ERBB-2 overexpression confers PI 3′ kinase-dependent invasion capacity on human mammary epithelial cells

KM Woods Ignatoski¹, T Maehama², SM Markwart³, JE Dixon², DL Livant³ and SP Ethier¹

Departments of ¹Radiation Oncology, ²Biological Chemistry and ³Anatomy and Cell Biology, University of Michigan Medical School, Ann Arbor, MI 48109–0616, USA

Summary Amplification and overexpression of ERBB-2 in human breast cancer is thought to play a significant role in the progression of the disease; however, its precise role in the etiology of altered phenotypes associated with human breast cancer is unknown. We have previously shown that exogenous overexpression of ERBB-2 conferred growth factor independence on human mammary epithelial cells. In this study, we show that ERBB-2 overexpression also causes the cells to acquire other characteristics exhibited by human breast cancer cells, such as anchorage-independent growth and invasion capabilities. ERBB-2-induced invasion is dependent on fibronectin and correlates with the down-regulation of cell surface α4 integrin. In addition ERBB-2 co-immunoprecipitates with focal adhesion kinase (FAK) in these cells. We have also shown, by use of exogenously expressed PTEN and by treatment with the PI3′-kinase inhibitor LY294002, that ERBB-2-induced invasion is dependent on the PI3′-kinase pathway; however, PTEN does not dephosphorylate FAK in these cells. © 2000 Cancer Research Campaign

Keywords: ERBB-2; invasion; P13′ kinase; human breast cancer; human mammary epithelial cells

Amplification and overexpression of genes that regulate growth and differentiation of normal cells is a common mechanism for oncogene activation in human solid cancers. Among the oncogenes known to be amplified and overexpressed in human breast cancer (HBC) are ERBB-2 (Kraus et al, 1987; Slamon et al, 1987; Berger et al, 1988; Guerin et al, 1989) (also known as HER-2/neu), C-MYC (Bonilla et al, 1988; Borg et al, 1992; Watson et al, 1993), CCND1 (Gillet et al, 1994), FGFR 1, 2 and 4 (Luqmani et al, 1992; Jaakkola et al, 1993; Jacquemier et al, 1994), as well as other less well characterized genes (Bieche et al, 1996; Anzick et al, 1997; Sen et al, 1997). In addition to gene amplification, HBC cells often acquire other genetic alterations that may play a role in altered phenotypes expressed by cancer cells. And, while it is often possible to determine the signalling pathways activated as a result of gene amplification, it is much more difficult to discern the role of a specific genetic change in the expression of altered growth phenotypes exhibited by breast cancer cells. Previously, we demonstrated that HBC cells with an amplification of ERBB-2 express high levels of constitutively activated p185ERBB-2 and exhibit varying degrees of growth factor independence in vitro that is directly related to the level of overexpression of the gene (Ram et al, 1996). More recently, we have been able to recapitulate the growth factor independence phenotype in immortalized human mammary epithelial (HME) cells by overexpressing ERBB-2 to levels equivalent to that expressed by breast cancer cells, indicating that ERBB-2 itself can drive factor independent growth when expressed to high enough levels (Woods Ignatoski et al, 1999). In the present studies, we investigated the role of ERBB-2 overexpression in the acquisition of other altered phenotypes exhibited by breast cancer cells. We found that high level overexpression of ERBB-2 in the immortalized HME cell line H16N2 results, not only in the acquisition of anchorage-independent growth capacity, but also in the ability to invade naturally occurring basement membranes in a manner similar to breast cancer cells. The ability of ERBB-2 to induce invasion is linked to its ability to activate PI 3′-kinase, involves down-regulation of α,β integrins, and may involve signalling from focal adhesion kinase, which is associated with ERBB-2 in the overexpressing cells. Thus, when expressed to very high levels, ERBB-2 becomes associated with signalling complexes not detectable in normal cells, and is directly involved in the acquisition of a number of phenotypes characteristic of transformed cells.

MATERIALS AND METHODS

Cell culture

The base medium for H16N2-PTP, H16N2-ERBB2 and SUM-190PT cells was Ham’s F12 media supplemented with 0.1% bovine serum albumin, 0.5 μg ml⁻¹ fungizone, 5 μg ml⁻¹ gentamycin, 5 mM ethanolamine, 10 mM HEPES, 5 μg ml⁻¹ transferrin, 10 μM T₃, 50 μM selenium and 1 μg ml⁻¹ hydrocortisone. SUM-190PT cell medium was further supplemented with 5 μg ml⁻¹ insulin, and H16N2-PTP cell medium was further supplemented with 10 ng ml⁻¹ epidermal growth factor (EGF). The medium for SUM-52PE, SUM-I49PT, and SUM-225CWN was Ham’s F12 supplemented with 5% fetal bovine serum (FBS), 0.5 μg ml⁻¹ fungizone, 5 μg ml⁻¹ gentamycin and 5 μg ml⁻¹ insulin and further supplemented with 1 μg ml⁻¹ hydrocortisone. SKBr3 cells were grown in Dulbecco’s modified Eagles medium (DMEM) supplemented with 10% FBS, 0.5 μg ml⁻¹ fungizone and 5 μg ml⁻¹ gentamycin. Cells infected with retroviral expression vectors were selected in 100 μg ml⁻¹ geneticin (G418) for 2 weeks. All cell culture reagents were
obtained from Sigma Chemical Co. (St Louis, MO, USA). Detailed descriptions of all of the SUM cell lines can be found at http://p53.cancer.med.umich.edu/clines/clines.html on the worldwide web.

Retrovirus construction

pTPerbB-2 retrovirus construction was previously described (Harris et al, 1998; Woods Ignatowski et al, 1999). pTPPTEN (wt) and pTPPTEN (C124S) were constructed by excising the FLAG-tagged PTEN genes from FLAG-PTEN/CMV5 (Machama and Dixon, 1998) with EcoRI and BamHI. The ends of each PTEN gene were blunt and ligated into pTP 2000 which was digested with XhoI and blunt ended. Five micrograms of DNA were transfected into φNX-A cells by CaPO₄ precipitation and virus was collected 48 h after transfection. Cells were infected with the appropriate virus and selected in 100 μg ml⁻¹ gentamycin for 2 weeks. Because erbB-2 overexpressing cells already contained a construct that was G418-resistant, we tested the quality of infectious pTPPTEN virus by infecting non-G418-resistant MCF-10A cells and selecting in 100 μg ml⁻¹ gentamycin for 2 weeks. Each pTPPTEN virus infected approximately 75% of the MCF-10A cells.

Soft agar assays

Six-well dishes were coated with a 1:1 mix of the appropriate 2 × medium for the cell line being studied and 1% Bactoagar. Cells were plated at 1 × 10⁴, 1 × 10⁵ and 1 × 10⁶ cells per well in a mixture of appropriate medium and 0.3% Bactoagar. Cells were fed 3 times per week for 3 weeks, stained with 500 μg ml⁻¹ P=Rhoidonitrotetrazolium violet (Catalog no. I-8377, Sigma Chemical Co., St Louis, MO, USA) overnight and photographed. Each cell line was plated in duplicate in three separate dilutions. The experiment was repeated three times.

FAK immunoprecipitations and protein blots

Cells were lysed in a buffer containing 20 mM Tris–HCl, pH 8.0, 137 mM sodium chloride (NaCl), 1% NP-40, 10% glycerol, 1 mM Na₃VO₄, 1 mM phenylmethyl sulphoxide (PMSF), 1% aprotinin, 20 μg ml⁻¹ leupeptin. Protein concentrations were equalized to 100 μg for whole cell lysates or to 1 mg for immunoprecipitations using the Lawry method (Bradford, 1976). For whole cell lysates, Laemmli sample buffer (Sambrook et al, 1989) was added and the samples were boiled. For immunoprecipitations, 1 μg of α-FAK antibody (Catalog no. sc-558, Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA) was added to equivalent protein amounts of sample and incubated at 4°C for 1 h. Immune complexes were then bound to protein A/G beads for 1 h at 4°C. Immunoprecipitates were washed once in lysis buffer, twice in phosphate-buffered saline (PBS) containing 1% Triton X-100 and twice in PBS. Laemmli sample buffer was added and the samples were boiled. Equal amounts of protein were separated in 7.5% sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE). Proteins were blotted to polyvinylidene difluoride (PVDF) membrane and probed with α-PyrGlu antibody (Catalog no. 05-321, Upstate Biotechnology, Inc., Lake Placid, NY, USA) or with α-FAK antibody. These experiments were repeated three times.

PTEN immunoprecipitates and protein blots

Infected cells were lysed in a buffer containing 10 mM Tris–HCl, pH 8.0, 50 mM NaCl, 5 mM EDTA, 50 mM sodium fluoride (NaF), 1% NP-40, 1 mM NaVO₄, 1 mM PMSF. Anti-FLAG M2 beads (Catalog no. A-1205, Sigma Chemical Co.) were added to lysates and the mix was incubated for 2 h at 4°C. Immunoprecipitates were washed 3 x with lysis buffer and once with Tris-buffered saline (TBS)-0.03% Brij-35. Proteins were separated in 10% SDS-PAGE and blotted to PVDF membrane. Blots were probed with α-PTEN antibody (Catalog no. 06-894, Upstate Biotechnology, Inc.). The experiment was repeated twice.

FACs analysis

Cells were removed from tissue culture plastic with a room temperature incubation with 10 mM EDTA in Hank’s balanced salt solution, washed twice in PBS, then incubated with anti-integrin α₅ antibody (Catalog no. 12077-012, Gibco) or with anti-integrin α₅ antibody (Catalog no. CP12L, Oncogene Research Products, Cambridge, MA, USA) for 1 h at 22°C. Cells were then washed twice in PBS and incubated with a fluorescein isothiocyanate (FITC)-conjugated secondary antibody for 1 h at 37°C. After two washes in PBS, cells were fixed with 70% ethanol and subjected to fluorescence activated cell sorting.

Invasion

Cells were suspended in 0.23% trypsin–EDTA (Catalog no. 15050–057, Gibco Life Technologies, Grand Island, NY) and placed on SU-basement membranes with or without fetal calf serum (FCS) according to established methods (Livant et al, 1995) for 4 h at 37°C, the time required to observe maximal invasion percentages for normal and metastatic cells (Livant et al, 1995, and data not shown). The percentages of spread and adherent cells were evaluated in each assay to check viability prior to fixation in 2% formaldehyde and scored at 400 × magnification using phase contrast optics. Viability ranged from 90 to 98% in all assays. Mean invasion percentages resulted from three independent determinations involving the scoring of all cells in contact with the invasion substrates. Fn-FCS was prepared by repeated affinity chromatography of FCS on gelatin sepharose as previously described (Catalog no. 17095601, Pharmacia Biotech, Piscataway, NJ, USA) (Akiyama and Yamada, 1985). The pFn depletion was confirmed by immunoblotting with α-Fn antibody (Catalog no. 12112–017, Gibco) or with anti-integrin α₅ antibody (Catalog no. CP12L, Oncogene Research Products, Cambridge, MA, USA) (Hayman et al., 1982, data not shown). LY294002 was added to an identical plate as a control.

RESULTS

ErbB-2 overexpression results in anchorage-independent survival and growth of HME cells

In previous work, we developed a series of cell lines by retroviral infection of the immortalized HME cell line, H16N2, with a vector...
containing the full-length human ERBB-2 gene. We found that, when selected for proliferation in serum-free, growth factor-free medium, the transduced cells expressed very high levels of p185\textsuperscript{ERBB-2} that were constitutively tyrosine phosphorylated. Importantly, these cells expressed levels of activated p185\textsuperscript{ERBB-2} that are equivalent to levels expressed by HBC with an ERBB-2 gene amplification. Thus, as predicted from studies using ERBB-2 amplified HBC cells (Ram et al, 1996), high level ERBB-2 overexpression and activation results in multiple growth factor independence (Woods Ignatoski et al, 1999).

To begin to examine other altered phenotypes that result from ERBB-2 overexpression, H16N2 cells expressing different levels of ERBB-2 protein (infected 1 x with pTPRBB2 or infected 3 x with pTPRBB2-infected) were tested for their ability to grow in soft agar. Figure 1 demonstrates that, like breast cancer cells with an ERBB-2 gene amplification, H16N2 cells overexpressing ERBB-2 have the ability to survive under anchorage-independent conditions. Furthermore, H16N2 cells expressing the highest levels of p185\textsuperscript{ERBB-2} (3 x pTPRBB2-infected) grew to form large colonies in agar. Thus, progressive overexpression of ERBB-2 resulted in increasing ability of HME cells to grow in an anchorage-independent manner. These results are consistent with those reported previously by Pierce et al (1991).

### ERBB-2 overexpressing cells acquire invasive potential

To assess the invasive capacity of H16N2-ERBB2 cells, we used a novel assay that utilizes naturally occurring sea urchin (SU) embryo basement membranes. This experimental system allows one to assess the invasive capacity of human cancer cells and transformed cells in a completely defined system, because, as they are obtained from an invertebrate, SU-basement membranes are naturally serum-free. This technique has been described previously and accurately reflects the in vivo invasive and metastatic capacity of a variety of cancer and normal cell types (Livant et al, 1995).

We first assessed the invasive capacity of human breast cancer cell lines developed in our laboratory, and compared that to the invasive capacity of H16N2-ERBB2 cells, and vector transduced controls. Figure 2A shows that three human breast cancer cell lines, all of which were derived from aggressive breast cancers, but which have different molecular etiologies, expressed significant invasive capacity in this system. By contrast, HME cells failed to invade under these same conditions. H16N2-ERBB2 cells exhibited invasive capacity similar to that expressed by breast cancer cells, whereas control vector-transduced cells were not invasive. These results indicate that high level overexpression of p185\textsuperscript{ERBB-2} results in important transformed phenotypes beyond growth factor independence and anchorage independent growth.

It was demonstrated previously that invasion of prostate cancer cells through SU-basement membranes is dependent on the presence of plasma fibronectin, and that the PHSRN sequence of the fibronectin cell-binding domain is sufficient to stimulate invasion (DL Livant, manuscript submitted). The PHSRN sequence is a known ligand for the \( \alpha_5 \beta_1 \) integrin (Aota et al, 1994). It has been shown previously that \( \alpha_5 \beta_1 \) integrin binding to plasma fibronectin fragments containing the cell-binding domain but not the C-terminal domain, which contains the binding site for \( \alpha_5 \beta_1 \) integrin, results in the activation of metalloproteinase gene expression in fibroblasts which is essential for invasion (Huhtala et al, 1995).

We therefore examined the dependence on plasma fibronectin for the invasion of breast cancer cells and H16N2-ERBB2 cells, as well as their expression of \( \alpha_5 \beta_1 \) and \( \alpha_4 \beta_1 \) integrins. Table 1 shows that breast cancer cells express dramatically lower levels of cell surface \( \alpha_5 \) integrins than do normal HME cells, consistent with results obtained previously with prostate epithelial and invasive prostate cancer cells (Witkowski et al, 1993; Roklin and Cohen, 1995).

To examine integrin expression in these cells, FACs analysis was performed using the cell lines described above following incubation of intact, non-fixed cells with fluorescently tagged antibodies against the \( \alpha_5 \) and \( \alpha_4 \) integrins. Table 1 shows that breast cancer cells express dramatically lower levels of cell surface \( \alpha_5 \) integrins than do normal HME cells, consistent with results obtained previously with prostate epithelial and invasive prostate cancer cells (Witkowski et al, 1993; Roklin and Cohen, 1995).

Similarly, control vector transduced H16N2 cells, which are non-invasive, express \( \alpha_4 \) whereas, this integrin was down-regulated from the cell surface in invasive H16N2-ERBB2 cells.
Interestingly, \( \alpha_c \) expression was increased in the invasive H16N2-ERBB2 cells as compared to the vector transduced controls. These results are consistent with the hypothesis that invasion by transformed breast epithelial cells through naturally occurring basement membranes is mediated by interactions between specific integrins and particular domains of plasma fibronectin. Thus, like invasive breast and prostate cancer cells, H16N2-ERBB2 cells have reduced expression of cell surface \( \alpha_c \beta_1 \) integrins, which bind to a specific region of plasma fibronectin, and which may be functionally relevant to their capacity to invade basement membranes.

Experiments are underway to evaluate the role of cell surface \( \alpha_c \) integrin in the invasive behaviour of HME cells.

### Signalling pathways mediating the invasive phenotype in ERBB-2 overexpressing cells

We and others have shown previously that an important consequence of ERBB-2 overexpression is a dramatic increase in the level of activation of phosphatidylinositol 3'-kinase (PI 3' kinase) and its downstream signalling pathways (Fedi et al, 1994; Soltoff et al, 1994; Ram and Ethier, 1996). Therefore, a series of experiments were performed to assess the role of this pathway in the invasive potential of H16N2-ERBB2 cells.

The tumour suppressor gene, PTEN, is a phosphatase that regulates the activity of the PI 3'-kinase pathway by virtue of its ability to directly dephosphorylate PIP3 (Maehama and Dixon, 1998). Therefore, we prepared retroviral expression vectors containing full-length human PTEN, or a mutant PTEN that has the ability to bind substrate but which is defective in both its lipid and protein phosphatase activities. Figure 3A illustrates the constructs used, and Figure 3B shows the expression of the transduced PTEN proteins in H16N2-ERBB2 cells following infection with the

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**Table 1** Cell surface integrin expression

| Cell line          | \( \alpha_c \) expression | \( \alpha_5 \) expression |
|--------------------|---------------------------|--------------------------|
| HME                | 54                        | 62                       |
| SUM-149PT          | 0                         | 10                       |
| SUM-190PT          | 14                        | 27                       |
| H16N2-PTP          | 55                        | 47                       |
| H16N2-ERBB-2       | 11                        | 86                       |
| H16N2-ERBB2/PTENWT | 12                        | 21                       |

*As determined by FACs analysis. *Per cent increase in amount of staining for \( \alpha_c \) or \( \alpha_5 \) integrins over negative control.

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**Figure 2** HBC cells and H16N2-ERBB-2 cells undergo fibronectin-dependent invasion. (A) Invasion assays for HME, HBC, and H16N2-ERBB-2 cells. Percent invasion relative to SUM-149PT invasion given. (B) Invasion assays for HBC and H16N2-ERBB-2 cells were performed in serum-free medium or serum-free medium supplemented with FCS or pFN, as indicated. Per cent invasion relative to 5% FCS invasion was graphed.
overexpressing cells. (CS, 5. Per cent invasion relative to H16N2-ERBB-2 invasion was graphed
H16N2-ERBB-2/PTP, 3; H16N2-ERBB-2/PTEN, 4; H16N2-ERBB-2/PTEN CS, 5.

exogenous PTEN. Lanes: H16N2-PTP, 1; H16N2-ERBB-2, 2; H16N2-ERBB-
H16N2-ERBB-2/PTEN and H16N2-ERBB-2/PTEN CS were expressing
on anti-FLAG immunoprecipitates using anti-FLAG antibody showing that
ligated into the biscistronic vector pTP2000. FL = FLAG Tag. (C)

2/PTP, 3; H16N2-ERBB-2/PTEN, 4; H16N2-ERBB-2/PTEN CS, 5. Relative % invasion
H16N2-ERBB-2 cells with a vector containing the cs-PTEN

To confirm that signalling from PI 3'-kinase is involved in the invasion of HBC and
ERBB-2 overexpressing cells. HBC cells and H16N2-ERBB-2 cells were
incubated with LY294002 then tested for invasion. Per cent invasion relative
to SUM-149PT invasion given

(1998) suggests that, under conditions that are relevant to the in
vivo situation, pp125FAK is a poor substrate for PTEN. Therefore,
we performed experiments to assess the pp125FAK protein activation
status in invasive H16N2-ERBB2 cells versus non-invasive control
cells, and to assess the role of PTEN in modulating pp125FAK
tyrosine phosphorylation in these cells.

To determine if the level of tyrosine phosphorylated pp125FAK
was elevated in invasive H16N2-ERBB2 cells compared to
controls, FAK immunoprecipitates were used to prepare Western
blots that were probed with phosphotyrosine antibodies. The data
in Figure 5A shows that overexpression of p185ERBB-2 in H16N2-
ERBB2 cells did not result in significant changes in the levels of
tyrosine phosphorylated pp125FAK. However, in these blots, we did
detect a tyrosine phosphorylated p185 that co-immunoprecipitated
with FAK in the ERBB-2 overexpressing cells, but not in control
cells. Reprobing of an identical blot with ERBB-2 specific anti-
bodies confirmed the identity of the tyrosine phosphorylated
protein as p185ERBB-2. The data in Figure 5A also shows that the
association of pp125FAK with p185ERBB-2, detected in the H16N2-
ERBB-2 cells, is also detectable in the SkBr-3 breast cancer cell
line which expresses high levels of p185ERBB-2 as a result of a gene
amplification. Thus, whereas levels of tyrosine phosphorylated
FAK are not altered in invasive ERBB-2 overexpressing cells,
p185ERBB-2 can be detected in immunocomplexes with pp125FAK
when ERBB-2 is expressed to high levels.

Experiments were next performed to examine the influence of
PTEN overexpression on levels of tyrosine phosphorylated
pp125FAK and its association with p185ERBB-2. The results shown in
Figure 5B indicate that PTEN overexpressing cells that have lost
their invasive capacity express the same levels of tyrosine phosphorylated
pp125FAK as non-PTEN overexpressing cells with full
invasive capacity. Furthermore, PTEN expression did not influence
the association of p185ERBB-2 with pp125FAK, and did not result
in the re-expression of α5 integrin on the cell surface (Fig. 5 and

Figure 4 The PI 3'-kinase inhibitor, LY294002, blocks invasion of HBC and
ERBB-2 overexpressing cells. HBC cells and H16N2-ERBB-2 cells were
incubated with LY294002 then tested for invasion. Per cent invasion relative
to SUM-149PT invasion given
acquisition of altered phenotypes typically expressed by aggressive cancer cells. ERBB-2 overexpressing cells are not only growth factor-independent and anchorage independent for growth, these cells can invade naturally occurring basement membranes in a pFn-dependent manner. When expressed to very high levels, p185<sub>ERBB-2</sub> can be detected in immunocomplexes with pp125<sub>FAK</sub>, and blocking PI 3'-kinase signalling in these cells completely abrogates the invasive potential of H16N2-ERBB2 cells.

**DISCUSSION**

ERBB-2 is an important oncogene in breast and other cancers, and overexpression and constitutive activation of its tyrosine kinase is essential to its oncogenic properties (DiFiore et al, 1987, 1990a, 1990b; Hudziak et al, 1987; Lonardo et al, 1990; Pierce et al, 1991). Despite the clear role of ERBB-2 in the aetiology of some human cancers, the specific cellular phenotypes expressed by cancer cells that are the direct result of ERBB-2 overexpression have not been completely defined.

In previous work, we demonstrated that breast cancer cells that overexpress p185<sub>ERBB-2</sub> are independent of growth factors required by normal HME cells for continuous proliferation in serum-free medium (Ram et al, 1996). More recently, we showed that overexpression of p185<sub>ERBB-2</sub> in HME cells to levels expressed by breast cancer cells with a ERBB-2 gene amplification is sufficient to induce the growth factor independence phenotype (Woods Ignatowski et al, 1999). The activation of signalling pathways, such as PI 3'-kinase and MAP kinase, in ERBB-2 overexpressing cells has been reported by a number of investigators and is consistent with the ability of the ERBB-2 kinase to stimulate growth of cancer cells (Benlevy et al, 1994; Marte et al, 1995). However, HBC cells exhibit other altered cellular phenotypes beyond growth factor independence. Therefore, we undertook experiments to determine if ERBB-2 overexpression in HME cells could induce other altered cellular phenotypes expressed by breast cancer cells.

H16N2 human mammary epithelial cells are a keratin-19-positive cell line that was immortalized by transduction of the entire HPV-16 genome (Band and Sager, 1989; Band et al, 1990). These cells are immortal in culture, are partially independent of insulin for growth in serum-free medium, but are still completely dependent on EGF for growth. In addition, H16N2 cells do not grow in soft agar, are not invasive in vitro, and do not form tumours in immunodeficient animals. H16N2-ERBB2 cells, which express levels of activated ERBB-2 equivalent to HBC cells with a ERBB-2 gene amplification, are independent of both insulin and EGF for growth in serum-free medium, have the ability to grow under anchorage-independent conditions, and can invade naturally occurring basement membranes. Thus, this is the first report to demonstrate that ERBB-2 overexpression can have a direct effect on the invasive potential of HME cells, and that progressive ERBB-2 overexpression results in the acquisition of phenotypes associated with increasing malignancy.

Based on the findings of our study, there are at least two pathways that are affected by overexpression of ERBB-2 which are important to the invasive potential of the cells. The first is the ERBB-2-mediated activation of PI 3' kinase signalling. The second is the plasma fibronectin dependence for the invasive phenotype, and the related down-regulation of cell surface α<sub>5</sub> integrin in the invasive cells.

Several laboratories, including our own, have demonstrated that overexpression of ERBB-2 results in high level activation of PI 3' internal Figure 5: ERBB-2 co-immunoprecipitates with FAK in H16N2-ERBB-2 cells. (A) Anti-FAK immunoprecipitates from control and ERBB-2 overexpressing cells probed with anti-PTYR<sub>152</sub> FAK and ERBB-2 are indicated; an identical blot was probed with anti-ERBB-2 to ensure that the tyrosine-phosphorylated p185 was ERBB-2. Anti-FAK immunoprecipitates from H16N2-ERBB-2 and SkBr3 cells were also probed with anti-PTYR<sub>152</sub> to show co-immunoprecipitating tyrosine-phosphorylated p185. (B) FAK is not dephosphorylated in PTEN-expressing cells. Anti-FAK immunoprecipitations from H16N2-ERBB-2 (Lane 1), H16N2-ERBB-2/PTP (Lane 2), H16N2-ERBB-2/PTEN WT (Lane 3), or H16N2-ERBB-2/PTEN CS (Lane 4) cells were probed with anti-PTYR<sub>152</sub> or an identical blot was probed with anti-FAK to show FAK levels.
kinase (Fedi et al, 1994; Soltoff et al, 1994; Ram and Ethier, 1996). There are at least three possible mechanisms by which ERBB-2 modulates the activity of this pathway (reviewed in Earp et al, 1986; Alroy and Yarden, 1997). First, the cytoplasmic domain of p185<sub>ERBB-2</sub> has a single YXXM binding site for the regulatory subunit of PI 3' kinase. Thus, either direct or indirect phosphorylation of this tyrosine residue can result in activation of the enzyme. Second, ERBB-2 activation often results in the concomitant activation of ERBB-3, which has six YXXM sites in its cytoplasmic domain. Thus, ERBB-3 is a potent activator of PI 3' kinase, and the H16N2-ERBB2 cells used in these studies express high levels of constitutively tyrosine phosphorylated ERBB-3. The third potential mechanism involves the association of the p85 subunit of PI 3' kinase with tyrosine phosphorylated pp125<sub>FAK</sub> (Bachelot et al, 1996). This mechanism is potentially significant in our experiments since, in invasive ERBB-2 overexpressing cells, p185<sub>ERBB-2</sub> was found to be associated with activated pp125<sub>FAK</sub> in immunocomplexes, and the p85 subunit of PI 3' kinase was detectable in the same immunocomplexes (not shown). Thus, it is possible that the association of p185<sub>ERBB-2</sub> with pp125<sub>FAK</sub> could potentiate PI 3' kinase activation beyond what occurs as a result of ERBB-2 homodimer and ERBB-2/ERBB-3 heterodimer interactions. The data obtained in our studies indicate that PI 3' kinase activation is a direct mediator of the invasive phenotype in that both LY294002 and exogenous expression of PTEN reversed the invasive capacity of H16N2-ERBB2 cells. The effect of PTEN is particularly interesting in that PTEN transduced cells proliferate as well in monolayer culture as control cells (data not shown). In contrast, the cells are growth inhibited by the concentration of LY204002 used to block invasion (data not shown).

The SU-basement membrane invasion assay used in these studies offers the ability to study cell invasion of naturally occurring basement membranes under serum-optional conditions. The defined nature of the assay allowed for the previous observation of the importance of plasma fibronectin for invasion of basement membranes by prostate cancer cells (Livant et al, 1995), and for the observation of its importance for basement membrane invasion by breast cancer cells, as described here. Like metastatic breast and prostate cancer cells, H16N2-ERBB2 cells also only invaded SU-basement membranes in the presence of plasma fibronectin. The importance of plasma fibronectin for invasion is particularly interesting because all cells examined thus far that are invasive in this assay have been found to express low levels of cell surface α<sub>4</sub> integrin, while expressing high levels of α<sub>5</sub>. Indeed, cell surface α<sub>4</sub> integrin expression, which is normal in H16N2 cells, was found to be down-regulated in ERBB-2 overexpressing cells with invasive potential. The down-regulation of α<sub>5</sub> detected by FACs analysis was not observed in Western blots (not shown) suggesting that integrin trafficking is altered in the invasive cells. These results are consistent with those recently reported by Weaver et al (1997) who not only demonstrated that the repertoire of integrin expression is functionally important to the transformed phenotype of the cells, but that changes in cell-surface localization of integrins can occur without changes in overall levels of transcription or translation. Our data are also consistent with those of Seftor et al (1998) who demonstrated that maspin, which can modulate the invasive capacity of MDA-435 cells, can directly modulate cell surface expression of a number of integrins.

The loss of cell surface α<sub>4</sub>β<sub>1</sub> integrin in the presence of α<sub>5</sub>β<sub>1</sub> is important because each of these integrin dimers bind to distinct domains of plasma fibronectin. In normal cells, the cell binding domain of pFn interacts with α<sub>5</sub>β<sub>1</sub>, whereas the C-terminal domain interacts with α<sub>4</sub>β<sub>1</sub> integrin. This dual interaction is important for the maintenance of normal cells in a non-motile, non-invasive state (Postlethwaite et al, 1981; Seppa et al, 1981; Clark et al, 1988). During wound healing, plasma fibronectin fragments are generated by proteolysis resulting in the formation of fragments containing the cell-binding domain. These fragments contain the PHSRN sequence, but do not contain the C-terminal domain that binds α<sub>5</sub>β<sub>1</sub>. The stimulation of fibroblasts by proteolytic fragments of plasma fibronectin is essential for the invasion and migration of cells during this process (Grinnell et al, 1992).

Indeed, Huhtala et al (1995), demonstrated that in fibroblasts, interaction of α<sub>4</sub>β<sub>1</sub> with pFn in the absence of α<sub>5</sub>β<sub>1</sub>, results in the up-regulation of metalloproteinases that are important for invasion across basement membranes. Similarly, Livant et al (submitted for publication) demonstrated that direct stimulation of α<sub>5</sub>β<sub>1</sub> integrin using the PHSRN peptide derived from the pFn cell-binding domain, was sufficient to induce migration and invasion by normal epithelial cells. Thus, down-regulation of cell surface α<sub>4</sub> integrin (and the consequent loss of α<sub>5</sub>β<sub>1</sub> plasma fibronectin receptors) in ERBB-2 overexpressing cells, as well as in breast and prostate cancer cells, is likely to be of functional significance to their ability to invade basement membranes and to metastasize.

Since integrins are involved in motility and invasion of cancer cells, and since integrin signalling is known to be mediated by activation of pp125<sub>FAK</sub> (Guan, 1997; Ilic et al, 1997), the observation that p185<sub>ERBB-2</sub> and pp125<sub>FAK</sub> interact and can be co-immunoprecipitated in invasive, ERBB-2 overexpressing cells, is particularly intriguing. This result suggests the possibility that the interaction of p185<sub>ERBB-2</sub> and pp125<sub>FAK</sub>, and the down-regulation of α<sub>5</sub>β<sub>1</sub> integrin from the cell surface, are functionally related. In addition, since the p85 subunit of PI 3' kinase is also detectable in pp125<sub>FAK</sub> immunoprecipitates, and blocking PI 3' kinase signalling abrogates invasiveness of the cells, the interaction of p185<sub>ERBB-2</sub> and pp125<sub>FAK</sub> could be important for the overall activation of PI 3' kinase signalling. Distinguishing the precise roles of p185<sub>ERBB-2</sub> and pp125<sub>FAK</sub>, PI 3' kinase activation, and cell surface α<sub>4</sub>β<sub>1</sub> integrin expression in mediating the invasive phenotype will require further experiments.

In summary, the data reported here, and previously, indicate that progressive overexpression of ERBB-2 in HME cells results in the step-wise acquisition of malignant properties. Moderate ERBB-2 overexpression is sufficient to liberate cells from their requirements of exogenous insulin-like growth factors for proliferation, but is not sufficient to drive EGF independence, growth in soft agar, or invasive capacity. However, expression of ERBB-2 to levels seen in breast cancer cells with a gene amplification not only results in complete growth factor autonomy, but also yields cells with robust anchorage-independent growth potential and with the ability to invade naturally occurring basement membranes. HME cells expressing high levels of p185<sub>ERBB-2</sub> exhibit many alterations in normal signalling patterns including; constitutive tyrosine phosphorylation of ERBB-2, ERBB-3 and EGFRI, constitutive activation of PI 3' kinase, association of ERBB-2 with pp125<sub>FAK</sub>, and down-regulation of cell surface α<sub>5</sub>β<sub>1</sub> integrins. Future studies will be aimed at dissecting the relative contributions of each of these signalling pathways in mediating the different altered phenotypes exhibited by ERBB-2 overexpressing mammary epithelial cells and breast cancer cells.
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