Abstract. We have previously shown that melanoma cells proliferate in response to the metabolic hormones TRH and TSH. The objective of the present study was to test the hypothesis that a third metabolic hormone, leptin, serves as a growth factor for melanoma. Using Western blotting, indirect immunofluorescence, and RT-PCR, leptin receptors were found to be expressed by human melanoma cells. In contrast, cultured melanocytes expressed message for the receptor without detectable protein. Melanoma cells responded to treatment with leptin by activating the MAPK pathway and proliferating. Melanoma cells but not melanocytes, also expressed leptin protein, creating a potential autocrine loop. Examination of human melanoma tumors by immunohistochemistry revealed that melanomas and nevi expressed leptin at a high frequency. Melanomas also strongly expressed the leptin receptor, whereas nevi expressed this receptor to a much lesser degree. We conclude that leptin is a melanoma growth factor and that a leptin autocrine-loop may contribute to the uncontrolled proliferation of these cells.

Introduction

Cutaneous melanoma, the most aggressive form of skin cancer, develops from the malignant transformation of melanocytes located in the dermal/epidermal junction. These normally quiescent cells originate embryologically from pleuripotent neural crest cells, thus sharing this progenitor with cells of the neuroendocrine system. The endocrine features of melanoma have been an active topic of research in our laboratory over the past several years. We have reported that melanoma cells produce thyrotropin releasing hormone (TRH) and respond to this peptide hormone by proliferation, suggesting the presence of an autocrine loop (1). We have also demonstrated the expression of functional thyroid stimulating hormone (TSH) receptors by human melanoma cells, which exhibit a proliferative response to TSH through activation of the mitogen-activated protein kinase (MAPK) pathway (2). These findings raise the possibility that an even wider spectrum of metabolic hormones may play a significant role in melanoma pathobiology.

Over the past decade, considerable interest has been generated in the metabolic hormone leptin. This 16-kDa protein, encoded by the Ob (obese) gene, exhibits a wide range of physiological functions, with roles in energy balance, reproduction, immunity, and inflammation. Best understood is the secretion of leptin by adipocytes as a satiety signal to the hypothalamus, where neurons of the paraventricular nucleus express leptin receptors (3). Multiple forms of the leptin receptor (ObR), resulting from alternative mRNA splicing, have been described and are generally categorized as the long form (ObRb), several short forms (ObRa, c, d, and f), and the soluble form (ObRe) (4). ObRb is thought to function as the major signaling isoform and dominates in the hypothalamus where binding of leptin initiates production of TRH. TRH induces TSH production and secretion by the pituitary gland, the first step in the initiation of metabolic processes involving energy expenditure.

Leptin receptors have been described in various types of tumor cells, including breast, prostate, colon, endometrium, and others, and leptin has been implicated as a growth factor for these cancers (5-9). Based on these reports and our previous data regarding the endocrine nature of melanoma, we hypothesized that melanoma cells express functional leptin receptors. Here, we report that melanoma cells do indeed express these receptors which, in the presence of leptin, signal through the MAPK pathway and induce proliferation. Furthermore, we demonstrate that melanomas also express leptin, providing a potential autocrine growth pathway.

Materials and methods

Patient material. Sections of melanomas and nevi were obtained from the M.D. Anderson Cancer Center (MDACC) Melanoma Tumor Bank. Procurement of all patient materials was conducted with IRB approval and in accordance with HIPAA guidelines.
RT-PCR. RT-PCR was carried out using the GeneAmp RNA PCR Kit (Applied Biosystems, Foster City, CA) and oligonucleotide primers prepared by Sigma Genosys (The Woodlands, TX). Total RNA was extracted using TRizol reagent (Invitrogen, Carlsbad, CA). Reverse transcription to cDNA, carried out using 2 μg of total RNA, took place at 42°C for 30 min. PCR primers and conditions are as follows: Leptin: forward, 5'-CACCAAAACCCTCATCAAGACA-3'; reverse, 5'-TCGAGCATCT-3'; reverse, 5'-CAAAAGCACACCCACTCT CTC-3'. Conditions: 94°C 60 sec; 58°C 60 sec; 72°C 60 sec, magnesium chloride 5.0 mM, 35 cycles.

DNA sequencing. Sequencing of PCR products was performed by the MDACC DNA Core Facility using an ABI Prism 3100 DNA Genetic Analyzer (Applied Biosystems), using Big Dye® v.3.1 dye terminator chemistry (Applied Biosystems). Forward and reverse DNA strands were sequenced for all samples.

Proliferation assay. Cells were plated at a density of 3x10^4 per chamber of a two-chamber tissue culture slide. The following day, the melanoma cells were placed in serum-free media and the melanocytes in a minimal media consisting of RPM-1640 with 10 μg/ml insulin, 5 mg/ml transferrin, and 10 nm hydrocortisone. Leptin was added at the desired concentration after 6 h for an overnight incubation. Effects on proliferation were examined after 24 h, by means of a commercially available BrdU immunocytochemistry assay (BrdU Staining Kit, Zymed, Carlsbad, CA). BrdU labeling reagent (Zymed) was added at a 1:100 dilution to the cultures the evening prior to the assay. On the following day, the cells were fixed with 70% ethanol and incubated in 3% H2O2 in methanol to block endogenous peroxidase activity. After permeabilization and blocking, the cells were incubated with biotinylated anti-BrdU antibody, followed by streptavidin peroxide, the chromogen diaminobenzidine (DAB), and a hematoxylin counterstain. To calculate the percentage of cells positive for BrdU, the stained sections were examined for brown DAB-stained nuclei. A minimum of 100 cells was counted from two different representative areas of the slide, and the mean and standard deviation calculated from these counts.

Immunohistochemistry. Tissue sections were deparaffinized in xylenes, and rehydrated in graded concentrations (100-85%) of ethanol. Sections were placed in Antigen Unmasking Solution (Vector Laboratories, Burlingame, CA) and microwave intermittently for a total of 10 min, to maintain boiling temperature. After cooling, the slides were placed in 3% H2O2 in cold methanol for 15 min to block endogenous peroxidase activity. This step was followed by permeabilization with 0.05% Triton X-100 (Sigma, St. Louis, MO) in phosphate-buffered saline for 5 min. An avidin-biotin-peroxidase complex (ABC) kit (Vectastain, Vector Laboratories) was then used for antigen detection. After a 30-min incubation with blocking serum, the primary antibody was applied and incubated for 2 h at room temperature. The slides were then washed, incubated for 30 min with secondary biotinylated antibody, followed by a 30-min incubation with the ABC reagent. The immunolabeling was developed with the chromogen 3-amino-9-ethylcarbazole (AEC). Hematoxylin was applied as a counterstain.

Immunolabeling was scored separately for two variables: first, for number of positive cells; second, for the overall intensity of immunoreactivity of the positive cells. Scoring for the number of positive tumor cells was defined as follows: '0', <5% positive cells; '1', 5-25% positive cells; '2', 26-75% positive cells; and '3', >75% positive cells. Intensity scoring consisted of: '0', no staining; '1', weak staining; '2', moderate staining; and '3', intense staining. The slides were interpreted by two readers (A.H.D. and J.A.E.).
Statistical analysis. Differences in the mean percentage of BrdU positive cells at different leptin concentrations were examined for significance using the unpaired t-test. IHC scores were compared between melanomas and nevi using the Mann-Whitney test. Minitab software was used for all computations and significance was set at p≤0.05.

Results

Expression of leptin receptors by melanocytic cells. Melanoma cell lines were first examined for ObR expression. Antibodies from a variety of sources were tested by immunoblotting, yielding positive results, but of relatively poor quality. The result of a representative blot, examining four human melanoma cell lines, is shown in Fig. 1, revealing bands of the appropriate size for ObRb (~125 kDa) and ObRa (~100 kDa). To better demonstrate the presence of ObR expression, indirect immunofluorescence (IF) was performed using non-permeabilized cells grown on chamber slides. The MCF-7 breast cancer cell line, which has been previously reported to express ObR, was included as a positive control (8). As shown in Fig. 2A, both of the melanoma cell lines included in this experiment expressed ObR whereas expression was notably absent or below the level of detection in melanocytes. For further confirmation of these findings, RT-PCR was performed using RNA from the same lines. Unlike the antibody used in the above experiments, which does not distinguish between the various isoforms, the PCR primers were specific for the long isoform ObRb. All lines, including the melanocytes, expressed message for ObRb (Fig. 2B), which was confirmed by DNA sequencing. Taken together, these findings convincingly demonstrate the expression of ObR by human melanoma cells. Although cultured melanocytes also express ObR message, it appears that translation is inhibited, as protein is not detected.

Activation of proliferation and pathway signaling in response to leptin. To explore the hypothesis that melanoma leptin receptors are functional and that leptin is a growth factor for these cells, cultured melanoma cells and melanocytes were exposed to increasing concentrations of leptin from 0 to 1000 ng/ml and examined for induction of proliferation by BrdU uptake. The melanoma cells were serum-starved for 6 h prior to an overnight incubation with leptin. Because the melanocytes did not survive serum starvation, they were assayed instead in a minimal medium consisting of RPMI-1640 with 10 μg/ml insulin, 5 μg/ml transferrin, and 10 nm hydrocortisone (2). Both melanoma lines included in these experiments proliferated in response to leptin, with an optimal concentration of 100 ng/ml, which is a physiologically relevant level (Fig. 3). Melanocytes also responded to leptin with a proliferative response. However, in contrast to the melanoma cells, this response did not appear to be dose-dependent, suggesting the presence of sparse surface receptors that were saturated at the lowest leptin concentration. Furthermore, the melanocyte proliferation rate was low compared to melanoma cells, remaining below 10%. The difference in melanocyte and melanoma cell proliferation was likely underestimated in these assays as the melanocytes had additional growth factors in their media as described above.

Activation of the MAPK and signal transducer and activator of transcription 3 (STAT3) pathways by leptin have been reported in colon and breast cancer cell lines (8,9). Both of these pathways by leptin have been reported in colon and breast cancer cell lines (8,9). Both of these pathways by leptin have been reported in colon and breast cancer cell lines (8,9).
to be the more prevalent of the two. To establish the pattern of leptin signaling in melanoma, WM793 cells were serum starved overnight, and then treated with leptin 100 ng/ml for 2, 5, 10, 15, and 30 min. As anticipated, the phosphorylated forms of ERK (pERK) and STAT3 (pSTAT3) were detected in the baseline samples. An increase in the level of pERK was seen at the earliest timepoint, and peaked at 5 min (Fig. 4). In contrast, no evidence for increased STAT3 phosphorylation was detected. Similar findings were seen with WM35 cells (data not shown). These results indicate that leptin preferentially activates the MAPK pathway in human melanoma cells, and that signaling through STAT3 is not a consequence of ObR activation in this system.

Figure 3. Proliferative response of melanoma cells to leptin. Melanoma cells and melanocytes were treated overnight with escalating concentrations of leptin. Both melanoma cell lines demonstrate a dose-dependent proliferative response. Data are representative of duplicate experiments. Charted are the mean of two separate counts of BrdU-positive cells; error bars represent the standard deviation. Asterisks indicate a significant difference compared to the '0' leptin concentration.

Figure 4. Pathway activation by leptin in melanoma cells. WM793 cells were treated with leptin 100 ng/ml and assayed for ERK and STAT3 phosphorylation at the indicated timepoints. (A) An increase in the amount of pERK is detected as early as 2 min after leptin exposure. In contrast, levels of pSTAT3 do not change over the time course. (B) Enhanced ERK phosphorylation was confirmed by densitometry.

Figure 5. Production of leptin by melanoma cells and melanocytes. (A) Leptin is demonstrated in lysates of three of four melanoma cell lines by Western blotting. (B) Leptin message is detected in WM793 and WM35 melanoma cells, as well as in melanocytes. Adipocyte mRNA serves as a positive control. (+), RT, followed by PCR; (-), PCR without RT, as a control for amplification of genomic DNA; M-cyte, melanocyte.

Production of leptin by melanoma cells. The most obvious source of leptin which might stimulate melanoma cells in vivo is the naturally circulating hormone produced by adipocytes in response to caloric intake. However, a potential alternative or complementary mechanism would be production and secretion of leptin by the tumor cells themselves as part of an autocrine growth stimulatory pathway. To explore the second alternative, the presence of leptin protein was examined in a panel of melanoma cell lines and melanocytes. Fig. 5A demonstrates that leptin is indeed present in lysates of three of the four melanoma cell lines examined and is absent in melanocytes. To rule out the possibility that the leptin detected was exogenous and simply taken up from the serum in the culture media, RT-PCR and sequencing were performed to
confirm the presence of leptin message in these cells. As demonstrated in Fig. 5B, leptin message was found in melanoma cells, confirming that the hormone was produced endogenously and supporting the hypothesis of a leptin autocrine loop. Similar to the case with ObR, melanocytes produce message for leptin which is not translated into the protein.

ObR and leptin expression by human melanoma and nevi. Finally, we examined ObR expression in 16 human primary melanomas and 15 benign nevi. This group of samples has been previously described in terms of maturation detected by the anti-leptin receptor antibody (11). Here, we have quantitated immunostaining for ObR and leptin, by both percent of positive cells and intensity of staining (Materials and methods), and compared expression in the two sample types (Fig. 6). As shown in Table 1, only one of the 16 nevi (6.7%), expressed ObR in a high percentage of cells (score of 2 or 3) and 60% were completely negative; in contrast, 63% of melanomas had a high percentage score and 31% were negative (p=0.009). The intensity scores were in agreement with these findings and bordered on significance (p=0.054). Additional sections were available from all 15 nevi and 11 of the melanomas for immunohistochemical study with antibody to leptin. In this regard, the two sample types were more similar. Although melanomas and nevi differed significantly in their percentage scores, expression was generally diffuse in both, with 73% of nevi and 100% of melanomas receiving a score of 2 or 3. Intensity scores for leptin did not significantly differ between the two sample types. These data suggest that while melanomas and nevi both produce a considerable amount of the leptin hormone,

Table 1. Comparison of IHC scores between nevi and melanomas.

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<th>Score</th>
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<th>Melanomas N (%)</th>
<th>Score</th>
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Figure 6. Immunostaining of melanomas and nevi for ObR and leptin. Representative sections are shown. The melanoma expresses leptin and its receptor. The nevus is positive for leptin, as well, but negative for the receptor. AEC, x20.
the receptor is found more frequently on the melanoma cells, providing the opportunity for dysregulated autocrine-stimulated growth in the malignant cells.

Discussion

In keeping with our previous data showing proliferative responses of melanoma cells to the metabolic hormones TRH and TSH, we now demonstrate a similar response to the hormone leptin. The unique aspect in this case is the fact that melanoma cells simultaneously express the leptin receptor and produce the hormone ligand, creating a potential autocrine loop by which dysregulated tumor growth can be supported. Furthermore, expression of leptin and ObR follows a progressive pattern across the spectrum of melanocytic cell types. Melanocytes express message for leptin and ObR, but little or no protein; nevi, as benign tumors, express high levels of leptin and low levels ObR; and melanoma cells express high levels of both. These findings lend credence to the concept that the proposed leptin autocrine loop is absent in normal cells, moderated in nevi, and uncontrolled in melanomas.

The pleuripotent nature of leptin is well-suited for exploitation by melanoma cells. Beyond its role in food intake and energy balance, leptin additionally functions as a pro-inflammatory ‘adipokine’ with a broad range of activities including cytokine production, cellular immunity, and inflammation (12). Inflammatory processes, in particular, are highly relevant to melanoma progression. We have reported a strong correlation of tumor-expressed inducible nitric oxide synthase (iNOS) with shortened survival of Stage III melanoma patients (13). iNOS catalyzes the production of the inflammatory mediator nitric oxide (NO), which has been shown to promote melanoma cells from apoptosis (14). A stimulatory effect of leptin on nitric oxide production, secondary to enhanced expression of iNOS, is well-established, best demonstrated in vascular and neuronal systems (15-17). Similarly, expression of the inflammatory enzyme cyclooxygenase-2 (COX-2) by human melanomas predicts poor patient outcomes, and leptin expression of iNOS, is well-established, best demonstrated in vascular and neuronal systems (15-17). Similarly, expression of the inflammatory enzyme cyclooxygenase-2 (COX-2) by human melanomas predicts poor patient outcomes, and leptin

The elevation of circulating leptin levels in obese individuals has generated interest in a role for leptin in the biology of melanoma, one of the few cancers associated with an increased risk for developing a melanoma, and the elevated leptin levels could not be attributed to obesity (25). One possibility is that individuals genetically predisposed to have higher circulating leptin levels, regardless of body mass, are at greater risk for the development of leptin-responsive tumors. Alternatively, the higher pre-diagnosis leptin levels seen in patients who develop these cancers may be a product of the developing tumor. If this is the case, serum leptin levels may serve as a screening tool and a marker of progression.

Taken together, there is mounting evidence for a role of leptin and ObR in cancer biology; our data now bring this process to the forefront in melanoma. Important issues for future research include the involvement of leptin signaling in melanoma inflammatory pathways and the possibility that tumor-derived leptin actually enters the circulation, elevating measurable levels and potentially functioning as a melanoma biomarker.

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References


