Abstract. Leptin, which is encoded by the obese gene, is a multifunctional neuroendocrine peptide that regulates appetite, bone formation, reproductive function and angiogenesis. The aims of the present study were to investigate the expression of leptin in 80 patients with colorectal cancer (CRC) and to determine the effects of leptin on the malignant properties of CRC cells. We evaluated the expression of leptin in tissues of 80 patients with CRC. Suspension cultures were used to isolate CRC stem cells following pretreatment with leptin. We analyzed the effects of leptin on the adhesion and invasive capacities of CRC cell lines. The effects of leptin on JAK and ERK activation were examined using western blotting. Leptin expression was associated with CRC progression and increased the number and size of spheroid formation by CRC cell lines. Leptin enhanced cell invasion and adhesion and activated JAK and ERK signaling in the CRC cell lines. The present study demonstrated that leptin influences the growth and survival of CRC stem cells and regulates adhesion and invasion of colorectal carcinoma through activation of the JAK and ERK signaling pathways.

Introduction

Colorectal cancer (CRC) is one of the most common neoplasms worldwide (1). A large body of epidemiological evidence suggests that obesity increases CRC risk in men (relative risks of ~1.5-2.0) and women (relatively risks of ~1.2-1.5) (2,3). Although the molecular mechanisms underlying this association are unknown, data acquired from experiments performed in vitro suggest the direct involvement of fat tissue in CRC development. Adipocytes and preadipocytes stimulate the growth of CRC cells (4), and the principal hormone synthesized by adipocytes is leptin. Leptin is encoded by the obese gene and functions as a neuroendocrine hormone that has attracted attention since its identification in 1995 (5). Leptin regulates appetite, bone formation, reproduction and angiogenesis (6). These biological activities suggest that it plays an important role in proliferation, invasion and metastasis of cancer cells (7). In humans, circulating leptin levels correlate with body mass index and are significantly elevated in obese individuals (8). Recent data clearly indicate that the mitogenic, anti-apoptotic, proinflammatory and angiogenic properties of leptin promote the development and progression of different types of cancers (9). Several reports have described the mitogenic effects of leptin on gastric (10), breast (11), ovarian (12), prostate (13) and endometrial cancer cells (14). Two studies demonstrated that increased leptin levels are associated with greater risk of CRC development, particularly in males (15,16). Furthermore, in colon epithelial cells, leptin was found to induce chemokine production associated with macrophage activation similar to that observed in an adenomatous polyposis coli genotype (17,18). The aims of the present study were to investigate the expression of leptin in 80 patients with CRC and to determine the effects of leptin on the malignant properties of CRC cell lines.

Materials and methods

Case selection and immunohistochemical assessment. For immunohistochemical staining, we obtained formalin-fixed, paraffin-embedded tissue samples from 80 patients diagnosed with CRC at Chonnam National University Hospital, Gwangju, Korea. The histology of the tumors was analyzed, and the pathological stage was estimated according to the TNM score. Patients (stage I, n=20; stage II, n=20; stage III, n=20 and stage IV, n=20) were randomly selected from each of the stage categories. The specimens were fixed in 10% neutral-buffered formalin, embedded in paraffin and stained with hematoxylin and eosin. The present study was approved by the Institutional Review Board of Chonnam National University Hospital.

For immunohistochemical staining, tissue sections were deparaffinized, rehydrated and subjected to epitope retrieval.
To block endogenous peroxidase activity, tissues were treated with Peroxidase-Blocking solution (Dako, Carpinteria, CA, USA) and incubated with a polyclonal rabbit anti-leptin antibody (A-20; Santa Cruz Biotechnology, Santa Cruz, CA, USA), which was diluted 1:100 using goat serum and incubated with the sections at room temperature for 1 h. After three 2-min washes with phosphate-buffered saline (PBS), the sections were incubated with a biotinylated goat anti-rabbit secondary antibody for 30 min (Dako). After three 2-min washes in PBS, horseradish peroxidase-streptavidin (Dako) was added to the sections for 30 min, followed by another three washes for 2 min in PBS. Reactions were detected using with 3,3′-diaminobenzidine substrate (Vector Laboratories, Burlington, ON, Canada) for 1 min, and the cells were counterstained using Mayer’s hematoxylin. Then slides were dehydrated following a standard procedure and sealed with coverslips.

Tissue specimens reacted with the anti-leptin antibody were examined at low and then at high magnification by two pathologists blinded to the identities of the samples. In cases of heterogeneous patterns in some sections, the classification was determined by the dominant pattern, and the intensity of stained cells was designated as negative, weak, moderate or strong.

Cell culture and leptin pretreatments. We selected three human CRC cell lines (LS174T, HCT 116 and CaCo-2) since they express leptin at high levels. They were obtained from the Korean Cell Line Bank (KCLB) and cultured at 37°C in a humidified atmosphere containing 5% CO₂. Cells were grown in Dulbecco’s modified Eagle’s medium (DMEM) containing 4,500 mg/l glucose, 100 mg/l streptomycin and 2 mM L-glutamine supplemented with 10% fetal bovine serum (FBS) (both from Gibco Invitrogen Inc., USA). After 24 h of serum starvation, the culture media were replaced with serum-free media containing the indicated treatments. After a further 24 h incubation, 10 ng/ml human recombinant leptin (Sigma-Aldrich Corp., St. Louis, MO, USA) was added for different times.

Spheroid formation assay. Cells were grown to 70% confluence, trypsinized and plated in 100 cm² diameter culture dishes at a density of 1,000 cells/ml in serum-free DMEM containing 10 ng/ml human recombinant basic fibroblast growth factor, (bFGF) and 10 ng/ml human epidermal growth factor (hEGF) (both from R&D Systems, Minneapolis, MN, USA). A density of at least 1,000 cells/ml was established for forming tumorspheres/colonospheres (serving as an in vitro model of cancer stem cells). All cultures were incubated at 37°C in a humidified atmosphere containing 5% CO₂. Cells were grown as suspension cultures for 1-2 weeks for tumorsphere formation. Colonies were counted in 10 randomly selected fields at x10 magnification using an Olympus IX50 inverted microscope.

Western blotting. To extract proteins, cells were lysed with RIPA buffer (1 M Tris-HCl, 150 mM NaCl, 1% Triton X-100 and 2 mM EDTA) containing 1 mM phenylmethylsulfonyl fluoride (PMSF) and Halt™ protease inhibitor cocktail (Thermo, Rockford, IL, USA). The protein concentrations of the cell lysates were quantitated using the BCA™ protein assay (Thermo) with bovine serum albumin (BSA) as a standard. The lysates (25 μg protein) were subjected to electrophoresis on 10% SDS-polyacrylamide gels and then electrophoretically transferred to polyvinylidene fluoride membranes (Millipore, Billerica, MA, USA). The membranes were incubated for 1 h in blocking solution [5% BSA in TBS-Tween-20 buffer (TBST)] and sequentially blotted with primary antibodies at 4°C overnight. Antibodies against Janus kinase 2 (JAK2), phospho-JAK2, AKT, phospho-AKT, ERK, phospho-ERK, MAPK, phospho-MAPK and β-actin were purchased from Cell Signaling Technology (Beverly, MA, USA). After rinsing in TBST, membranes were incubated with horse-radish peroxidase (HRP)-labeled anti-rabbit or anti-mouse immunoglobulin secondary antibodies (1:2,000 dilution) (Cell Signaling Technology) at room temperature for 1 h. Enhanced chemiluminescence was used to detect the bands, which were visualized using a Fuji LAS-3000 image analyzer (Fuji Film, Tokyo, Japan).

Adhesion assay. The 96-well plates were prepared. Three human CRC cell lines were detached from the surfaces of culture flasks with 5 mM EDTA in PBS, resuspended in culture medium containing 0.02% BSA to 2.4x10⁵ cells/ml, and 100 μl of cell suspension was added to each well. All cells were assayed in quadruplicate. After incubation for 12 h at 37°C in an atmosphere containing 5% CO₂, the supernatant from each well was removed. After washing out non-adherent cells, adherent cells were incubated for 4 h in medium containing 500 μg/ml 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) solution. The absorbance of the reaction product was measured at 550 nm. Adherent cells were counted in three random area of each well.

Invasion assay. Cell invasion assays were performed using Transwell filter chambers (8.0-µm pores) coated with 1% gelatin overnight and dried at room temperature. Human CRC cell lines were harvested, washed once in serum-free media, and seeded at 2x10⁵ cells in 120 μl of medium containing 0.2% BSA in the upper chamber. Then, 400 μl of 0.2% BSA medium containing 20 μg/ml of human plasma fibronectin (Calbiochem, La Jolla, CA, USA), a chemotactic factor, was loaded into the lower chamber. The Transwell apparatus was incubated for 24 h at 37°C. Cells that invaded the bottom surface of the upper chamber were fixed with 70% ethanol and stained with Diff-Quik solution (Sysmex, Kobe, Japan) following the manufacturer’s protocol. The non-invasive cells on the top surface were wiped off with cotton balls, and the stained cells on the bottom surface were counted in five selected fields (each 0.5 mm²) of six random squares using a hemocytometer placed on the stage of a light microscope. Results are expressed as the means ± standard error of the mean (SEM) of the number of cells/field of three individual experiments.

Statistical analysis. The statistical significance of differences between data sets was determined using the paired t-test. The χ² and Fisher’s exact test, where appropriate, were used to compare expression of leptin with various tumor stages. All reported p-values were two-sided and p<0.05 was considered to indicate a statistically significant result. The Statistical Package for the Social Sciences (SPSS)/PC 20.0 (Chicago, IL, USA) was used to perform the calculations.
Results

Association of leptin expression and tumor stage. We determined the level of expression of leptin in 80 CRCs of different stages. Leptin was clearly expressed in the cytoplasm of the CRC cells. Leptin expression was ‘undetectable’ in 19/20 (95%) patients with stage I CRC and 5/20 (25%) patients with stage IV CRC. In contrast, leptin was ‘moderately to strongly’ expressed in 0/20 patients with stage I CRC and in 10/20 tissue samples with stage IV CRC. Expression of leptin was significantly associated with tumor stage (p<0.0001).

Spheroid formation induced by leptin. Cancer cells can be cultured in suspension to form spheres in serum-free replacement medium. To test whether human CRC cell lines could form spheres, CaCo-2, LS174T and HCT 116 cells were cultured in a suspension-culture system. Tumorsphere formation by CaCo-2, LS174T and HCT 116 cells was observed on day 5, accounting for 3.73±0.25, 2.80±0.26 and 3.13±0.32% of the total cell population by day 11, respectively. A greater number of large aggregates and colonies formed in the leptin-treated cultures relative to the controls (p<0.05). Leptin exposure increased the average colony size formed by each cell line (Fig. 2).

Induction of adhesion and invasion in leptin-treated cells. In the adhesion assays, leptin treatment for 12 h enhanced cell-cell adhesion of HCT 116 cells compared with the untreated HCT 116 cells (p=0.006); however, there was no statistically significant difference between the treated and untreated LS174T (p=0.140) and CaCo-2 cells (p=0.147, Fig. 3).

We next determined whether leptin treatment influences the invasiveness of the colon cancer cell lines The number of invading HCT 116 cells treated with leptin (10 ng/ml) was 574.3±562.5 when compared with the control value of 502.6±502.2 (Fig. 4A, p=0.178). The number of invading CaCo-2 cells treated with leptin (10 ng/ml) was 3,395.6±1571.2 when compared with the control value of 1,512.6±850.9 (Fig. 4B, p<0.001).

Activation of JAK2 and ERK signaling in cells treated with leptin. In the CRC CaCo-2 cells, leptin (10 ng/ml) stimulated the phosphorylation of JAK2 at the different treatment times. Increased phosphorylation of JAK2 was observed within
30 min after leptin treatment followed by a decline. In the LS174T and HCT 116 cells, leptin stimulated the phosphorylation of ERK. The levels of total (T) JAK2, ERK, MAPK and AKT were not altered in the three CRC cell lines after leptin treatment (Fig. 5).

Discussion

In the present study, we demonstrated that leptin enhanced the malignant phenotypes of three cell lines derived from human CRCs. Epidemiologic studies suggest that obesity is a risk factor for colon cancer development (16,19). In contrast, Stattin et al. suggested that leptin is simply an innocent bystander, serving merely as a correlate of other obesity-induced adverse alterations in metabolism that may be the true cause of CRC (16). In human CRC tissue, but not in normal mucosa or adenomas, leptin and its receptor are overexpressed, which correlates with the expression of the proneoplastic transcriptional regulator, hypoxia-inducible factor 1, and a more advanced tumor phenotype (21,22). Moreover, leptin is gradually expressed in the progression of normal cells to adenomas and subsequently to carcinomas, suggesting that leptin may be involved in multistep colorectal carcinogenesis (3).

In the present study, we determined the expression of leptin in tissue samples from 80 Korean patients with CRC and the associated clinicopathologic factors. We showed that patients with more advanced stage tumors expressed higher levels of leptin. Expression levels of leptin and its receptor were previously found to correlate with the grade of tumor differentiation, depth of bowel wall invasion, Dukes’ stage and distant metastasis in CRC patients, suggesting that the binding of leptin to its cellular receptor promotes the proliferation of CRC cells (23). However, data for the leptin concentration in the sera of patients with CRC are discrepant. For example, three studies report that decreased serum leptin levels are associated with tumor aggressiveness (21,24,25).

Cancer stem cells are rare, but play an important role in the maintenance of cancer homeostasis (26). Stem cells of the gastrointestinal tract may serve as a principal target of tumorigenic mutations due to their naturally long life-span and capacity for self-renewal. The concept of stem cell-driven tumorigenesis in CRC is supported by the identification and phenotypic characterization of a subpopulation of colon cancer cells capable of initiation and sustained reproduction.
of human colon carcinomas in immune-compromised mice (tumor-initiating cells or colorectal tumor stem cells) (27). Cancer stem cells (CSCs) form spheres when cultured in vitro in serum-free suspension cultures. This sphere-formation technique was used to isolate putative CSCs from freshly isolated brain (28), breast (29) and colon tumors (30). Furthermore, spheroid culture (or colonospheres) generated from a limited number of human CRC-derived cell lines are enriched for cells that express colonic CSC markers (31,32). Increasing evidence suggests that stem cells play a decisive role in the progression and metastasis of CRC.

CSCs possess the ability to self-renew and differentiate into different cell types. Moreover, CSCs play an important role in the maintenance of cancer homeostasis, metastasis, resistance to therapy and subsequent tumor recurrence (26,33). In the present study, leptin increased the number and size of tumorspheres formed by HCT 116, LS174T and CaCo-2 cells. Increased phosphorylation of JAK2 was observed within 30 min after leptin treatment (10 ng/ml) followed by a decline. (B) In LS174T and (C) HCT 116 cells, leptin stimulated the phosphorylation of ERK. The levels of total (T) JAK2, ERK, MAPK and AKT were not altered in three CRC cell lines following leptin treatment.

Figure 4. Cell invasion assay. (A) The number of invading leptin-treated (10 ng/ml) HCT 116 cells was 574.3±562.5, whereas that of the negative control was 502.6±502.2 (p=0.178). (B) The number of invading leptin-treated (10 ng/ml) CaCo-2 cells was 3395.6±1571.2, whereas that for the negative control was 1512.6±850.9 (p<0.001).

Figure 5. Analysis of JAK and ERK phosphorylation. (A) Leptin stimulated the phosphorylation (P) of JAK2 at different treatment times in CaCo-2 cells. Increased phosphorylation of JAK2 was observed within 30 min after leptin treatment (10 ng/ml) followed by a decline. (B) In LS174T and (C) HCT 116 cells, leptin stimulated the phosphorylation of ERK. The levels of total (T) JAK2, ERK, MAPK and AKT were not altered in three CRC cell lines following leptin treatment.
Cancer progression is a multistep process that enables tumor cells to migrate to points far from a given primary tumor mass (20). Here we showed that leptin enhanced the invasive potential of CaCo-2 cells and the adhesive potential of HCT 116 cells. These effects are consistent with the ability of leptin to enhance the malignant phenotypes of CRC cells, such as local invasiveness and the formation of distant metastasis. Moreover, these results are consistent with our confirmation here of leptin expression in human CRC tissue, which was associated with advanced tumor stage.

To elucidate the signaling pathways involved in leptin-mediated induction of malignant properties of CRC cells, we showed that leptin rapidly induced the phosphorylation of JAK2 and ERK, thus activating these key signal transduction pathways associated with cell growth. These results are consistent with those of others that demonstrated the enhanced activation of the JAK/STAT signaling pathway and elevated expression of genes that are targeted by the STAT3 signaling pathway in colorectal adenomas compared with normal colorectal tissues (34). Although we did not study the pharmacologic inhibition of this pathway here, our results suggest that leptin-mediated JAK and ERK signaling may control invasion and migration. Further investigations are required to clarify the role of CSC in the invasion and migration of CRC cells.

Our results were inconsistent regarding adhesion and invasion, signaling pathways and spheroid formation in the three CRC cell lines. These cell lines mirror the features of the original, individual, diverse tumors from which they were derived and reflect different stages of tumors occurring in the same organ as well as their metastatic features. Therefore, each of these three colon cancer cell lines may possess distinct combinations of unique oncogenes and tumor-suppressor mutations (35).

In conclusion, in the present study, we demonstrated that leptin affected the spheroid formation of colorectal cancer cell lines and regulated adhesion and invasion of colorectal carcinomas through activation of the JAK and ERK signaling pathways.

References