Energy Balance Modulates Mouse Skin Tumor Promotion through Altered IGF-1R and EGFR Crosstalk

Tricia Moore1, Linda Beltran1, Steve Carbajal1, Stephen D. Hursting2, and John DiGiovanni1,2

Abstract

Obesity, an established risk factor for epithelial cancers, remains prevalent in the United States and many other countries. In contrast to positive energy balance states (overweight, obesity), calorie restriction (CR) has been shown to act as a universal inhibitor of tumorigenesis in multiple animal models of human cancer. Unfortunately, the mechanisms underlying the enhancing effects of obesity or the inhibitory effects of CR on cancer etiology remain elusive. Here, we evaluated the impact of dietary energy balance manipulation on epithelial carcinogenesis and identified several potential mechanisms that may account for the differential effects of obesity and CR on cancer. Obesity enhanced tumor promotion during epithelial carcinogenesis, in part, due to altered insulin-like growth factor-1 receptor (IGF-1R)/EGF receptor (EGFR) crosstalk and downstream signaling to effectors such as Akt/mTOR. Obesity-induced changes in cellular signaling subsequently led to altered levels of cell-cycle proteins that favored enhanced epidermal proliferation during tumor promotion. In contrast, CR reduced susceptibility to tumor promotion, attenuated IGF-1R/EGFR crosstalk and downstream signaling, and altered levels of cell-cycle proteins that favored reduced epidermal proliferation during tumor promotion. Collectively, these findings suggest potential targets for the prevention of epithelial cancers, as well as for reversal of obesity-mediated cancer development and progression. Cancer Prev Res; 1–11. ©2012 AACR.

Introduction

Energy balance refers to the relationship between caloric consumption and energy expenditure (1). Epidemiologic and animal studies have established a direct correlation between positive energy balance states (e.g., overweight, obesity) and risk of developing multiple cancers (2–5). In contrast, negative energy balance states, such as calorie restriction (CR), have been shown to consistently inhibit tumorigenesis in animal models regardless of mode of tumor induction (1, 6–8). Previous studies have evaluated the impact of fat consumption and/or CR on 2-stage skin carcinogenesis (8–10). This model of chemically induced epithelial carcinogenesis enables mechanistic evaluation of dietary manipulation during all stages of tumor development, including tumor initiation, promotion, and progression (11). Boutwell (8) found that CR inhibited skin tumor development using the 2-stage carcinogenesis protocol, and Birt and colleagues (9, 10) showed that 40% CR consistently inhibited tumor promotion by 12-O-tetradecanoylphorbol-13-acetate (TPA), with little or no effects on either tumor initiation or tumor progression. Although these findings indicated a role for energy balance in the modulation of 2-stage skin carcinogenesis, especially during tumor promotion, the effect of diet-induced obesity (DIO) per se has not been adequately studied in this model.

Several mechanisms have been proposed to explain the inhibitory effects of CR on tumor development (12, 13) including elevated serum corticosterone (6, 14) and alterations in cellular signaling. TPA-induced epidermal AP-1 activation and extracellular signal–regulated kinase (ERK) phosphorylation was reduced in mice subject to 40% CR (15–17). Xie and colleagues (18) reported a reduction in TPA-mediated activation of epidermal phosphoinositol 3-kinase (PI3K) and Ras signaling following 20% CR. Recently, we reported that both CR and DIO modulated steady-state growth factor signaling pathways in mouse epidermis, liver, and prostate (19). Additional studies suggested that these effects may be mediated by diet-induced changes in circulating insulin-like growth factor-1 (IGF-1) levels and altered IGF-1 receptor (IGF-1R) signaling (19, 20).

Diets of varying caloric density were used to induce changes in body mass and body fat content to determine the impact of both positive and negative energy balance on the promotion stage of multistage epithelial carcinogenesis in mouse skin. We provide novel evidence showing that dietary energy balance affects susceptibility to epithelial carcinogenesis, in part, through diet-induced changes in

1Division of Pharmacology and Toxicology, College of Pharmacy, and 2Department of Nutritional Sciences, College of Natural Sciences, The University of Texas at Austin, Austin, Texas

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Corresponding Author: John DiGiovanni, Dell Pediatric Research Institute, The University of Texas at Austin, 1400 Barbara Jordan Blvd. Austin, TX 78723. Phone: 512-495-4726; Fax: 512-495-4945; E-mail: John digsiovanni@austin.utexas.edu
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IGF-1R and EGF receptor (EGFR) crosstalk and downstream signaling during tumor promotion.

Materials and Methods

Animals

ICR female mice (3–4 weeks of age, Harlan Teklad) were group housed; however, mice maintained on a 30% CR regimen were separated for 1 hour for feeding. Mice were weighed before randomization and then every 2 weeks for the duration of the experiments.

Diet

Four diets ranging in caloric density were used (Research Diets) and have been previously described: 30% CR diet, 15% CR diet, 10 kcal% fat (ad libitum), and 60 kcal% fat (ad libitum; ref. 1). CR was achieved by administering a daily aliquot equivalent to 70% or 85% of the daily energy consumed by the 10 kcal% fat group, as previously detailed (19).

Serum analysis and body fat composition

Blood was collected immediately following CO₂ asphyxiation, serum was prepared, and serum IGF-1, insulin, leptin, and adiponectin levels were determined as previously described (19, 21). Corticosterone levels were determined by ELISA (Kamiya Biomedical Co.). Percentage of body fat was determined as previously described (22).

Two-stage skin carcinogenesis

ICR female mice were placed on the 10 kcal% fat control diet at 7 weeks of age and initiated with 25 nmol of 7,12-dimethylbenz[a]anthracene (DMBA; Eastman Kodak Co.). Four weeks following initiation, mice were randomized and placed on the 4 diets (n = 30 per group). Four weeks later, mice received twice weekly topical treatments of 3.4 nmol of TPA (Alexis Biochemicals) for 50 weeks. Tumor incidence, average carcinomas per mouse, carcinomas (SCC) were confirmed by histopathology.

Epidermal hyperproliferation

Female ICR mice were maintained on the diets described above for 15 weeks, after which they were treated twice weekly for 2 weeks with either acetone vehicle or 3.4 nmol of TPA (n = 3 per group). Mice were injected with bromodeoxyuridine (BrdUrd; Sigma Aldrich; 100 μg/g body weight) 30 minutes before sacrifice. Whole skin sections were excised, processed, and evaluated for epidermal thickness and labeling index as previously described (20).

Epidermal protein lysates

For short-term in vivo experiments, ICR mice were maintained on the diets for 15 weeks, after which they received a single application of either acetone (vehicle) or 3.4 nmol of TPA. Mice were killed either 6 (acetone, TPA) or 18 hours (TPA) after treatment, epidermis was scraped and protein lysates were prepared as previously described (20).

Cell culture

C50 cells are a nontumorigenic keratinocyte cell line derived from spontaneously immortalized normal mouse keratinocytes. The cells were obtained from Dr. Susan Fischer (UT MDACC, Houston, TX) and were cultured as previously described (23) with no further characterizations. When cells reached approximately 80% confluency, they were serum and growth factor starved for 24 hours. Plates were then stimulated with either 25 ng/mL recombinant human IGF-1 (rhIGF-1; Sigma Aldrich) or 10 ng/mL EGF (BD Biosciences) and harvested at multiple time points for protein or RNA isolation.

Immunoprecipitation and Western blot analysis

Immunoprecipitation and Western blot analyses were conducted using lysates prepared from either epithelium or cultured keratinocytes. For co-immunoprecipitation experiments, lysates were precipitated with IGF-1R (Cell Signaling) or EGFR (Millipore) antibodies using the Dynabead Protein G IP Kit (Invitrogen). Western blot analyses were conducted as previously described (20).

Real-time quantitative reverse transcriptase PCR

For in vivo experiments, ICR mice were maintained on the diets for 15 weeks, after which they received a single application of either acetone vehicle or 3.4 nmol of TPA. Mice were killed either 6 (acetone, TPA) or 18 hours (TPA) after treatment and epidermis was scraped for RNA isolation (n = 3 per group). For in vitro experiments, C50 cells were stimulated with either IGF-1 or EGF, harvested for RNA, and RNA was isolated using the Qiagen RNeasy Protect Mini Kit. RNA quality was assessed using the Agilent 2100 Bioanalyzer (Agilent Technologies, Inc.), and quantitative PCR (qPCR) was carried out, as previously described, using assays on demand specific to TGF-α, HB-EGF, amphiregulin, and EGF (21). RNA was normalized to glyceraldehyde-3-phosphate dehydrogenase (GAPDH). All measurements were duplicated.

Results

Dietary energy balance effects on skin tumor promotion

A 2-stage skin carcinogenesis experiment was conducted using 4 experimental diet groups to generate lean (30% CR), normal (15% CR), overweight (10 kcal% fat), and obese (DIO, 60 kcal% fat) body phenotypes, as previously described (5, 21, 24). Groups were initiated with DMBA, maintained on 10 kcal% diet for 4 weeks, randomized into the experimental groups and promoted twice weekly with 3.4 nmol of TPA for 50 weeks. Tumor incidence and multiplicity (average number of papillomas per mouse) were calculated weekly until the latter reached a plateau at 29 weeks. CR (15% and 30%) significantly inhibited
papilloma multiplicity when compared with both the overweight control and DIO groups \((P < 0.05; \text{Mann–Whitney } U\) test); however, no significant differences were observed between the overweight control and DIO groups (Fig. 1A; Supplementary Table S1). Dietary energy balance manipulation had no significant effect on papilloma incidence under these experimental conditions (Fig. 1B; Supplementary Table S1).

Treatments continued for an additional 21 weeks to evaluate the impact of dietary energy balance on tumor progression. As shown in Fig. 1C and D, the number and incidence of SCCs was significantly reduced in the 30% CR group \((P < 0.05, \text{Mann–Whitney } U\) test and \(P < 0.05, \chi^2\) test, respectively), compared with the 15% CR, overweight control and the DIO groups; however, dietary manipulation did not affect the overall rate of malignant conversion (see also Supplementary Table S1). A second 2-stage skin carcinogenesis experiment was carried out that yielded nearly identical findings (data not shown).

**Effect of long-term dietary manipulation on body profiles during two-stage skin carcinogenesis**

Body mass, percentage of body fat, and circulatory protein levels (i.e., IGF-1, insulin, leptin, adiponectin)

![Figure 1. Effect of dietary energy balance on 2-stage skin carcinogenesis and TPA-induced epidermal proliferation. Two-stage skin carcinogenesis was conducted using lean (30% CR), normal (15% CR), overweight control (10 kcal% fat), and obese (60 kcal% fat) mice \((n = 30\) per group). A, tumor multiplicity; CR (30%, 15%) significantly reduced tumor multiplicity \((P < 0.05, \text{Mann–Whitney } U\) test). B, tumor incidence. C, average carcinomas per mouse; 30% CR significantly reduced, whereas DIO (60 kcal%) significantly increased the average number of carcinomas per mouse \((P < 0.05, \text{Mann–Whitney } U\) test). D, carcinoma incidence. CR (30%, 15%) significantly reduced carcinoma incidence \((P < 0.05, \chi^2\) test). E and F, ICR female mice were maintained on the four diets for 15 weeks and were treated with either acetone (white bars) or 3.4 nmol TPA (black bars), twice weekly for 2 weeks \((n = 3\) per group). E, representative BrdUrd-stained skin sections. F, epidermal thickness (top) and percent BrdUrd incorporation (bottom). *\(a\), statistically different from all similarly treated groups; *\(a\), significantly different from similarly labeled values \((P < 0.05, \text{Mann–Whitney } U\) test).}
Dietary energy balance modulates TPA-induced epidermal hyperproliferation

To evaluate potential mechanisms whereby dietary energy balance modulates 2-stage skin carcinogenesis, additional mice were maintained on the 4 diets for 15 weeks, thus generating lean, normal, overweight, and obese body phenotypes. Body mass, percentage of body fat, and levels of circulatory proteins were then determined at this time point (Supplementary Table S2.II). Body mass and percentage of body fat, as well as levels of circulatory proteins were observed between the overweight control and DIO groups at the end of the 50-week experimental period.

Additional analyses were conducted to determine the impact of dietary energy balance on levels of negative cell-cycle regulatory proteins. Levels of both p27 and p21 were significantly higher in 30% CR mice (vehicle control), as compared with DIO mice (Fig. 2C). Diet-induced changes in cyclin A and c-myc levels following TPA treatment were also significant at the 6-hour time point (P < 0.05, Student t test).

Additional analyses were conducted to determine the impact of dietary energy balance on levels of negative cell-cycle regulatory proteins. Levels of both p27 and p21 were significantly higher in 30% CR mice (vehicle control), as compared with DIO mice (Fig. 2C). Following TPA treatment (6 hours), both p27 and p21 levels were significantly higher in mice maintained on the 30% CR diet, as compared with mice maintained on the DIO diet (P < 0.05, Student t test). A similar trend was observed at 18 hours,
although these differences were not significantly different. Preliminary experiments (data not shown) showed increased nuclear localization of both p27 and p21 in the epidermis of CR mice, as compared with mice maintained on the DIO regimen, suggesting that dietary energy balance may also modulate these negative cell-cycle regulatory proteins through additional mechanisms.

**Activation of the IGF-1R stimulates IGF-1R/EGFR crosstalk in cultured mouse keratinocytes**

Both our current and previous data suggested a role for altered IGF-1 levels in modulating the effect of dietary energy balance on epithelial carcinogenesis. A reduction in circulating IGF-1, either due to genetic (LID mouse model) or dietary manipulation (CR) attenuated both IGF-1R and EGFR signaling (refs. 19, 20; Fig. 2), indicating a potential for crosstalk between these 2 receptors. C50 cells were used to examine the role of IGF-1 in modulating both IGF-1R and EGFR signaling and crosstalk, as well as erbB2 activation. Cultured cells were stimulated with IGF-1 (25 ng/mL), harvested at multiple time points (0–120 minutes), and analyzed for IGF-1R activation (measured by IRS-1 phosphorylation) and for EGFR and erbB2 activation (measured by phosphorylation) using Western blot analysis. IGF-1 treatment significantly induced activation of all 3 cell surface receptors (Fig. 3). EGFR phosphorylation was rapid, with maximal activation occurring within 5 minutes and phosphorylation was maintained above the basal level up to 120 minutes. EGFR phosphorylation correlated directly with IGF-1–induced activation of the IGF-1R. Rapid phosphorylation of erbB2 also paralleled both IGF-1R and EGFR activation; however, erbB2 activation was maintained at a
consistent level across all time points examined. Stimulation of C50 cells with EGF (10 ng/mL) over the same time course led to rapid activation of both the EGFR and erbB2; however, there was no significant effect on IGF-1R activation (Fig. 3).

Previous studies have identified several potential mechanisms for IGF-1R and EGFR crosstalk (26–30). The ability of IGF-1 to induce heterodimerization between the IGF-1R and the EGFR was evaluated in co-immunoprecipitation experiments carried out using lysates prepared from C50 cells treated with either IGF-1 or EGF. As shown in Fig. 4A, IGF-1 treatment induced a statistically significant increase in IGF-1R/EGFR association from 5 to 60 minutes. In contrast, EGF had no observable effect. Additional experiments were carried out to determine the impact of IGF-1 stimulation on levels of mRNA for the EGFR and EGFR ligands. qPCR was carried out using RNA extracted from C50 cells (stimulated with IGF-1 or EGF and harvested at 0–120 minutes) to determine the relative expression of EGFR, TGF-α, HB-EGF, and amphiregulin. Both IGF-1 and EGFR stimulation significantly increased expression of HB-EGF and amphiregulin, with a greater induction occurring with EGF treatment (Fig. 4B). These relative increases in mRNA expression, however, were observed only at later time points (≥ 60 minutes). No increases in TGF-α and EGFR mRNA expression were observed following stimulation of C50 cells with either IGF-1 or EGF (Fig. 4B).

**Dietary energy balance alters IGF-1R/EGFR crosstalk in vivo**

Finally, we evaluated the impact of dietary energy balance manipulation on IGF-1R/EGFR crosstalk during tumor promotion in mouse epidermis in vivo. Female ICR mice were maintained on the previously described 30% CR and DIO diets for 15 weeks. These lean and obese mice were then treated with a single application of acetone (vehicle) or 3.4 nmol of TPA and killed 6 or 18 hours later. Epidermal lysates and epidermal RNA were prepared for Western blot and qPCR analyses, respectively. Co-immunoprecipitation experiments were carried out to evaluate diet-induced changes in IGF-1R/EGFR heterodimerization. As shown in Fig. 5A, there was a greater degree of association between the IGF-1R and EGFR in obese mice relative to CR mice in the absence of TPA treatment. Furthermore, TPA (6 hours) induced a significant increase in IGF-1R and EGFR association in both the CR and obese mice although the relative increase in heterodimerization was significantly greater in obese mice (P < 0.05, Student t test).
Dietary manipulation did not modulate mRNA levels of EGFR and EGFR ligands in the absence of TPA treatment; however, DIO significantly increased levels of mRNA for EGFR ligands compared with 30% CR, following treatment with TPA (Fig. 5B). Specifically, DIO significantly increased mRNA levels of TGF-α (18 hours), HB-EGF, amphiregulin, and EGFR following TPA treatment. EGFR mRNA levels were not affected by diet or treatment with TPA. Collectively, the in vitro and in vivo results suggest that dietary energy balance modulates IGF-1R/EGFR crosstalk, at least in part, due to diet-induced changes in mRNA levels of EGRF ligands and changes in IGF-1R and EGFR heterodimerization.

Discussion

In this study, potential mechanism(s) underlying the effects of dietary energy balance manipulation on skin tumor formation during the promotion phase of 2-stage skin carcinogenesis were examined. Consistent with previously published data, both 15% and 30% CR significantly inhibited the promotion of papillomas by TPA compared with both the overweight control and DIO groups but had no significant effect on tumor progression (8–10). Notably, no significant difference in tumor response was seen between the DIO and overweight control groups, although the DIO regimen significantly increased both body mass and percentage of body fat. Previous mammary carcinogenesis studies also reported that despite significant differences in body mass, no significant increase in tumor incidence occurred when dietary fat consumption exceeded 20 Kcal% to 30 Kcal% (31–33). These data support the hypothesis that in some model systems, there is an upper limit to the effects of a positive energy balance state above which further increases in total calories do not further increase tumor response.

Genetic reduction of circulating IGF-1 levels (i.e., LID mouse model) attenuated TPA-induced epidermal hyperproliferation, which correlated with reduced responsiveness to skin tumor promotion by TPA (20). In comparison to the overweight control and DIO groups, both the 30% and 15% CR groups had significantly reduced epidermal thickness and LI in the absence of TPA. Following TPA treatment, we...
observed a significantly progressive effect of dietary energy balance across all the diet groups on the epidermal proliferative response, with 30% CR exhibiting the lowest response and DIO exhibiting the highest. Several other studies have shown that energy balance impacts cellular proliferation in both normal and tumorigenic tissue (e.g., mammary, colon, liver, epidermis, and bladder; refs. 34–38). Collectively, our results suggest that manipulating dietary energy balance impacts TPA-mediated epidermal hyperproliferation, thus modulating epithelial carcinogenesis in mouse skin.

Previous studies have indicated an important role for IGF-1R and EGFR activation in TPA-induced epidermal hyperproliferation and tumor promotion in mouse skin (reviewed in ref. 39). Inhibition of TPA-induced EGFR activation in vivo, using either RG13022 (tyrosine kinase inhibitor) or GW2974 (dual specific erbB2/EGFR inhibitor) significantly reduced TPA-induced epidermal hyperproliferation as well as skin tumor promotion (GW2974; refs. 40, 41). In addition, epidermal IGF-1 overexpression increased epidermal proliferation, both in the presence and absence of TPA, and increased sensitivity to 2-stage skin carcinogenesis (42, 43). Furthermore, dietary energy balance induced changes in steady-state activation of these critical growth factor receptors in epidermis and other epithelial tissues of FVB/N and C57BL/6 mice (19). As shown in Fig. 2A, 30% CR reduced activation and/or phosphorylation of the IGF-1R, EGFR, and downstream signaling following treatment with TPA compared with the DIO group. Xie and colleagues reported a similar reduction in TPA-mediated epidermal Akt activation in CR mice (18).

The impact of dietary manipulation on levels of positive and negative cell-cycle regulatory proteins was also evaluated. Notably, in mice treated with TPA, the levels of positive cell-cycle regulatory proteins (cyclin D1, cyclin E, cyclin A, c-myc) correlated directly with caloric consumption, whereas the levels of negative cell-cycle regulatory proteins (i.e., p21, p27) inversely correlated with caloric consumption. As reported earlier, elevated IGF-1R activation or increased Akt activity, respectively, leads to increased levels of positive cell-cycle regulatory proteins (cyclins D, A, E, c-myc) during tumor promotion, providing a mechanism for the upregulation of cellular proliferation observed in mouse epidermis after treatment with TPA (25, 44). Together, these observations suggest that dietary energy balance–induced alterations in epidermal proliferation seen during tumor promotion result from altered upstream growth factor signaling which modulates levels of both positive and negative cell-cycle regulatory proteins.

Using C50 cells, we established that IGF-1 treatment specifically induced activation of not only the IGF-1R but also the EGFR and erbB2. Similar to previous studies (29,
30), EGF stimulation did not activate the IGF-1R, suggesting that IGF-1R/EGFR crosstalk may be unidirectional. Thus, IGF-1 modulated activation of the EGFR in cultured keratinocytes through multiple mechanisms, including induction of IGF-1R/EGFR heterodimerization and heightened expression of EGFR ligand mRNA. Both mechanisms likely contribute to the sustained activation of the EGFR observed at later time points; however, they do not fully explain the rapid induction of EGFR and erbB2 phosphorylation observed following IGF-1 stimulation. EGFR phosphorylation may also occur as a result of IGF-1–mediated ectodomain shedding of preformed, membrane-bound EGFR ligands (29, 30, 45), which may explain the early activation of the EGFR by IGF-1 observed in our studies. The activation (phosphorylation) of erbB2 seen in IGF-1–treated cells is likely due to EGFR-erbB2 heterodimerization, although we cannot completely rule out the possibility that other mechanisms may be involved, including heterodimerization between IGF-1R and erbB2 (46, 47). Finally, we conducted in vivo experiments to determine whether dietary energy balance modulates IGF-1R/EGFR crosstalk. CR (30%) not only reduced IGF-1R/EGFR heterodimerization in epidermis following TPA treatment but also significantly reduced mRNA expression of EGFR ligands, as compared with DIO. To our knowledge, this is the first report showing that dietary energy balance impacts IGF-1R and EGFR crosstalk. These diet-related changes in receptor crosstalk account, at least in part, for the observed reduction (CR) or increase (DIO) in epidermal signaling that controls the proliferative response to TPA during tumor promotion.

In the current study, the 2-stage skin carcinogenesis model was used primarily as a general model of epithelial carcinogenesis to evaluate mechanisms associated with dietary energy balance manipulation. Nevertheless, the observed changes may also be relevant to UVB-mediated skin carcinogenesis. In this regard, diet-induced obesity has been shown to increase UVB-mediated inflammation as well as inflammation and cell survival signaling (48). Akt activation (phosphorylation) was found to be elevated in the epidermis of obese mice exposed to UVB compared with mice on a control diet in this study. In a model of leptin deficiency-induced obesity (i.e., ob/ob mice), similar increases in UVB-induced inflammation signaling (NF-κB, COX-2, TNFα) and survival signaling (Akt activation) were observed compared with wild-type mice (49). Elevated levels of proliferating cell nuclear antigen (PCNA) and cyclin D1 were also observed in epidermis of the obese mice following exposure to UVB. Thus, a number of changes similar to those seen in our current study have been observed in skin/epidermis of obese mice (either diet-induced or genetically induced obesity) following exposure to UVB. In a study by Hopper and colleagues (50), CR was reported to inhibit UVB-induced upregulation of AP-1-DNA binding and UVB-induced alteration in AP-1 constituent protein levels suggesting that CR inhibited events associated with UVB-mediated skin tumor promotion. Only limited information is available on the potential link between dietary energy balance and non-melanoma skin cancer (NMSC) in humans (reviewed in ref. 51); however, stronger evidence points to a relationship between obesity and melanoma skin cancer (reviewed in ref. 51). Further studies evaluating the role of dietary energy balance in both NMSCs and melanoma skin cancer seem warranted on the basis of the data presented in our current studies as well as those published on UV effects in mouse models of obesity as noted above.

In conclusion, we have shown that CR reduced, whereas DIO increased, signaling through the IGF-1R and EGFR following TPA treatment. Both in vitro and in vivo studies (20) suggest that levels of circulating IGF-1, which are modulated by dietary energy balance, regulate activation of the IGF-1R which in turn regulates crosstalk with the EGFR (and erbB2). These diet-induced changes in IGF-1R and EGFR activation subsequently altered downstream signaling to Akt and mTOR (and other signaling pathways), thus modulating levels and activity of cell-cycle regulatory proteins. These changes in cell-cycle regulatory proteins correlated directly with diet-induced changes in TPA-induced epidermal proliferation. Supplementary Figure S1 summarizes the effects of dietary energy balance on signaling pathways and cell-cycle proteins observed in the present study in relation to altered epidermal proliferation and tumor promotion. Taken together, these effects of dietary energy balance on IGF-1R/EGFR signaling and crosstalk, cell-cycle regulation, and epidermal proliferation provide a plausible mechanism for the inhibitory effects of negative energy balance and the enhancing effects of positive energy balance on susceptibility to skin tumor promotion during 2-stage skin carcinogenesis. These studies provide new information about potential molecular targets for prevention of epithelial cancers and for reversing the effects of obesity on cancer development.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

Authors’ Contributions
Conception and design: T. Moore, S.D. Hursting, J. DiGiovanni
Development of methodology: T. Moore, J. DiGiovanni
Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): T. Moore, L. Beltran, S. Carbajal, J. DiGiovanni
Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): T. Moore, S.D. Hursting, J. DiGiovanni
Writing, review, and/or revision of the manuscript: T. Moore, L. Beltran, S. Carbajal, S.D. Hursting, J. DiGiovanni
Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases): T. Moore, J. DiGiovanni
Study supervision: J. DiGiovanni

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