properties in C-81 LNCaP cells

The metastatic potential of human prostate cancer cells was analyzed by *in vitro* Matrigel Boyden chamber assay. As compared to C-33 LNCaP cells, the C-81 cells showed a significantly higher level of motility through polyethylene terephthalate (Fig. 1A) and a higher degree of invasion through matrigel (Fig. 1B).

MECA-79 epitope is located in mucin-type glycans

The MECA-79 epitope was detected in two protein bands at 77 and 131 kDa from lysates of both C-33 and C-81 cells. The high passage LNCaP cells showed greater intensity (2.6-fold of 131 kDa and 1.8-fold of 77 kDa) than low passage LNCaP cells. The biochemical properties of MECA-79 epitopes present in C-33 and C-81 cells were characterized by digestion of cell lysates with *N*-glycanase, KS-II, Ch-ABC and OSGP. The MECA-79 epitope was resistant to *N*-glycanase, KS-II and Ch-ABC (Data not shown). However, the two protein bands carrying MECA-79 epitope were digested by OSGP, which specifically cleaves the peptide backbone of glycoproteins carrying sialylated mucin type glycans (Fig. 2) [17].

L-Selectin interaction with LNCaP cells

Flow cytometry analysis showed that 70% more C-81 LNCaP cells bound to L-selectin than C-33 cells (Fig. 3).

Analysis of glycozymes involved in the synthesis of L-selectin ligand in LNCaP cells

To understand the difference in metastatic potential as related to L-selectin ligand expression between C-33 and C-81 LNCaP cells, a combination of western blotting and RT-PCR analysis of key glycozymes involved in the synthesis of L-selectin ligand was performed. Western blot analysis showed a 70% increase of the C1 β GalT-1 enzyme (core 1 synthase) in C-81 cells as compared to C-33 cells (Fig. 4A). RT-PCR analysis of the expression of glycosyltransferase and sulfotransferase genes involved in L-selectin ligand synthesis showed a significantly higher expression of the following glycozymes in C-81 LNCaP cells as compared to those in C-33 cells: β 3GnT-3 (120%), FUT7 (20%), GlcNAc6ST-1 (50%), ST3Gal-III (30%) and ST3Gal-IV (30%) (Fig. 4B). There was no difference in the expression of C2GnT-1, β 4GalT-1,

β 4GalT-4 and iGnT genes between these two LNCaP cells although their expression levels were relatively high in both cells.

Discussion

Cell–cell interactions involving selectins and their carbohydrate ligands present on opposite cells play a key role in a variety of biological events, such as trafficking and homing of lymphocytes to specific tissues or lymphoid organs in the body [18]. For example, the L-selectin ligand in the high endothelial venues of the lymphoid organs is responsible for the arrest of the L-selectin-carrying leukocytes that traffic through these organs. Current study demonstrates that LNCaP cells express L-selectin ligand, which may facilitate metastasis of these cells to the regional lymph nodes by hitching on the circulating leukocytes. This finding is consistent with the fact that these cells are originally derived from the lymph nodes of a patient with metastatic prostate cancer [19].

The C-33 and C-81 LNCaP cell model has been shown to mimic the clinical progression of prostate cancer metastasis [14,15] and has been widely used for studying the mechanisms of prostate cancer progression [20–24]. We find that C-81 cells express more L-selectin ligand and exhibit higher levels of *in vitro* migratory and invasive properties as compared with C-33 cells. The results indicate that increased expression of L-selectin ligand correlates with elevated metastatic potential of LNCaP cells. Although elevation of L-selectin ligand can be used to explain the increased *in vivo* tumorigenicity, it is not clear how this property can be used to explain the enhanced *in vitro* metastatic potential of LNCaP cells. One possible explanation is that this property has nothing to do with the *in vitro* tumorigenicity, *i.e.* the observation is simply a co-incidental phenomenon. However, we can not exclude the possibility that this L-selectin ligand may facilitate *in vitro* migration of LNCaP cells because these cells express not only L-selectin ligand but also L-selectin (Radhakrishnan, unpublished observation).

6-sulfo-sLe^x is the L-selectin ligand, which also can be recognized by MECA-79 antibody. This antibody is specific for mucin core 1-associated 6-sulfo-*N*-acetylglucosamine [Gal β 1–4 (6-sulfo)GlcNAc β 1–3Gal β 1–3GalNAc α] [25] with or without further modifications, which include α 2–3 sialylation of β 1–4Gal, α 1–3 fucosylation of GlcNAc, and/or C6 sulfation of β 1–3Gal [6,7]. MECA-79 antibody specifically binds to luminal surface of HEV and inhibits binding of lymphocytes to HEVs *in vivo* and *in vitro* [26]. MECA-79 epitope can be found in mucin-type [17] and N-linked [27] glycans. But, they are present only in mucin-type glycans in LNCaP cells (Fig. 2).

Mucin glycan-associated 6-sulfo-sLe^x can be found on extended core 1 and core 2 branch as shown in Fig. 5. Synthesis of this epitope involves several glycosyl- and sulfo-transferases working in a sequential ordered manner [25]. Mucin glycan synthesis is initiated by the formation of GalNAc α Ser/Thr as catalyzed by peptidyl GalNAc transferases, which is followed by the formation of core 1 [Gal β 1–3GalNAc] as catalyzed by core 1 synthase (C1 β 3GalT-1). After the formation of core 2 [GlcNAc β 6 (Gal β 1–3)GalNAc α 1R] and/or extended core 1 [GlcNAc β 1–3Gal β 1–3GalNAc α 1R], GlcNAc 6-sulfate is generated before β 1–4 galactosylation of GlcNAc. This enzymatic step is followed by sialylation and then fucosylation to complete the synthesis of the L-selectin ligand. L-selectin ligand can be located at core 1 and core 2, but only core 1-associated L-selectin ligand can be recognized by MECA-79 antibody [25]. The increase in MECA-79 epitope and L-selectin ligand in C-81 LNCaP cells as compared to C-33 cells can be explained by upregulation of the expression of several glycosyltransferases involved in the synthesis of these glycotopes. C-81 cells express 70% more C1 β 3GalT-1 than C-33 cells. Since core 1 is the obligatory precursor for core 2 [28], more core 1 may lead to the synthesis of more core 2 branch. In addition, C81 cells express 120% more core 1 extension enzyme (β 3GnT-3), which may lead to the production of more extended

core 1 structure. Furthermore, C-81 cells express more sulfotransferase (GlcNAc6ST-1), sialyltransferases (ST3Gal-III and -IV) and fucosyltransferase (FUT7). The expression pattern of these glycosyltransferases positively correlates with higher level expression of MECA-79 epitope and L-selectin ligand on C-81 cells than in C-33 cells. Furthermore, it is of interest to note that alteration of the expression of selective glycosyltransferase and sulfo-transferase genes is correlated with the metastatic potential of LNCaP cells, suggesting the important role played by these glycogenes in carcinogenesis. The role of each of these glycogenes plays in this process will be examined *in vivo* in an animal model. Further, some of these genes may serve as prognostic markers for cancer progression. Modulation of the expression of these genes may be used as a potential therapeutic approach to suppress cancer metastasis.

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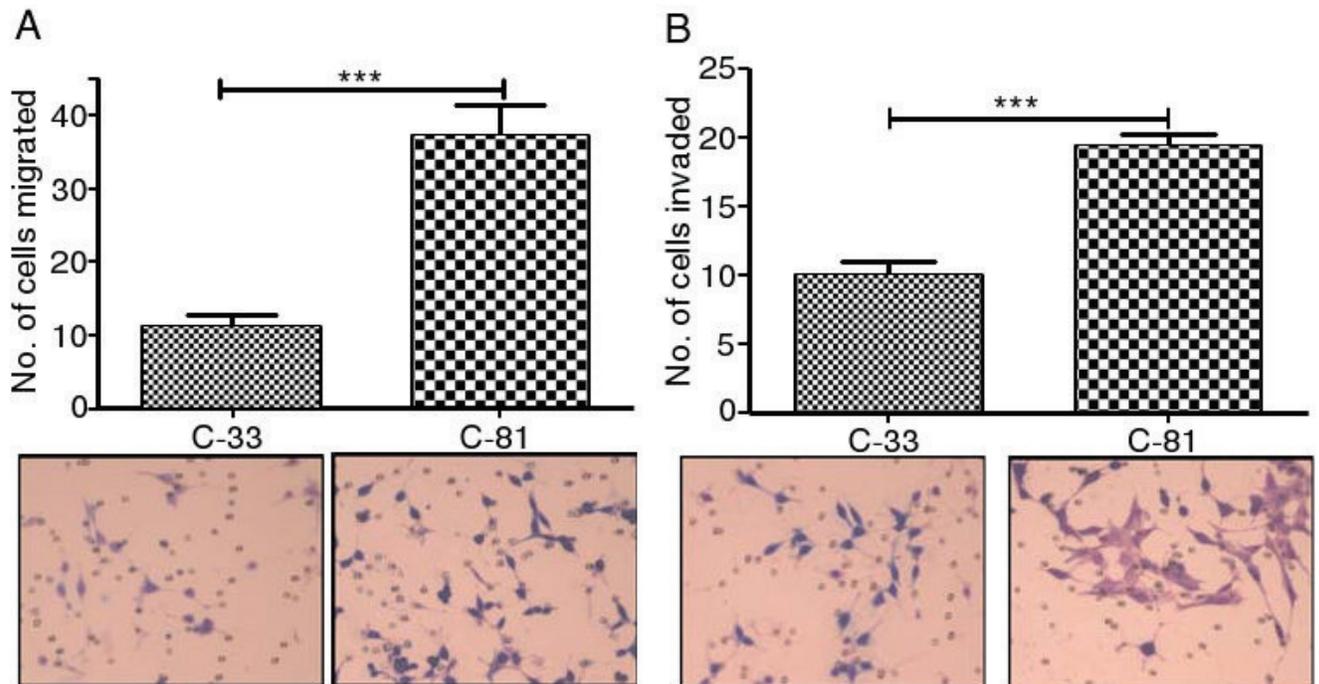


Fig. 1. Migration and invasion analysis of LNCaP cells. C-81 and C-33 LNCaP cells of 5×10^4 cells/ml medium were inoculated in the upper chambers of 24-well migration and matrigel invasion chamber plates. After 22–24 h, the cells that transversed to bottom side of the chamber plates were fixed, stained, and then counted under a phase contrast microscope (40 \times) in three independent fields for each insert. C-81 cells showed highly significant level of migration (a) and invasion (b) than C-33 cells through polyethylene teraphthalate membrane and matrigel invasion chambers, respectively. The data, which were expressed as mean \pm SD, were obtained from three independent experiments (** $p < 0.05$ and *** $p < 0.001$)

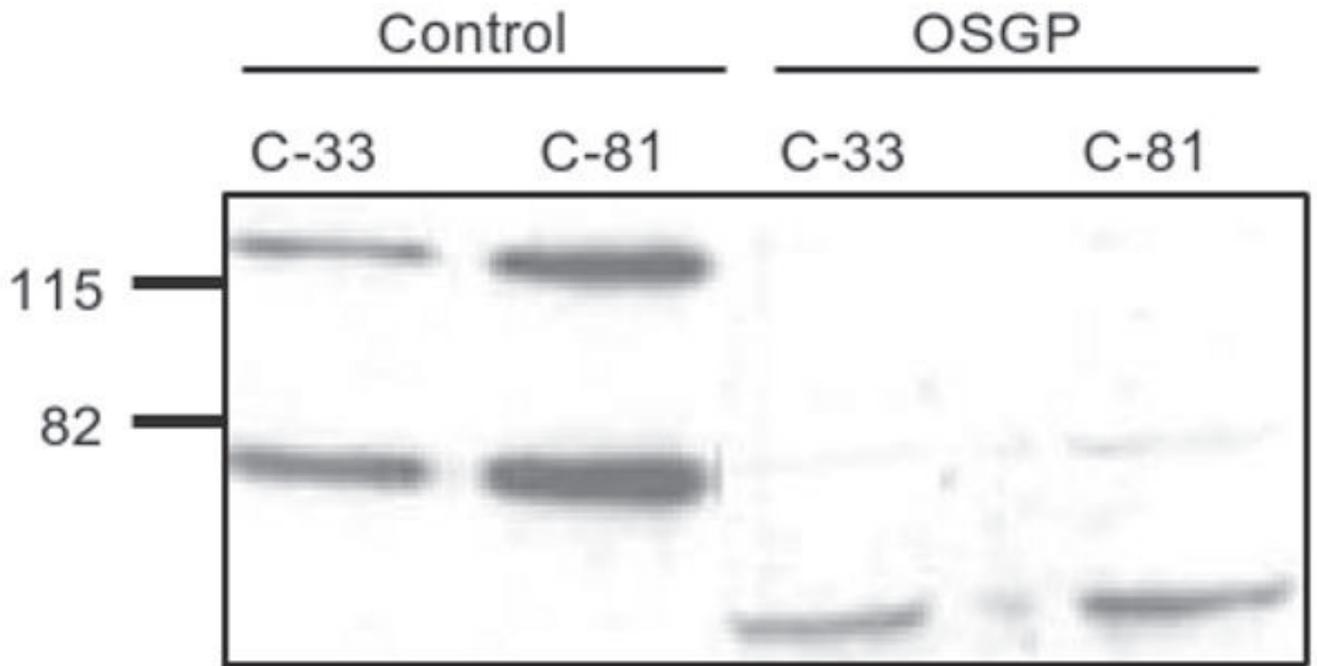


Fig. 2. Western blot analysis and characterization of MECA-79 epitopes on C-33 and C-81 LNCaP cells. Two bands of ~77 and 131 were recognized as MECA-79 epitopes. The high passage LNCaP cells showed more intense bands than low passage LNCaP cells. The lysates treated with OSGP showed loss of MECA-79 epitopes

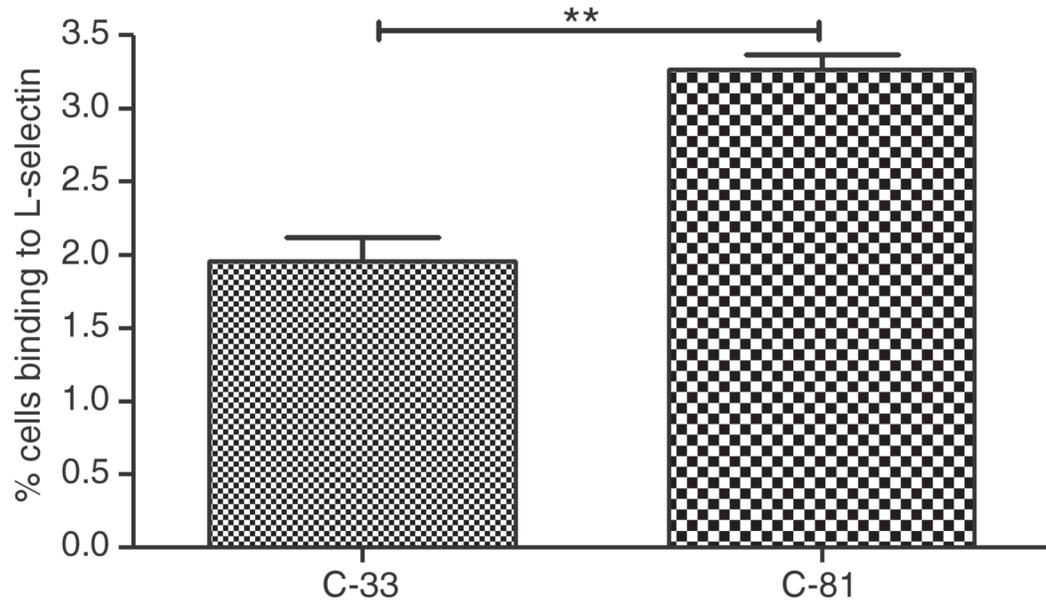


Fig. 3. Selectin-ligand interaction on LNCaP cells. The high passage LNCaP cells showed 1.7 fold higher numbers of cells that binding to L-selectin than low passage LNCaP cells. The data represented an average of three independent experiments (** $p < 0.01$)

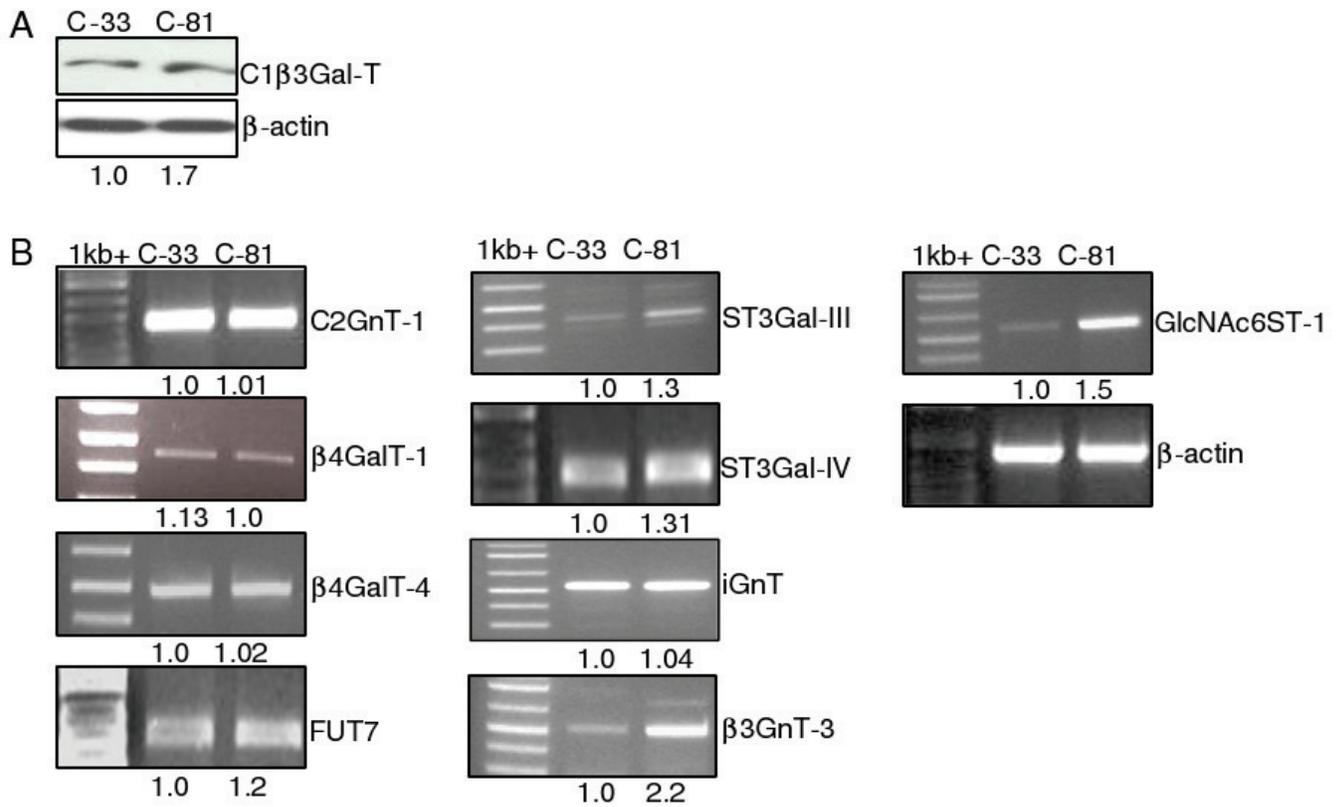
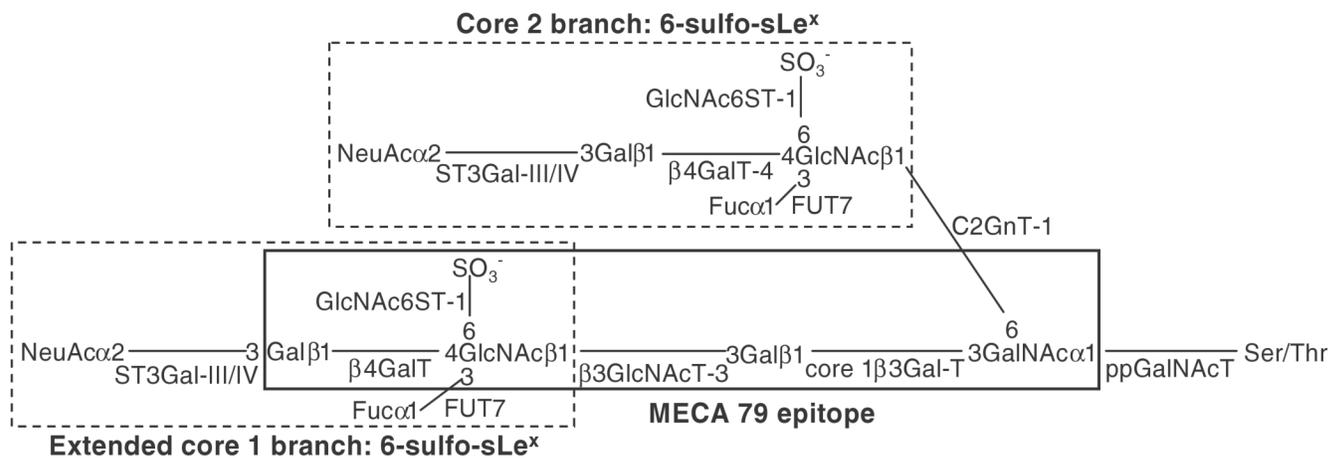


Fig. 4. Western blotting of C1 β GalT-1 (a) and RT-PCR analysis of the expression of sulfotransferase and other glycosyltransferase genes involved in the biosynthesis of mucin-type L-selectin ligand (b). β 3GnT-3, GlcNAc6ST-1, FUT7, ST3Gal-III and -IV genes were expressed higher in C-81 than C-33 LNCaP cells. There was no difference in the expression of C2GnT-1, β 4galT-1, β 4galT-4 and iGnT genes between these two LNCaP cells

**Fig. 5.**

Schematic diagram of the biosynthetic pathway of 6-sulfo-sLe^x located on biantennary mucin glycans. Key glycosyltransferases and sulfotransferase involved in the biosynthesis of 6-sulfo-sLe^x on extended core 1 and core 2 branch structures are shown. [ppGalNAcTF, peptidyl *N*-acetylgalactosaminyltransferase; β 3Gal-T1, β 3-Galactosyltransferase-1; C2GnT-1, β 1-6*N*-acetylglucosaminyltransferase-1; GlcNAc6ST1, *N*-acetylglucosaminyl-6-*O*-sulfotransferase-1; β 4GalT-4, β 4-Galactosyltransferase-4; FUT-7, α 1-3 Fucosyltransferase-7; ST3Gal-III/IV, α 2-3 sialyltransferase-III/IV; β 3GlcNAcT-3, β 3-*N*-acetylglucosaminyltransferase-3; β 4GalT, β 4-galactosyltransferase]