

World Journal of *Gastroenterology*

World J Gastroenterol 2018 February 21; 24(7): 767-876



**MINIREVIEWS**

- 767 Epidemiology, determinants, and management of AIDS cholangiopathy: A review
Naseer M, Dailey FE, Juboori AA, Samiullah S, Tahan V

ORIGINAL ARTICLE**Basic Study**

- 775 Glucose transporter expression in the human colon
Merigo F, Brandolese A, Facchin S, Missaggia S, Bernardi P, Boschi F, D'Inca R, Savarino EV, Sbarbati A, Sturniolo GC
- 794 Translational pancreatic cancer research: A comparative study on patient-derived xenograft models
Rubio-Manzanares Dorado M, Marín Gómez LM, Aparicio Sánchez D, Pereira Arenas S, Praena-Fernández JM, Borrero Martín JJ, Farfán López F, Gómez Bravo MÁ, Muntané Relat J, Padillo Ruiz J
- 810 Cryopreservation for delayed circulating tumor cell isolation is a valid strategy for prognostic association of circulating tumor cells in gastroesophageal cancer
Brungs D, Lynch D, Luk AW, Minaei E, Ranson M, Aghmesheh M, Vine KL, Carolan M, Jaber M, de Souza P, Becker TM
- 819 Metformin attenuates motility, contraction, and fibrogenic response of hepatic stellate cells *in vivo* and *in vitro* by activating AMP-activated protein kinase
Li Z, Ding Q, Ling LP, Wu Y, Meng DX, Li X, Zhang CQ
- 833 Fish oil alleviates liver injury induced by intestinal ischemia/reperfusion *via* AMPK/SIRT-1/autophagy pathway
Jing HR, Luo FW, Liu XM, Tian XF, Zhou Y

Retrospective Cohort Study

- 844 Elderly patients had more severe postoperative complications after pancreatic resection: A retrospective analysis of 727 patients
Chen YT, Ma FH, Wang CF, Zhao DB, Zhang YW, Tian YT

Retrospective Study

- 852 Predictors of functional benefit of hepatitis C therapy in a 'real-life' cohort
Steinebrunner N, Stein K, Sandig C, Bruckner T, Stremmel W, Pathil A
- 862 Predictors of post-treatment stenosis in cervical esophageal cancer undergoing high-dose radiotherapy
Kim JW, Kim TH, Kim JH, Lee IJ

CASE REPORT

- 870** Esophageal metastasis of stem cell-subtype hepatocholangiocarcinoma: Rare presentation of a rare tumor
Salimon M, Chapelle N, Matysiak-Budnik T, Mosnier JF, Frampas E, Touchefeu Y

CORRECTION

- 876** Correction for "Evaluation of a multiplex PCR assay for detection of cytomegalovirus in stool samples from patients with ulcerative colitis" (*World J Gastroenterol* 2015; 21: 12667-12675)
Hokama A

ABOUT COVER

Editorial board member of *World Journal of Gastroenterology*, Serdar Topaloglu, MD, Associate Professor, Department of Surgery, School of Medicine, Karadeniz Technical University, Trabzon 61080, Turkey

AIMS AND SCOPE

World Journal of Gastroenterology (*World J Gastroenterol*, *WJG*, print ISSN 1007-9327, online ISSN 2219-2840, DOI: 10.3748) is a peer-reviewed open access journal. *WJG* was established on October 1, 1995. It is published weekly on the 7th, 14th, 21st, and 28th each month. The *WJG* Editorial Board consists of 642 experts in gastroenterology and hepatology from 59 countries.

The primary task of *WJG* is to rapidly publish high-quality original articles, reviews, and commentaries in the fields of gastroenterology, hepatology, gastrointestinal endoscopy, gastrointestinal surgery, hepatobiliary surgery, gastrointestinal oncology, gastrointestinal radiation oncology, gastrointestinal imaging, gastrointestinal interventional therapy, gastrointestinal infectious diseases, gastrointestinal pharmacology, gastrointestinal pathophysiology, gastrointestinal pathology, evidence-based medicine in gastroenterology, pancreatology, gastrointestinal laboratory medicine, gastrointestinal molecular biology, gastrointestinal immunology, gastrointestinal microbiology, gastrointestinal genetics, gastrointestinal translational medicine, gastrointestinal diagnostics, and gastrointestinal therapeutics. *WJG* is dedicated to become an influential and prestigious journal in gastroenterology and hepatology, to promote the development of above disciplines, and to improve the diagnostic and therapeutic skill and expertise of clinicians.

INDEXING/ABSTRACTING

World Journal of Gastroenterology (*WJG*) is now indexed in Current Contents[®]/Clinical Medicine, Science Citation Index Expanded (also known as SciSearch[®]), Journal Citation Reports[®], Index Medicus, MEDLINE, PubMed, PubMed Central and Directory of Open Access Journals. The 2017 edition of Journal Citation Reports[®] cites the 2016 impact factor for *WJG* as 3.365 (5-year impact factor: 3.176), ranking *WJG* as 29th among 79 journals in gastroenterology and hepatology (quartile in category Q2).

EDITORS FOR THIS ISSUE

Responsible Assistant Editor: Xiang Li
Responsible Electronic Editor: Yan Huang
Proofing Editor-in-Chief: Lian-Sheng Ma

Responsible Science Editor: Ze-Mao Gong
Proofing Editorial Office Director: Jin-Lei Wang

NAME OF JOURNAL
World Journal of Gastroenterology

ISSN
ISSN 1007-9327 (print)
ISSN 2219-2840 (online)

LAUNCH DATE
October 1, 1995

FREQUENCY
Weekly

EDITORS-IN-CHIEF
Damian Garcia-Olmo, MD, PhD, Doctor, Professor, Surgeon, Department of Surgery, Universidad Autonoma de Madrid; Department of General Surgery, Fundacion Jimenez Diaz University Hospital, Madrid 28040, Spain

Stephen C Strom, PhD, Professor, Department of Laboratory Medicine, Division of Pathology, Karolinska Institutet, Stockholm 141-86, Sweden

Andrzej S Tarnawski, MD, PhD, DSc (Med), Professor of Medicine, Chief Gastroenterology, VA Long Beach Health Care System, University of California, Irvine, CA, 5901 E. Seventh Str., Long Beach,

CA 90822, United States

EDITORIAL BOARD MEMBERS
All editorial board members resources online at <http://www.wjgnet.com/1007-9327/editorialboard.htm>

EDITORIAL OFFICE
Ze-Mao Gong, Director
World Journal of Gastroenterology
Baishideng Publishing Group Inc
7901 Stoneridge Drive, Suite 501,
Pleasanton, CA 94588, USA
Telephone: +1-925-2238242
Fax: +1-925-2238243
E-mail: editorialoffice@wjgnet.com
Help Desk: <http://www.f6publishing.com/helpdesk>
<http://www.wjgnet.com>

PUBLISHER
Baishideng Publishing Group Inc
7901 Stoneridge Drive, Suite 501,
Pleasanton, CA 94588, USA
Telephone: +1-925-2238242
Fax: +1-925-2238243
E-mail: bpgoffice@wjgnet.com
Help Desk: <http://www.f6publishing.com/helpdesk>
<http://www.wjgnet.com>

PUBLICATION DATE
February 21, 2018

COPYRIGHT
© 2018 Baishideng Publishing Group Inc. Articles published by this Open-Access journal are distributed under the terms of the Creative Commons Attribution Non-commercial License, which permits use, distribution, and reproduction in any medium, provided the original work is properly cited, the use is non commercial and is otherwise in compliance with the license.

SPECIAL STATEMENT
All articles published in journals owned by the Baishideng Publishing Group (BPG) represent the views and opinions of their authors, and not the views, opinions or policies of the BPG, except where otherwise explicitly indicated.

INSTRUCTIONS TO AUTHORS
Full instructions are available online at <http://www.wjgnet.com/bpg/gerinfo/204>

ONLINE SUBMISSION
<http://www.f6publishing.com>

Basic Study

Glucose transporter expression in the human colon

Flavia Merigo, Alessandro Brandolese, Sonia Facchin, Silvia Missaggia, Paolo Bernardi, Federico Boschi, Renata D'Incà, Edoardo Vincenzo Savarino, Andrea Sbarbati, Giacomo Carlo Sturniolo

Flavia Merigo, Silvia Missaggia, Paolo Bernardi, Andrea Sbarbati, Department of Neuroscience, Biomedicine and Movement, Human Anatomy and Histology Section, University of Verona, Verona I-37134, Italy

Alessandro Brandolese, Department of Medicine, Gastroenterology Section, University of Verona, Verona I-37134, Italy

Sonia Facchin, Renata D'Incà, Edoardo Vincenzo Savarino, Giacomo Carlo Sturniolo, Department of Surgery, Oncology and Gastroenterology, Gastroenterology Section, University Hospital of Padua, Padua I-35128, Italy

Federico Boschi, Department of Computer Science, University of Verona, Verona I-37134, Italy

ORCID number: Flavia Merigo (0000-0002-6919-7169); Alessandro Brandolese (0000-0003-2977-992X); Sonia Facchin (0000-0002-6774-590X); Silvia Missaggia (0000-0002-8437-379X); Paolo Bernardi (0000-0002-1225-9525); Federico Boschi (0000-0001-9623-6859); Renata D'Incà (0000-0003-1169-1890); Edoardo Vincenzo Savarino (0000-0002-3187-2894); Andrea Sbarbati (0000-0002-0954-2119); Giacomo Carlo Sturniolo (0000-0002-2398-110X).

Author contributions: Merigo F was involved in experimental design, immunohistochemistry, data analyses, and manuscript writing; Brandolese A contributed to clinical sample documentation, presentation of the results, and manuscript writing; Facchin S, D'Incà R and Savarino EV contributed to clinical sample collection and documentation; Missaggia S contributed to immunohistochemistry analysis; Bernardi P performed SEM analysis; Boschi F analyzed the data; Sbarbati A and Sturniolo GC were involved in experimental design, final critical reading of the manuscript before submission; all authors contributed to the final manuscript.

Institutional review board statement: The study was reviewed and approved by the Ethics Committee of Padua University Hospital (study n° 2813P) Institutional Review Board.

Conflict-of-interest statement: All authors declare no conflicts of interest.

Data sharing statement: Technical appendix and data set are

available from the corresponding author at flavia.merigo@univr.it. Participants gave informed consent for data sharing. The present data are anonymized and the risk of identification is low.

Open-Access: This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

Manuscript source: Unsolicited manuscript

Correspondence to: Flavia Merigo, BSc, Human Anatomy and Histology Section, Department of Neuroscience, Biomedicine and Movement, University of Verona, Strada Le Grazie 8, Verona I-37134, Italy. flavia.merigo@univr.it
Telephone: +39-45-8027155
Fax: +39-45-8027163

Received: December 6, 2017

Peer-review started: December 6, 2017

First decision: December 12, 2017

Revised: December 13, 2017

Accepted: December 19, 2017

Article in press: December 19, 2017

Published online: February 21, 2018

Abstract

AIM

To investigate by immunostaining glucose transporter expression in human colorectal mucosa in controls and patients with inflammatory bowel disease (IBD).

METHODS

Colorectal samples were obtained from patients undergoing lower endoscopic colonoscopy or recto-sigmoidoscopy. Patients diagnosed with ulcerative

colitis ($n = 18$) or Crohn's disease ($n = 10$) and scheduled for diagnostic colonoscopy were enrolled. Patients who underwent colonoscopy for prevention screening of colorectal cancer or were followed-up after polypectomy or had a history of lower gastrointestinal symptoms were designated as the control group (CTRL, $n = 16$). Inflammatory status of the mucosa at the sampling site was evaluated histologically and/or endoscopically. A total of 147 biopsies of colorectal mucosa were collected and processed for immunohistochemistry analysis. The expression of GLUT2, SGLT1, and GLUT5 glucose transporters was investigated using immunoperoxidase labeling. To compare immunoreactivity of GLUT5 and LYVE-1, which is a marker for lymphatic vessel endothelium, double-labeled confocal microscopy was used.

RESULTS

Immunohistochemical analysis revealed that GLUT2, SGLT1, and GLUT5 were expressed only in short epithelial portions of the large intestinal mucosa. No important differences were observed in glucose transporter expression between the samples obtained from the different portions of the colorectal tract and between the different patient groups. Unexpectedly, GLUT5 expression was also identified in vessels, mainly concentrated in specific areas where the vessels were clustered. Immunostaining with LYVE-1 and GLUT5 antibodies revealed that GLUT5-immunoreactive (-IR) clusters of vessels were concentrated in areas internal to those that were LYVE-1 positive. GLUT5 and LYVE-1 did not appear to be colocalized but rather showed a close topographical relationship on the endothelium. Based on their LYVE-1 expression, GLUT5-IR vessels were identified as lymphatic. Both inflamed and non-inflamed mucosal colorectal tissue biopsies from the IBD and CTRL patients showed GLUT5-IR clusters of lymphatic vessels.

CONCLUSION

Glucose transporter immunoreactivity is present in colorectal mucosa in controls and IBD patients. GLUT5 expression is also associated with lymphatic vessels. This novel finding aids in the characterization of lymphatic vasculature in IBD patients.

Key words: Ulcerative colitis; Colon; Crohn's disease; Glucose transporter; LYVE-1, Immunohistochemistry

© **The Author(s) 2018.** Published by Baishideng Publishing Group Inc. All rights reserved.

Core tip: Our study demonstrates that GLUT2, SGLT1, and GLUT5 glucose transporters are expressed in the colorectal mucosa in controls and patients with inflammatory bowel disease (IBD). In addition, it provides first evidence that GLUT5 expression is associated with lymphatic vessels in controls and IBD patients. GLUT5-immunoreactive vessels were isolated or clustered in specific areas. We interpreted the presence of clusters as a pattern related to proliferative

zones. As GLUT5 is the main fructose transporter, fructose may have a role in the atypical aggregation of lymphatic vessels. This novel finding yields further insight into the characterization of lymphatic vasculature, whose dysfunction is a long-recognized feature in humans with IBD.

Merigo F, Brandolese A, Facchin S, Missaggia S, Bernardi P, Boschi F, D'Inca R, Savarino EV, Sbarbati A, Sturniolo GC. Glucose transporter expression in the human colon. *World J Gastroenterol* 2018; 24(7): 775-793 Available from: URL: <http://www.wjgnet.com/1007-9327/full/v24/i7/775.htm> DOI: <http://dx.doi.org/10.3748/wjg.v24.i7.775>

INTRODUCTION

Facilitative monosaccharide transport across the cell membrane is mediated by the GLUT family of membrane proteins encoded by the solute carrier transporter 2 (SLC2) gene. The GLUT protein family comprises 14 isoforms that differ by expression pattern and by affinity for sugars, polyols, and other carbon compounds^[1]. The GLUT5 and GLUT2 (also known as SLC2A5 and SLC2A2, respectively) isoforms are predominantly but not exclusively distributed in the intestine.

GLUT5 is a sugar transporter that facilitates fructose uptake. In mice, GLUT5 mRNA is abundantly detected by in situ hybridization and PCR in the small intestine but not in the large intestine. It is expressed in the small intestine and other organs such as the brain, testes, skeletal muscle, adipocytes, and kidneys^[2,3]. In enterocytes, it is located on the apical membrane. In the small intestine of rats, GLUT5 expression may be increased by exposure to fructose (or its metabolites), steroids, and thyroid hormones depending on age and weaning stage^[4-7]. GLUT5 expression differs according to insulin-resistance in adipocytes and duodenal enterocytes^[8,9]. It is also implicated in the development of metabolic diseases such as fructose-induced hypertension and non-alcoholic fatty liver disease^[10].

GLUT2 expression is widely present throughout the digestive system^[11], greatest in the small intestine and least in the terminal ileum. Molecular biology studies indicate that it is not expressed in the large intestine in mice^[12]. In humans, GLUT2 expression has been observed in enterocytes and enteroendocrine L-cells. In enterocytes, it is mainly located in the basolateral membrane where it facilitates the passage of glucose towards the interstitial space and bloodstream. In mice, it has been observed in the apical region of enterocytes after a sugar-rich meal or when insulin-resistance develops^[13-16]. GLUT2 is also expressed in enteroendocrine L-cells of the distal small intestine and large intestine in humans, and it is responsible for the secretion of glucagon-like peptide 1 (GLP-1)^[17-19].

Because of the gradient of glucose concentration between the cytosol and the intestinal lumen, glucose uptake by enterocytes depends mostly on SGLT1, which is a Na⁺/glucose co-transporter of the human family SLC5^[20]. SGLT1 is located on the apical membrane of enterocytes^[21]. In mice, SGLT1 expression was found to be intensely present from the duodenum to the end of the ileum where its levels tend to decrease, as demonstrated by in situ hybridization and PCR techniques. In the large intestine, SGLT1 expression remains controversial because SGLT1 mRNA in the proximal colon has been detected by in situ hybridization but not by PCR^[12]. SGLT1 is also known to be implicated in glucose-mediated GLP1 exocytosis through depolarization of the L-cell membrane^[22].

Further implications of these transporters other than sugar uptake and metabolism have been proposed for inflammation, malignancy, and gut microbiota regulation. Research has shown that GLUT5 activity and expression are regulated under normal conditions but they are altered in pathological conditions^[7]. GLUT5 is over-expressed in malignant but not in normal human tissues, including colonic and hepatic tissues, and it can be considered an important diagnostic or therapeutic target in tumors relying on fructose for proliferation such as breast cancer and pancreatic cancer^[23-25]. GLUT5 mRNA levels in non-inflamed small intestine of rats with iodacetamide-induced colitis were observed to be lower than in controls, suggesting a possible link between this fructose transporter and inflammatory cytokines. On the other hand, fructose absorption in rabbit was found to be inversely proportional to TNF- α expression and bacterial lipopolysaccharide (LPS) exposure^[26,27]. In mice models, GLUT5 deficiency has been reported to be associated with severe malabsorption due to dilation of the cecum and colon^[28].

GLUT2 has a high affinity for glucosamine, an amino-monosaccharide with anti-inflammatory activity against sodium dextran sulfate-induced colitis, which is a model for inflammatory bowel disease (IBD) in rats^[29]. Moreover, it has also been reported to have a role in modulating gut microbiota: after intestinal-specific deletion of GLUT2 in mice, an increase in commensal bacteria was observed in association with an improvement in inflammatory status^[19].

SGLT1 (like GLUT2) mRNA and protein levels were found to be significantly increased in intestinal biopsies of diabetic human subjects^[9]. Some mechanisms of cellular survival linked to SGLT1 have been demonstrated in Caco-2 cells models when exposed to bacterial LPS^[30]. Furthermore, SGLT1 expression in these models was found to be inversely proportional to TNF- α expression, implicating its potential involvement in colonic inflammatory diseases^[31]. In addition, in vivo studies on mice have shown that oral glucose and 3-O-methyl-d-glucose (a non-metabolizable glucose analogue) administration promotes survival in endotoxic shock induced by LPS^[32]. These studies

suggest a protective role for SGLT1 in inflammatory processes or sepsis, possibly by maintaining intestinal barrier integrity. SGLT1 acts through the inhibition of intestinal epithelial cell activation mediated by phosphorylation of the signaling molecule NF- κ B, a protein complex involved in the gene regulation of several cellular responses, which is a known mechanism in the epithelium of patients with Crohn's disease (CD) and ulcerative colitis (UC)^[33-35].

These findings raise expectations for future discoveries; however, most studies to date on sugar transporters in the gastrointestinal tract have been performed using molecular biology in murine or in vitro models. Since topographical variations in glucose transporters have been detected mainly by biochemical rather than standard immunohistochemical analysis also in humans, the localization and distribution of glucose transporters in the large intestinal mucosa of the colon have not yet been clarified.

For this reason, the aim of this study was to investigate by immunostaining at light and confocal microscopy the expression of the major intestinal glucose transporters (GLUT5, GLUT2, SGLT1) in human colonic mucosa in control subjects and subjects with IBD.

MATERIALS AND METHODS

The study protocol was approved by the Ethics Committee of Padua University Hospital (study No. 2813P). Written, informed consent was obtained from all participants prior to inclusion in the study. The study was prospectively conducted from December 2013 to November 2015.

Patients

Patients diagnosed with UC or CD and scheduled for diagnostic colonoscopy were enrolled. Patients who underwent colonoscopy for prevention screening of colorectal cancer or were followed-up after polypectomy or lower gastrointestinal symptoms were designated as the control group (CTRL). Inclusion criteria are presented in Table 1.

The UC patient group ($n = 18$) included 8 men and 10 women (mean age 47 years, range 30-66; average body-mass index (BMI, weight in kg divided by height in m squared) 24 kg/m², range 18-32.9 kg/m²).

The CD patient group ($n = 10$) included 5 men and 5 women (mean age 37 years, range 19-53; average BMI 23.6 kg/m², range 19.6-26.1 kg/m²).

The CTRL patient group ($n = 16$) included 6 men and 10 women (mean age 56 years, range 27-72, average BMI 24.85 kg/m², range 18.1-31.1 kg/m²). All CTRL patients were evaluated based on histological and/or endoscopic examination of intestinal biopsies; none was affected by UC or CD.

Patients were classified by their BMI into one of three groups: normal weight for BMI < 24.9 kg/m²,

Table 1 Criteria of inclusion for patients in each group

Patient group	Periodic Follow-up (n)	New stadiation of IBD (n)	Abdominal pain (n)	Chronic diarrhea (n)	Colorectal cancer screening (n)	Stipsis (n)	Hemorrhoids (n)
UC	16	2	-	-	-	-	-
CD	2	8	-	-	-	-	-
CTRL	3 ¹	-	5	4	2	1	1

¹Follow up after polypectomy. CD: Crohn's disease; UC: Ulcerative colitis; CTRL: Controls; IBD: Inflammatory bowel disease.

Table 2 Number of biopsies analyzed by immunohistochemistry in each tract of large intestine

Patient group	Biopsy status	Large Intestine					
		Proximal Tract			Distal Tract		
		Cecum colon (n)	Ascending colon (n)	Transverse colon (n)	Descending colon (n)	Sigmoid colon (n)	Rectum (n)
UC	Inflamed	2	6	4	6	7	7
	Non-Inflamed	7	9	5	6	3	3
CD	Inflamed	0	4	3	4	4	4
	Non-Inflamed	1	1	0	1	2	1
CTRL	Inflamed	2	4	3	2	4	4
	Non-Inflamed	7	5	6	6	7	7

CD: Crohn's disease; UC: Ulcerative colitis; CTRL: Controls.

overweight for 25.0-29.9, and obese for > 30 kg/m².

Colorectal samples

Colorectal samples were obtained from patients undergoing lower endoscopic colonoscopy or recto-sigmoidoscopy (2 in the CTRL group). Biopsies of portions of the colonic tract were taken for diagnostic purposes according to the endoscopist's judgment, and for immunohistochemistry (IHC). The biopsies were collected from adjacent sites to compare the level of inflammation in independent samples. In 14 of the 44 patients who underwent a complete colonoscopy, biopsies were obtained of all 6 portions of the colon-rectum (cecum, ascending colon, transverse, descending, sigmoid colon, rectum). In the remaining 30 patients, biopsies were obtained only from the endoscopically examined portions.

A total of 147 biopsies of colonic mucosa were collected for IHC analysis. Inflammatory status of the mucosa at the sampling site was evaluated endoscopically in all biopsies and histologically in 127 out of 147 biopsies by an experienced pathologist who evaluated the mononuclear and polymorphonuclear cell infiltration of the mucosal layer. For the endoscopic findings, inflammatory status was graded according to the Mayo endoscopic score in the UC patients (Mayo score of 0 indicates normal colonic mucosa, > 0 evidence of macroscopic active inflammation), and according to the Rutgeerts score in previously resected CD (3 out of 10) patients (Rutgeerts score of 0-1 indicates normal ileocolic anastomosis, and > 1 macroscopic relapse of CD). For the non-resected CD and CTRL patients, inflammatory mucosal status was graded as documented in the endoscopic report. Based on histological and endoscopic grading, their status towards inflammation and classification as

inflamed or non-inflamed was determined. When the endoscopic grade differed from the histological grade, the final biopsy status was decided on the basis of the pathologist's assessment. The complete list of all biopsies analyzed by IHC, classified according to inflammation status, is shown in Table 2.

Tissue preparation

After sampling, specimens were fixed in 40 g/L formaldehyde and processed by embedding in paraffin using standard methods. Sections were cut to 7 µm thickness, mounted on polylysine-coated microscope slides, and processed for immunoperoxidase and double immunofluorescence labeling. Primary and secondary antibodies are listed in Table 3.

Peroxidase-immunohistochemistry

Sections were deparaffinized in xylene, rehydrated through passages in alcohol with decreasing concentrations to water, and then microwaved at 800 W for three five-minute cycles in 0.01 mol/L citrate buffer (pH 6) to optimize antigen retrieval. Endogenous peroxidase activity was blocked by incubating the slides in 30 mL/L H₂O₂ in methanol for 30 min. After rinsing in PBS, the slides were first incubated with blocking solution (3 mL/L Triton X-100, 10 g/L bovine serum albumin, and 10 mL/L normal swine or rabbit serum); the solution was used to dilute the antibodies. Subsequently, the sections were incubated with primary antibodies overnight at 4 °C and then reacted with biotinylated secondary antibody for 1 h at room temperature. The immunoreaction was detected using a Vectastain Elite ABC kit (Vector Laboratories, Burlingame, CA, United States) and then visualized with 3,3'-diaminobenzidine tetrahydrochloride (Dako, Milan, Italy) for 5-10 min. Finally, the sections were mounted in Entellan (Merck,

Table 3 Primary and secondary antibodies for immunohistochemical analysis

Antibody	Host	Dilution	Source
Anti-GLUT2	Goat	1:200	cat#ab111117, abcam, Cambridge, United Kingdom
Anti-GLUT5	Rabbit	1:400	cat#ab36057, abcam
Anti-SGLT1	Rabbit	1:200	cat#ab14686, abcam
Anti-LYVE-1	Rabbit	1:500	cat#Bs-1311R, Bioss, Woburn, MA, United States
Anti-VEGF	Rabbit	1:200	cat#RB-222, Thermo Scientific, Fremont, CA, United States
Polyclonal anti-rabbit IgG/biotinylated	Swine	1:400	cat#E0353, Dako, Milan, Italy
Polyclonal anti-goat IgG/biotinylated	Rabbit	1:400	cat#E0466, Dako
Cy TM 3-Fab fragment anti-rabbit IgG	Goat	1:100	cat#111-167, Jackson Lab., Baltimore, MD, United States
FITC-fab fragment anti-rabbit IgG	Goat	1:100	cat#111-097, Jackson Lab.
Unconjugated fab fragment anti-rabbit IgG	Goat	1:30	cat#111-007, Jackson Lab.

Milan, Italy).

Sections processed as above, but without the primary antibody, were used as negative control. Control sections for SGLT1 were also prepared by preabsorbing the primary antibodies with the corresponding peptide (5 µg/mL of antibody; SGLT1 peptide, cat#ab99447, abcam). Sections of human small intestine were used as positive controls.

Light microscopy

Sections were examined with an Olympus BX51 microscope (Olympus, Tokyo, Japan) equipped with a digital camera (DKY-F58 CCD JVC, Yokohama, Japan). Digital images were analyzed and processed with Image-ProPlus 7.0 software (Media Cybernetics, Silver Spring, MD, United States). Images were composed with Adobe Photoshop software ver. 6.0 (Adobe Systems, Mountain View, CA, United States) to regulate contrast and brightness.

Immunofluorescence microscopy

Double immunofluorescent staining was used to compare immunolabeling of GLUT5 with that of lymphatic vessel endothelial hyaluronan receptor 1 (LYVE-1), a marker for lymphatic vessel endothelium in humans and rodents, and with that vascular endothelial growth factor (VEGF), a marker of vascular endothelial cells.

The double-label assay was carried out sequentially using a method that relied on the use of secondary monovalent Fab fragments because all primary antibodies are raised in the same species^[36,37]. Paraffin sections (processed as described above) were used for double staining, which was performed as described in Merigo *et al.*^[38]. After completing the staining protocol, the sections were immersed 1 g/L Sudan Black B (Merck) in 70% ethanol for 20 min at room temperature to reduce tissue autofluorescence. After rinsing, the sections were mounted with fluorescent mounting medium (DAKO) and observed with a Zeiss LSM 510 confocal microscope equipped with argon (488 nm) and helium/neon (543 nm) excitation beams. Sequential acquisition, *i.e.*, one color at a time, was utilized on double-labeled tissues to avoid side-band excitation of the inappropriate fluorophore. All images for publication were composed using Adobe Photoshop

software, adjusting only brightness and contrast.

Control sections were prepared using one of the following methods: (1) replacing the second primary antibody with normal rabbit serum; or (2) exchanging the fluorophores of the secondary antibodies; or (3) omitting the primary antibody; or (4) changing the sequence of secondary antibody application.

Scanning electron microscopy

Sections of colonic mucosa, previously immunostained for GLUT5 antibody, were reused for scanning electron microscopy (SEM) analysis. After removal of the coverslip, the samples were brought to absolute alcohol, processed by critical point drying (CPD 030, Balzers, Vaduz, Liechtenstein), mounted on stubs with colloidal silver, sputtered with gold by means of an MED 010 coater (Balzers), and examined under an FEI XL30 scanning electron-microscope (FEI Company, Eindhoven, NL).

RESULTS

The results of immunohistochemical analysis of samples from the cecum, ascending and transverse colon were grouped together as data of the proximal tract; the results of samples from the descending, sigmoid colon, and rectum were grouped as data of the distal portion of the large intestine.

Immunohistochemical analysis of GLUT2, SGLT1, and GLUT5 expression in large intestine

Immunostaining for GLUT2 was detected on the apical cell pole and/or basolateral membrane of epithelial intestinal cells. Only small portions of the epithelium were positive for GLUT2 (Figure 1A-D). A few samples in each group of patients were immunostained for GLUT2 in both the proximal and the distal tract (Table 4).

Immunoreactivity for SGLT1 was concentrated mainly on the apical pole of epithelial cells, which were often observed in small clusters of 2 or 3 (Figure 1E and F). SGLT1 expression was observed in the samples from the distal tract; no substantial differences between the three patient groups were noted (Table 4).

Two distinct expression patterns of GLUT5 immunoreactivity were observed throughout the portions of the large intestine. GLUT5 expression was found on

Table 4 Percentage of samples with GLUT2, SGLT1 and GLUT5 expression in epithelial cells of large intestine biopsies

Patient group	Biopsy status	Large			Biopsy status	Intestine		
		Proximal	Tract			Distal	Tract	
			GLUT2, %	SGLT1, %			GLUT5, %	GLUT2, %
UC	Inflamed (<i>n</i> = 12)	8	9	17	Inflamed (<i>n</i> = 20)	0	15	15
	Non-inflamed (<i>n</i> = 21)	5	5	33	Non-inflamed (<i>n</i> = 12)	8	8	17
CD	Inflamed (<i>n</i> = 7)	0	0	0	Inflamed (<i>n</i> = 12)	8	8	0
	Non-inflamed (<i>n</i> = 2)	0	0	0	Non-inflamed (<i>n</i> = 4)	0	0	0
CTRL	Inflamed (<i>n</i> = 9)	11	0	33	Inflamed (<i>n</i> = 10)	30	10	0
	Non-inflamed (<i>n</i> = 18)	17	0	17	Non-inflamed (<i>n</i> = 20)	20	1	15

CD: Crohn's disease; UC: Ulcerative colitis; CTRL: Controls.

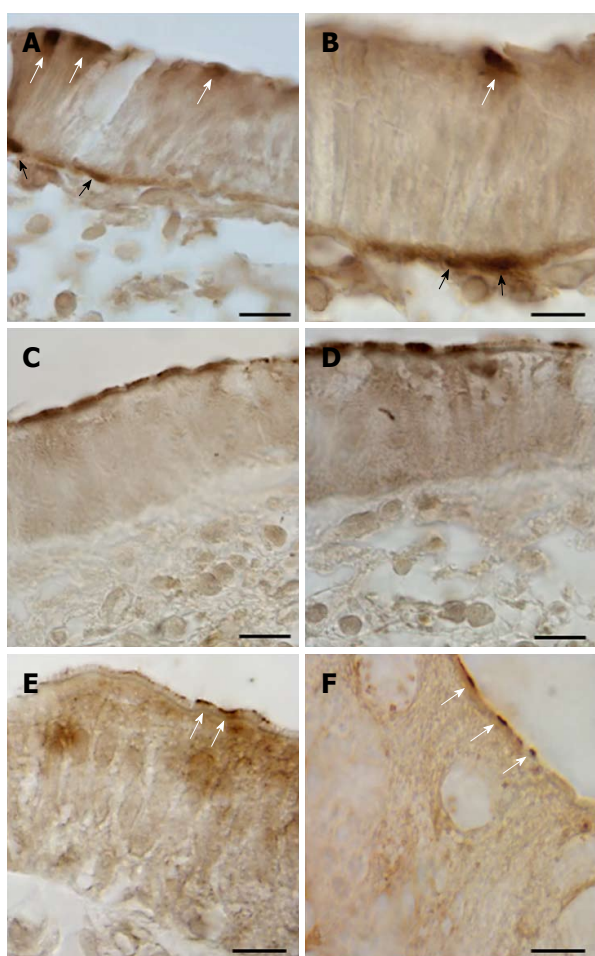


Figure 1 Immunoperoxidase staining showing GLUT2 (A-D) and SGLT1 (E and F) immunoreactivity in epithelial cells of the cecum (A and B), descending (C and D), and sigmoid (E and F) colon of human intestine. White arrows indicate staining on the apical pole, black arrows on the basolateral membrane of epithelial intestinal cells. Scale bar: (A, C, D) 50 μ m; (B, E, F) 5 μ m.

the apical membrane of epithelial cells, most likely representing labeling of the brush border membrane (Figure 2A-L). Only short portions of the epithelium showed positive epithelial cells. This staining was observed from the cecum to the rectum in both the UC and CTRL groups, but without significant differences between them (Table 4).

GLUT5 staining was also seen in vascular structures, where it displayed a heterogeneous pattern of

expression and distribution. GLUT5 immunoreactivity was observed in the endothelial cells of small vessels, with well-defined endothelium, and slit-like or rounded luminal spaces. The vessels were scattered separately just below the epithelium or around the glands in the intestinal mucosa (Figure 3A-C). This pattern was defined as normal GLUT5 expression in vessels because no samples lacking this characteristic were observed.

In some samples, however, this immunoreactivity pattern was detected in small, numerous vessels which, though remaining individually distinct, tended to aggregate. They were mainly concentrated below the epithelium or around the glands (Figure 3D-I).

No specific labeling was seen in the control sections when IHC was performed without the primary antibody (Figure 3J).

Aggregation of GLUT5-immunoreactive (-IR) vessels was much more evident in other samples, where GLUT5 immunostaining was observed to be mainly concentrated in specific areas formed by clusters of GLUT5-IR vessels scattered throughout the intestinal tract (Figure 4A-F). The clusters varied in number (number ranged from 1 to 12 *per* section) and size (min area 262 μ m²; max area 10695 μ m²) and were predominantly localized beneath the epithelium (Figure 5A and B) or around the glands (Figure 5C and D). They consisted of many non-rounded vessels which appeared to be strongly dilated, with only short portions lining the lumen covered by endothelial cells. The flat endothelial cells were stained for GLUT5 (Figure 5B). The epithelium overlying these aggregates often appeared damaged by the presence of vacuolar structures (Figure 5E and F). The epithelium was missing in some cases.

GLUT5-IR clusters in samples from UC, CD, and CTRL patients

GLUT5-IR clusters were observed throughout the entire tract of the large intestine in the samples of both inflamed and non-inflamed mucosal tissue from the UC, CD, and CTRL patients. The percentage of samples with GLUT5-IR clusters is shown in Table 5.

In the UC patients, the expression of GLUT5-IR clusters was higher in the samples from the distal than from the proximal tract, especially in the samples of non-inflamed tissue (83.3% vs 47.6%). The

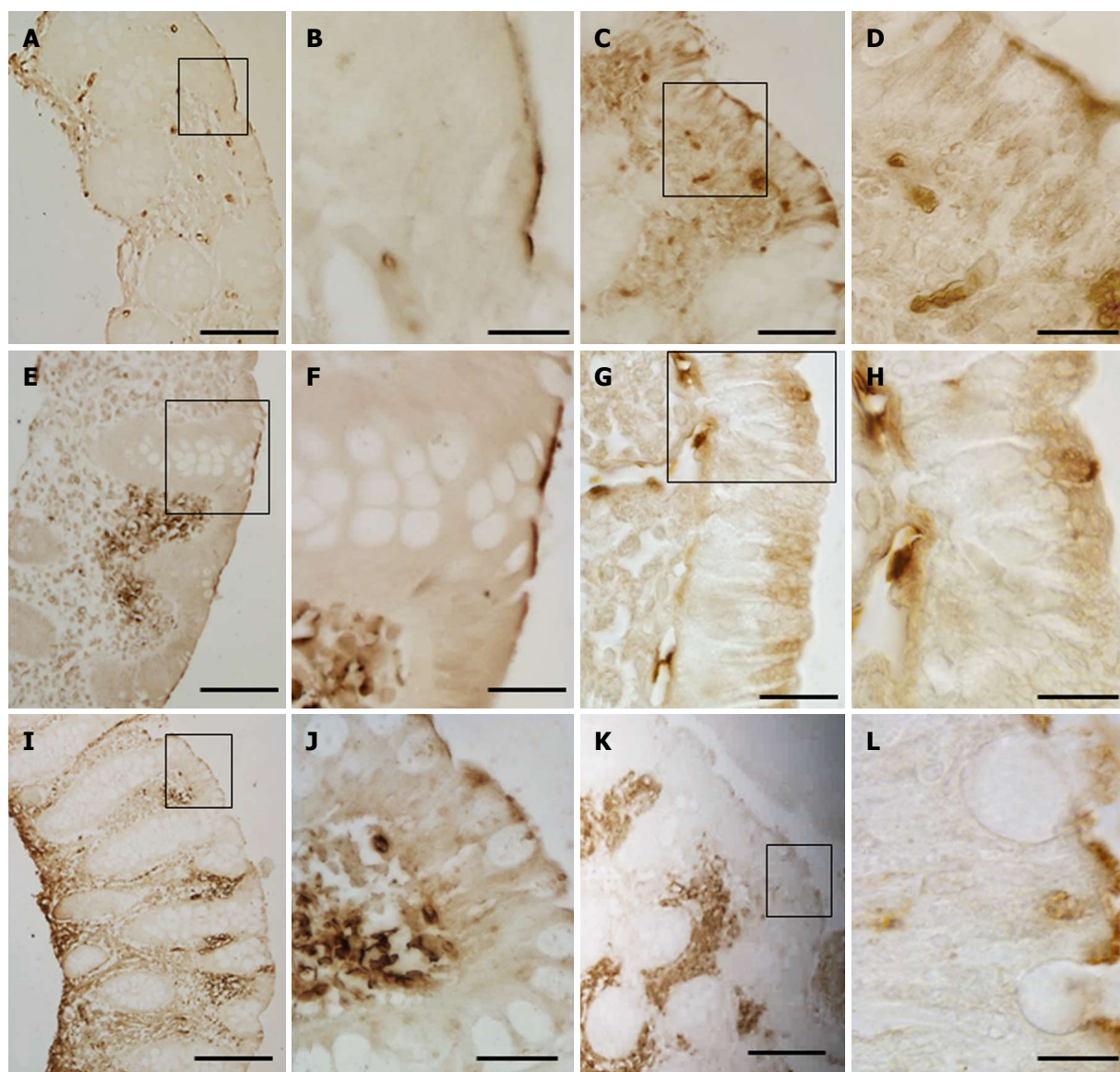


Figure 2 Immunoperoxidase staining showing GLUT5 immunoreactivity in the brush border membrane of epithelial cells in lumen of cecum (A and B), ascending (C and D), transverse (E and F), descending (G and H), sigmoid (I and J) colon, and rectum (K and L). The boxed area in (A, C, E, G, I, K) is shown at higher magnification in (B, D, F, H, J, L, respectively). Scale bar: (A, E, G, I) 50 μ m; (C, K) 25 μ m; (D, F) 10 μ m; (B, H, J, L) 5 μ m.

percentage of inflamed and non-inflamed samples was similar in the proximal tract (50% vs 47.6%), whereas in the distal tract the percentage of non-inflamed samples was higher (83.3% vs 55%).

In the CD patients, the percentage of GLUT5-IR clusters was higher in inflamed samples obtained from the distal than from the proximal tract (83% vs 71%), whereas the percentage was higher in non-inflamed samples from the proximal tract than from the distal tract (50% vs 25%). However, it was always higher in the inflamed than in the non-inflamed samples from both the proximal (70% vs 50%) and the distal tract (83% vs 25%).

In the CTRL group, the percentage of GLUT5-IR clusters in both the inflamed (60% vs 55.5%) and non-inflamed tissue samples (85% vs 44.4%) was higher in those obtained from the distal tract than from

the proximal tract. The percentage was similar for the inflamed samples and non-inflamed samples from the proximal tract (55.5% vs 44.4%) and higher in the non-inflamed as compared to the inflamed tissue samples from the distal tract (85% vs 60%). Figure 6 reports the percentages of samples with GLUT5-IR clusters as bar graphs, for both the proximal (Panel A) and the distal (Panel B) tract.

The expression of GLUT5-IR clusters was also evaluated in a sample group obtained by classifying the patients according to their BMI (Table 6). In the UC and CTRL patients, the expression of GLUT5-IR clusters was higher in the inflamed samples of patients with normal weight as compared with non-inflamed samples from the proximal tract; whereas in the distal tract the percentage of GLUT5-IR clusters was always higher in the non-inflamed samples as compared

Table 5 Percentage of samples with GLUT5 expression in clusters of lymphatic vessels in large intestine biopsies

Patient group	Large		Intestine	
	Proximal	Tract	Distal	Tract
	Biopsy status	GLUT5, %	Biopsy status	GLUT5, %
UC	Inflamed (<i>n</i> = 12)	50.0	Inflamed (<i>n</i> = 20)	55.0
	Non-Inflamed (<i>n</i> = 21)	47.6	Non-Inflamed (<i>n</i> = 12)	83.3
CD	Inflamed (<i>n</i> = 7)	71.0	Inflamed (<i>n</i> = 12)	83.0
	Non-Inflamed (<i>n</i> = 2)	50.0	Non-Inflamed (<i>n</i> = 4)	25.0
CTRL	Inflamed (<i>n</i> = 9)	55.5	Inflamed (<i>n</i> = 10)	60.0
	Non-Inflamed (<i>n</i> = 18)	44.4	Non-Inflamed (<i>n</i> = 20)	85.0

CD: Crohn's disease; UC: Ulcerative colitis; CTRL: Controls.

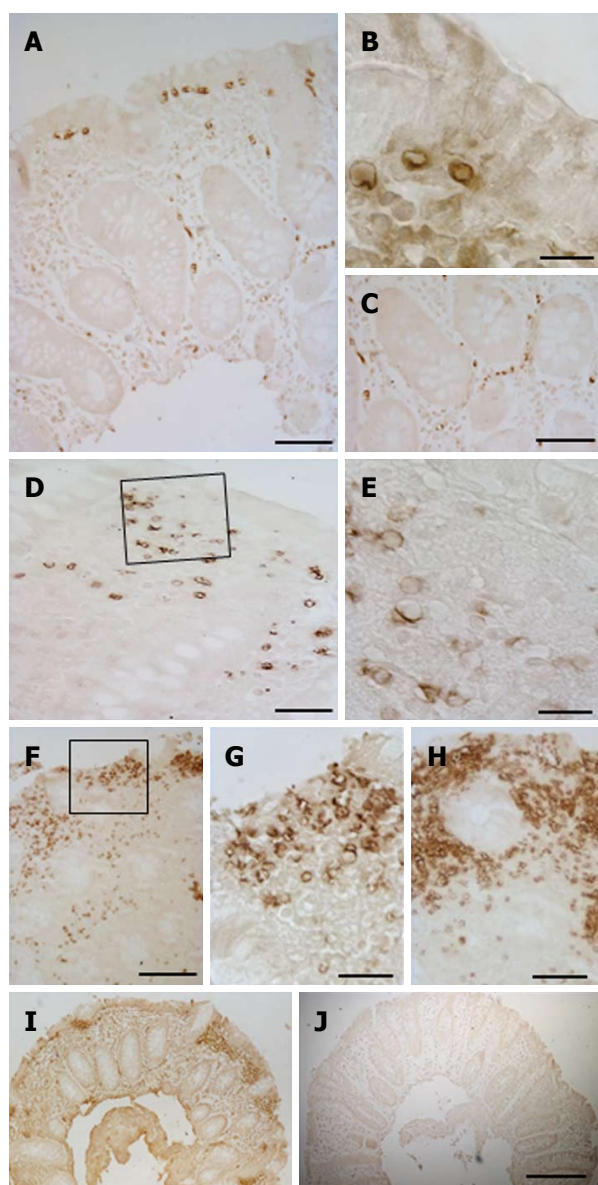


Figure 3 Immunoperoxidase staining showing GLUT5 immunoreactivity in small, rounded vessels in cecum (A and C), ascending (B and I), transverse (D and E), and sigmoid (F-H) colon of human intestine. No specific staining is observed in adjacent sections (I and J) when GLUT5 antibody was omitted (J). The boxed area in (D and F) is shown at higher magnification in (E and G, respectively). Scale bar: (I and J) 125 μ m; (A, C, D, F) 50 μ m; (B, E, G, H) 10 μ m.

to the inflamed samples from all three groups. The percentage was higher in the inflamed than in the non-inflamed tissue samples from both the proximal and distal tract in the normal weight and overweight CD patients.

SEM analysis of GLUT5 immunoreactivity

The GLUT5 positive sections were also observed at SEM for a more detailed evaluation of the labeled areas. Comparison between the SEM (Figure 7A and C) and the light microscope (Figure 7B and D) images showed that the area with GLUT5 immunoreactivity (Figure 8A-E) was characterized by large spaces and numerous dilated vessels, probably lymphatic because of the absence of red blood cells inside the lumen (Figure 8G and H). The vessels had a non-continuous endothelium, as observed at light microscopy. In the non-labeled areas, there were vessels containing cells inside the lumen that, judging by their rounded appearance, were probably lymphocytes or plasma cells (Figure 8F).

Comparison of GLUT5 immunoreactivity with LYVE-1 and VEGF expression in large intestine

GLUT5 expression was compared with expression of LYVE-1 and VEGF by light immunohistochemistry in contiguous sections and by laser-scanning confocal microscopy in double staining. Light immunohistochemistry revealed the presence of GLUT5 immunoreactivity in areas that were also LYVE-1 positive (Figure 9A, B, G and H). GLUT5 positivity was limited to areas usually located under the epithelium (Figure 9C and I), whereas LYVE-1 staining was distributed in large areas of the lamina propria (Figure 9D and J). The GLUT5 expression pattern was generally much more marked than LYVE-1 labeling, which was formed by short spots (Figure 9E and F). However, some corresponding areas in contiguous sections showed GLUT5 immunoreactivity (Figure 9G and K) but lacked LYVE-1 staining (Figure 9H and L).

Double immunofluorescence microscopy was consistent with light-microscopic evaluation. GLUT5-IR clusters were found distributed inside LYVE-1 positive

Table 6 Percentage of samples with GLUT5-immunoreactive clusters of vessels in biopsy groups subdivided by patient body mass index

Patient group	Biopsy status	Large			Intestine		
		Proximal	Tract		Distal	Tract	
		Patient	BMI		Patient	BMI	
		Normal weight, %	Over-weight, %	Obese, %	Normal weight, %	Over-weight, %	Obese, %
UC	Inflamed	71	20	0	70	43	33
	Non-inflamed	57	25	50	87	100	0
CD	Inflamed	80	50	0	87	75	0
	Non-inflamed	100	0	0	50	0	0
CTRL	Inflamed	80	25	0	67	50	0
	Non-inflamed	22	50	100	82	80	100

BMI: Body mass index; CD: Crohn's disease; UC: Ulcerative colitis; CTRL: Controls.

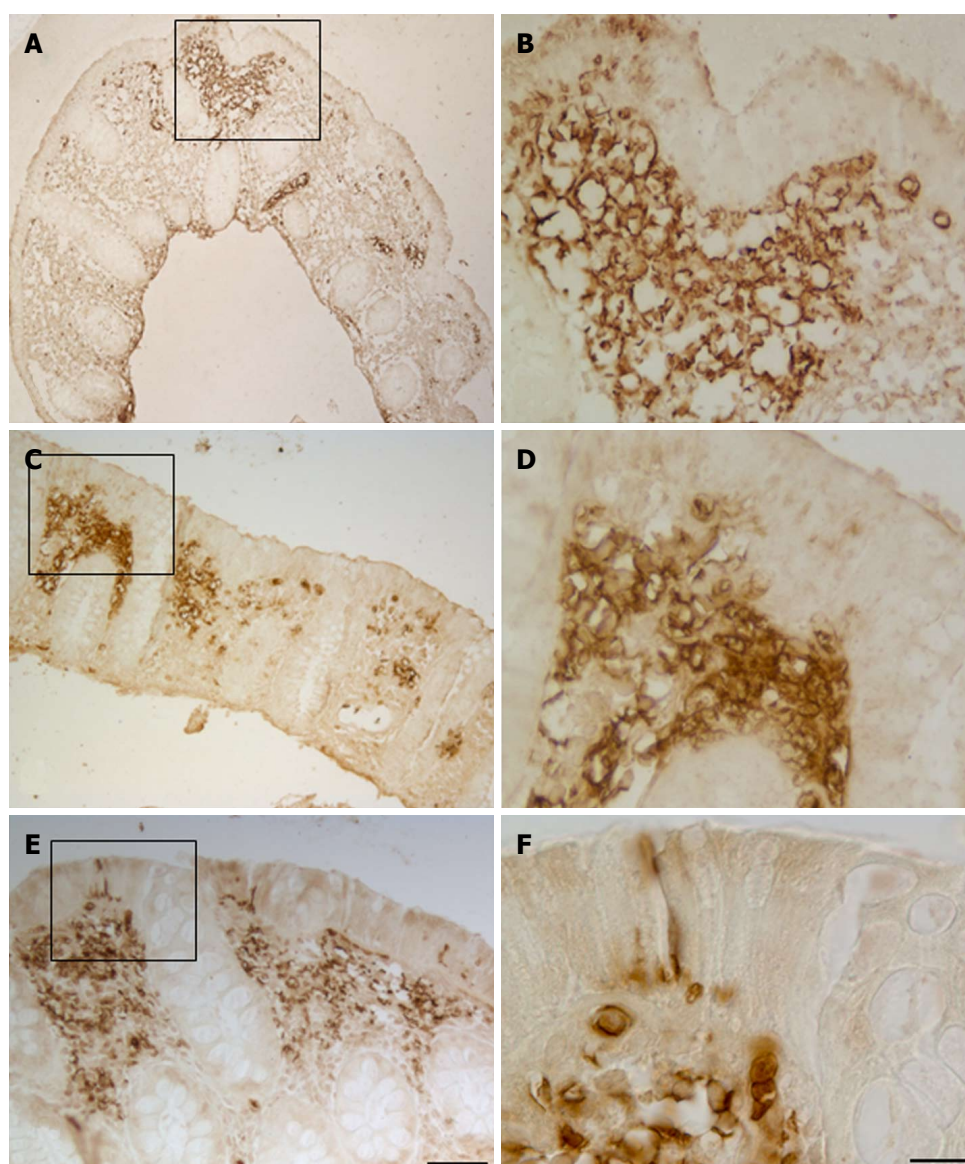


Figure 4 Immunoperoxidase staining showing GLUT5 immunoreactivity in vessel clusters in ascending (A and B), descending (C and D) colon, and rectum (E and F). The boxed area in (A, C, E) is shown at higher magnification in (B and D, F respectively). Scale bar: (A, C, E) 125 μ m; (B, D, F) 25 μ m.

areas and formed by numerous vessels, most of which were irregularly shaped and with ill-defined *lumina* (Figure 10A-F and J-L). GLUT5 and LYVE-1 expression

were never colocalized on the morphologically undefined wall but rather were expressed in close proximity to some areas of the endothelium (Figure 10M-O). Some

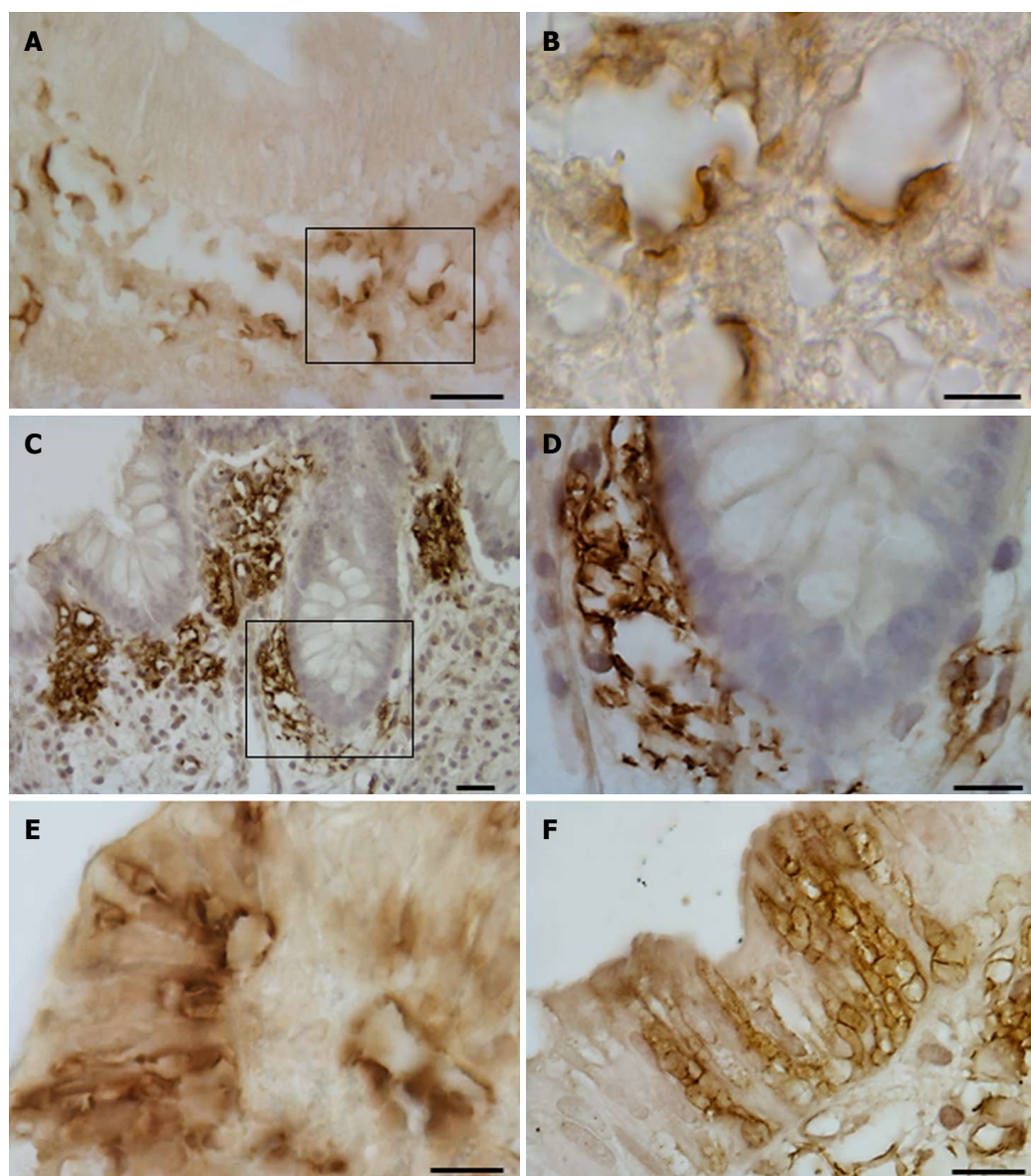


Figure 5 GLUT5 immunoreactivity is shown in vessel clusters localized beneath the epithelium (A and B) or around the glands (C and D), and in vacuolar structures in the epithelium (E and F). The boxed area in (A and C) is shown at higher magnification in (B, D, respectively). Scale bar: (A, C, E and F) 25 μ m; (D) 10 μ m; (B) 5 μ m.

vessels showed only GLUT5 expression (Figure 10G-I).

In addition to staining the lymphatic vessels, LYVE-1 also labeled the brush border membrane of some epithelial cells which was observed in both immunohistochemistry (Figure 9D) and double-immunofluorescent experiments, in the latter also seen colocalized with GLUT5 expression (Figure 10P-R).

To further discriminate between lymphatic vessels and blood vessels, expression of GLUT5 and VEGF antibodies was compared in contiguous sections using light immunohistochemistry and immunofluorescence. Both analyses revealed that VEGF expression was absent in GLUT5-IR clusters, confirming that GLUT5-immunoreactivity is selective for lymphatic vessels (data not shown).

In control sections, no specific double-labeling was seen when the second antibody was replaced with

normal rabbit serum.

DISCUSSION

Here we show that GLUT2, SGLT1, and GLUT5 glucose transporters are expressed in the epithelial cells of tissue samples of the large intestine, and that they are mainly located in the brush border membrane from IBD and control patients. Also in the colonic mucosa their location reflects the canonical expression of the small intestine and confirms the tissue-specific expression of GLUT isoforms also in humans. However, unlike the small intestine, their expression is present only in short epithelial portions, involving a limited number of cells. We observed no important differences in glucose transporter expression between the samples obtained from the proximal and distal tracts and

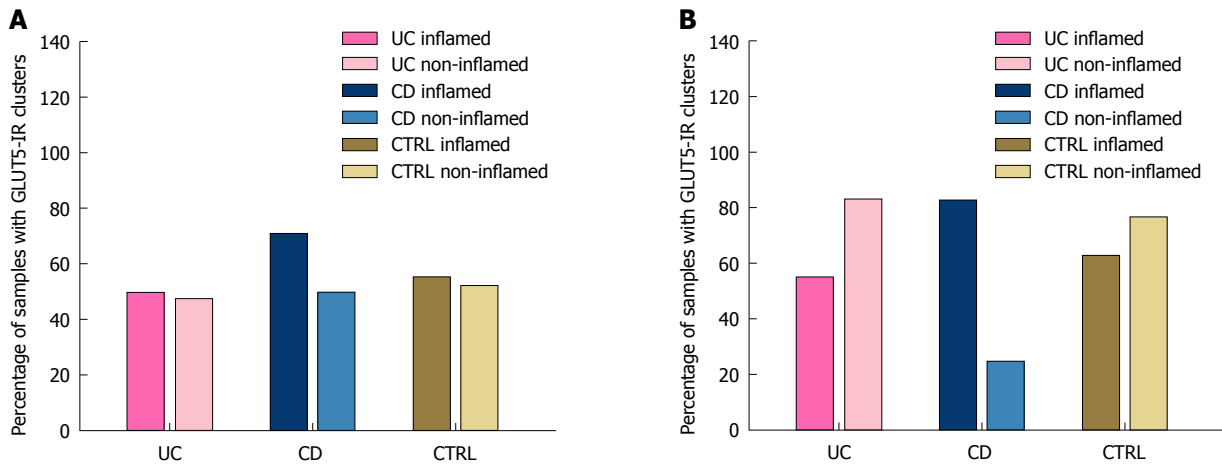


Figure 6 Bar graphs of the percentage of samples with GLUT5-immunoreactive clusters of lymphatic vessels in the proximal tract (Panel A) and distal tract (Panel B) of large intestine of patients with ulcerative colitis, Crohn's disease, and controls. Inflamed and non-inflamed mucosa can be distinguished for each patient group. In the UC and CTRL groups, the percentage is similar for inflamed and non-inflamed mucosa in the proximal tract, and higher for non-inflamed than inflamed mucosa in the distal tract. In the CD group, the percentage is higher for inflamed than non-inflamed mucosa in both the proximal and the distal tract. UC: Ulcerative colitis; CD: Crohn's disease; CTRL: Controls.

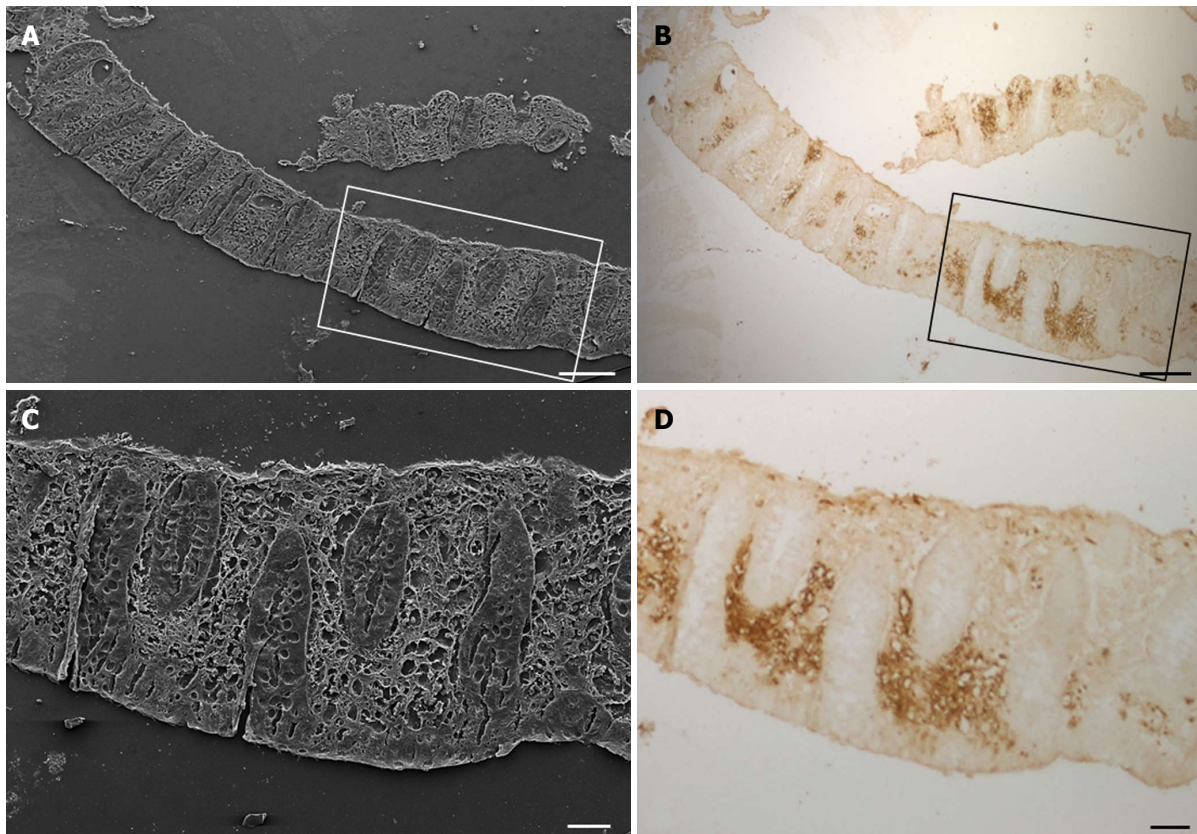


Figure 7 Scanning electron (A and C) and light (B and D) microscopy images of a sample of descending colon with non-inflamed mucosa after immunoperoxidase GLUT5 staining from a ulcerative colitis patient. Scale bar: (A and B) 250 μm ; (C and D) 100 μm .

between the different patient groups. Nevertheless, we consider this finding important as it provides the first evidence for the distribution of glucose transporter immunoreactivity throughout the epithelial tract of the human large intestine. Because of the limited expression of the glucose transporters, we were unable to compare their expression across the three patient groups. However, we believe that our data support

previous findings. In humans, GLUT5 expression has been observed in normal colonic cells; expression of both GLUT2 and GLUT5 has been reported in colon carcinoma and expression of GLUT5 with higher levels of staining than in normal colon cells^[23]. GLUT2 and GLUT5 have been found to be overexpressed in many other types of tumors. Since both transporters are involved in fructose uptake, it was suggested that

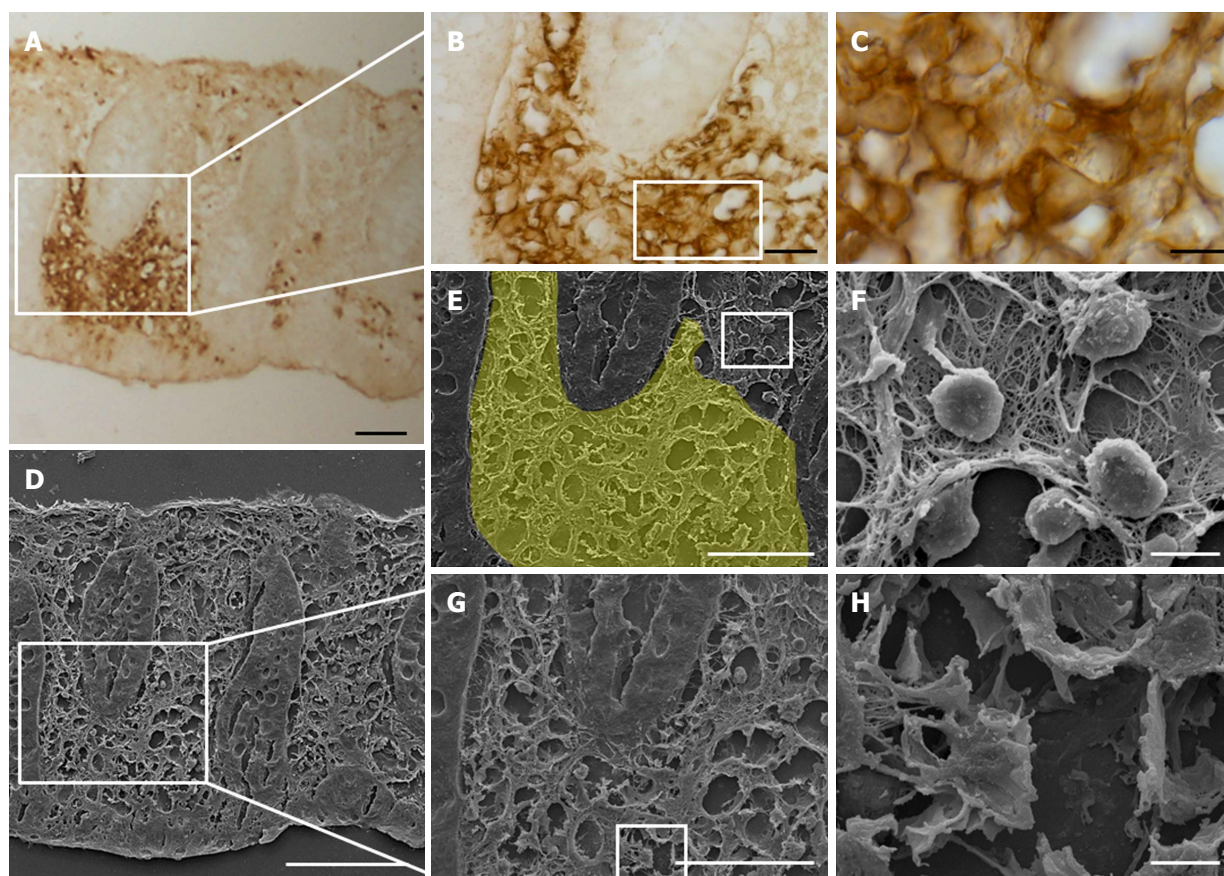


Figure 8 Light (A-C) and scanning electron (D-H) microscopy images of a sample of descending colon with non-inflamed mucosa after immunoperoxidase GLUT5 staining from a UC patient. The boxed areas in (A) and (D) are corresponding areas and are shown enlarged in (B) and (G), respectively. The yellow area in (E) represents the immunostaining area shown in (B). The boxed area in (B, E, G) is shown at higher magnification in (C, F, H, respectively). Scale bar: (D) 100 μ m; (A, E, G) 50 μ m; (B) 25 μ m; (C, F, H) 5 μ m.

tumor cells utilize fructose as an energy substrate^[23].

On the other hand, while excessive, chronic consumption of the GLUT5 and GLUT2 substrate fructose has been associated with numerous diseases and syndromes, including hypertension, obesity, diabetes, hyperinsulinemia, and non-alcoholic fatty liver disease, the role of fructose transporters in causing or contributing to the diseases is still unclear^[10].

In addition, we also identified GLUT5 expression in vessels, mainly concentrated in specific areas where the vessels were clustered. This staining was detected in the samples from IBD and control patients. Furthermore, immunohistochemical and SEM evaluation revealed that the vessels in the GLUT5-IR clusters had a reticular architecture, indistinct and dilated lumen, and were labeled on the endothelium that appeared to be fragmented. Large spaces between the vessels were present. Immunostaining with LYVE-1, a specific marker of lymphatic endothelium, and GLUT5 antibodies in contiguous sections revealed that GLUT5-IR clusters of vessels were concentrated in areas that were well-circumscribed but internal to those that were LYVE-1 positive. This finding was also confirmed in double staining studies. GLUT5 and LYVE-1 did not appear to be colocalized but rather showed a

close topographical relationship on the endothelium lining the lumen. Based on their LYVE-1 expression, GLUT5-IR vessels were unambiguously identified as lymphatic. We interpreted the presence of clusters as an impaired pattern of lymphatic capillary networks, probably related to proliferative zones, and consisting of vessels with morphological characteristics resembling those of immature lymphatics. They differed markedly from the vessels that we classified as "normal", which were characterized by a well-defined lumen lined by endothelial cells, a rounded appearance, and a non-aggregated distribution.

To the best of our knowledge, this is the first study to provide evidence that GLUT5 expression is associated with lymphatic vessels in controls and UC and CD patients. This novel finding yields further insight into the characterization of lymphatic vasculature, whose dysfunction is a long-recognized feature in humans with IBD.

There is considerable evidence for the proliferative expansion of lymphatic vessels during the course of inflammatory disease^[39-41]. Alteration and remodeling of the lymphatic system, besides blood vessel angiogenesis, are implicated in IBD^[42-44]. Increased lymphatic vessels density is a well-established feature in biopsy

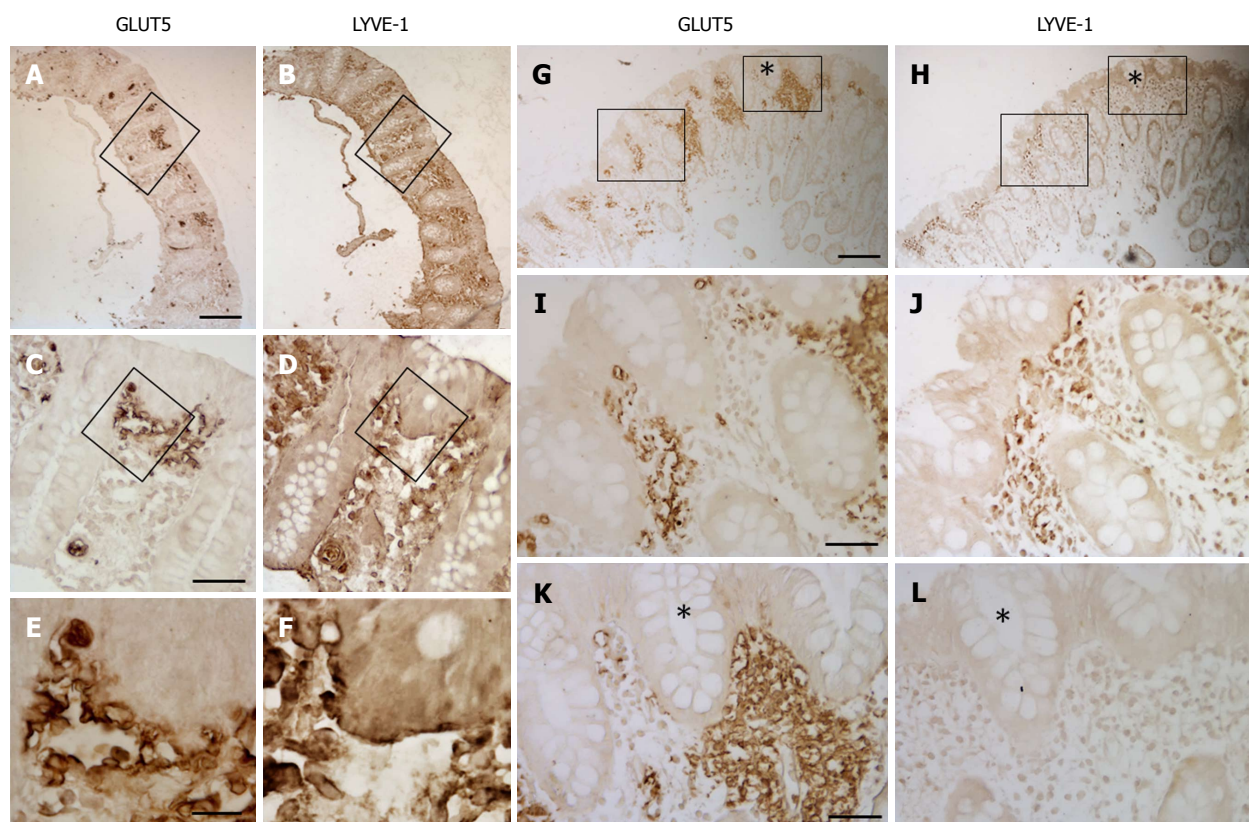


Figure 9 Immunoperoxidase staining showing GLUT5 and LYVE-1 immunoreactivity in samples of the distal tract of large intestine from UC (A-F) and CD (G-L) patients. In both samples, the mucosa was non-inflamed. The boxed area in (A and B) is enlarged in (C and D, respectively), the area in (C and D) is enlarged in (E and F, respectively), the area without an asterisk in (G and H) is enlarged in (I and J, respectively), the area with an asterisk in (G and H) is enlarged in (K and L, respectively). Scale bar: (A, B, G, H) 100 μ m; (C, D, I-L) 25 μ m; (E and F) 10 μ m.

samples from CD and UC patients and is correlated with disease severity^[42,44-47]. Similarly, a reduced density of lymphatic vessels was demonstrated to be associated with CD recurrence^[46].

Moreover, studies have reported that lymphatic vessel dysfunction contributes to perpetuating intestinal inflammation in both UC and CD patients and that the intestinal lymphatic system can profoundly influence gut immune homeostasis^[48,49]. However, it is still unclear whether expansion of the lymphatic network in IBD is a defensive mechanism or contributes to worsening the condition. It can be interpreted as a protective/adaptive, local response linked to the necessity to reduce fluid accumulation and to remove infiltrated immune cells, thus limiting further tissue injury. But lymphatic vessels also undergo morphological and functional changes in inflammatory conditions that impair their ability to drain fluid^[50].

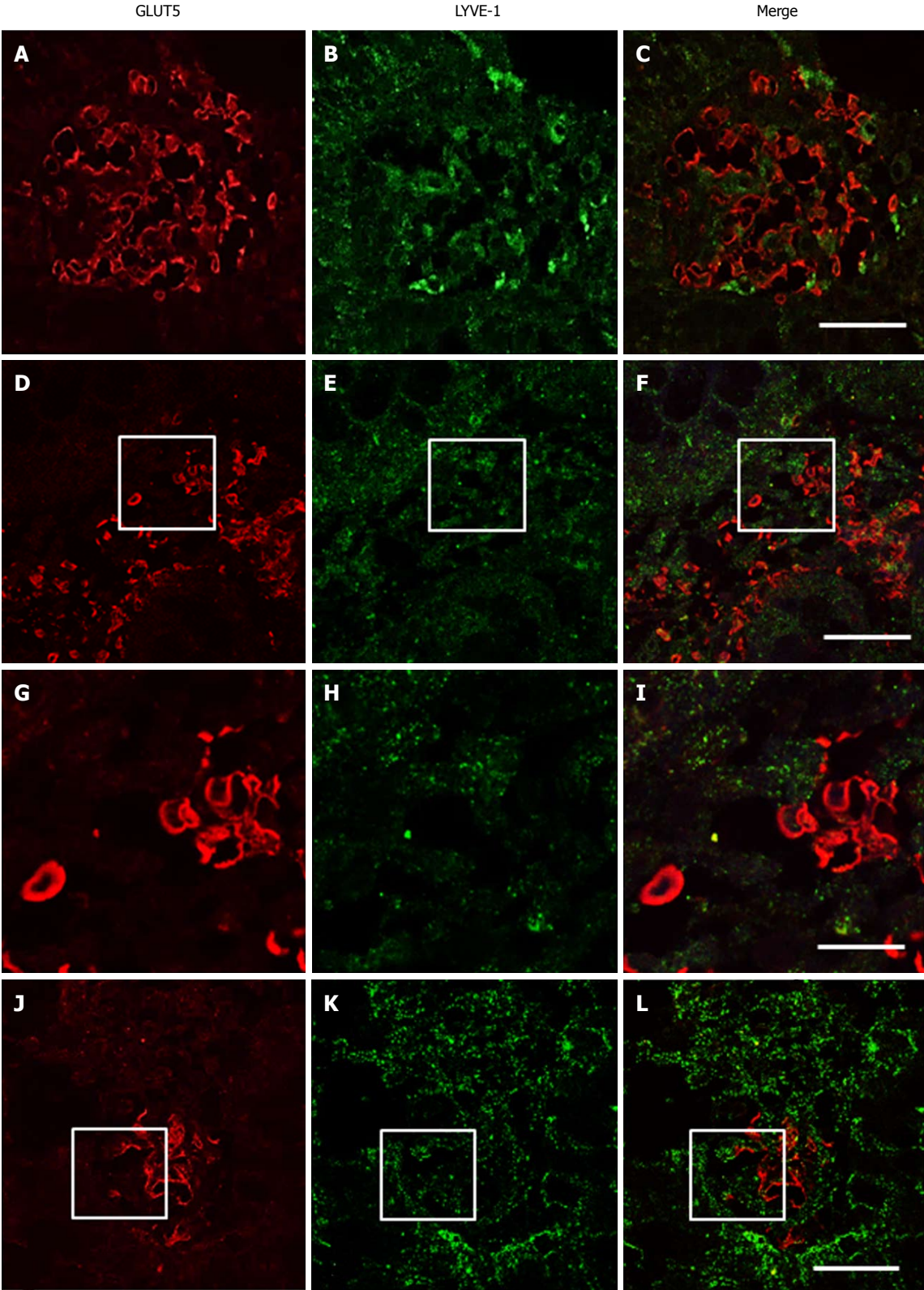
Lymphangiectasia is typically observed in the colonic mucosa of CD and UC patients^[44,45], and impaired intestinal lymphatic transport function was found to play a role in the early-stages and development of IBD^[51,52]. The relevance of the lymphatic transport function in IBD was also demonstrated in an experimental murine model that showed that acute colitis was aggravated by compromised lymphatic function; in contrast, immune cell clearance, fluid transport, and course of the disease

were improved by induction of lymphangiogenesis^[53,54].

While it is beyond doubt that increased lymphatic vessel density is closely linked to inflammatory disease, whether this process in IBD is an event that precedes inflammation or is its direct consequence remains to be elucidated.

Here we demonstrated a relevant presence of GLUT5-IR clusters of lymphatic vessels in both inflamed and non-inflamed tissue samples of large intestine from IBD and CTRL patients. The intestinal distal tract was, by far, the area most affected by their presence, especially in the UC and CTRL groups. This finding is consistent with the abundance of lymphatic tissue in the most distal regions of the intestine^[50].

To our surprise, we found more non-inflamed than inflamed tissue samples expressing GLUT5-IR clusters in the distal tract of the UC patients and controls: these percentages were similar across all three patient groups when classified by BMI. Differently, in the CD patients, similar results were found for the proximal and the distal intestinal tract, and the number of inflamed samples with GLUT5-IR clusters was higher than non-inflamed tissue samples, also in relation to the BMI of patients. Our data indicate that GLUT5-IR clusters are present in both UC patients and controls, and that their presence does not appear to be conditioned by the level of inflammation of the mucosal area where the



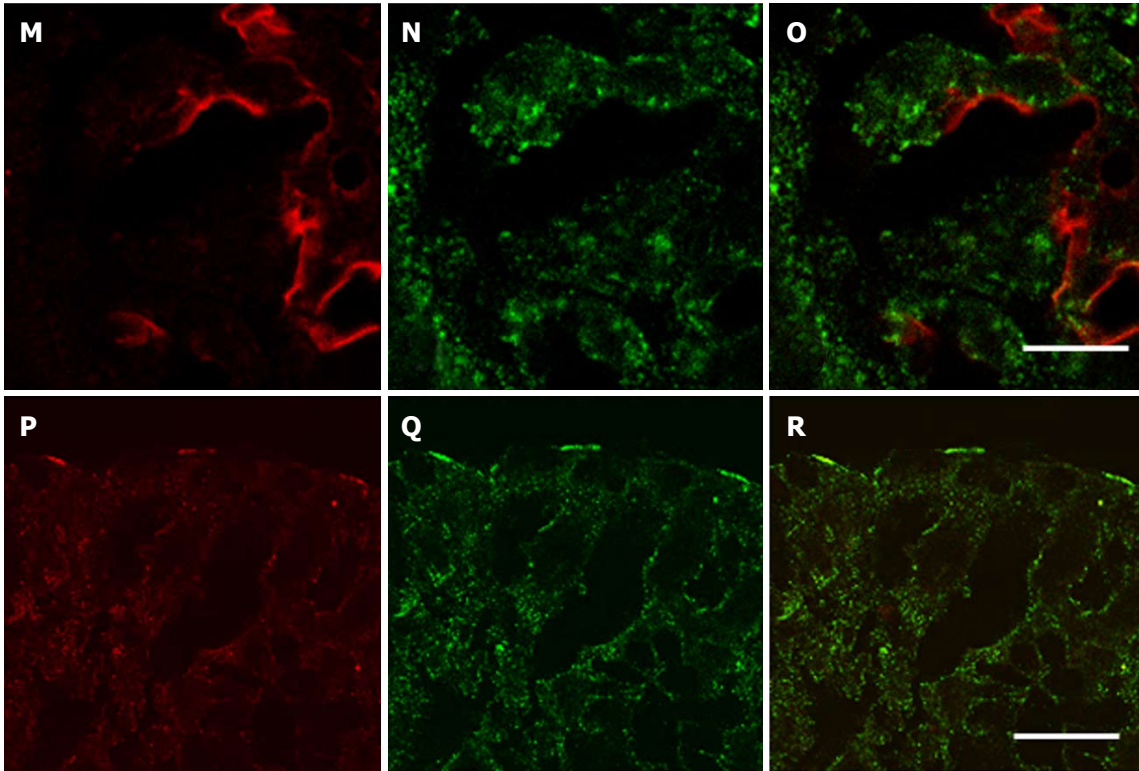


Figure 10 Double-immunofluorescent confocal microscopy showing expression of GLUT5 (red) with LYVE-1 (green) in samples of descending colon from UC (A-C, J-R) and CD (D-I) patients. The mucosa was inflamed in the UC sample and non-inflamed in the CD sample. The boxed area in (D-F) is shown at higher magnification in (G-I, respectively); the boxed area in (J-L) is shown at higher magnification in (M-O, respectively). Scale bar: (C, F, L, R) 20 μ m; (I, O) 10 μ m.

clusters are found. The opposite finding in the CD group may indicate a particular aspect of this condition. Since the amount of tissue analyzed in the CD patients was limited, it may not be completely representative of the whole group, however.

The few studies that have investigated the relationship between lymphoproliferation and inflammation of the area where it occurs have shown that lymphatic proliferation and inflammation do not occur in close association. Several explanations for this have been proposed. Increased lymphatic vessel density has been documented as a persistent feature in samples from IBD patients, in which it was also observed in fibrotic specimens from end-stage disease characterized by low inflammation^[42]. Similarly, the increased lymphatic vessel density detected in both the inflamed and non-inflamed tissue samples from the UC and CD patients has been interpreted as a process that occurs before the disease becomes clinically active, and therefore can manifest itself independent of inflammation^[46]. No obvious correlation between increased lymphatic vessel density and inflammation was demonstrated in the colonic mucosa of the UC patients, where it has been predicted that structural changes in the mucosa, including “*muscularis* mucosa expansion, infiltration by muscle fibers, and filiform epithelial changes” are an essential prerequisite to justify lymphatic proliferation^[45].

Both these interpretations may also be consistent with our data: the first is especially valid for the controls

where the presence of vessel clusters can be a sub-clinical aspect, at times not necessarily related to a particular pathology. The second can be valid for IBD: we sometimes observed vacuolar structures in the epithelium over the areas where lymphatic vessel clusters were present and altered epithelial cell integrity. Recent studies have shown that epithelial barrier dysfunction in IBD can contribute to disease progression by activating immunoregulatory processes, probably following increased exposure of the mucosa to the luminal microbiota or their products^[55-58]. Although the underlying mechanism is unclear, defects in intestinal barrier function are now attributed a direct or indirect role in the onset or progression of IBD. We observed morphological epithelial damage limited to certain areas of the epithelium in some samples and only at the light microscope, in which the real damage may be underestimated.

Other mechanisms/factors have been investigated to explain the stimulation of lymphatic vessels in IBD, given that it may occur via different processes in UC and CD. Lymphatic system expansion during inflammation in peripheral tissues progresses through the expansion and proliferation of an existing vascular network or sprouting of new vessels (lymphangiogenesis)^[40,59]. However, the expression of lymphangiogenic factors (*e.g.*, VEGF-C, VEGF-D) and cytokines (*e.g.*, TNF and IL-1b), which have been found to be increased in inflammatory conditions, were decreased in the

inflamed and non-inflamed colonic tissues of the IBD patients^[46,60].

LYVE-1 has also been demonstrated to be selectively expressed in inflammation. In cultured primary lymphatic endothelial cells, LYVE-1 uptake and degradation and inhibition of its gene expression were seen to occur under the stimulus of pro-inflammatory cytokines, leading to a subsequent reversible loss of the surface expression of LYVE-1^[61]. Subsequently, quantification of LYVE-1 by RT-PCR in human colonic mucosa revealed a lower mRNA level in both the non-inflamed and inflamed samples from the UC patients than in the controls, but no differences in LYVE-1 mRNA levels were seen in the CD patients^[46]. In the UC patients, the reduction was more evident in the non-inflamed than the inflamed samples.

In this study, double-immunofluorescence staining revealed that GLUT5-IR clusters were present in zones with LYVE-1 positivity, however, the appearance of the vessels inside the clusters was very heterogeneous, as was their phenotypic expression: GLUT5-positive but LYVE-1 negative or GLUT5-negative but LYVE-1 positive vessels were often identified, whereas the simultaneous expression of LYVE-1 and GLUT5 was limited to restricted endothelial areas of some clustered vessels. In the cases where LYVE-1 was missing, we saw the expression of GLUT5, which appeared in close continuity with LYVE-1, without ever being colocalized with it. These data strongly suggest a reduced expression of LYVE-1, which may depend on the different degrees of vessel differentiation within the clusters. There is considerable evidence that immature vessels lack LYVE-1 expression, particularly intratumoral lymphatic vessels, that differ morphologically from lymphatic vessels found within normal tissue^[62,63]. The reason for this difference is unclear; however, it has been also demonstrated that altered expression of lymphatic endothelial markers such as LYVE-1 may compromise lymphatic transport function^[50].

Collectively, our results regarding GLUT5 expression in IBD patients are consistent with previous data obtained for lymphatic proliferation in IBD. The presence of GLUT5 alone does not imply that it is involved in lymphatic proliferation because it was found to be expressed in both normal and clustered lymphatic vessels. However, its expression indicates that GLUT5 may play a role in controlling the formation of lymphatic vessels. It is plausible that aberrant lymphatic growth in clusters may result from the effect of GLUT5 on cellular mechanisms underlying the differentiation of new lymphatic vessels. It may also be that fructose, GLUT5 being its main transporter, may be the energy substrate that endothelial cells use for lymphatic expansion.

In this regard, we speculate that fructose present in the lumen may have some effect on the proliferation of these vessels. We found that a percentage of samples expressed GLUT5 in the brush border membrane of epithelial cells from both the UC and CTRL patients.

This finding merits further investigation because it could be a clear example of crosstalk between the cells projecting into the intestinal lumen and underlying environments.

In conclusion, our study provides evidence that GLUT2, SGLT1, and GLUT5 glucose transporters are expressed in the epithelial cells of the mucosa of the large intestine of IBD and control patients. Furthermore, both inflamed and non-inflamed mucosal colorectal tissue biopsies from the IBD and control patients showed GLUT5-IR lymphatic vessel clusters. This finding aids in the characterization of lymphatic vessels in IBD. As GLUT5 is the main fructose transporter in the human intestine, it is conceivable that fructose is a metabolic substrate that has a role in the atypical aggregation of lymphatic vessels.

Quantification of lymphatic vessel density in the gastrointestinal tract, particularly in cancer and chronic inflammatory disease, holds interest because it can be predictive of the early stages of disease. In this regard, GLUT5 expression on endothelial lymphovascular cells may have implications for routine use in the histopathological evaluation of lymphangiogenesis, also in combination with LYVE-1, a marker of lymphatic endothelium that can be down-modulated under inflammatory conditions.

A future area of focus is the precise function of GLUT5 expression in lymphatic vessels and the factors that interfere with the development or maintenance of IBD.

ARTICLE HIGHLIGHTS

Research background

Glucose transporter expression is present throughout the digestive system, greatest in the small intestine and least in the terminal ileum. Most studies to date on glucose transporters in the gastrointestinal tract have been performed using biochemical rather than standard immunohistochemical analysis also in humans.

Research motivation

The localization and distribution of glucose transporters in the intestinal mucosa of the human colon has not yet been clarified despite evidences that support the role of glucose transporters in broad areas of pure glucose absorption and metabolism, such as inflammation, malignancy, and gut microbiota regulation.

Research objectives

The aim of this study was to investigate by immunostaining at light and confocal microscopy the expression of the major intestinal glucose transporters (GLUT2, SGLT1, GLUT5) in human colonic mucosa in control subjects and subjects with IBD.

Research methods

Patients diagnosed with ulcerative colitis or Crohn's disease and scheduled for diagnostic colonoscopy were enrolled. Patients who underwent colonoscopy for prevention screening of colorectal cancer or were followed-up after polypectomy or lower gastrointestinal symptoms were designated as the control group. Colorectal samples were obtained from patients undergoing lower endoscopic colonoscopy or recto-sigmoidoscopy. Biopsies of portions of the colonic tract (cecum, ascending colon, transverse, descending, sigmoid colon, rectum) were taken for diagnostic purposes according to the endoscopist's judgment and for

immunohistochemistry. Inflammatory status of the mucosa at the sampling site was evaluated endoscopically and histologically.

Samples were fixed in formaldehyde and embedded in paraffin. The expression of GLUT2, SGLT1, and GLUT5 glucose transporters was investigated using immunoperoxidase labeling. Immunoreactivity of GLUT5 was compared with that of LYVE-1, which is a marker for lymphatic vessel endothelium, using double-labeled confocal microscopy.

Research results

GLUT2, SGLT1, and GLUT5 glucose transporter immunostaining was found in short epithelial portions of the large intestine from IBD and control patients. No difference in glucose transporter expression was observed between the samples obtained from the proximal and distal tracts and between the different patient groups. GLUT5 immunostaining was also detected in vessels, which were mainly concentrated in specific areas. In double fluorescent-labeled sections with GLUT5 and LYVE-1, GLUT5-immunoreactive clusters of vessels were concentrated in areas internal to those that were LYVE-1 positive. The GLUT5 and LYVE-1 labeling patterns were never colocalized but rather showed a close topographical relationship on the endothelium lining the lumen. Based on their LYVE-1 expression, GLUT5 immunoreactive vessels were identified as lymphatic and observed both in inflamed and non-inflamed mucosal colorectal tissue biopsies from the IBD and CTRL patients. This novel finding yields further insight into the characterization of lymphatic vasculature, whose dysfunction is a long-recognized feature in humans with IBD.

Research conclusions

This study provides evidence that GLUT2, SGLT1, and GLUT5 glucose transporters are expressed in the colorectal mucosa in controls and IBD patients. Furthermore, it provides first evidence that GLUT5 expression is associated with lymphatic vessels in controls and IBD patients. Its expression indicates that GLUT5 may play a role in controlling the formation of lymphatic vessels. GLUT5 expression on endothelial lymphovascular cells may have implications for routine use in the histopathological evaluation of lymphangiogenesis, also in combination with LYVE-1, a marker of lymphatic endothelium that can be down-modulated under inflammatory conditions.

Research perspectives

A future area of focus is the precise function of GLUT5 expression in lymphatic vessels and the factors that interfere with the development or maintenance of IBD.

REFERENCES

- Mueckler M, Caruso C, Baldwin SA, Panico M, Blench I, Morris HR, Allard WJ, Lienhard GE, Lodish HF. Sequence and structure of a human glucose transporter. *Science* 1985; **229**: 941-945 [PMID: 3839598 DOI: 10.1126/science.3839598]
- Kayano T, Burant CF, Fukumoto H, Gould GW, Fan YS, Eddy RL, Byers MG, Shows TB, Seino S, Bell GI. Human facilitative glucose transporters. Isolation, functional characterization, and gene localization of cDNAs encoding an isoform (GLUT5) expressed in small intestine, kidney, muscle, and adipose tissue and an unusual glucose transporter pseudogene-like sequence (GLUT6). *J Biol Chem* 1990; **265**: 13276-13282 [PMID: 1695905]
- Blakemore SJ, Aledo JC, James J, Campbell FC, Lucocq JM, Hundal HS. The GLUT5 hexose transporter is also localized to the basolateral membrane of the human jejunum. *Biochem J* 1995; **309** (Pt 1): 7-12 [PMID: 7619085]
- Douard V, Cui XL, Soteropoulos P, Ferraris RP. Dexamethasone sensitizes the neonatal intestine to fructose induction of intestinal fructose transporter (Slc2A5) function. *Endocrinology* 2008; **149**: 409-423 [PMID: 17947353 DOI: 10.1210/en.2007-0906]
- Matosin-Matekalo M, Mesonero JE, Laroche TJ, Lacasa M, Brot-Laroche E. Glucose and thyroid hormone co-regulate the expression of the intestinal fructose transporter GLUT5. *Biochem J* 1999; **339** (Pt 2): 233-239 [PMID: 10191252]
- David ES, Cingari DS, Ferraris RP. Dietary induction of intestinal fructose absorption in weaning rats. *Pediatr Res* 1995; **37**: 777-782 [PMID: 7651763 DOI: 10.1203/00006450-199506000-00017]
- Douard V, Ferraris RP. Regulation of the fructose transporter GLUT5 in health and disease. *Am J Physiol Endocrinol Metab* 2008; **295**: E227-E237 [PMID: 18398011 DOI: 10.1152/ajpendo.90245.2008]
- Litherland GJ, Hajdich E, Gould GW, Hundal HS. Fructose transport and metabolism in adipose tissue of Zucker rats: diminished GLUT5 activity during obesity and insulin resistance. *Mol Cell Biochem* 2004; **261**: 23-33 [PMID: 15362482]
- Dyer J, Wood IS, Palejwala A, Ellis A, Shirazi-Beechey SP. Expression of monosaccharide transporters in intestine of diabetic humans. *Am J Physiol Gastrointest Liver Physiol* 2002; **282**: G241-G248 [PMID: 11804845 DOI: 10.1152/ajpgi.00310.2001]
- Douard V, Ferraris RP. The role of fructose transporters in diseases linked to excessive fructose intake. *J Physiol* 2013; **591**: 401-414 [PMID: 23129794 DOI: 10.1113/jphysiol.2011.215731]
- Thorens B, Cheng ZQ, Brown D, Lodish HF. Liver glucose transporter: a basolateral protein in hepatocytes and intestine and kidney cells. *Am J Physiol* 1990; **259**: C279-C285 [PMID: 1701966]
- Yoshikawa T, Inoue R, Matsumoto M, Yajima T, Ushida K, Iwanaga T. Comparative expression of hexose transporters (SGLT1, GLUT1, GLUT2 and GLUT5) throughout the mouse gastrointestinal tract. *Histochem Cell Biol* 2011; **135**: 183-194 [PMID: 21274556 DOI: 10.1007/s00418-011-0779-1]
- Kellett GL, Brot-Laroche E, Mace OJ, Leturque A. Sugar absorption in the intestine: the role of GLUT2. *Annu Rev Nutr* 2008; **28**: 35-54 [PMID: 18393659 DOI: 10.1146/annurev.nutr.28.061807.155518]
- Gouyon F, Caillaud L, Carriere V, Klein C, Dalet V, Citadelle D, Kellett GL, Thorens B, Leturque A, Brot-Laroche E. Simple-sugar meals target GLUT2 at enterocyte apical membranes to improve sugar absorption: a study in GLUT2-null mice. *J Physiol* 2003; **552**: 823-832 [PMID: 12937289 DOI: 10.1113/jphysiol.2003.049247]
- Kellett GL, Helliwell PA. The diffusive component of intestinal glucose absorption is mediated by the glucose-induced recruitment of GLUT2 to the brush-border membrane. *Biochem J* 2000; **350** Pt 1: 155-162 [PMID: 10926839 DOI: 10.1042/0264-6021:3500155]
- Ait-Omar A, Monteiro-Sepulveda M, Poitou C, Le Gall M, Cotillard A, Gilet J, Garbin K, Houllier A, Château D, Lacombe A, Veyrie N, Hugol D, Tordjman J, Magnan C, Serradas P, Clément K, Leturque A, Brot-Laroche E. GLUT2 accumulation in enterocyte apical and intracellular membranes: a study in morbidly obese human subjects and ob/ob and high fat-fed mice. *Diabetes* 2011; **60**: 2598-2607 [PMID: 21852673 DOI: 10.2337/db10-1740]
- Reimann F, Habib AM, Tolhurst G, Parker HE, Rogers GJ, Gribble FM. Glucose sensing in L cells: a primary cell study. *Cell Metab* 2008; **8**: 532-539 [PMID: 19041768 DOI: 10.1016/j.cmet.2008.11.002]
- Depoortere I. Taste receptors in the gut tune the release of peptides in response to nutrients. *Peptides* 2015; **66**: 9-12 [PMID: 25683908 DOI: 10.1016/j.peptides.2015.01.013]
- Schmitt CC, Aranas T, Viel T, Chateau D, Le Gall M, Waligora-Dupriet AJ, Melchior C, Rouxel O, Kapel N, Gourcerol G, Tavitian B, Lehuen A, Brot-Laroche E, Leturque A, Serradas P, Grosfeld A. Intestinal invalidation of the glucose transporter GLUT2 delays tissue distribution of glucose and reveals an unexpected role in gut homeostasis. *Mol Metab* 2016; **6**: 61-72 [PMID: 28123938 DOI: 10.1016/j.molmet.2016.10.008]
- Wright EM, Turk E. The sodium/glucose cotransport family SLC5. *Pflügers Arch* 2004; **447**: 510-518 [PMID: 12748858 DOI: 10.1007/s00424-003-1063-6]
- Hediger MA, Coady MJ, Ikeda TS, Wright EM. Expression cloning and cDNA sequencing of the Na⁺/glucose co-transporter. *Nature* 1987; **330**: 379-381 [PMID: 2446136 DOI: 10.1038/330379a0]
- Gorboulev V, Schürmann A, Vallon V, Kipp H, Jaschke A, Klessen D, Friedrich A, Scherneck S, Rieg T, Cunard R, Veyhl-Wichmann M, Srinivasan A, Balen D, Breljak D, Rexhepaj R, Parker HE, Gribble FM, Reimann F, Lang F, Wiese S, Sabolic I, Sendtner M, Koepsell H. Na⁺(+)-D-glucose cotransporter SGLT1

- is pivotal for intestinal glucose absorption and glucose-dependent incretin secretion. *Diabetes* 2012; **61**: 187-196 [PMID: 22124465 DOI: 10.2337/db11-1029]
- 23 **Godoy A**, Ulloa V, Rodríguez F, Reinicke K, Yañez AJ, García Mde L, Medina RA, Carrasco M, Barberis S, Castro T, Martínez F, Koch X, Vera JC, Poblete MT, Figueroa CD, Peruzzo B, Pérez F, Nualart F. Differential subcellular distribution of glucose transporters GLUT1-6 and GLUT9 in human cancer: ultrastructural localization of GLUT1 and GLUT5 in breast tumor tissues. *J Cell Physiol* 2006; **207**: 614-627 [PMID: 16523487 DOI: 10.1002/jcp.20606]
- 24 **Wuest M**, Trayner BJ, Grant TN, Jans HS, Mercer JR, Murray D, West FG, McEwan AJ, Wuest F, Cheeseman CI. Radiopharmacological evaluation of 6-deoxy-6-[18F]fluoro-D-fructose as a radiotracer for PET imaging of GLUT5 in breast cancer. *Nucl Med Biol* 2011; **38**: 461-475 [PMID: 21531283 DOI: 10.1016/j.nucmedbio.2010.11.004]
- 25 **Liu H**, Huang D, McArthur DL, Boros LG, Nissen N, Heaney AP. Fructose induces transketolase flux to promote pancreatic cancer growth. *Cancer Res* 2010; **70**: 6368-6376 [PMID: 20647326 DOI: 10.1158/0008-5472.CAN-09-4615]
- 26 **García-Herrera J**, Marca MC, Brot-Laroche E, Guillén N, Acín S, Navarro MA, Osada J, Rodríguez-Yoldi MJ. Protein kinases, TNF- α , and proteasome contribute in the inhibition of fructose intestinal transport by sepsis in vivo. *Am J Physiol Gastrointest Liver Physiol* 2008; **294**: G155-G164 [PMID: 17962360 DOI: 10.1152/ajpgi.00139.2007]
- 27 **García-Herrera J**, Abad B, Rodríguez-Yoldi MJ. Effect of lipopolysaccharide on D-fructose transport across rabbit jejunum. *Inflamm Res* 2003; **52**: 177-184 [PMID: 12755384]
- 28 **Barone S**, Fussell SL, Singh AK, Lucas F, Xu J, Kim C, Wu X, Yu Y, Amlal H, Seidler U, Zuo J, Soleimani M, Slc2a5 (Glut5) is essential for the absorption of fructose in the intestine and generation of fructose-induced hypertension. *J Biol Chem* 2009; **284**: 5056-5066 [PMID: 19091748 DOI: 10.1074/jbc.M808128200]
- 29 **Uldry M**, Ibberson M, Hosokawa M, Thorens B. GLUT2 is a high affinity glucosamine transporter. *FEBS Lett* 2002; **524**: 199-203 [PMID: 12135767 DOI: 10.1016/S0014-5793(02)03058-2]
- 30 **Yu LC**, Flynn AN, Turner JR, Buret AG. SGLT-1-mediated glucose uptake protects intestinal epithelial cells against LPS-induced apoptosis and barrier defects: a novel cellular rescue mechanism? *FASEB J* 2005; **19**: 1822-1835 [PMID: 16260652 DOI: 10.1096/fj.05-4226com]
- 31 **Barrenetxe J**, Sánchez O, Barber A, Gascón S, Rodríguez-Yoldi MJ, Lostao MP. TNF α regulates sugar transporters in the human intestinal epithelial cell line Caco-2. *Cytokine* 2013; **64**: 181-187 [PMID: 23910014 DOI: 10.1016/j.cyt.2013.07.004]
- 32 **Palazzo M**, Gariboldi S, Zanolio L, Salleri S, Dusio GF, Mauro V, Rossini A, Balsari A, Rumio C. Sodium-dependent glucose transporter-1 as a novel immunological player in the intestinal mucosa. *J Immunol* 2008; **181**: 3126-3136 [PMID: 18713983 DOI: 10.4049/jimmunol.181.5.3126]
- 33 **Yomogida S**, Kojima Y, Tsutsumi-Ishii Y, Hua J, Sakamoto K, Nagaoka I. Glucosamine, a naturally occurring amino monosaccharide, suppresses dextran sulfate sodium-induced colitis in rats. *Int J Mol Med* 2008; **22**: 317-323 [PMID: 18698490 DOI: 10.3892/ijmm.00000025]
- 34 **Reed KL**, Fruin AB, Gower AC, Gonzales KD, Stucchi AF, Andry CD, O'Brien M, Becker JM. NF-kappaB activation precedes increases in mRNA encoding neurokinin-1 receptor, proinflammatory cytokines, and adhesion molecules in dextran sulfate sodium-induced colitis in rats. *Dig Dis Sci* 2005; **50**: 2366-2378 [PMID: 16416193 DOI: 10.1007/s10620-005-3066-y]
- 35 **Visekruna A**, Joeris T, Seidel D, Kroesen A, Loddenkemper C, Zeitz M, Kaufmann SH, Schmidt-Ullrich R, Steinhoff U. Proteasome-mediated degradation of IkappaBalpha and processing of p105 in Crohn disease and ulcerative colitis. *J Clin Invest* 2006; **116**: 3195-3203 [PMID: 17124531 DOI: 10.1172/JCI28804]
- 36 **Lewis Carl SA**, Gillette-Ferguson I, Ferguson DG. An indirect immunofluorescence procedure for staining the same cryosection with two mouse monoclonal primary antibodies. *J Histochem Cytochem* 1993; **41**: 1273-1278 [PMID: 7687266 DOI: 10.1177/41.8.7687266]
- 37 **Negoescu A**, Labat-Moleur F, Lorimier P, Lamarcq L, Guillemet C, Chambaz E, Brambilla E. F(ab) secondary antibodies: a general method for double immunolabeling with primary antisera from the same species. Efficiency control by chemiluminescence. *J Histochem Cytochem* 1994; **42**: 433-437 [PMID: 7508473 DOI: 10.1177/42.3.7508473]
- 38 **Merigo F**, Benati D, Cristofolletti M, Osculati F, Sbarbati A. Glucose transporters are expressed in taste receptor cells. *J Anat* 2011; **219**: 243-252 [PMID: 21592100 DOI: 10.1111/j.1469-7580.2011.01385.x]
- 39 **Jackson DG**. Biology of the lymphatic marker LYVE-1 and applications in research into lymphatic trafficking and lymphangiogenesis. *APMIS* 2004; **112**: 526-538 [PMID: 15563314 DOI: 10.1111/j.1600-0463.2004.apm11207-0811.x]
- 40 **Alitalo K**, Tammela T, Petrova TV. Lymphangiogenesis in development and human disease. *Nature* 2005; **438**: 946-953 [PMID: 16355212 DOI: 10.1038/nature04480]
- 41 **Kerjaschki D**. The lymphatic vasculature revisited. *J Clin Invest* 2014; **124**: 874-877 [PMID: 24590271 DOI: 10.1172/JCI74854]
- 42 **Geleff S**, Schoppmann SF, Oberhuber G. Increase in podoplanin-expressing intestinal lymphatic vessels in inflammatory bowel disease. *Virchows Arch* 2003; **442**: 231-237 [PMID: 12647212 DOI: 10.1007/s00428-002-0744-4]
- 43 **Danese S**, Fiocchi C. Etiopathogenesis of inflammatory bowel diseases. *World J Gastroenterol* 2006; **12**: 4807-4812 [PMID: 16937461 DOI: 10.3748/wjg.v12.i30.4807]
- 44 **Pedica F**, Ligorio C, Tonelli P, Bartolini S, Baccarini P. Lymphangiogenesis in Crohn's disease: an immunohistochemical study using monoclonal antibody D2-40. *Virchows Arch* 2008; **452**: 57-63 [PMID: 18040712 DOI: 10.1007/s00428-007-0540-2]
- 45 **Kaiserling E**, Kröber S, Geleff S. Lymphatic vessels in the colonic mucosa in ulcerative colitis. *Lymphology* 2003; **36**: 52-61 [PMID: 12926829]
- 46 **Rahier JF**, De Beauce S, Dubuquoy L, Erdual E, Colombel JF, Jouret-Mourin A, Geboes K, Desreumaux P. Increased lymphatic vessel density and lymphangiogenesis in inflammatory bowel disease. *Aliment Pharmacol Ther* 2011; **34**: 533-543 [PMID: 21736598 DOI: 10.1111/j.1365-2036.2011.04759.x]
- 47 **Fogt F**, Pascha TL, Zhang PJ, Gausas RE, Rahemtulla A, Zimmerman RL. Proliferation of D2-40-expressing intestinal lymphatic vessels in the lamina propria in inflammatory bowel disease. *Int J Mol Med* 2004; **13**: 211-214 [PMID: 14719125 DOI: 10.3892/ijmm.13.2.211]
- 48 **Vranova M**, Halin C. Lymphatic Vessels in Inflammation. *J Clin Cell Immunol* 2014; **5**: 250 [DOI: 10.4172/2155-9899.1000250]
- 49 **Becker F**, Yi P, Al-Kofahi M, Ganta VC, Morris J, Alexander JS. Lymphatic dysregulation in intestinal inflammation: new insights into inflammatory bowel disease pathomechanisms. *Lymphology* 2014; **47**: 3-27 [PMID: 25109166]
- 50 **Alexander JS**, Chaitanya GV, Grisham MB, Boktor M. Emerging roles of lymphatics in inflammatory bowel disease. *Ann N Y Acad Sci* 2010; **1207** Suppl 1: E75-E85 [PMID: 20961310 DOI: 10.1111/j.1749-6632.2010.05757.x]
- 51 **Von Der Weid PY**, Rehal S. Lymphatic pump function in the inflamed gut. *Ann N Y Acad Sci* 2010; **1207** Suppl 1: E69-E74 [PMID: 20961308 DOI: 10.1111/j.1749-6632.2010.05715.x]
- 52 **Linares PM**, Gisbert JP. Role of growth factors in the development of lymphangiogenesis driven by inflammatory bowel disease: a review. *Inflamm Bowel Dis* 2011; **17**: 1814-1821 [PMID: 21744436 DOI: 10.1002/ibd.21554]
- 53 **D'Alessio S**, Correale C, Tacconi C, Gandelli A, Pietrogrande G, Vetrano S, Genua M, Arena V, Spinelli A, Peyrin-Biroulet L, Fiocchi C, Danese S. VEGF-C-dependent stimulation of lymphatic function ameliorates experimental inflammatory bowel disease. *J Clin Invest* 2014; **124**: 3863-3878 [PMID: 25105363 DOI: 10.1172/JCI72189]

- 54 **Becker F**, Potepalov S, Shehzahdi R, Bernas M, Witte M, Abreo F, Traylor J, Orr WA, Tsunoda I, Alexander JS. Downregulation of FoxC2 Increased Susceptibility to Experimental Colitis: Influence of Lymphatic Drainage Function? *Inflamm Bowel Dis* 2015; **21**: 1282-1296 [PMID: 25822012 DOI: 10.1097/MIB.0000000000000371]
- 55 **Edelblum KL**, Turner JR. The tight junction in inflammatory disease: communication breakdown. *Curr Opin Pharmacol* 2009; **9**: 715-720 [PMID: 19632896 DOI: 10.1016/j.coph.2009.06.022]
- 56 **Berry D**, Reinisch W. Intestinal microbiota: a source of novel biomarkers in inflammatory bowel diseases? *Best Pract Res Clin Gastroenterol* 2013; **27**: 47-58 [PMID: 23768552 DOI: 10.1016/j.bpg.2013.03.005]
- 57 **Klag T**, Stange EF, Wehkamp J. Defective antibacterial barrier in inflammatory bowel disease. *Dig Dis* 2013; **31**: 310-316 [PMID: 24246980 DOI: 10.1159/000354858]
- 58 **Lanis JM**, Kao DJ, Alexeev EE, Colgan SP. Tissue metabolism and the inflammatory bowel diseases. *J Mol Med (Berl)* 2017; **95**: 905-913 [PMID: 28528514 DOI: 10.1007/s00109-017-1544-2]
- 59 **Wirzenius M**, Tammela T, Uutela M, He Y, Odorisio T, Zambruno G, Nagy JA, Dvorak HF, Ylä-Herttua S, Shibuya M, Alitalo K. Distinct vascular endothelial growth factor signals for lymphatic vessel enlargement and sprouting. *J Exp Med* 2007; **204**: 1431-1440 [PMID: 17535974 DOI: 10.1084/jem.20062642]
- 60 **Flister MJ**, Wilber A, Hall KL, Iwata C, Miyazono K, Nisato RE, Pepper MS, Zawieja DC, Ran S. Inflammation induces lymphangiogenesis through up-regulation of VEGFR-3 mediated by NF-kappaB and Prox1. *Blood* 2010; **115**: 418-429 [PMID: 19901262 DOI: 10.1182/blood-2008-12-196840]
- 61 **Johnson LA**, Prevo R, Clasper S, Jackson DG. Inflammation-induced uptake and degradation of the lymphatic endothelial hyaluronan receptor LYVE-1. *J Biol Chem* 2007; **282**: 33671-33680 [PMID: 17884820 DOI: 10.1074/jbc.M702889200]
- 62 **Beasley NJ**, Prevo R, Banerji S, Leek RD, Moore J, van Trappen P, Cox G, Harris AL, Jackson DG. Intratumoral lymphangiogenesis and lymph node metastasis in head and neck cancer. *Cancer Res* 2002; **62**: 1315-1320 [PMID: 11888898]
- 63 **Padera TP**, Kadambi A, di Tomaso E, Carreira CM, Brown EB, Boucher Y, Choi NC, Mathisen D, Wain J, Mark EJ, Munn LL, Jain RK. Lymphatic metastasis in the absence of functional intratumor lymphatics. *Science* 2002; **296**: 1883-1886 [PMID: 11976409 DOI: 10.1126/science.1071420]

P- Reviewer: Huerta-Franco MR, Maric I, Nakase H, Yu LCH

S- Editor: Gong ZM **L- Editor:** A **E- Editor:** Huang Y





Published by **Baishideng Publishing Group Inc**
7901 Stoneridge Drive, Suite 501, Pleasanton, CA 94588, USA
Telephone: +1-925-223-8242
Fax: +1-925-223-8243
E-mail: bpgoffice@wjgnet.com
Help Desk: <http://www.f6publishing.com/helpdesk>
<http://www.wjgnet.com>



ISSN 1007-9327

