

Case Control Study

Detection of fucosylated haptoglobin using the 10-7G antibody as a biomarker for evaluating endoscopic remission in ulcerative colitis

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Abstract**BACKGROUND**

Inflammatory bowel disease (IBD) is a chronic, relapsing inflammation of the digestive tract. Although fecal and serum biomarkers have been extremely important and supportive for monitoring of IBD, their low sensitivity and high variability characteristics limit clinical efficacy. Thus, the establishment of better biomarkers is expected. Fucosylation is one of the most important glycosylation modifications of proteins. Fucosylated haptoglobin (Fuc-Hpt) is used as a biomarker for several cancers and inflammation-related diseases. We recently established a novel glycan monoclonal antibody (mAb), designated 10-7G, which recognizes Fuc-Hpt. We developed an enzyme-linked immunosorbent assay (ELISA) to measure serum levels of Fuc-Hpt (10-7G values).

AIM

To investigate the usefulness of the serum 10-7G values as a potential biomarker

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for monitoring disease activity in IBD.

METHODS

This was a case control study. Intestinal tissues of IBD patients ($n = 10$) were examined immunohistochemically using the 10-7G mAb. We determined 10-7G values using serum from patients with ulcerative colitis (UC, $n = 110$), Crohn's disease ($n = 45$), acute enteritis (AE, $n = 11$), and healthy volunteers (HVs) who exhibited normal ($n = 20$) or high ($n = 79$) C-reactive protein (CRP) levels at medical check-up. We investigated the correlation between the 10-7G value and various clinical parameters of IBD patients by correlation analysis. Receiver operating characteristic (ROC) curve analysis was performed to evaluate the usefulness of the 10-7G values as a biomarker for clinical and endoscopic remission of UC compared to conventional serum biomarkers.

RESULTS

In the immunohistochemical analysis, positive 10-7G mAb staining was observed in lymphocytes infiltrating into inflammatory sites of the mucosal layer and lymphoid follicles. The 10-7G values were significantly higher in patients with IBD ($P < 0.001$) and AE ($P < 0.05$) compared with HVs. In addition, 10-7G values were correlated with clinical examination parameters related to inflammation in patients with UC, particularly the CRP level ($r_s = 0.525$, $P = 0.003$) and clinical activity index score ($r_s = 0.435$, $P = 0.038$). However, there was no correlation between 10-7G values and CRP in HVs with high CRP levels, suggesting that the 10-7G values is not the same as a general inflammation biomarker. ROC curve analysis showed that area under the curve (AUC) value of 10-7G values for the diagnosis of endoscopic remission was higher than other biomarkers (AUC value = 0.699).

CONCLUSION

The serum 10-7G value is a novel biomarker for evaluating intestinal inflammation and endoscopic mucosal healing in UC.

Key Words: Inflammatory bowel disease; Biomarkers; Glycosylation; Fucosylated haptoglobin; Prohaptoglobin; Endoscopic remission

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Core Tip: Fucosylation is one of the most important glycosylation involved in cancer and inflammation. Recently, we have succeeded to establish a novel glycan antibody 10-7G mAb which directly recognizes fucosylated haptoglobin (Fuc-Hpt), and developed its enzyme-linked immunosorbent assay (ELISA) system. In this study, we demonstrated the serum 10-7G values (Fuc-Hpt levels determined with 10-7G mAb ELISA) were significantly higher in patients with inflammatory bowel disease. We also found that Fuc-Hpt was produced by lymphocytes infiltrating into inflammatory sites of intestine. Moreover, statistical analyses showed that the 10-7G value could evaluate intestinal inflammation and endoscopic mucosal healing in ulcerative colitis.

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INTRODUCTION

Inflammatory bowel disease (IBD), which comprises Crohn's disease (CD) and

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ulcerative colitis (UC), is a chronic recurrent intestinal inflammatory disease of unknown etiology, and the number of individuals affected is increasing worldwide^[1]. Although the etiology of IBD remains unknown, the disease is known to involve complex interactions between genetic, environmental, and microbial factors and the immune response^[2,3]. For proper management and selection of appropriate treatment strategy, it is essential to evaluate disease activity of IBD accurately. Endoscopy is the gold standard for the diagnosis and monitoring status of IBD patients, but cannot be frequently performed because of the cost and invasiveness. Although fecal and serum biomarkers, such as fecal calprotectin and serum C-reactive protein (CRP), have been important and supportive for monitoring of IBD, their low sensitivity and high variability characteristics limit clinical efficacy^[4,5]. Thus, the establishment of better biomarkers is expected.

Haptoglobin, an acute-phase protein produced primarily in the liver, is rapidly upregulated following stimulation with cytokines such as tumor necrosis factor alpha, interleukin (IL)-6, and IL-1^[6]. Haptoglobin modulates inflammation *via* its systemic anti-inflammatory activity^[7]. A major function of haptoglobin is binding free plasma hemoglobin produced due to hemolysis, thereby preventing damage associated with oxidative stress^[8]. Haptoglobin is an α - and β -chain heterodimer linked by disulfide bonds, with all four N-glycans located in the β chain and no potential N-glycosylation sites in the α chain^[9,10].

Fucosylation is one of the most important glycosylation modifications of proteins involved in inflammation and cancer^[11]. We previously reported that fucosylated haptoglobin (Fuc-Hpt) is found in the serum of patients with pancreatic cancer^[12] and established a lectin-antibody enzyme-linked immunosorbent assay (ELISA) to determine serum Fuc-Hpt levels^[13]. Recently, we established a novel glycan antibody [10-7G monoclonal antibody (mAb)] that directly recognizes Fuc-Hpt^[14]. Epitope analysis of the 10-7G mAb indicated that the peptide recognized by this antibody is in the haptoglobin α -chain, and the epitope appears as a result of aberrant glycosylation, which induces a conformational change in mature haptoglobin^[15]. In human serum, the 10-7G mAb recognizes not only fucosylated mature haptoglobin (Fuc-mHpt) but also fucosylated prohaptoglobin (Fuc-pHpt), a precursor molecule. Moreover, affinity chromatography using the 10-7G mAb followed by lectin blotting and mass spectrometry analysis revealed that the 10-7G mAb predominantly recognizes both Fuc-mHpt and Fuc-pHpt^[14]. The novel ELISA we developed using the 10-7G mAb enables measurement of both types of Fuc-Hpt in the serum. Here, we use Fuc-Hpt to describe both Fuc-mHpt and Fuc-pHpt, and Fuc-Hpt levels determined using the 10-7G mAb ELISA are reported as 10-7G values (Figure 1). As expected, 10-7G values are increased in pancreatic cancer patients compared with healthy volunteers (HVs)^[14,16].

We recently found that the 10-7G mAb can be used for immunohistochemical staining of lymphocytes and macrophages associated with pancreatic cancer tissues. In immunoblotting analyses using the 10-7G mAb, we demonstrated that some immune cell lines produce Fuc-Hpt and that several inflammatory cytokines induce its secretion (manuscript submitted). These results suggest that the production of Fuc-Hpt by immune cells is related to local immune conditions, such as those associated with cancer and inflammation. With respect to inflammatory diseases, IBD is one of the most representative diseases in terms of the relationship between immune cells and local immunity.

In general, serum haptoglobin levels increase during acute inflammation, as do levels of CRP^[6], and fucosylation of serum proteins increases dramatically in chronic inflammation^[11]. As described above, the 10-7G mAb we developed recognizes characteristic types of haptoglobin, including Fuc-mHpt and Fuc-pHpt. In the present study, we investigated the usefulness of the 10-7G value as a potential serum biomarker for monitoring disease activity of IBD, comparing 10-7G values with several conventional serum biomarkers.

MATERIALS AND METHODS

Clinical samples

Serum samples were obtained from patients with CD ($n = 45$), UC ($n = 110$), and acute enteritis (AE; $n = 11$) at Osaka University and Osaka University Hospital. Serum samples were also obtained from HVs with normal serum CRP levels (≤ 0.3 mg/dL; $n = 20$) and high serum CRP levels (> 0.3 mg/dL; $n = 79$) who underwent medical health examinations at aMs New Otani Clinic, Osaka, Japan. Detailed characteristics of HVs and UC, CD, and AE patients are presented in Table 1. Resected intestinal tissue

Table 1 Characteristics of patients with ulcerative colitis, Crohn's disease, and acute enteritis and healthy volunteers

Characteristic	Patients with UC	Patients with CD	Patients with AE	HVs	CRP-high HVs
Number (male:female)	110 (63:47)	45 (36:9)	11 (5:6)	20 (14:6)	79 (56:26)
Age, yr, mean (SD)	48.79 (15.08)	46.24 (11.75)	42.89 (20.20)	53.05 (6.58)	56.01 (11.23)
Age at IBD diagnosis, yr, mean (SD)	34.74 (16.64)	26.89 (8.36)			
UC disease location, extensive colitis/left-sided colitis/proctitis	52/40/18				
CD location, colic/ileal/ileo-colic		5/17/23			
Alb, g/dL, mean (SD)	4.14 (0.31)	3.71 (0.51)	3.84 (0.41)	4.41 (0.19)	4.28 (0.24)
ALT, U/L, mean (SD)	20.27 (17.49)	16.00 (6.04)		37.00 (22.19)	16.96 (5.22)
ALP, U/L, mean (SD)	212.58 (69.81)	244.79 (88.12)		214.35 (54.46)	209.84 (56.33)
T-Bil, mg/dL, mean (SD)	0.51 (0.36)	0.45 (0.24)		0.77 (0.22)	0.63 (0.23)
γ -GTP, U/L, mean (SD)	30.75 (23.66)	22.67 (15.61)		42.78 (25.20)	28.97 (17.30)
CRP, mg/dL, mean (SD)	0.38 (0.61)	0.68 (1.16)	8.15 (2.46)	0.10 (0.08)	1.35 (0.85)
Hb, g/dL, mean (SD)	12.79 (1.87)	12.78 (1.75)	13.08 (1.75)	14.53 (1.05)	13.96 (1.32)
Ht, %, mean (SD)	38.91 (7.58)	39.65 (4.77)	39.11 (4.59)	42.6 (2.68)	40.79 (3.66)
WBC, $10^3/\mu\text{L}$, mean (SD)	7.35 (2.48)	6.75 (2.17)	12.50 (5.18)		
Plt, $10^4/\mu\text{L}$, mean (SD)	27.58 (7.07)	28.90 (6.21)	27.34 (6.22)	21.60 (5.62)	23.43 (6.45)
Mayo, mean (SD)	3.27 (2.83)				
CDAI, mean (SD)		129.18 (70.48)			

UC: Ulcerative colitis; CD: Crohn's disease; AE: Acute enteritis; HVs: Healthy volunteers; Alb: Albumin; ALT: Alanine aminotransferase; ALP: Alkaline phosphatase; T-Bil: Total-bilirubin; γ -GTP: γ -Glutamyl transpeptidase; CRP: C-reactive protein; Hb: Hemoglobin; Ht: Hematocrit; WBC: White blood cells; Plt: Platelets; Mayo: Full Mayo score; CDAI: Crohn's disease activity index; SD: Standard deviation.

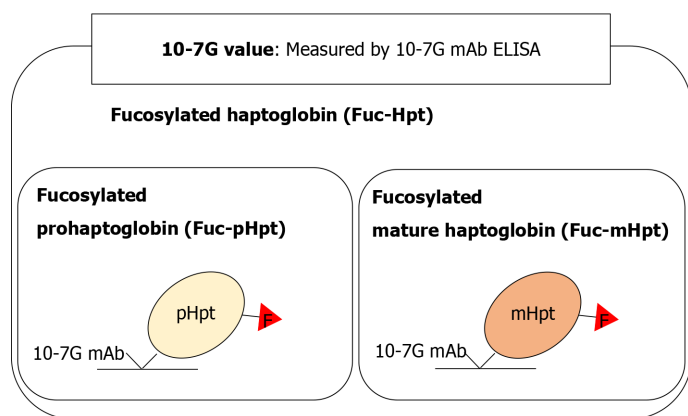


Figure 1 Schematic illustration of fucosylated haptoglobin detection using the 10-7G mAb. ELISA: Enzyme-linked immunosorbent assay.

sections from patients with severe CD ($n = 5$) or UC ($n = 5$) were embedded in paraffin at the Department of Pathology of Osaka University Graduate School of Medicine. This study was approved by the ethics committee of Osaka University Hospital and conducted in accordance with approved guidelines (approval number 14107-10). Written informed consent was obtained from all subjects in this study at the time of enrollment or blood sampling.

Immunohistochemistry

Tissue sample sections (4- μm thickness) were fixed in 10% formaldehyde and embedded in paraffin. Sections were then deparaffinized using a xylene alternative, Hemo-De (Falma, Tokyo, Japan), and then rehydrated in 100% ethanol (FUJIFILM Wako Pure Chemical Corporation, Osaka, Japan). The sections were rinsed with

phosphate-buffered saline [PBS; 137 mmol/L NaCl, 10 mmol/L Na₂HPO₄, 2.7 mmol/L KCl, 1.76 mmol/L KH₂PO₄ (pH 7.4)]. For antigen retrieval, sections were submerged in antigen-retrieval buffer [10 mmol/L Na₃C₆H₅O₇ (pH 6.0)] for 30 s at 125 °C and then submerged for 10 s at 90 °C in a Pascal decloaking chamber (Dako, Glostrup, Denmark). The sections were cooled for 10 min in the chamber and 10 min at room temperature (RT) following antigen retrieval. After rinsing in Tris-buffered saline containing Tween 20 [TBST; 136 mmol/L NaCl, 2.6 mmol/L KCl, 24 mmol/L Tris, 0.005% Tween 20 (pH 7.4)], endogenous peroxidase activity was blocked by incubation in Dako REAL peroxidase-blocking solution (Dako) for 5 min at RT. The sections were incubated overnight at 4 °C with 0.22 µg/mL of the 10-7G mAb diluted in Dako REAL antibody diluent (Dako). Next, the sections were incubated with Dako ChemMate ENVISION kit/HRP (DAB) (Dako) for 30 min at RT. After each incubation step, the sections were washed with TBST three times for 5 min each. Subsequently, the sections were incubated with DAB+ chromogen solution (Dako) diluted with DAB substrate buffer (Dako) for 3 min to develop the stain. Finally, the slides were counterstained with hematoxylin (CHROMA, Nordrhein-Westfalen, Germany), dehydrated, cleared, and mounted. Images were acquired using a BZ-9000 microscope (KEYENCE, Osaka, Japan).

ELISAs

The 10-7G mAb ELISA was performed according to our previously reported method, with slight modifications^[14]. 3,3',5,5'-Tetramethylbiphenyl-4-4'-diamine solution was used for color development, as described previously^[17]. The chromogen and the optical density were measured at 450 nm using an iMark™ microplate absorbance reader (Bio-Rad, Hercules, CA, United States).

A lectin-antibody ELISA was used to determine serum Fuc-mHpt levels, as described previously^[13]. The Fab fragment of anti-haptoglobin polyclonal antibody (pAb) (Dako) was coated onto an ELISA plate to capture serum haptoglobin, and fucosylation of haptoglobin was detected using biotinylated *Aleuria aurantia* lectin (AAL). Serum Fuc-mHpt levels determined using this lectin-antibody ELISA are reported as AAL-mHpt.

Serum levels of leucine-rich alpha-2 glycoprotein (LRG) were determined using a human LRG assay kit (IBL, Gunma, Japan), according to the manufacturer's instructions.

pHpt plasmid construction

A pcDNA3.1-Hyg (+) vector (Invitrogen, Carlsbad, CA, United States) encoding the human haptoglobin transcript variant 1 (Hpt2) gene was prepared as previously described^[18]. To construct the human pHpt expression vector, site-directed mutagenesis was employed to replace arginine with glutamine at residue 161, which is important for digestion by complement C1r subcomponent-like protein (C1RL). Amino acid substitution was carried out by polymerase chain reaction (PCR). In brief, the cycle parameters were as follows: initial denaturation at 95 °C for 3 min, one cycle of 98 °C for 10 s, 50 °C for 2 min, and 68 °C for 8 min, followed by 35 cycles of 10 s of denaturation at 98 °C, 15 s annealing at 57 °C, and an 8-min extension at 68 °C. The resulting DNA fragments were ligated with T4 DNA ligase (Promega, Madison, WI, United States). Primer sequences were as follows: sense, CCAAGAATCCGGCAA-CCCAGTGCAGCAGATCCTGGGTGGAC; anti-sense, GGCATCCAGGTGTCCACCC-AGGATCTGCTGCACTGGGTTTGCC. The mutated sequences were confirmed by sequencing analysis.

Purification of Fuc-pHpt from HEK293T cells

Human embryonic kidney cells (HEK293T) were obtained from the American Type Culture Collection (Manassas, VA, United States) and maintained at 37 °C in a humidified atmosphere with 5% CO₂ and 95% air in DMEM high-glucose (Nacalai Tesque Inc., Kyoto, Japan) medium supplemented with 10% heat-inactivated fetal bovine serum (HyClone, Logan, UT, United States), 100 units/mL of penicillin, and 100 µg/mL of streptomycin (Nacalai Tesque Inc.).

HEK293T cells were transfected with the expression vector pcDNA3.1-Hyg(+) containing the mutated Hpt2 gene. After 24 h of incubation, transfected cells were selected using hygromycin B (Wako) for 2-3 wk to obtain the pHpt-overexpressing stable transfectants.

Fuc-pHpt products were collected from the conditioned medium of pHpt-overexpressing HEK293T cells and purified using an affinity chromatography column coupled with 10-7G mAb, as previously described^[14]. The concentration of Fuc-pHpt

was determined using a Nano Drop 1000 spectrophotometer (Thermo Fisher Scientific, Waltham, MA, United States).

Fuc-pHpt semi-quantification by Western blotting

A 0.25- μ L aliquot of serum in 10 μ L of PBS was electrophoresed on a 10% polyacrylamide gel and transferred onto a polyvinylidene difluoride membrane. After blocking with 5% skim milk for 1 h, the membrane was incubated with 1/7500-diluted 10-7G mAb at 4 °C overnight. After three 10-min washes with TBST, the membrane was incubated with peroxidase-conjugated anti-mouse immunoglobulin G (IgG) for 1 h. After three 10-min washes with TBST, bound immunoglobulins were visualized using ImmunoStar® Zeta (Wako), according to standard protocols. Chemiluminescence signal detection and quantification were using a FUSION chemiluminescence imaging system (Vilber-Lourmat, Collegien, France). The quantity of Fuc-pHpt in each serum sample was calculated based on the ratio of the chemiluminescence signal intensity of the sample to that of the positive control. A fixed amount (5.0 ng) of purified Fuc-pHpt was used as a positive control sample for every Western blot analysis.

Haptoglobin phenotyping

For haptoglobin phenotyping, serum samples were subjected to Western blotting using anti-human haptoglobin pAb, as described previously^[16]. Haptoglobin phenotypes for each sample were determined after antibody signals were developed using ImmunoStar® Zeta (Wako) and detection of α -1 and/or α -2.

Statistical analysis

Statistical analyses were performed using JMP Pro 14.0 software (SAS Institute Inc., Cary, NC, United States). Data were analyzed using the Wilcoxon test, Spearman rank correlation analysis, and receiver operating characteristic (ROC) curve analysis. As the 10-7G value, AAL-mHpt, Fuc-pHpt amount, CRP levels, hemoglobin (Hb), and albumin (Alb) data did not exhibit a Gaussian distribution, these parameters were common log-transformed before correlation diagrams were drawn. In the statistical analyses, samples of the Hpt1-1 phenotype were excluded because the 10-7G values were below the level of detection. *P* values < 0.05 were considered indicative of a statistically significant difference.

RESULTS

Immunohistochemical identification of Fuc-Hpt-producing cells in IBD patient tissues

To identify Fuc-Hpt-producing cells in UC and CD intestinal tissues collected from IBD patients, we performed immunohistochemical analyses using the 10-7G mAb. As shown in **Figure 2A** and **B**, positive 10-7G mAb staining was observed in lymphocytes infiltrating into inflammatory sites of the mucosal layer and lymph nodules in the intestinal tissues of UC patients. Similar but much more diffuse 10-7G mAb staining was observed in comparable cells in CD tissues (**Figure 2C** and **D**).

Determination of the serum 10-7G value in patients with IBD and AE

We determined serum 10-7G values for HVs and patients with UC, CD, or AE. As expected, 10-7G values were significantly higher in patients with UC, CD, and AE as compared to HVs (**Figure 3A**). To determine whether the 10-7G mAb primarily recognized Fuc-mHpt or Fuc-pHpt, we assessed serum Fuc-pHpt levels by immunoblot analysis using the 10-7G mAb, whereas serum Fuc-mHpt levels were determined using the lectin-antibody ELISA. Data regarding the semi-quantification of serum Fuc-pHpt levels by immunoblot analysis using the 10-7G mAb are described as Fuc-pHpt amount, whereas serum Fuc-mHpt levels determined using the lectin-antibody ELISA are described as AAL-mHpt.

Data regarding 10-7G values and Fuc-pHpt amount or AAL-mHpt were statistically analyzed using Spearman's rank correlation coefficient (*r_s*). As shown in **Figure 3B**, a marked positive correlation was observed between the 10-7G value and Fuc-pHpt amount (*r_s* = 0.695, *P* < 0.0001). In contrast, a lower positive correlation was observed between serum AAL-mHpt levels and 10-7G values (*r_s* = 0.386, *P* = 0.0002) (**Figure 3C**). These results indicate that the 10-7G value depends primarily on serum Fuc-pHpt and partly reflects serum Fuc-mHpt in patients with intestinal inflammatory diseases.

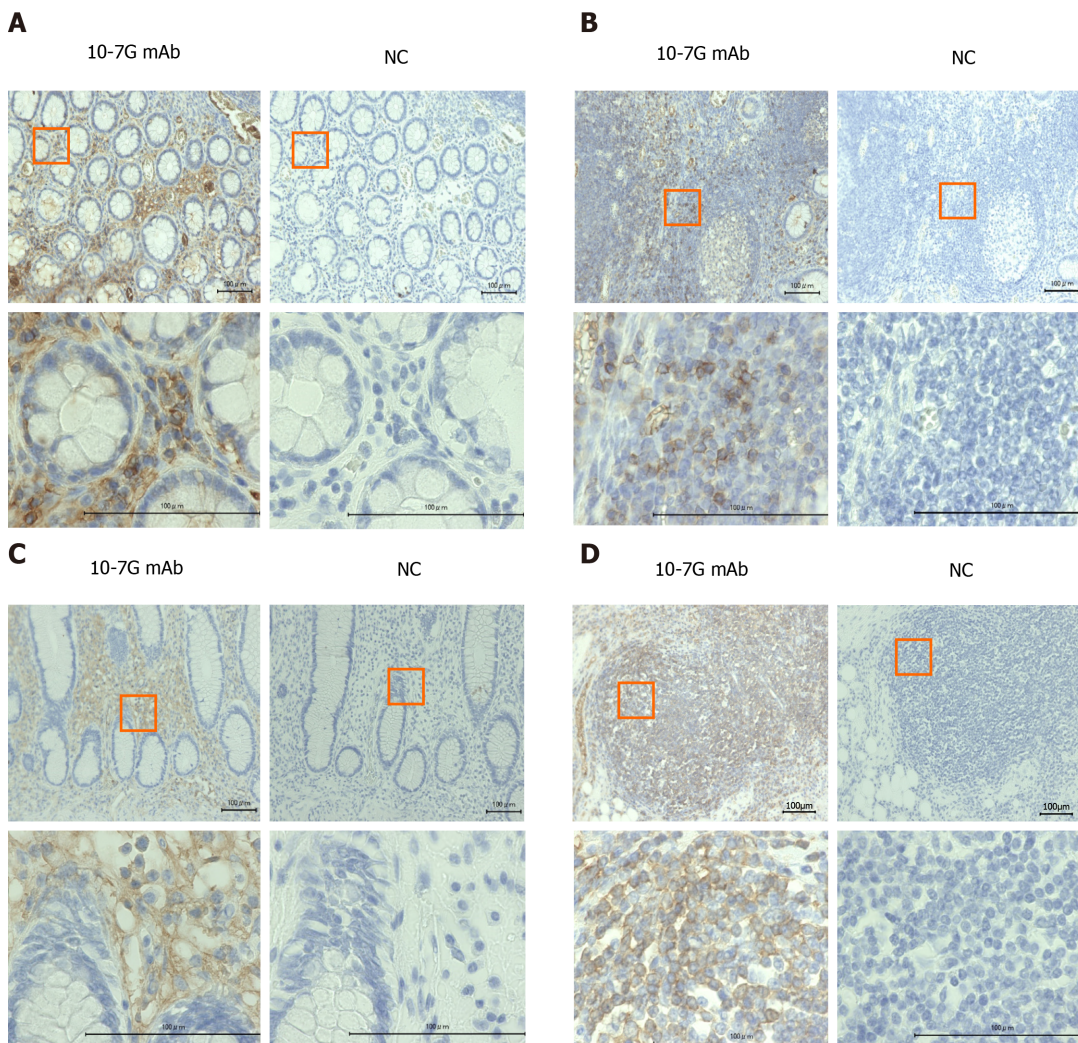


Figure 2 Immunohistochemical study of ulcerative colitis and Crohn's disease intestinal tissues using the 10-7G mAb. A and B: Lymphocytes infiltrating into inflammatory sites of the mucosal layer (A) and lymph nodules in intestinal tissues (B) of patients with ulcerative colitis ($n = 5$) were stained using the 10-7G mAb; C and D: Lymphocytes infiltrating into inflammatory sites of the mucosal layer (C) and lymph nodules in intestinal tissues (D) of patients with Crohn's disease ($n = 5$) were also stained. Photographs were acquired using a 20 × (upper) or 100 × (lower) objective. Scale bar, 100 μm. Positive staining was judged by comparison to the negative control. NC: Negative control.

Correlation between the 10-7G value and various clinical parameters

To elucidate the pathophysiologic significance of 10-7G values in IBD, we investigated the correlation between the 10-7G value and various clinical parameters of IBD patients, including Alb, alanine aminotransferase, alkaline phosphatase, total-bilirubin, γ -glutamyl transpeptidase, CRP, Hb, hematocrit, white blood cell count, platelet count, IBD activity index score, and full Mayo score (Mayo) for UC or Crohn's disease activity index (CDAI) for CD (Table 2). As shown in Figure 4A-C the 10-7G values of UC patients were negatively correlated with Hb ($r_s = -0.308, P = 0.0419$) and positively correlated with CRP ($r_s = 0.525, P = 0.0003$) and Mayo ($r_s = 0.435, P = 0.038$). In contrast, as shown in Figure 4D and E, the 10-7G values of CD patients were negatively correlated with Alb ($r_s = -0.376, P = 0.014$) and positively correlated with CRP ($r_s = 0.595, P < 0.0001$). However, no significant correlation was observed between 10-7G values and CDAI (Figure 4F). These results indicate that 10-7G values are correlated with some inflammation-related biomarkers, especially CRP, in both UC and CD patients and reflect an inflammatory condition.

To determine whether the 10-7G value is a suitable inflammation marker similar to CRP, we also determined 10-7G values of HVs who were found to have elevated CRP levels during routine medical examinations (CRP-high HVs, $n = 79$). As shown in Figure 5A, there was no significant difference in 10-7G values in CRP-high or normal HVs. In addition, there was no significant correlation between the 10-7G value and CRP level in CRP-high HVs (Figure 5B). In contrast, the 10-7G values of CRP-high IBD patients were significantly higher than those of CRP-normal IBD patients, and there

Table 2 Correlation coefficients between 10-7G value and clinical parameters in ulcerative colitis and Crohn's disease patients

	UC			CD		
	<i>r</i> s	<i>P</i> value	<i>n</i>	<i>r</i> s	<i>P</i> value	<i>n</i>
Alb	-0.182	0.254	41	-0.376 ^a	0.014	42
ALT	-0.214	0.339	22	-0.148	0.51	22
ALP	0.453	0.052	19	0.329	0.169	19
T-Bil	-0.419	0.059	21	-0.081	0.728	21
γ-GTP	-0.06	0.801	20	0.392	0.079	21
CRP	0.525 ^a	0.0003	44	0.595 ^a	< 0.0001	44
Hb	-0.308 ^a	0.0419	44	-0.287	0.072	40
Ht	-0.304	0.064	38	-0.261	0.091	43
WBC	0.289	0.057	44	0.215	0.177	41
Plt	0.06	0.712	40	0.145	0.387	38
Mayo	0.435 ^a	0.038	23			
CDAI				0.134	0.392	43

*r*s: Spearman's rank correlation coefficient.

^a*P* < 0.05. UC: Ulcerative colitis; CD: Crohn's disease; Alb: Albumin; ALT: Alanine aminotransferase; ALP: Alkaline phosphatase; T-Bil: Total-bilirubin; γ-GTP: γ-Glutamyl transpeptidase; CRP: C-reactive protein; Hb: Hemoglobin; Ht: Hematocrit; WBC: White blood cells; Plt: Platelets; Mayo: Full Mayo score; CDAI: Crohn's disease activity index.

was a significant correlation between 10-7G values and CRP in IBD patients (Figure 5C and D). These data suggest that the 10-7G value mainly reflects intestinal inflammation, whereas CRP reflects systemic inflammation.

Usefulness of the 10-7G value as a biomarker of UC disease activity

As the 10-7G value was found to correlate with IBD activity index score in a small number of UC patients (Figure 4C and Table 2), we investigated the usefulness of the 10-7G value as an indicator of UC disease activity using a larger number of serum samples (*n* = 110). We compared the 10-7G value with CRP and LRG, a novel serum biomarker for UC disease activity^[19]. We first analyzed the correlation between the 10-7G value, LRG, CRP, and three clinical activity index scores: the partial Mayo score (pMayo), an evaluation of symptomatic severity; the Mayo endoscopic subscore (MES), an evaluation of endoscopic severity; and Mayo, the sum of pMayo and MES, which is an indicator of clinical severity (Supplementary Table 1). Although the 10-7G value was significantly correlated with three of the scores (pMayo: *r*s = 0.2433, *P* = 0.0224, MES: *r*s = 0.3540, *P* = 0.0064, Mayo: *r*s = 0.2273, *P* = 0.0440), the correlation coefficients were slightly lower than those for CRP (pMayo: *r*s = 0.2947, *P* = 0.0053, MES: *r*s = 0.4375, *P* = 0.0006, Mayo: *r*s = 0.2597, *P* = 0.0208) and LRG (pMayo: *r*s = 0.3108, *P* = 0.0032, MES: *r*s = 0.3665, *P* = 0.0047, Mayo: *r*s = 0.3010, *P* = 0.0070).

The sensitivity and specificity of the serum 10-7G value, LRG, and CRP for diagnosis of endoscopic remission and clinical remission based on MES and Mayo were determined using ROC curve analysis. Endoscopic remission (*i.e.*, mucosal healing) was defined as MES ≤ 1^[20], and clinical remission was defined as Mayo ≤ 2 (pMayo ≤ 1 and MES ≤ 1)^[21]. For the diagnosis of endoscopic remission, the area under the curve (AUC) of the 10-7G value (MES ≤ 1: 0.699) (Figure 6A) was higher than that of both LRG (MES ≤ 1: 0.659) (Figure 6B) and CRP (MES ≤ 1: 0.663) (Figure 6C). In terms of clinical remission, the AUC value of the 10-7G value (Mayo ≤ 2: 0.626) (Supplementary Figure 1A) was lower than that of both LRG (Mayo ≤ 2: 0.714) (Supplementary Figure 1B) and CRP (Mayo ≤ 2: 0.678) (Supplementary Figure 1C). These results indicate that the 10-7G value is a useful biomarker for evaluating endoscopic mucosal healing in UC patients.

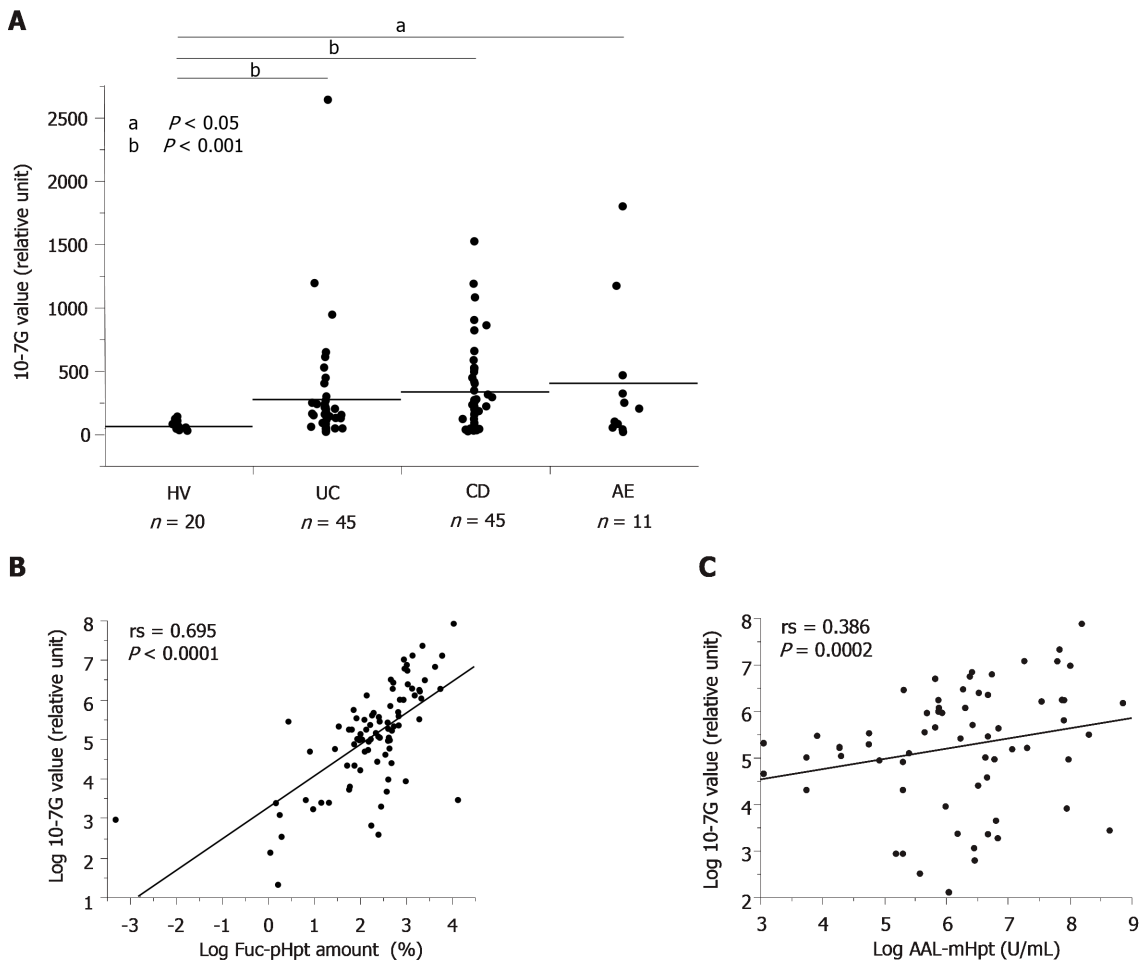


Figure 3 Measurement of the serum 10-7G value and fucosylated prohapptoglobin amount in patients with inflammatory bowel disease. **A:** Serum 10-7G value of healthy volunteers and patients with ulcerative colitis, Crohn's disease, or acute enteritis. Values are shown in relative units; **B** and **C:** Coefficients of correlation (**B**) between the 10-7G value and Fuc-pHpt amount and (**C**) between the 10-7G value and AAL-mHpt level. HV: Healthy volunteer; UC: Ulcerative colitis; CD: Crohn's disease; AE: Acute enteritis.

DISCUSSION

In this study, we demonstrated that the 10-7G value reflects intestinal inflammation. We found that the 10-7G value was not increased in the serum of HVs with elevated CRP levels, but it was significantly increased in the serum of patients with IBD and AE (Figures 3A and 5A). In addition, there was a significant correlation between 10-7G values and CRP in IBD patients, whereas no significant correlation in HVs with elevated CRP levels (Figure 5B and D). We previously reported agalactosyl IgG as a diagnostic biomarker of IBD^[22]. Agalactosyl IgG is also an indicator of chronic inflammation associated with conditions such as rheumatoid arthritis^[23]. The 10-7G value, on the other hand, is associated with acute and/or chronic intestinal inflammation.

We also demonstrated that the 10-7G value is a useful biomarker of disease activity in IBD patients, especially those with UC. Correlation analyses revealed that the 10-7G value is correlated with UC activity index score, non-CD Mayo score, and CDAI (Figure 4C and F). We concluded that this difference was due to the specificity of the 10-7G value for intestinal inflammation. Whereas the inflammatory lesions associated with UC are localized in the large intestine, those associated with CD can occur throughout the gastrointestinal tract, from the oral cavity to the anal canal^[24]. The CDAI reflects not only intestinal inflammation but also lesions related to extraintestinal complications, such as arthritis, stomatitis, and anal fistulas^[25]. Furthermore, calculation of the Mayo includes endoscopic findings that enable accurate evaluation of the intestinal condition, but the CDAI does not consider these findings. Therefore, the 10-7G value may not be correlated with the CDAI.

We also compared the 10-7G value and the LRG level, a promising tool for monitoring disease activity in IBD patients that was recently accepted for insurance

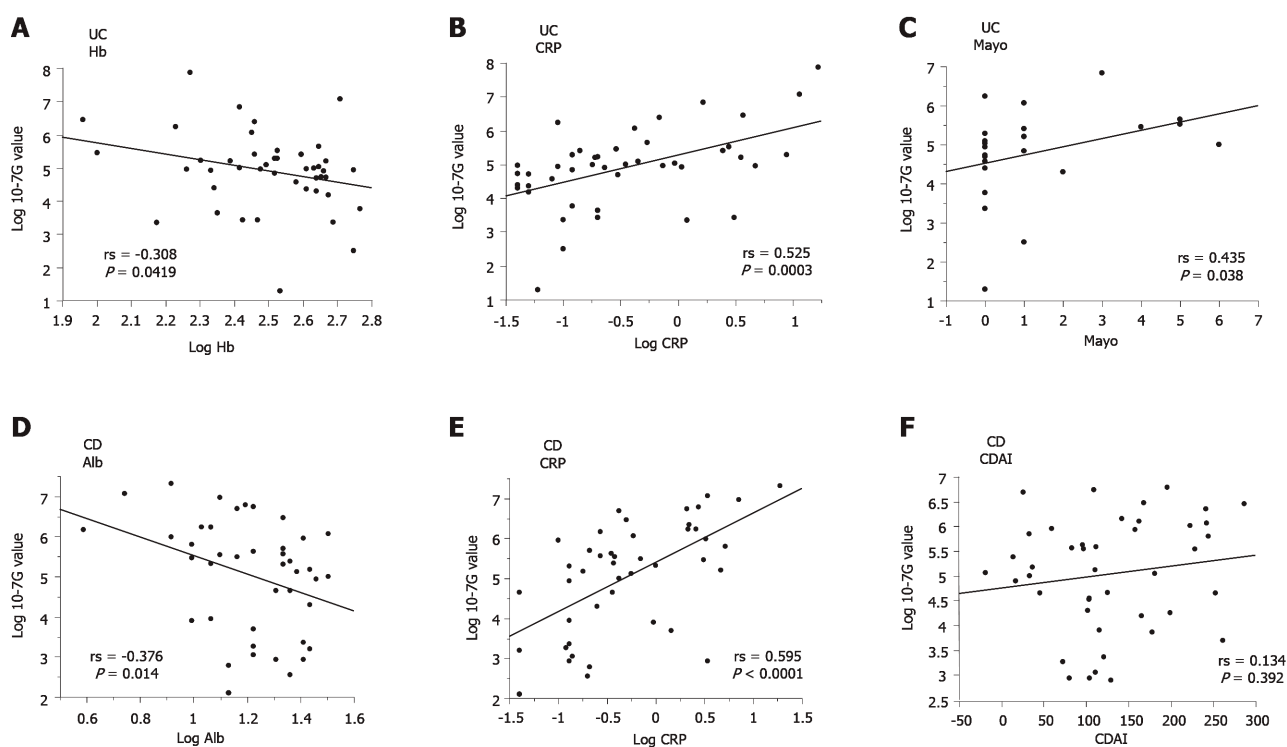


Figure 4 Correlations between 10-7G values and clinical parameters in inflammatory bowel disease patients. A-C: In ulcerative colitis patients, the 10-7G value was negatively correlated with hemoglobin and positively correlated with C-reactive protein (CRP) level and full Mayo score; D and E: In Crohn's disease patients, the 10-7G value was negatively correlated with albumin and positively correlated with CRP level; F: The 10-7G value was not correlated with the Crohn's disease activity index. UC: Ulcerative colitis; CD: Crohn's disease; Hb: Hemoglobin; CRP: C-reactive protein; Mayo: Mayo score; Alb: Albumin; CDAI: Crohn's disease activity index.

coverage in Japan. As indicated in [Figure 6](#), the AUC of the 10-7G value for a diagnosis of endoscopic remission ($MES \leq 1$) was higher than that of LRG. In contrast, the AUC of LRG for the diagnosis of symptomatic remission ($pMayo \leq 1$) was higher than that of the 10-7G value ([Supplementary Figure 1D and E](#)). In other words, the 10-7G value is superior to LRG for evaluating the endoscopic mucosal condition, and LRG is superior to the 10-7G value for evaluating general symptoms such as stool frequency and hematochezia. As indicated in [Figure 3B](#), the 10-7G value of IBD patients primarily reflects Fuc-pHpt levels, and some studies have reported that pHpt increases epithelial permeability^[26]. Based on our results and other reports, we attributed the usefulness of the 10-7G value for evaluating the mucosal condition of UC patients to the detection of Fuc-pHpt.

To identify Fuc-Hpt-producing cells in intestinal tissues of IBD patients, we performed immunohistochemical staining with the 10-7G mAb. As shown in [Figure 2](#), lymphocytes infiltrating into areas of inflammation in the mucosal layer were stained with the 10-7G mAb. The staining signal was thought to depend primarily on Fuc-pHpt rather than Fuc-mHpt. Cleavage of pHpt by C1RL is required for haptoglobin maturation. C1RL is a serine protease that cleaves $\alpha\beta$ -junctions within pHpt^[27]. C1RL expression is reportedly markedly elevated in the liver^[28]. Therefore, most haptoglobin produced in hepatocytes matures before it is secreted into the blood. The expression of C1RL in other tissues is typically much lower than in hepatocytes; therefore, synthesized haptoglobin does not fully mature. In addition, in an immunoblotting analysis using 10-7G mAb, we recently demonstrated that some lines of non-adherent cells produce Fuc-pHpt and that several inflammatory cytokines facilitate Fuc-pHpt secretion (manuscript submitted). These data support our hypothesis that infiltrating lymphocytes produce Fuc-pHpt in sites of intestinal inflammation and that the 10-7G value is an indicator of Fuc-pHpt production. Further studies are needed to fully elucidate the mechanism underlying the secretion of Fuc-pHpt by immune cells, however.

Considering our findings in this study, we proposed the flow chart of non-invasive detection methods of clinical activity of IBD, especially UC, using 10-7G value based on the cutoff value of 211 defined as ROC curve analysis ([Figure 6A](#)), the point of the best split of endoscopic remission ([Figure 7](#)). Since clinical symptoms appear after

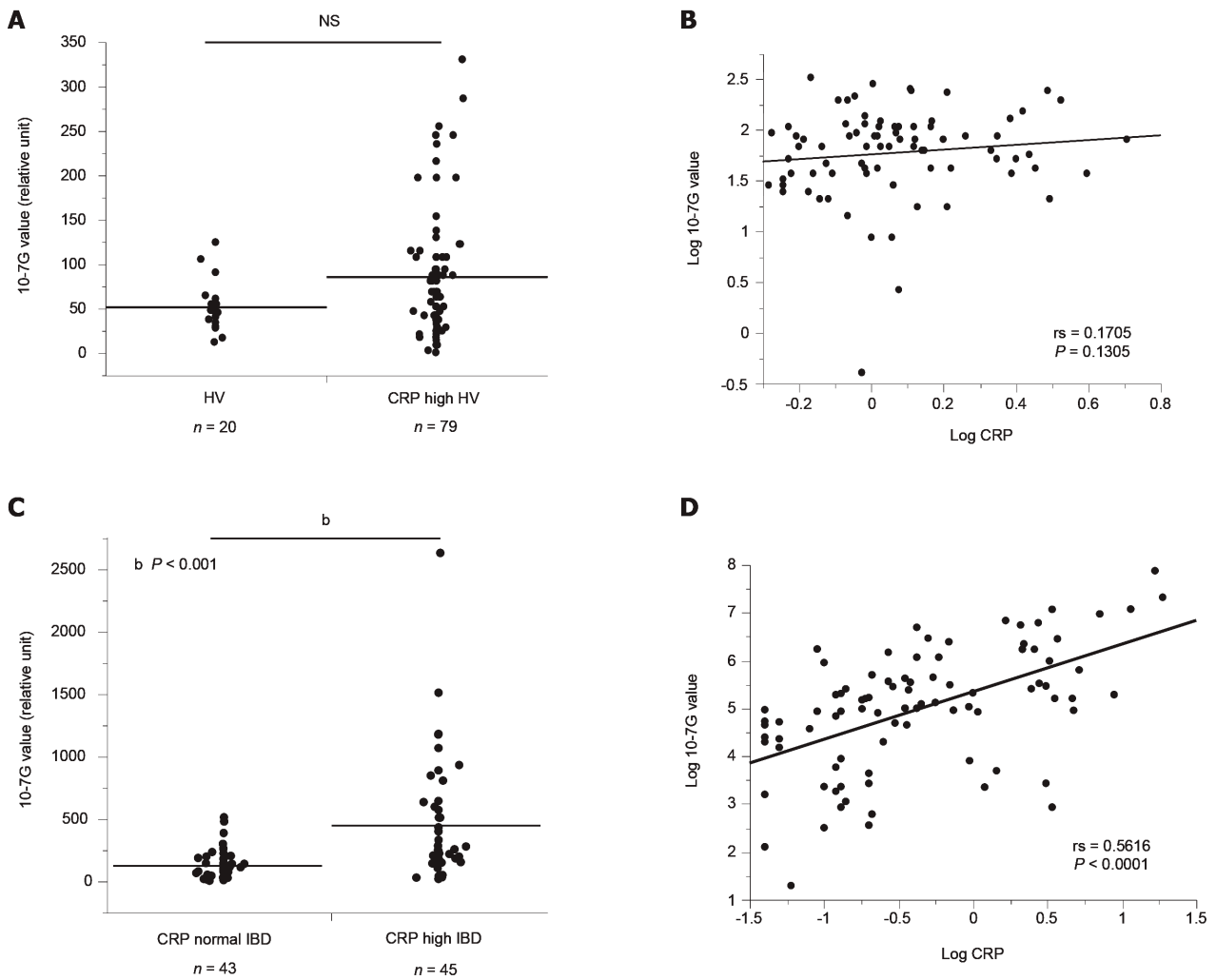


Figure 5 Measurement of serum 10-7G values and correlation between 10-7G values and C-reactive protein in healthy volunteers who underwent a medical examination. A: The 10-7G value of healthy volunteers (HVs) who had elevated C-reactive protein (CRP) levels (> 0.3 mg/dL) diagnosed during routine medical examination (CRP-high HVs); B: Coefficient of correlation between the 10-7G value and CRP level in CRP-high HVs; C: 10-7G values of inflammatory bowel disease (IBD) patients with normal CRP level (≤ 0.3 mg/dL) (CRP-normal IBD) or high CRP level (> 0.3 mg/dL) (CRP-high IBD); D: Coefficient of correlation between the 10-7G value and CRP level in IBD patients. HV: Healthy volunteer; CRP: C-reactive protein; IBD: Inflammatory bowel disease; NS: Not significant.

changes in mucosal inflammation status of the intestine and colonoscopy itself has a risk of exacerbating mucosal inflammation, non-invasive biomarkers such as 10-7G value are required to follow up the patients with IBD.

CONCLUSION

In conclusion, we demonstrated that the serum 10-7G value reflects intestinal inflammation and is a useful biomarker for evaluating endoscopic mucosal healing in UC, as Fuc-Hpt detected by the 10-7G mAb is produced by lymphocytes infiltrating into sites of inflammation in the intestinal mucosal layer.

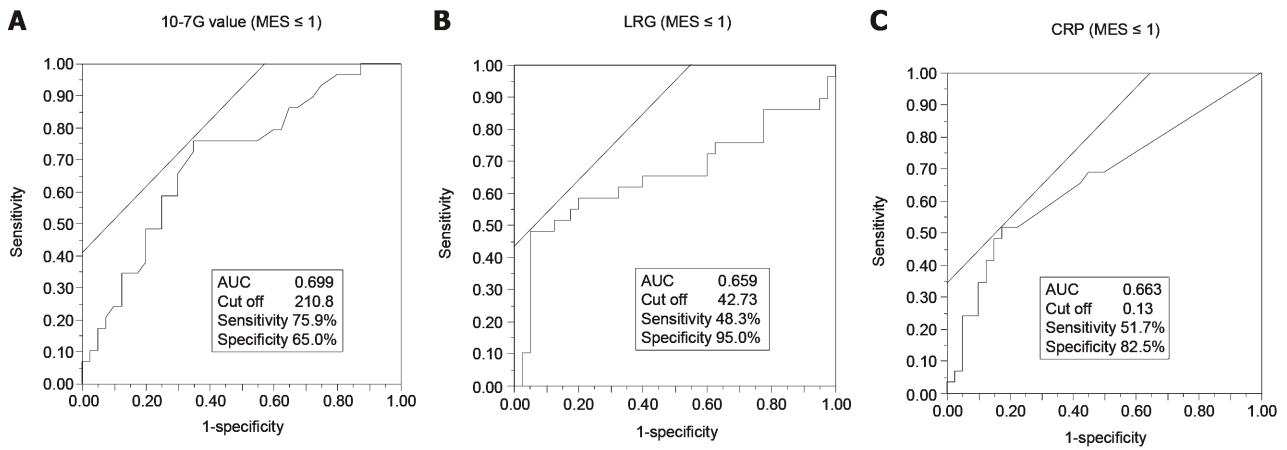


Figure 6 Evaluation of the usefulness of the 10-7G value for pathologic diagnosis of ulcerative colitis compared to C-reactive protein and leucine-rich alpha-2 glycoprotein. A-C: Analysis of receiver operating characteristic curves for 10-7G value, leucine-rich alpha-2 glycoprotein, and C-reactive protein for differentiation between ulcerative colitis (UC) patients with endoscopic mucosal remission by Mayo endoscopic subscore (MES). AUC: Area under the curve; MES: Mayo endoscopic subscore; LRG: Leucine-rich alpha-2 glycoprotein; CRP: C-reactive protein.

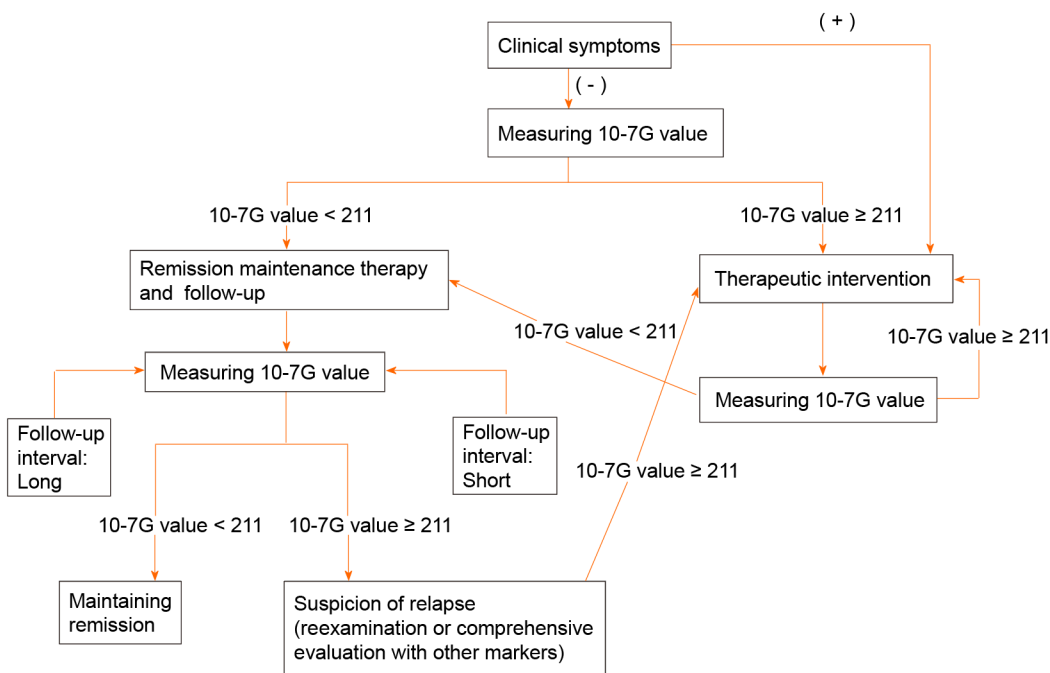


Figure 7 Non-invasive follow-up chart of patients with ulcerative colitis using 10-7G value.

ARTICLE HIGHLIGHTS

Research background

Fucosylation is one of the most important glycosylation involved in cancer and inflammation. Recently, we have succeeded to establish a novel glycan antibody 10-7G mAb which directly recognizes fucosylated haptoglobin (Fuc-Hpt), and developed its enzyme-linked immunosorbent assay system.

Research motivation

Although fecal and serum biomarkers have been extremely important and supportive for monitoring of Inflammatory bowel disease (IBD), their low sensitivity and high variability characteristics limit clinical efficacy. Thus, the establishment of better biomarkers is expected.

Research objectives

This research aimed to investigate the usefulness of the serum levels of Fuc-Hpt (10-7G values) as a potential biomarker for monitoring disease activity in IBD.

Research methods

Intestinal tissues of IBD patients were examined immunohistochemically using the 10-7G mAb. The 10-7G values were measured using serum from patients with IBD, acute enteritis (AE), and healthy volunteers. Correlation analysis and Receiver operating characteristic curve analysis were performed to evaluate the usefulness of the 10-7G values as a biomarker for IBD compared to conventional serum biomarkers and various clinical parameters.

Research results

Positive 10-7G mAb staining was observed in lymphocytes infiltrating into inflammatory sites of the mucosal layer. The 10-7G values were significantly higher in patients with IBD and AE. Statistical analyses showed that the 10-7G values could evaluate intestinal inflammation and endoscopic mucosal healing in ulcerative colitis (UC).

Research conclusions

Fuc-Hpt is produced by lymphocytes infiltrating into sites of inflammation in the intestine and the serum 10-7G value is a non-invasive biomarker for evaluating endoscopic remission in UC.

Research perspectives

Further studies are needed to fully elucidate the mechanism underlying the secretion of Fuc-Hpt by immune cells. In addition, we will measure serum 10-7G values of same patients with UC in follow-up study.

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