

The Great Escape: When Cancer Cells Hijack the Genes for Chemotaxis and Motility

John Condeelis,^{1,2} Robert H. Singer,¹
and Jeffrey E. Segall¹

¹Anatomy and Structural Biology and ²Analytical Imaging Facility, Albert Einstein College of Medicine, Bronx, New York 10461-1975; email: condeeli@aecom.yu.edu; rhsinger@aecom.yu.edu; segall@aecom.yu.edu

Annu. Rev. Cell Dev. Biol.
2005. 21:695–718

First published online as a
Review in Advance on
July 5, 2005

The *Annual Review of
Cell and Developmental
Biology* is online at
<http://cellbio.annualreviews.org>

doi: 10.1146/
annurev.cellbio.21.122303.120306

Copyright © 2005 by
Annual Reviews. All rights
reserved

1081-0706/05/1110-
0695\$20.00

Key Words

actin, cofilin, N-WASP, capping protein, Arp2/3 complex

Abstract

The combined use of the new technologies of multiphoton-based intravital imaging, the chemotaxis-mediated collection of invasive cells, and high sensitivity expression profiling has allowed the correlation of the behavior of invasive tumor cells in vivo with their gene expression patterns. New insights have resulted including a gene expression signature for invasive cells and the tumor microenvironment invasion model. This model proposes that tumor invasion and metastasis can be studied as a problem resembling normal morphogenesis. We discuss how these new insights may lead to a better understanding of the molecular basis of the invasive behavior of tumor cells in vivo, which may result in new strategies for the diagnosis and treatment of metastasis.

Contents

INTRODUCTION.....	696
WHAT MOTILITY BEHAVIORS CONTRIBUTE TO INVASION AND INTRAVASATION?	696
Intravital Imaging of Tumor Cell Behavior in Tumors In Vivo....	696
Motility Behaviors Contributing to Invasion and Intravasation	697
IS THERE AN EXPRESSION SIGNATURE THAT CORRELATES WITH THESE BEHAVIORS?	698
The Concept of an Invasion Signature	698
HOW DO THE GENES OF THE INVASION SIGNATURE CONTRIBUTE TO INVASION?	700
Coordinate Regulation of Motility Pathways in Invasion	700
Tests of Function of Genes of the Invasion Signature in Chemotaxis, Invasion, and Metastasis.....	709
NEW INSIGHTS INTO TUMOR INVASION AND METASTASIS.	711
The Tumor Microenvironment Invasion Model.....	712

INTRODUCTION

The ability of tumor cells to spread from primary tumors (and metastatic tumors) is the major cause of death in cancer patients. Spreading of tumor cells relies upon cell motility, which results in the invasion of neighboring connective tissue and entry into lymphatics and blood vessels (intravasation) (Clark et al. 2000, Condeelis & Segall 2003, Woodhouse et al. 1997). We focus on tumor invasion and metastasis as a problem in cell motility. In this context, the motility behavior of tumor cells inside the tumor must be analyzed as carefully as the gene expression patterns displayed by invasive cancer cells. Dur-

ing motility, the microenvironment becomes a determinant in the success or failure of a cancer cell in its attempt to traverse the tumor and enter blood and lymphatic vessels (Liotta & Kohn 2001). Subtle changes in the cancer cell's interactions with extracellular matrix and gradients of growth factors and cytokines define whether a cell becomes invasive or remains stationary in the tumor mass. Only by understanding the basic biology of how the motility of cells inside the tumor is influenced by, and influences, the gene expression patterns of cancer cells and their microenvironment will it be possible to define strategies to impede the spread of cancer cells from tumors.

In this review three questions are considered: What motility behaviors contribute to invasion and intravasation? Is there an expression signature that correlates with these behaviors, that is, an invasion signature? How do the genes of the invasion signature contribute to invasion? Answering these questions to date has suggested a novel model for tumor invasion and metastasis, which is discussed at the end of the review.

WHAT MOTILITY BEHAVIORS CONTRIBUTE TO INVASION AND INTRAVASATION?

Intravital Imaging of Tumor Cell Behavior in Tumors In Vivo

An attempt to understand the behavior of tumor cells at single-cell resolution in vivo predates the introduction of green fluorescent protein (GFP) and its derivatives. Tumor cells were transiently labeled with vital dyes and observed with conventional transmitted and fluorescence microscopy (Chambers et al. 1995, Scherbarth & Orr 1997, Suzuki et al. 1996, Vajkoczy et al. 1999, Wood 1958, Yuan et al. 1995). This required the use of short-lived preparations in thin regions of tissue where light could pass efficiently and sometimes the use of viewing windows (Chambers et al. 1995, Wood 1958). These approaches

usually limited the analysis of tumors to artificial locations and introduced the potential for artifact resulting from the viewing method. A major step was the introduction of stable GFP expression, which allowed genetic labeling of cells in tumors with tissue- and cell-type specificity without rejection of the GFP-tagged cells (Chishima et al. 1997, Farina et al. 1998). Thus more clinically relevant tumor models were developed that could be imaged in the location in the animal where the tumor naturally forms and progresses to different stages (Ahmed et al. 2002, Brown et al. 2001, Yang et al. 2000).

The introduction of the laser-scanning confocal microscope was an essential advance that made optical sectioning and single-cell resolution possible, essential capabilities for relating cell behavior to mechanisms of invasion (Chantrain et al. 2004, Farina et al. 1998). However, conventional one-photon laser-scanning confocal microscopy is limited by the relatively poor optical depth of penetration of short wavelength excitation light, photobleaching, and phototoxic damage to the whole tissue and not just at the focal point. The recent introduction of multiphoton microscopy, which uses 800–900 nm light from a pulsed laser, has largely solved the problems of photobleaching and toxicity and extended the depth of penetration by 20-fold (Condeelis & Segall 2003, Helmchen & Denk 2002, Jain et al. 2002, Wang et al. 2002, Williams et al. 2001, Zipfel et al. 2003b). In addition, the multiphoton excitation of tissue causes second harmonic scattering of photons from α -helix-containing proteins, thus allowing the imaging of extracellular matrix proteins such as collagen without the need for fluorescent labeling of the tissue matrix (Campagnola et al. 2001, Condeelis & Segall 2003, Zipfel et al. 2003a). This benefit of multiphoton excitation can be used to analyze cell-extracellular matrix interactions and matrix remodeling directly in live tissue (Condeelis & Segall 2003, Masters et al. 1997). The application of gradient index (GRIN) lens technology to multiphoton imaging holds the promise of extend-

ing intravital imaging to any depth within live mice, making systemic analysis of tumor invasion and metastasis possible (Levene et al. 2004). The tumor cell behaviors in live tumors discussed in the next section are derived from studies using laser-scanning confocal and multiphoton imaging in rats and mice.

Motility Behaviors Contributing to Invasion and Intravasation

An understanding of the motility of cancer cells and its contribution to metastasis has begun to emerge from intravital imaging of cells, at single-cell resolution, inside tumors within living animals. An important outcome of studying tumor cells within their normal tumor environment is that the behavior observed is an indication of what tumor cells actually do *in vivo*, not what they can do as inferred from *in vitro* and ectopic models *in vivo*. That is, artificial models such as ectopic growth of tumor cells in tissues in which tumors do not normally form, e.g., growing breast tumor cells in dermis instead of mammary gland, can lead to the observation of cell behaviors that do not occur in real breast tumors. Tumor models where tumor cells are grown in tissue that is the natural site for the tumor, e.g., breast in breast, are called orthotopic models. The behaviors discussed next are those seen in orthotopic models.

A number of behaviors have been observed in orthotopic models *in vivo* that relate to metastatic potential. These form a pattern common to a variety of tumor types and provide insight into mechanisms of invasion and intravasation (Condeelis & Segall 2003, Farina et al. 1998, Friedl & Wolf 2003, Sahai et al. 2005, Wang et al. 2002, Wyckoff et al. 2000a).

Tumor cells in primary mammary tumors move as solitary cells at up to 10 times their velocity *in vitro*. Tumor cell motility is characterized as solitary amoeboid movement, but it can also occur as cell streams and linear files suggesting the use of common

paths on extracellular matrix (ECM) fibers (Farina et al. 1998, Friedl & Wolf 2003, Sahai et al. 2005, Wang et al. 2002, Wyckoff et al. 2000a). In fact, the highest velocities are observed for carcinoma cells in metastatic tumors that are moving along linear paths in association with ECM fibers, in particular, collagen fibers. These high-velocity linear excursions are unrestricted by networks of ECM in mammary tumors, except around blood vessels (Condeelis & Segall 2003).

Intravasation and invadopodia. Tumor cell motility is restricted at the basement membrane of blood vessels, where the cells must squeeze through small pores in the basement membrane/endothelium to gain access to the blood space. The degree to which the basement membrane of blood vessels represents a barrier has been documented by direct observations of cell behavior during intravasation. Carcinoma cells in nonmetastatic tumors are fragmented during intravasation as they squeeze across the basement membrane/endothelium indicating that the cell must be highly distended and under tension as it crosses. Remarkably, carcinoma cells in metastatic tumors cross this restriction as intact cells, possibly in large measure owing to the high levels of expression of cytokeratins in metastatic cells (Wang et al. 2002) and their ability to extend invadopodia (Condeelis & Segall 2003, Wang et al. 2002, Wyckoff et al. 2000a, Yamaguchi et al. 2005).

Chemotaxis to blood vessels. Carcinoma cells in metastatic tumors are attracted to blood vessels, where they form a layer of cells that are morphologically polarized toward the vessel. Chemotaxis to epidermal growth factor by carcinoma cells has been demonstrated in vivo (Wyckoff et al. 2004) and resembles that observed for tumor cells in vitro (Wyckoff et al. 2000b). Chemotaxis ability of tumor cells is highly correlated with their potential for invasion, intravasation, and metastasis and appears responsible for the attraction of carcinoma cells to blood vessels (Wyckoff

et al. 2000a). Cell polarity toward blood vessels is correlated with increased intravasation and metastasis, indicating a local blood vessel associated source of chemoattractants (Pollard 2004, Wyckoff et al. 2004).

Blood vessel-associated macrophages are a source of EGF and other chemoattractants.

An in vivo invasion assay (Wyckoff et al. 2000b) has been used to study the mechanism of chemotaxis in primary mammary tumors of rats and mice. These studies demonstrate that macrophages form a paracrine loop with invasive tumor cells (Wyckoff et al. 2004). Expression analysis of tumor cells and macrophages caught invading together indicates how these cells are attracted to each other and can invade jointly: Tumor cells express CSF-1, which stimulates macrophage chemotaxis, whereas macrophages express epidermal growth factor (EGF), which stimulates tumor cell chemotaxis (Wyckoff et al. 2004). Because metastatic mammary tumors contain large numbers of rapidly moving macrophages with many clustered near blood vessels (Condeelis & Segall 2003, Wyckoff et al. 2000a), they are a local source of chemotactic cytokines and chemotactic growth factors, such as EGF, within the tissue and near blood vessels (Lin et al. 2001, Pollard 2004, Wyckoff et al. 2004). The in vitro assay of invasive cell motility inside collagen matrices demonstrates that macrophages and tumor cells and the activity of their CSF-1 and EGF receptors, respectively, are necessary and sufficient for enhancement of invasion (Goswami et al. 2005).

IS THERE AN EXPRESSION SIGNATURE THAT CORRELATES WITH THESE BEHAVIORS?

The Concept of an Invasion Signature

Gene expression profiling has been used extensively in an attempt to sort tumors into

subtypes that might be diagnosed and treated more effectively (Ramaswamy et al. 2003, van 't Veer et al. 2002). In addition, expression profiling has been used in an attempt to identify invasion- and metastasis-specific genes that might predict the metastatic potential of tumors and to gain insight into the mechanisms of invasion and metastasis. In general, studies involving (a) entire primary tumors, (b) laser capture microdissection of fixed primary tumors, and (c) cells isolated from metastases of bone marrow, lymphatics and distant solid organs have identified candidate genes that might be important for tumor cell invasion (reviewed in Wang et al. 2005). However, such approaches have had limitations:

- Expression analysis of whole primary tumors provides bulk tumor expression patterns, in which case the specific patterns of expression typical of invasive cells might be diluted.
- Laser capture microdissection studies must rely on morphology and histological location, an uncertain exercise, to select cells that might have been invading, thereby making the relevance to invasion of expression profiles from such cells questionable.
- Isolating cells from metastases are likely to produce expression profiles that are relevant to successful growth at the new site but not necessarily profiles indicative of invasion potential from the primary tumor.

In an alternative approach, an *in vivo* invasion assay, based on the chemotaxis of tumor cells to blood vessels seen *in vivo* (Wyckoff et al. 2000b), was used to collect invasive cells from live primary tumors in mice and rats. Because the *in vivo* invasion assay employs microneedles containing chemoattractants such as EGF and extracellular matrix that mimic conditions around blood vessels that are involved in chemotaxis (Wyckoff

et al. 2000b, 2004), the invasive cells collected are likely also to be the cells involved in intravasation. Invasive tumor cells collected by this method can then be interrogated directly relative to the tumor cells that remain behind in the primary tumor and, after subtraction of gene expression changes occurring in response to EGF and other collection conditions, this reveals the expression pattern unique to invasive tumor cells, an “invasion signature.” The invasion signature shown in **Tables 1, 2, and 3** is derived from invasive cells collected in rat mammary tumors generated from carcinoma cell lines (Goswami et al. 2004; Wang et al. 2003, 2004). A similar invasion signature has been derived from mouse mammary tumors resulting from expression of the PyMT oncogene *in situ* (W. Wang, personal communication). This indicates that the invasion signature is common to mammary tumors in rats and mice regardless of the origins of the mammary tumor. The invasion signature shown in **Tables 1, 2, and 3** indicates that invasive cells are a population that is neither proliferating nor apoptotic but is highly motile (Goswami et al. 2004, Wang et al. 2004). The reduction in apoptosis is consistent with tumor cells having a survival advantage owing to suppression of apoptosis genes and up-regulation of pro-survival genes (**Table 2**). Furthermore, the pattern of expression of genes involved in proliferation suggests that invasive tumor cells are not proliferating (**Table 1**). This predicts that cancer treatments targeting cell growth may not be very effective at killing invasive tumor cells. This was tested by exposing invasive cells collected using the *in vivo* invasion assay to conventional chemotherapy that is directed at dividing cells. As predicted, the invasive cells survived better compared with non-invasive cells from the same tumor (Goswami et al. 2004).

Several of the genes of the invasion signature have been identified in clinical and conventional gene expression profiling studies. Clinical studies of bladder, breast,

Table 1 Genes of the cell proliferation part of the invasion signature^a

Gene symbol	Gene description	Fold change ^b
Suppression of cell proliferation		
Psmc5	Protease (prosome, macropain) 26S subunit, ATPase 5	5.5
Rad9	Cell cycle checkpoint control protein (Rad9) mRNA	4.0
Hmg1	High mobility group protein 1	3.5
CKS2	Cyclin-dependent kinases regulatory subunit 2	3.4
Cks1	Cyclin-dependent kinase regulatory subunit 1	3.2
Fmo5	Flavin containing monooxygenase 5	3.0
GAS6	GAS 6 mRNA associated with growth arrest	2.8
Phb	Prohibitin	2.8
Mad2	Mitotic checkpoint component Mad2 mRNA	2.4
Madh3	MAD homolog 3	2.4
Hmg14	High mobility group protein 14	2.3
Enhancement of cell proliferation		
CGMC	Carcinoembryonic antigen CGM6 precursor	0.5
CPR2	Cell cycle progression 2 protein (CPR2)	0.4
Ask	Activator of S phase kinase	0.2

^aTo determine the significance of changes in gene expression in each of the functional categories of the genes represented in microarrays, the Student's t test, Chi-square, or SAM analysis were performed. The fold changes in gene expression of the invasion signature shown in **Tables 1–3** were found to be statistically significant in the invasive cells by Chi-square or SAM analysis. In addition, in all cases, $P < 0.05$. Random sets of equal numbers of genes did not generate the same pattern of up- and down-regulation, indicating that the pattern was not observed by chance ($P < 0.05$). Similarly, clustering the results from all genes of the general population in the same space of all genes on the microarray did not yield an outcome similar to the invasion signature. All results are from Goswami et al. 2004 and Wang et al. 2004.

^bThe fold change indicates the level of expression in the invasive tumor cells compared with the general population of tumor cells of the primary tumor.

and colorectal cancers have implicated Rho A, Rock (Kamai et al. 2003), Mena (Di Modugno et al. 2004), and the Arp2 and 3 subunits of the Arp2/3 complex (Otsubo et al. 2004), respectively, as up-regulated in these cancers. Studies have suggested that the elevated expression of Arp2/3 complex by both neoplastic and stromal cells contributes to the increased motility of both cell types and thus provides suitable conditions for invasion (Otsubo et al. 2004). In addition, LIM-kinase 1 is up-regulated in metastatic breast and prostate tumors (Davila et al. 2003, Yoshioka et al. 2003). Hence, the pathways involved in actin polymerization at the leading edge are implicated in invasion by different approaches (**Figure 1**).

HOW DO THE GENES OF THE INVASION SIGNATURE CONTRIBUTE TO INVASION?

Coordinate Regulation of Motility Pathways in Invasion

An important insight into the special motility properties of invasive cells, the high speeds of locomotion, chemotaxis, and invadopod formation, comes from the motility pathways portion of the invasion signature (**Table 3**). That is, the finding that the genes coding for the key effectors of the minimum motility machine (Loisel et al. 1999), i.e., the cofilin, capping protein, and Arp2/3 pathways, that regulate β -actin polymerization at the leading edge, are dramatically up-regulated

Table 2 Genes of the apoptosis and survival part of the invasion signature^a

Gene	Description	Fold change ^b
Anti-apoptotic genes		
Ier3	Immediate early response 3	4.9
Ubl1a2	Ubiquitin-like 1 (sentrin) activating enzyme subunit 2	4.7
Txn	Thioredoxin	3.7
Hsp105	Heat shock protein, 105 kDa	3.5
Odc	Ornithine decarboxylase, structural	3.0
Dad1	Defender against cell death 1	2.7
Trp53	Transformation related protein 53	2.5
Hsp60	Heat shock protein, 60 kDa	2.4
Api4	Apoptosis inhibitor 4	2.3
Cldn3	Claudin 3	2.3
Api5	Apoptosis inhibitor 5	2.3
Hsp86-1	Heat shock protein, 86 kDa 1	2.1
Api1	Apoptosis inhibitor 1	2.0
Adam17	A disintegrin and metalloproteinase domain 17	2.0
Pro-apoptotic genes		
Pdcd4	Programmed cell death 4	0.1
Fem1b	Feminization 1 b homolog (<i>C. elegans</i>)	0.4
Apaf1	Apoptotic protease activating factor 1	0.6
Pdcd8	Programmed cell death 8 (apoptosis inducing factor)	0.8
	Cellular apoptosis susceptibility protein	1.1
	ESTs, highly similar to apoptosis specific protein	1.1
	Apoptosis-associated speck-like protein containing CARD	1.2
AIF	Apoptosis-inducing factor AIF	1.3

^aFor details on methodology and results, please see footnote a to **Table 1**.

^bFor more details, please see footnote b to **Table 1**.

(**Figure 1**) (Wang et al. 2004). Furthermore, the genes of the motility portion of the invasion signature (**Table 3**) can be organized into a series of converging pathways based on the known functions of the proteins for which they code (**Figure 2**) (Wang et al. 2004). The functions of each of these pathways and how they may contribute to the behavior of tumor cells during invasion and intravasation is considered next.

Cofilin pathway. The cofilin family of proteins in vertebrates consists of cofilin/ADF. The cofilin pathway for invasive carcinoma cells is summarized in **Figure 2a**, with the genes whose expression is altered in invasive tumor cells highlighted. The invasion signa-

ture indicates that the cofilin activity cycle has been impacted at several levels of regulation in invasive cells. Cofilin is the more abundant isoform of the family in carcinoma cells, and its expression is highly up-regulated in invasive cells (**Figure 2a**).

Cofilin's severing and depolymerization activities are inhibited by phosphorylation, G-actin binding, and binding to phosphatidylinositol (4,5)-bisphosphate (PIP₂) (Bamburg 1999, DesMarais et al. 2004a, Paaivilainen et al. 2004). Changes in pH can also regulate the level of activity of cofilin, but over the physiological range of pH found in vertebrate cells (6.6–7.4) (Bernstein et al. 2000), the activities of cofilin are only graded, not inactivated, suggesting that pH may act more like a

Table 3 Genes of the motility part of the invasion signature^a

Accession	Gene description	Fold change ^b
AA414612	Capping protein $\alpha 1^c$	4.00
AW556230	Cell division cycle 42 ^c	3.96
AU015486	Capping protein $\alpha 2$	3.89
C79581	Moesin ^c	3.67
C86972	Arp 2/3 complex subunit p16 ^c	3.52
AW538432	Rho interactin protein 3 ^c	3.33
AU015879	LIM-kinase 1 ^c	3.24
AA285584	Palladin	3.12
AW555565	Zyxin	2.93
W10023	Catenin β	2.88
C76867	Tropomyosin α chain	2.86
AU023806	Rho-associated coiled-coil forming kinase 1 ^c	2.71
AW536576	Testis expressed gene 9	2.67
AI324089	Phosphatidylinositol-4-phosphate 5-kinase type II α^c	2.60
AI427644	Epidermal growth factor receptor ^c	2.59
AW541453	Capping protein (actin filament), gelsolin-like	2.53
C86107	Actinin $\alpha 3^c$	2.52
AW543636	Annexin A5	2.47
AA052404	CRIP1 protein	2.32
AA014771	Protein kinase C, ζ^c	2.30
AW546733	Arp 2/3 complex subunit p21 ^c	2.22
AA538228	RAB25, member RAS oncogene family	2.19
AA275245	Vinculin	2.16
AA386680	Kinesin family member 5B	2.13
AW536843	Chaperonin subunit 4 (δ)	2.06
AW536183	Chaperonin subunit 3 (γ)	2.06
AI326287	Tubulin alpha-4 chain	2.05
AW553280	Integrin $\beta 1$ (fibronectin receptor β)	2.00
AW536098	Cofilin 1, nonmuscle ^c	2.00
AU017992	Kinectin 1	2.00
AW557123	Downstream of tyrosine kinase 1	2.00
AW549817	Burkitt lymphoma receptor 1	2.00
AA272097	Fibroblast growth factor receptor 1	0.54
AA073514	Zipcode-binding protein 1 ^c	0.11

^aFor details on methodology and results, please see footnote a to **Table 1**.

^bFor more details, please see footnote b to **Table 1**.

^cThese results have been validated by quantitative real-time-PCR.

rheostat to regulate the amplitude of activity without acting like an on-off-switch.

The consequence of regulating cofilin by phosphorylation appears to differ by cell type. In some cell types, cofilin is almost 100%

phosphorylated in resting cells and motility is stimulated by dephosphorylation (Kanamori et al. 1995, Okada et al. 1996). In carcinoma cells in serum, phospho-cofilin is less than half of the total cofilin at steady state (Zebda et al.

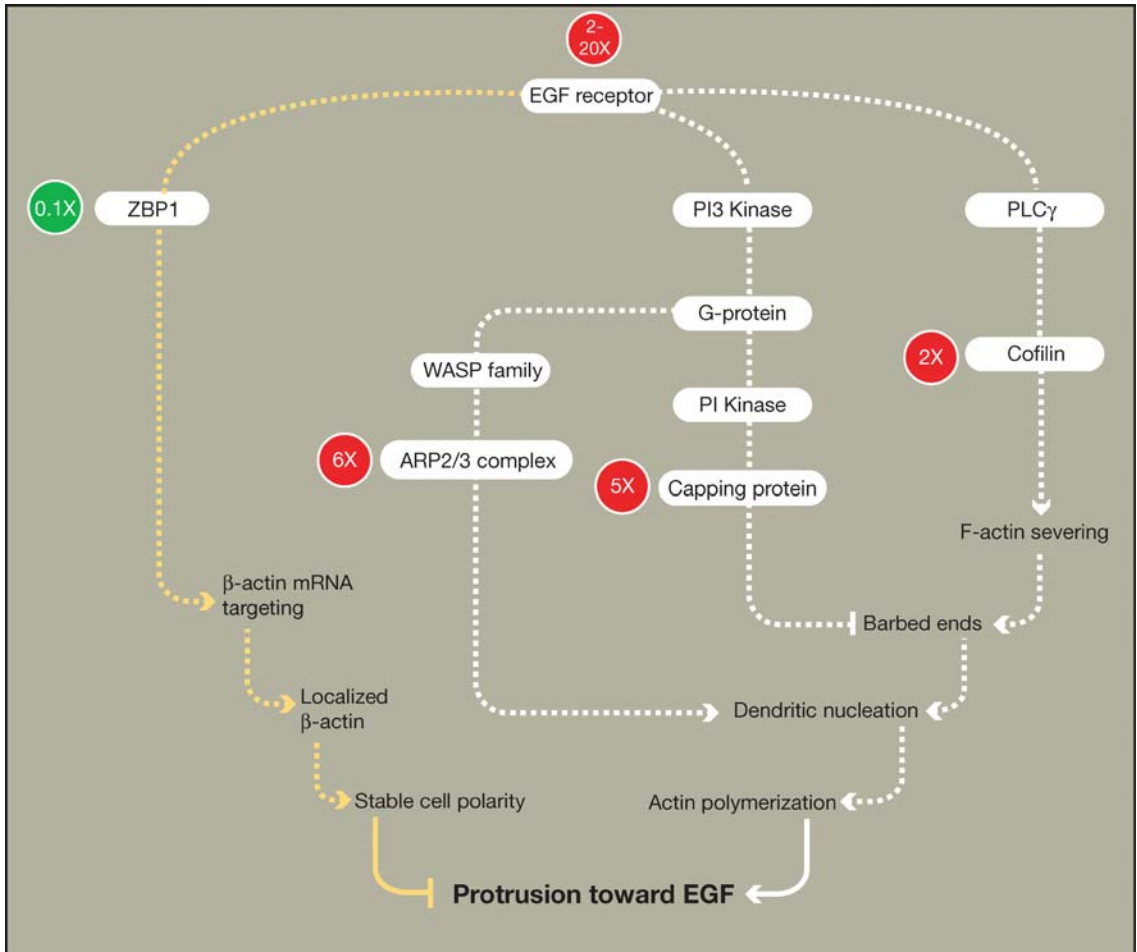


Figure 1

The four pathways leading to β -actin polymerization at the leading edge of carcinoma cells in response to EGF. The major effectors responsible for leading-edge polymerization are cofilin, capping protein, and the Arp2/3 complex. Zip-code-binding protein 1 (ZBP1) regulates chemotaxis to EGF through β -actin mRNA targeting. The fold changes in gene expression in this and **Figure 2** were determined by quantitative real-time PCR and are indicated as (*nx*). Cofilin and Arp2/3 complex are synergistic in the production of free barbed ends leading to dendritic nucleation and protrusive force. Capping protein funnels the available G-actin onto productive elongating barbed ends by capping nonproductive barbed ends. The four pathways therefore coordinately generate protrusions that act to steer the cells during chemotaxis and invasion.

2000), and in serum-starved cells, phospho-cofilin is as little as 10% of the total cofilin (X. Song & R. Eddy, personal communication). Even so, cofilin in both cases is mostly inactive (Chan et al. 2000), indicating that a mechanism other than phosphorylation must be at work to inhibit cofilin activity in carcinoma cells.

Another function of phosphorylation of cofilin in carcinoma cells is the recycling of cofilin from G-actin. Cofilin binds to G-actin with submicromolar affinity and the heterodimer is inactive in both severing and depolymerization (Bamburg 1999, Paavilainen et al. 2004). The release of cofilin from this heterodimer is crucial to the

recycling of cofilin activity. Cyclase-associated protein (CAP) is capable of releasing cofilin from the heterodimer through a direct interaction with actin (Bertling et al. 2004, Paavilainen et al. 2004). In addition, because phospho-cofilin cannot bind to actin, LIM-kinase may also be involved in breaking the G-actin-cofilin heterodimer *in vivo*. Phosphorylation may also function to put limits on the amplitude, location, and duration of cofilin activity after its activation by EGF. Hence, while the phosphorylation/

dephosphorylation cycle of cofilin may not be directly involved in the activation of cofilin in carcinoma cells by EGF, LIM-kinase, along with CAP, may be crucial in regulating the localization and recycling of cofilin activity.

Four different kinases that appear to be downstream of the Rho-family GTPases have been shown to phosphorylate cofilin, LIM-kinase 1 and 2, and TES-kinase 1 and 2 (Arber et al. 1998, Dan et al. 2001, Rosok et al. 1999, Toshima et al. 2001, Yang et al. 1998). In invasive carcinoma cells, LIM-kinase 1 is

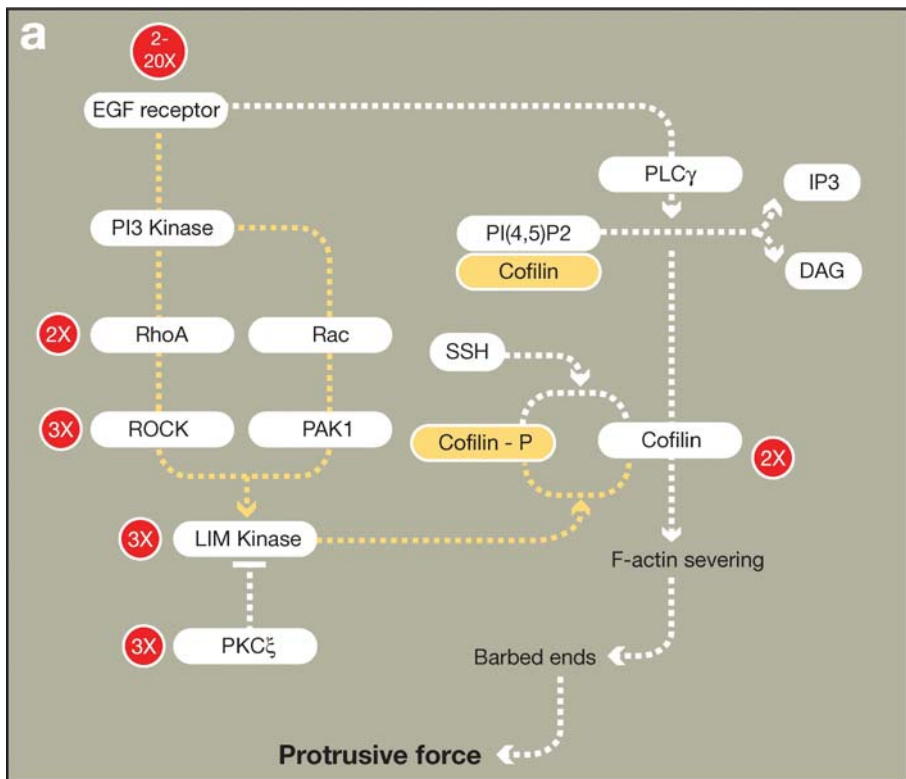


Figure 2

The pathways to barbed end generation and protrusive force. The fold changes in gene expression are indicated as (*nx*). (a) The cofilin pathway leading to barbed end production in response to EGF. Gene for both inhibitory (PAK, ROCK, LIM kinase) and stimulatory (PLC and PKC ξ) inputs to cofilin are more highly expressed in invasive cells; these regulate the location, timing, and sharpness of cofilin-dependent actin polymerization transients that are required for chemotaxis. For *a*, the inhibitory parts are in yellow and the stimulatory parts in white. (b) The capping protein pathway leading to barbed end capping. Genes for both inhibition (Mena and PI5K) and stimulation (capping protein) of the capping activity of this pathway are more highly expressed in invasive cells. (c) The Arp2/3 complex pathway leading to dendritic nucleation in response to EGF. Genes coding for Arp2/3 complex subunits and upstream activators are more highly expressed in invasive cells.

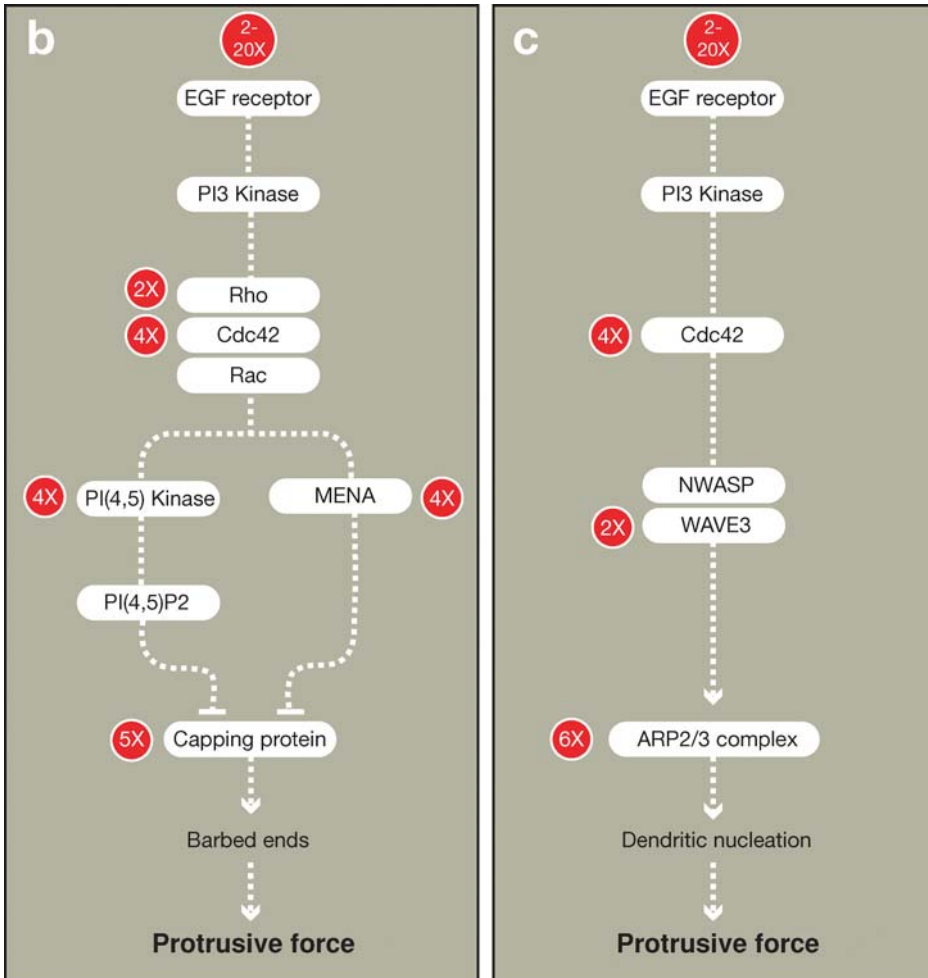


Figure 2

(Continued)

most prominently expressed, and its expression is up-regulated in invasive cells (Wang et al. 2004) (**Figure 2a**). Furthermore, the activation of LIM-kinase 1 occurs through the PI3K-induced activation of Rho-family G-proteins, which activate PAK and ROCK. Rho is highly expressed in invasive cells. Either PAK (Edwards et al. 1999) or ROCK (Ohashi et al. 2000) can phosphorylate LIM-kinase at threonine 508 thereby activating it to increase cofilin phosphorylation. Both kinases are also up-regulated in invasive cells (**Figure 2a**).

Inhibition of LIM-kinase activity is PKC dependent, and this involves one of the atypical PKC isoforms (Djafarzadeh & Niggli 1997, Kuroda et al. 1996). LIM-kinase and

PKC ζ tightly associate via the interaction through the second LIM domain of LIM-kinase, which indicates direct phosphorylation of LIM-kinase (Kuroda et al. 1996). Additional studies have implicated the δ isoform of PKC as a negative regulator of LIM-kinase (Martiny-Baron et al. 1993). The expression of PKC ζ is up-regulated in invasive cells (**Figure 2a**).

The general pattern of regulation in the cofilin pathway indicates that genes coding for proteins that both increase and decrease the activity of cofilin are coordinately up-regulated along with cofilin itself. This pattern may result from the toxicity of elevated cofilin expression (reviewed in Ghosh

et al. 2004), where expression of inhibitory genes is essential to maintain higher levels of cofilin. Alternatively, the significance of this paradoxical pattern may be understood when one considers that the cofilin pathway is directly involved in sensing during chemotaxis of carcinoma cells to EGF (Mouneimne et al. 2004), and cofilin is sufficient to set the direction of cell movement (Ghosh et al. 2004). Directional sensing of EGF requires an early transient of free, actin filament barbed ends resulting from cofilin severing that causes localized actin polymerization (Chan et al. 2000, Mouneimne et al. 2004). If the free barbed ends of the early transient are either inhibited or sustained, then directional protrusion in response to EGF fails (Chan et al. 2000, Mouneimne et al. 2004, Zebda et al. 2000). That is, it is the generation of a transient of free barbed ends that is essential in directional sensing, not sustained polymerization. The up-regulation of genes that both increase and decrease cofilin severing activity, as seen in **Figure 2a**, is consistent with the enhanced ability of invasive cells to generate an early transient that is essential for chemotaxis. In addition, the localization and timing of the stimulatory and inhibitory branches of the cofilin pathway are believed to determine the precise location and duration of cofilin activity and its recycling to compartments where cofilin is inhibited in resting cells (DesMarais et al. 2004a).

Capping protein pathway. Capping protein binds to the growing barbed ends of actin filaments to prevent further elongation and regulate filament length. The patterns of regulation of genes of the capping protein pathway exhibit the same antagonistic relationships as seen in the cofilin pathway where expression of stimulatory and inhibitory branches are up-regulated together. Expression of both the α and β -subunits of capping protein is dramatically increased, suggesting higher capping protein activity in the pathway. However, the expression of genes that code for proteins that are inhibitory to capping protein activity, the

type II- α isoform of PI4, 5 kinase (Cooper & Schafer 2000) and Mena (Bear et al. 2002), are also up-regulated (**Figure 2b**). Capping protein, like cofilin, is essential for viability, and large changes in its expression level may not be tolerated by cells over time (Cooper & Schafer 2000). Therefore, a more interesting interpretation of these results is that the amplitude and sharpness of capping protein activity as a transient is increased in invasive cells because of this antagonistic pattern of expression. The combination of heightened transient capping protein activity and changes in its timing and location could synergize with the barbed end generating activities of the cofilin and Arp2/3 pathways (**Figure 1**) to cause intense focal bursts of actin polymerization, as observed in *in vitro* experiments with purified proteins (Carlier 1998, Loisel et al. 1999).

Arp2/3 complex pathway. Both the cofilin and capping protein pathways converge on the Arp2/3 complex. Because the expression of key components of both pathways is up-regulated, it is interesting that the expression of several subunits of the Arp2/3 complex are also greatly up-regulated in invasive cells, as is the expression of upstream stimulators of the Arp2/3 complex, WAVE 3 and Cdc42 (**Figure 2c**).

Cofilin and Arp2/3 complex synergistically contribute to the nucleation of a dendritic array both *in vitro* (Ichetovkin et al. 2002) and *in vivo* (DesMarais et al. 2004b). This synergy results from the amplification of the Arp2/3 complex's nucleation activity by cofilin's severing activity, which creates barbed ends that elongate to form newly polymerized actin filaments (Ichetovkin et al. 2002). The newly polymerized filaments are the preferred filament type for Arp2/3 complex-mediated branching (DesMarais et al. 2004a,b; Ichetovkin et al. 2002). This synergistic amplification of the Arp2/3 complex activity has been proposed to explain the ability of cofilin to determine sites of protrusion and cell direction in uncaging

Stimulated protrusion (G-actin abundant)

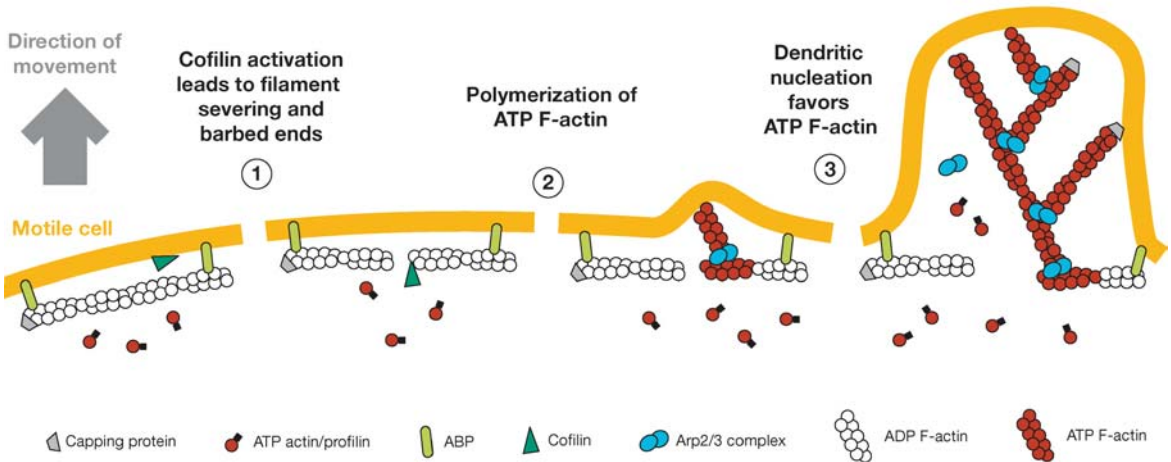


Figure 3

The stimulated protrusion model showing the role of cofilin severing in determining the site of dendritic nucleation, protrusion, and cell direction. Severing of actin filaments in the cortical actin cytoskeleton by cofilin creates free barbed ends that bias the location and the amount of dendritic nucleation by the Arp2/3 complex. Polymerization proceeds from a pool of pre-existing actin monomers, allowing the initiation of polymerization to occur without being tightly coupled to depolymerization. Redrawn from DesMarais et al. (2004a).

experiments (DesMarais et al. 2004a, Ghosh et al. 2004) (**Figure 3**).

Capping protein funnels actin monomers onto newly created free barbed ends by capping older filaments, thereby enhancing the formation of the short, branched filaments characteristic of Arp2/3 complex-nucleated dendritic arrays (Carlier 1998).

WAVE 3 is believed to activate the Arp2/3 complex, as do its relatives WAVEs 1 and 2 (Takenawa & Miki 2001). Both WAVEs 1 and 2 are regulated by Rac 1, which regulates their interaction with Arp2/3 complex to cause stimulation of the Arp2/3 complex's nucleation activity (Eden et al. 2002, Miki et al. 2000, Steffen et al. 2004). This in turn causes lateral (WAVE 2-dependent) and dorsal (WAVE 1-dependent) protrusions (Suetsugu et al. 2003). However, the molecular mechanism of regulation of WAVE 3 is unknown, as are the phenotypic consequences

on cell behavior of stimulating WAVE 3. More work will need to be done on the consequences of WAVE 3 expression and activation in carcinoma cells to understand its significance for tumor cell invasion.

An interesting finding is the coordinated up-regulation of expression of genes for several subunits of the Arp2/3 complex and Cdc42. Cdc42 regulates N-WASP, a ubiquitous member of the WASP family, which induces actin polymerization by activating Arp2/3 complex (Ho et al. 2004). Regulation of the activity of N-WASP involves an intramolecular interaction by which the VCA (verproline/cofilin/acidic) domain, the active site that binds to Arp2/3 complex, is masked by the N-terminal regulatory region of N-WASP (Kim et al. 2000, Rohatgi et al. 2000). The known regulators of N-WASP activity appear to operate by either stabilizing or destabilizing this autoinhibitory

conformation (Ho et al. 2004). Elevated expression of Cdc42, as observed in invasive cells, in combination with the elevated expression of Arp2/3 complex (**Figure 2c**), would be expected to enhance the activity of the N-WASP-Arp2/3 complex pathway, thereby leading to increased invadopod production and cell invasion (Mizutani et al. 2002, Yamaguchi et al. 2005). Cofilin also has been found to amplify and stabilize N-WASP generated invadopods, suggesting that the synergistic interaction between the cofilin and Arp2/3 complex pathways described above is at work during invasion (Ghosh et al. 2004, Yamaguchi et al. 2005).

An additional consequence of increased Cdc42 expression might be its effect on the ability of tumor cells to acquire polarity to blood vessels, as observed during intravasation. Carcinoma cells in metastatic tumors are attracted to blood vessels, where they form a layer of cells that are morphologically polarized toward the vessel. This vessel-directed polarization is believed to be important for intravasation (Condeelis & Segall 2003, Wyckoff et al. 2000a). Chemotaxis undoubtedly contributes to the accumulation of cells around the vessels, but the acquisition of vessel-directed polarity might require additional steps in the reorganization of the cytoskeleton.

A consensus has been building that Cdc42 is involved in determining the direction of cell movement and cell polarity. Inhibition of Cdc42 prevents macrophage migration toward a chemotactic signal (Allen et al. 1998) and directional migration in astrocytes (Etienne-Manneville & Hall 2001). The development of stable cell polarity in astrocytes involves Cdc42, the orientation of the microtubule organizing center, and depends on microtubule dynamics (nocodazole-sensitive) but not on actin polymerization (Etienne-Manneville & Hall 2001, Gundersen et al. 2004). In microtubule-dependent cell polarization in astrocytes, Cdc42 activation involves recruitment of a GEF, FGD-1 and appears to operate through

Par6 to recruit PKC ζ to inactivate GSK3 β (Etienne-Manneville & Hall 2003). This pathway requires stimulation of integrins because arginine-glycine-aspartate (RGD) peptides inhibit the activation of Cdc42, PKC, and protrusion (Etienne-Manneville & Hall 2001).

However, chemotaxis by tumor cells requires actin polymerization and is unaffected by concentrations of nocodazole (100 nM) sufficient to block microtubule dynamics and inhibit cell polarity in astrocytes (Segall et al. 1996). In addition, N-WASP, a major effector of Cdc42, is required for invadopod formation (Yamaguchi et al. 2005) and chemotaxis in tumor cells (L. Soon, personal communication), indicating that cell polarization toward EGF in these cells is dependent on Cdc42 through N-WASP and actin polymerization. During N-WASP- and actin-dependent events at the leading edge of lamellipods, the Cdc42 GEF, intersectin 1, binds to and is activated by N-WASP (Hussain et al. 2001). Therefore, N-WASP may recruit, through intersectin 1, GDP-Cdc42 and activate it locally at the leading edge, making N-WASP function in the polarization to EGF of tumor cells analogous to that of Par6 in cell polarity during wounding. Furthermore, PKC ζ may be involved in the regulation of LIM-kinase activity, which may regulate the amount of active cofilin at the leading edge and its synergy with N-WASP/Arp2/3 complex-mediated protrusion activity (**Figure 1**). These results illustrate parallels between how cells polarize (microtubule-dependent) and chemotax (actin-dependent) and suggest that N-WASP is involved in assembly of a compartment at the leading edge, analogous to the polarity complex in polarizing cells (Etienne-Manneville & Hall 2001), that is required for cell polarity during chemotaxis.

ZBP1 pathway. A gene whose expression is strongly down-regulated in invasive cells is ZBP1 (**Figure 1**). ZBP1 is a member of a family of RNA-binding proteins that contain four C-terminal hnRNP-K homology domains

and two N-terminal RNA recognition motifs (Yaniv & Yisraeli 2002). ZBP1 is a 68-kDa RNA-binding protein that binds to the mRNA zip-code of β -actin mRNA and functions to localize β -actin mRNA to the leading edge of crawling cells. Because β -actin is the preferred isoform of actin for the polymerization of filaments at the leading edge of cells, it is acted on by the cofilin, capping protein, and Arp2/3 pathways (Shestakova et al. 2001). β -actin mRNA localization is required for the maintenance of stable cell polarity as observed in the absence of exogenous signals such as that seen in normal primary fibroblasts, epithelial cells, and tumor cells with differing metastatic potential in which actin polymerization is nucleated at only one pole of the cell in normal and nonmetastatic tumor cells (Shestakova et al. 1999). Disruption of ZBP1-mediated β -actin mRNA targeting in cultured cells leads to cells without cell polarity that are able to nucleate actin polymerization globally and exhibit amoeboid movement (Shestakova et al. 2001). Therefore, ZBP1 may determine the sites in cells where the Arp2/3 complex, capping protein, and cofilin pathways converge to determine the leading edge and cell polarity by controlling the sites of targeting of β -actin mRNA and the location of β -actin protein that is the common downstream effector of these pathways.

Tests of Function of Genes of the Invasion Signature in Chemotaxis, Invasion, and Metastasis

The genes of the motility part of the invasion signature can be organized into three converging pathways based on the known functions of the proteins for which they code (Figure 2). The functions of key gene products in these pathways and how they affect chemotaxis, invasion, and metastasis by carcinoma cells have been tested. The results of these tests are described next.

Cofilin and LIM-kinase. Direct tests of cofilin function are complicated by the fact

that cofilin is required for viability, which makes genetic approaches in carcinoma cells difficult to interpret. However, the acute inhibition of cofilin activity in carcinoma cells inhibits the generation of barbed ends and actin polymerization at the leading edge in response to EGF (Chan et al. 2000, DesMarais et al. 2004b). Inhibition of cofilin activity, through either the inhibition of PLC γ or direct inhibition using acute siRNA suppression of cofilin expression and cofilin function blocking antibodies, inhibits the early barbed end transient that is essential for the chemotaxis of carcinoma cells to EGF (Mouneimne et al. 2004). Furthermore, cofilin is required for the formation of the stable invadopods by carcinoma cells that are important in the invasion of dense extracellular matrix (Mullins et al. 1998, Yamaguchi et al. 2005), particularly that found around blood vessels (Condeelis & Segall 2003). Finally, the local activation of cofilin in carcinoma cells is sufficient to generate protrusive activity and determine cell direction (Ghosh et al. 2004). All these results indicate that cofilin is essential for the chemotaxis and invasion of mammary carcinoma cells to EGF through a mechanism involving the localized generation of barbed ends that causes the localized protrusion, which defines cell direction (Figure 3).

The effects of altering LIM-kinase expression have been studied in tumor cells by several groups, who have shown that overexpression of LIM-kinase 1 in tumor cell lines increases their motility and invasiveness in vitro (Davila et al. 2003, Yoshioka et al. 2003). Experimental reduction in the expression of LIM-kinase 1 in metastatic prostate cell lines decreased invasiveness in matrigel invasion assays. To study the effect of LIM-kinase 1 on metastasis in vivo, an experimental metastasis model was used where cells were injected directly into the left ventricle of mice (Yoshioka et al. 2003). In this case, the ability of cancer cells to survive in the blood, extravasate from blood vessels, and grow at metastatic sites all contribute to the metastasis score, so it is not clear how these

results relate to invasion in the primary tumor. In general, these results are consistent with the observed overexpression of LIM-kinase 1 in invasive cells in mammary tumors and their invasion signature (**Table 3, Figure 2a**).

In a separate set of studies, the overexpression of either full-length-regulated LIM-kinase 1 or its constitutively active kinase domain has been reported to inhibit cofilin activity *in vivo* by phosphorylation. Overexpression also inhibits EGF-induced barbed end production, in particular the early barbed end transient, and lamellipod extension in culture (W. Wang, G. Mouneimne, J. Wyckoff, X. Chen, M. Sidani, and J. Condeelis, unpublished data; Zebda et al. 2000). Furthermore, the overexpression of full-length LIM-kinase 1 in carcinoma cells without altering cofilin expression is correlated with the inhibition of chemotaxis, invasion, intravasation, and metastasis of tumor cells in mammary tumors prepared from these carcinoma cells (W. Wang, G. Mouneimne, J. Wyckoff, X. Chen, M. Sidani, and J. Condeelis, unpublished data). Although these results appear contradictory to those described above, in fact they are consistent with the invasion signature associated with the cofilin pathway (**Figure 2a**). That is, highly invasive cells up-regulate LIM-kinase 1, cofilin, and their stimulatory and inhibitory effectors together (**Figure 2a**), consistent with the hypothesis that the up-regulation of both inhibitory and stimulatory branches of the cofilin pathway increases the amplitude and sharpness of cofilin-dependent actin polymerization transients that are essential for chemotaxis and invasion in carcinoma cells (Mouneimne et al. 2004; W. Wang, G. Mouneimne, J. Wyckoff, X. Chen, M. Sidani, and J. Condeelis, unpublished data). Therefore, to compare studies in which the expression of LIM-kinase, cofilin, or other members of this pathway are experimentally altered, it is essential to measure the output of the cofilin pathway as the timing and amplitude of cofilin-dependent barbed end production during

chemotaxis. Manipulations that increase the cofilin-dependent barbed end production of the early transient during chemotaxis are predicted to increase invasiveness, and this predicts that studies in which cells are more invasive and metastatic after overexpression of LIM-kinase have associated compensatory increases in the expression of other members of the cofilin pathway so as to increase barbed end production in response to EGF. Additional work will be required to investigate this possibility.

N-WASP. N-WASP has been implicated in invasion of extracellular matrix in a number of studies. The invasion of Madin-Darby canine kidney cells during tubulogenesis in collagen gels is inhibited by expression of dominant-negative N-WASP (Yamaguchi et al. 2002). Furthermore, N-WASP, in cooperation with cofilin, is required for the formation of invadopods (Yamaguchi et al. 2005), and its activity is localized to nascent invadopods during the invasion of fibronectin gels (Lorenz et al. 2004). In particular, the depletion of N-WASP or the p34arc subunit of Arp2/3 complex by siRNA interference suppresses invadopod formation. In addition, siRNA interference and dominant-negative mutant expression analyses revealed that cofilin and the N-WASP regulators, Nck1, Cdc42, and WIP, but not Grb2 and WISH, are necessary for invadopod formation (Yamaguchi et al. 2005). EGF receptor kinase inhibitors block the formation of invadopods by carcinoma cells in the presence of serum, and EGF stimulation of serum-starved cells induces invadopod formation. These results indicate that EGF receptor-activated N-WASP and cofilin are required for the formation of invadopods and that Nck1 and Cdc42 mediate the signaling pathway.

A phenomenon that may be related to chemotaxis, invadopod formation, and pathfinding is the observation that the localized stimulation of the EGF receptor on carcinoma cells using EGF-bound beads results in localized actin polymerization and protrusion

(Kempiak et al. 2003). This highly focal actin polymerization requires the activation of the Arp2/3 complex by N-WASP and cofilin and is regulated by Grb2 and Nck2 (Kempiak et al. 2005). This phenomenon may be relevant to how EGF receptor ligands, which can bind to extracellular matrix, stimulate focal protrusions, invadopod formation, and adhesion *in vivo* (Kempiak et al. 2005). Additional work will be required to determine the effects of altering N-WASP activity on invasion, intravasation, and metastasis *in vivo*.

ZBP1. The targeting of β -actin mRNA to the leading lamella is essential for stable cell polarity during locomotion, and ZBP1 is required for mRNA targeting (Condeelis & Singer 2005). Highly metastatic cell lines have reduced levels of ZBP1, and this is consistent with the reduction in ZBP1 expression seen in invasive cells (Wang et al. 2004). Decreased β -actin mRNA targeting seen in cells with reduced ZBP1 is correlated with the loss of cell polarity and increased amoeboid movement in metastatic carcinoma cell lines *in vitro* and *in vivo* (Shestakova et al. 1999, Wang et al. 2002) and increased chemotaxis (Wang et al. 2004). Increasing the level of expression of ZBP1 in invasive carcinoma cells rescues the localization of β -actin mRNA to one pole of the cell and results in the inhibition of chemotaxis to EGF both *in vitro* and *in vivo* in tumors. In addition, tumors prepared from cells re-expressing ZBP1 are significantly less invasive and metastatic than their parental cell-generated counterparts (Wang et al. 2004). However, tumor growth is not significantly affected by increasing the expression of ZBP1. This suggests that the suppression of invasion and metastasis by ZBP1 is not related to growth of the tumor. These results are consistent with the observation that mouse mammary tumors that overexpress the ZBP1 homologue CRD-BP are not metastatic (Tessier et al. 2004).

The invasion and metastasis suppression activity of ZBP1 may result from its ability to suppress the chemotaxis of cancer cells

by maintaining them in a polarized epithelial cell-like state. Cells that lack an intrinsic and stable polarity are more chemotactic to exogenous gradients, presumably because there is no intrinsic polarity to be overcome by the exogenous chemotactic signal and the cell can turn in any direction to respond to the gradient (Iijima et al. 2002, Parent & Devreotes 1999). This may account for the enhanced ability of invasive carcinoma cells to chemotax to blood vessels (Condeelis & Segall 2003, Wyckoff et al. 2000a). It also suggests that the generation of polarity in carcinoma cells that occurs around blood vessels is independent of ZBP1 activity, as discussed above.

NEW INSIGHTS INTO TUMOR INVASION AND METASTASIS

The identification of an invasion signature for mammary tumors that implicates the coordinate regulation of genes involved in functionally related activities presents a rich collection of targets for chemotherapy not previously detected in conventional expression profiling of whole tumors. The fact that the pathways are coordinately regulated in invasive cells suggests that combinations of therapeutics may be particularly effective.

An additional insight resulting from the study of invasive cells and their invasion signature comes from the comparison of expression profiles obtained from invasive cells with the conventional expression profiles of whole tumors. Gene expression profiles of whole tumors have shown promise in prognosis by identifying patterns of expression that are correlated with metastasis (Ramaswamy et al. 2003, van't Veer et al. 2002). However, unlike the invasion signature described for invasive cells, these patterns of expression appear as random sets of genes with unrelated functions and thus are difficult to interpret in terms of mechanisms of invasion and metastasis. This suggests that the invasion signature is either averaged out when interrogating the whole tumor because invasive cells are rare or that the changes in gene expression

that represent the invasion signature are largely transient. Therefore, it is interesting that the expression profiles of whole tumors demonstrate that the invasive and metastatic potential of the primary tumor can be encoded early in the development of the tumor and throughout the bulk of the tumor including the stroma (Ramaswamy et al. 2003, van 't Veer et al. 2002). These results suggest that metastasis could occur early in tumor progression and that most cells in the tumor are potentially metastatic, thus favoring a “transient expression” model rather than an “averaged-out model” to explain the discordance between expression profile results. This conclusion is surprising because the traditional view of tumor progression is that tumors develop through a succession of stable genetic changes acquired through selection pressures, a process analogous to Darwinian evolution. According to the traditional view of tumor progression, the cells selected to be metastatic are very rare, and metastases arise from progressive genetic changes in these rare cells within a primary tumor delaying metastasis to late stages of tumor progression (Bernards & Weinberg 2002, Hanahan & Weinberg 2000).

The Tumor Microenvironment Invasion Model

A new model, the tumor microenvironment invasion model (TMIM), has been proposed to explain the relationship between the expression pattern of invasive cells and expression patterns of whole tumors and how these relate to the traditional view of tumor progression (Wang et al. 2005). In this model, the transient changes in gene expression leading to invasion (the invasion signature) result from microenvironments in the tumor that are defined by stable genetic changes in both stromal and tumor cells. That is, tumor progression, as described by traditional models (Hanahan & Weinberg 2000), leads to the development of microenvironments encoded within the tumor, which elicit the tran-

sient gene expression patterns that support invasion. In this context, invasion is similar to a morphogenetic program involving the transient expression of genes that lead to a change in the location of cells, a program that can occur repeatedly during tumor development and in any location in the tumor that has the microenvironment that elicits the morphogenetic program. The expression of genes that are synergistic for inducing microenvironments causing invasion could lead to the random appearance, in time and location, of these microenvironments during tumor progression leading to repeated episodes of invasion and metastasis throughout tumor progression.

TMIM is consistent with the finding that genes encoding the tumor microenvironment for invasion and metastasis appear to be expressed throughout the bulk of the tumor. It is also consistent with the ability to collect invasive cells by chemotaxis using needles that are placed in random locations in tumors if the growth factors inside the needles mimic microenvironments inducing invasion, as claimed (Wang et al. 2004, Wyckoff et al. 2004). Furthermore, the TMIM hypothesis is supported by intravital imaging of experimental tumors where only a small proportion of tumor cells are motile, and moving cells are not uniformly distributed but are observed in localized areas of the tumor (Condeelis & Segall 2003, Wang et al. 2002), and the observation that micrometastases are often genetically heterogeneous, suggesting that invasive behavior is not stably specified (Klein 2002). Finally, the TMIM hypothesis is consistent with our current understanding of how the tumor microenvironment contributes to invasion and metastasis (Bissell & Radisky 2001).

The exciting new technologies reviewed here have brought us to the point where tumor invasion and metastasis can be studied as a problem in morphogenesis. The future will reveal if the new insights that are emerging will lead to new strategies for the diagnosis and treatment of metastasis.

LITERATURE CITED

- Ahmed F, Wyckoff J, Lin EY, Wang W, Wang Y, et al. 2002. GFP expression in the mammary gland for imaging of mammary tumor cells in transgenic mice. *Cancer Res.* 62:7166–69
- Allen WE, Zicha D, Ridley AJ, Jones GE. 1998. A role for Cdc42 in macrophage chemotaxis. *J. Cell Biol.* 141:1147–57
- Arber S, Barbayannis FA, Hanser H, Schneider C, Stanyon CA, et al. 1998. Regulation of actin dynamics through phosphorylation of cofilin by LIM-kinase. *Nature* 393:805–9
- Bailly M, Yan L, Whitesides GM, Condeelis JS, Segall JE. 1998. Regulation of protrusion shape and adhesion to the substratum during chemotactic responses of mammalian carcinoma cells. *Exp. Cell Res.* 241:285–99
- Bamburg JR. 1999. Proteins of the ADF/cofilin family: essential regulators of actin dynamics. *Annu. Rev. Cell Dev. Biol.* 15:185–230
- Bear JE, Svitkina TM, Krause M, Schafer DA, Loureiro JJ, et al. 2002. Antagonism between Ena/VASP proteins and actin filament capping regulates fibroblast motility. *Cell* 109:509–21
- Bernards R, Weinberg RA. 2002. A progression puzzle. *Nature* 418:823
- Bernstein BW, Painter WB, Chen H, Minamide LS, Abe H, Bamburg JR. 2000. Intracellular pH modulation of ADF/cofilin proteins. *Cell Motil. Cytoskelet.* 47:319–36
- Bertling E, Hotulainen P, Mattila PK, Matilainen T, Salminen M, Lappalainen P. 2004. Cyclase-associated protein 1 (CAP1) promotes cofilin-induced actin dynamics in mammalian nonmuscle cells. *Mol. Biol. Cell* 15:2324–34
- Bissell MJ, Radisky D. 2001. Putting tumors in context. *Nat. Rev. Cancer* 1:46–54
- Brown EB, Campbell RB, Tsuzuki Y, Xu L, Carmeliet P, et al. 2001. In vivo measurement of gene expression, angiogenesis and physiological function in tumors using multiphoton laser scanning microscopy. *Nat. Med.* 7:864–68
- Campagnola PJ, Clark HA, Mohler WA, Lewis A, Loew LM. 2001. Second-harmonic imaging microscopy of living cells. *J. Biomed. Opt.* 6:277–86
- Carlier MF. 1998. Control of actin dynamics. *Curr. Opin. Cell Biol.* 10:45–51
- Chambers AF, MacDonald IC, Schmidt EE, Koop S, Morris VL, et al. 1995. Steps in tumor metastasis: new concepts from intravital videomicroscopy. *Cancer Metastasis Rev.* 14:279–301
- Chan AY, Bailly M, Zebda N, Segall JE, Condeelis JS. 2000. Role of cofilin in epidermal growth factor-stimulated actin polymerization and lamellipod protrusion. *J. Cell Biol.* 148:531–42
- Chantrain CF, Shimada H, Jodele S, Groshen S, Ye W, et al. 2004. Stromal matrix metalloproteinase-9 regulates the vascular architecture in neuroblastoma by promoting pericyte recruitment. *Cancer Res.* 64:1675–86
- Chishima T, Miyagi Y, Wang X, Yamaoka H, Shimada H, et al. 1997. Cancer invasion and micrometastasis visualized in live tissue by green fluorescent protein expression. *Cancer Res.* 57:2042–47
- Clark EA, Golub TR, Lander ES, Hynes RO. 2000. Genomic analysis of metastasis reveals an essential role for RhoC. *Nature* 406:532–35
- Condeelis J, Segall JE. 2003. Intravital imaging of cell movement in tumours. *Nat. Rev. Cancer* 3:921–30
- Condeelis J, Singer R. 2005. How and why does β -actin mRNA target? *Biol. Cell* 97:97–110
- Condeelis J, Song X, Backer J, Wyckoff J, Segall J. 2003. *Chemotaxis of cancer cells during invasion and metastasis*. Presented at 5th Abercrombie Symp. Cell Behav., St. Catherine's College, Oxford, UK

- Cooper JA, Schafer DA. 2000. Control of actin assembly and disassembly at filament ends. *Curr. Opin. Cell Biol.* 12:97–103
- Dan C, Kelly A, Bernard O, Minden A. 2001. Cytoskeletal changes regulated by the PAK4 serine/threonine kinase are mediated by LIM kinase 1 and cofilin. *J. Biol. Chem.* 276:32115–21
- Davila M, Frost AR, Grizzle WE, Chakrabarti R. 2003. LIM kinase 1 is essential for the invasive growth of prostate epithelial cells: Implications in prostate cancer. *J. Biol. Chem.* 278:36868–75
- DesMarais V, Ghosh M, Eddy RE, Condeelis J. 2004a. Cofilin takes the lead. *J. Cell Sci.* 118:19–26
- DesMarais V, Macaluso F, Condeelis J, Bailly M. 2004b. Synergistic interaction between the Arp2/3 complex and cofilin drives stimulated lamellipod extension. *J. Cell Sci.* 117:3499–510
- Di Modugno F, Bronzi G, Scanlan MJ, Del Bello D, Cascioli S, et al. 2004. Human Mena protein, a serex-defined antigen overexpressed in breast cancer eliciting both humoral and CD8+ T-cell immune response. *Int. J. Cancer* 109:909–18
- Djafarzadeh S, Niggli V. 1997. Signaling pathways involved in dephosphorylation and localization of the actin-binding protein cofilin in stimulated human neutrophils. *Exp. Cell Res.* 236:427–35
- Eden S, Rohatgi R, Podtelejnikov AV, Mann M, Kirschner MW. 2002. Mechanism of regulation of WAVE 1. *Nature* 418:790–93
- Edwards DC, Sanders LC, Bokoch GM, Gill GN. 1999. Activation of LIM-kinase by Pak1 couples Rac/Cdc42 GTPase signalling to actin cytoskeletal dynamics. *Nat. Cell Biol.* 1:253–59
- Etienne-Manneville S, Hall A. 2001. Integrin-mediated activation of Cdc42 controls cell polarity in migrating astrocytes through PKC ζ . *Cell* 106:489–98
- Etienne-Manneville S, Hall A. 2003. Cdc42 regulates GSK-3 β and adenomatous polyposis coli to control cell polarity. *Nature* 421:753–56
- Farina KL, Wyckoff JB, Rivera J, Lee H, Segall JE, et al. 1998. Cell motility of tumor cells visualized in living intact primary tumors using green fluorescent protein. *Cancer Res.* 58:2528–32
- Friedl P, Wolf K. 2003. Tumour-cell invasion and migration: diversity and escape mechanisms. *Nat. Rev. Cancer* 3:362–74
- Ghosh M, Song X, Mouneimne G, Sidani M, Lawrence DS, Condeelis JS. 2004. Cofilin promotes actin polymerization and defines the direction of cell motility. *Science* 304:743–46
- Goswami S, Sahai E, Wyckoff J, Cammer M, Cox D, Pixley F, et al. 2005. Macrophages promote the invasion of breast carcinoma cells via a paracrine loop. *Cancer Res.* 65:5278–83
- Goswami S, Wang W, Wyckoff JB, Condeelis JS. 2004. Breast cancer cells isolated by chemotaxis from primary tumors show increased survival and resistance to chemotherapy. *Cancer Res.* 64:7664–67
- Gundersen GG, Gomes ER, Wen Y. 2004. Cortical control of microtubule stability and polarization. *Curr. Opin. Cell Biol.* 16:106–12
- Hanahan D, Weinberg RA. 2000. The hallmarks of cancer. *Cell* 100:57–70
- Helmchen F, Denk W. 2002. New developments in multiphoton microscopy. *Curr. Opin. Neurobiol.* 12:593–601
- Ho HY, Rohatgi R, Lebensohn AM, Le M, Li J, et al. 2004. Toca-1 mediates Cdc42-dependent actin nucleation by activating the N-WASP-WIP complex. *Cell* 118:203–16

- Hussain NK, Jenna S, Glogauer M, Quinn CC, Wasiak S, et al. 2001. Endocytic protein intersectin-1 regulates actin assembly via Cdc42 and N-WASP. *Nat. Cell Biol.* 3:927–32
- Ichetovkin I, Grant W, Condeelis J. 2002. Cofilin produces newly polymerized actin filaments that are preferred for dendritic nucleation by the Arp2/3 complex. *Curr. Biol.* 12:79–84
- Iijima M, Huang YE, Devreotes P. 2002. Temporal and spatial regulation of chemotaxis. *Dev. Cell* 3:469–78
- Jain RK, Munn LL, Fukumura D. 2002. Dissecting tumour pathophysiology using intravital microscopy. *Nat. Rev. Cancer* 2:266–76
- Kamai T, Tsujii T, Arai K, Takagi K, Asami H, et al. 2003. Significant association of Rho/ROCK pathway with invasion and metastasis of bladder cancer. *Clin. Cancer Res.* 9:2632–41
- Kanamori T, Hayakawa T, Suzuki M, Titani K. 1995. Identification of two 17-kDa rat parotid gland phosphoproteins, subjects for dephosphorylation upon beta-adrenergic stimulation, as destrin- and cofilin-like proteins. *J. Biol. Chem.* 270:8061–67
- Kempiaj SJ, Yamaguchi H, Sarmiento C, Sidani M, Ghosh M, et al. 2005. An N-Wasp mediated pathway for localized activation of actin polymerization which is regulated by cortactin. *J. Biol. Chem.* 280(7):5836–42
- Kempiaj SJ, Yip SC, Backer JM, Segall JE. 2003. Local signaling by the EGF receptor. *J. Cell Biol.* 162:781–87
- Kim AS, Kakalis LT, Abdul-Manan N, Liu GA, Rosen MK. 2000. Autoinhibition and activation mechanisms of the Wiskott-Aldrich syndrome protein. *Nature* 404:151–58
- Klein CA, Blankenstein T, Schmidt-Kittler O, Petronio M, Polzer B, et al. 2002. Genetic heterogeneity of single disseminated tumour cells in minimal residual cancer. *Lancet* 360:683–89
- Kuroda S, Tokunaga C, Kiyohara Y, Higuchi O, Konishi H, et al. 1996. Protein-protein interaction of zinc finger LIM domains with protein kinase C. *J. Biol. Chem.* 271:31029–32
- Levene MJ, Dombeck DA, Kasischke KA, Molloy RP, Webb WW. 2004. In vivo multiphoton microscopy of deep brain tissue. *J. Neurophysiol.* 91:1908–12
- Lin EY, Nguyen AV, Russell RG, Pollard JW. 2001. Colony-stimulating factor 1 promotes progression of mammary tumors to malignancy. *J. Exp. Med.* 193:727–40
- Liotta LA, Kohn EC. 2001. The microenvironment of the tumour-host interface. *Nature* 411:375–79
- Loisel TP, Boujemaa R, Pantaloni D, Carlier MF. 1999. Reconstitution of actin-based motility of *Listeria* and *Shigella* using pure proteins. *Nature* 401:613–16
- Lorenz M, Yamaguchi H, Wang Y, Singer RH, Condeelis J. 2004. Imaging sites of N-WASP activity in lamellipodia and invadopodia of carcinoma cells. *Curr. Biol.* 14:697–703
- Martiny-Baron G, Kazanietz MG, Mischak H, Blumberg PM, Kochs G, et al. 1993. Selective inhibition of protein kinase C isozymes by the indolocarbazole Go 6976. *J. Biol. Chem.* 268:9194–97
- Masters BR, So PT, Gratton E. 1997. Multiphoton excitation fluorescence microscopy and spectroscopy of in vivo human skin. *Biophys. J.* 72:2405–12
- Miki H, Yamaguchi H, Suetsugu S, Takenawa T. 2000. IRSp53 is an essential intermediate between Rac and WAVE in the regulation of membrane ruffling. *Nature* 408:732–35
- Mizutani K, Miki H, He H, Maruta H, Takenawa T. 2002. Essential role of neural Wiskott-Aldrich syndrome protein in podosome formation and degradation of extracellular matrix in src-transformed fibroblasts. *Cancer Res.* 62:669–74
- Mouneimne G, Soon L, DesMarais V, Sidani M, Song X, et al. 2004. Phospholipase C and cofilin are required for carcinoma cell directionality in response to EGF stimulation. *J. Cell Biol.* 166:697–708

- Mullins RD, Heuser JA, Pollard TD. 1998. The interaction of Arp2/3 complex with actin: nucleation, high affinity pointed end capping, and formation of branching networks of filaments. *Proc. Natl. Acad. Sci. USA* 95:6181–86
- Ohashi K, Nagata K, Maekawa M, Ishizaki T, Narumiya S, Mizuno K. 2000. Rho-associated kinase ROCK activates LIM-kinase 1 by phosphorylation at threonine 508 within the activation loop. *J. Biol. Chem.* 275:3577–82
- Okada K, Takano-Ohmuro H, Obinata T, Abe H. 1996. Dephosphorylation of cofilin in polymorphonuclear leukocytes derived from peripheral blood. *Exp. Cell Res.* 227:116–22
- Otsubo T, Iwaya K, Mukai Y, Mizokami Y, Serizawa H, et al. 2004. Involvement of Arp2/3 complex in the process of colorectal carcinogenesis. *Mod. Pathol.* 17:461–67
- Paavilainen V, Bertling E, Falck S, Lappalainen P. 2004. Regulation of cytoskeletal dynamics by actin-monomer-binding proteins. *Trends Cell Biol.* 14:386–94
- Parent CA, Devreotes PN. 1999. A cell's sense of direction. *Science* 284:765–70
- Pollard JW. 2004. Tumour-educated macrophages promote tumour progression and metastasis. *Nat. Rev. Cancer* 4:71–78
- Ramaswamy S, Ross KN, Lander ES, Golub TR. 2003. A molecular signature of metastasis in primary solid tumors. *Nat. Genet.* 33:49–54
- Rohatgi R, Ho HY, Kirschner MW. 2000. Mechanism of N-WASP activation by CDC42 and phosphatidylinositol 4,5-bisphosphate. *J. Cell Biol.* 150:1299–310
- Rosok O, Pedeutour F, Ree AH, Aasheim HC. 1999. Identification and characterization of TESK2, a novel member of the LIMK/TESK family of protein kinases, predominantly expressed in testis. *Genomics* 61:44–54
- Sahai E, Wyckoff J, Philippar U, Gertler F, Segall J, Condeelis J. 2005. Simultaneous imaging of GFP, CFP and collagen in tumors in vivo. *BMC Biotechnol.* 5:14
- Scherbarth S, Orr FW. 1997. Intravital videomicroscopic evidence for regulation of metastasis by the hepatic microvasculature: effects of interleukin-1alpha on metastasis and the location of B16F1 melanoma cell arrest. *Cancer Res.* 57:4105–10
- Segall JE, Tyrech S, Boselli L, Masseling S, Helft J, et al. 1996. EGF stimulates lamellipod extension in metastatic mammary adenocarcinoma cells by an actin-dependent mechanism. *Clin. Exp. Metastasis* 14:61–72
- Shestakova EA, Singer RH, Condeelis J. 2001. The physiological significance of beta-actin mRNA localization in determining cell polarity and directional motility. *Proc. Natl. Acad. Sci. USA* 98:7045–50
- Shestakova EA, Wyckoff J, Jones J, Singer RH, Condeelis J. 1999. Correlation of beta-actin messenger RNA localization with metastatic potential in rat adenocarcinoma cell lines. *Cancer Res.* 59:1202–5
- Steffen A, Rottner K, Ehinger J, Innocenti M, Scita G, et al. 2004. Sra-1 and Nap-1 link Rac to actin assembly driving lamellipodia formation. *EMBO J.* 23:749–59
- Suetsugu S, Yamazaki D, Kurisu S, Takenawa T. 2003. Differential roles of WAVE1 and WAVE2 in dorsal and peripheral ruffle formation for fibroblast cell migration. *Dev. Cell* 5:595–609
- Suzuki T, Yanagi K, Ookawa K, Hatakeyama K, Ohshima N. 1996. Flow visualization of microcirculation in solid tumor tissues: intravital microscopic observation of blood circulation by use of a confocal laser scanning microscope. *Front Med. Biol. Eng.* 7:253–63
- Takenawa T, Miki H. 2001. WASP and WAVE family proteins: key molecules for rapid rearrangement of cortical actin filaments and cell movement. *J. Cell. Sci.* 114:1801–9
- Tessier CR, Doyle GA, Clark BA, Pitot HC, Ross J. 2004. Mammary tumor induction in transgenic mice expressing an RNA-binding protein. *Cancer Res.* 64:209–14

- Toshima J, Toshima JY, Amano T, Yang N, Narumiya S, Mizuno K. 2001. Cofilin phosphorylation by protein kinase testicular protein kinase 1 and its role in integrin-mediated actin reorganization and focal adhesion formation. *Mol. Biol. Cell* 12:1131–45
- Vajkoczy P, Goldbrunner R, Farhadi M, Vince G, Schilling L, et al. 1999. Glioma cell migration is associated with glioma-induced angiogenesis in vivo. *Int. J. Dev. Neurosci.* 17:557–63
- van 't Veer LJ, Dai H, van de Vijver MJ, He YD, Hart AA, et al. 2002. Gene expression profiling predicts clinical outcome of breast cancer. *Nature* 415:530–36
- Wang W, Goswami S, Lapidus K, Wells AL, Wyckoff JB, et al. 2004. Identification and testing of a gene expression signature of invasive carcinoma cells within primary mammary tumors. *Cancer Res.* 64:8585–94
- Wang W, Goswami S, Sahai E, Wyckoff J, Segall J, Condeelis J. 2005. Tumor cells caught in the act of invading: How they revealed their strategy for enhanced cell motility. *Trends Cell Biol.* 15(3):138–45
- Wang W, Wyckoff JB, Frohlich VC, Oleynikov Y, Huttelmaier S, et al. 2002. Single cell behavior in metastatic primary mammary tumors correlated with gene expression patterns revealed by molecular profiling. *Cancer Res.* 62:6278–88
- Wang W, Wyckoff JB, Wang Y, Bottinger EP, Segall JE, Condeelis JS. 2003. Gene expression analysis on small numbers of invasive cells collected by chemotaxis from primary mammary tumors of the mouse. *BMC Biotechnol.* 3:13–25
- Williams RM, Zipfel WR, Webb WW. 2001. Multiphoton microscopy in biological research. *Curr. Opin. Chem. Biol.* 5:603–8
- Wood S. 1958. Pathogenesis of metastasis formation observed in vivo in the rabbit ear chamber. *Arch. Pathol.* 66:550–68
- Woodhouse EC, Chuaqui RF, Liotta LA. 1997. General mechanisms of metastasis. *Cancer* 80:1529–37
- Wyckoff J, Wang W, Lin EY, Wang Y, Pixley F, et al. 2004. A paracrine loop between tumor cells and macrophages is required for tumor cell migration in mammary tumors. *Cancer Res.* 64:7022–29
- Wyckoff JB, Jones JG, Condeelis JS, Segall JE. 2000a. A critical step in metastasis: in vivo analysis of intravasation at the primary tumor. *Cancer Res.* 60:2504–11
- Wyckoff JB, Segall JE, Condeelis JS. 2000b. The collection of the motile population of cells from a living tumor. *Cancer Res.* 60:5401–4
- Yamaguchi H, Lorenz M, Kempiak SJ, Sarmiento C, Coniglio S, et al. 2005. Molecular mechanism of invadopodium formation: The role of the N-WASP/Arp2/3 complex pathway and cofilin. *J. Cell Biol.* 168:441–52
- Yamaguchi H, Miki H, Takenawa T. 2002. Neural Wiskott-Aldrich syndrome protein is involved in hepatocyte growth factor-induced migration, invasion, and tubulogenesis of epithelial cells. *Cancer Res.* 62:2503–9
- Yang N, Higuchi O, Ohashi K, Nagata K, Wada A, et al. 1998. Cofilin phosphorylation by LIM-kinase 1 and its role in Rac-mediated actin reorganization. *Nature* 393:809–12
- Yang M, Baranov E, Jiang P, Sun FX, Li XM, et al. 2000. Whole body optical imaging of green fluorescent protein expressing tumors and metastasis. *PNAS* 97:1206–11
- Yaniv K, Yisraeli J. 2002. The involvement of a conserved family of RNA binding proteins in embryonic development and carcinogenesis. *Gene* 287:49–54
- Yoshioka K, Foletta V, Bernard O, Itoh K. 2003. A role for LIM kinase in cancer invasion. *Proc. Natl. Acad. Sci. USA* 100:7247–52
- Yuan F, Dellian M, Fukumura D, Leunig M, Berk DA, et al. 1995. Vascular permeability in a human tumor xenograft: molecular size dependence and cutoff size. *Cancer Res.* 55:3752–56

- Zebda N, Bernard O, Bailly M, Welti S, Lawrence DS, Condeelis JS. 2000. Phosphorylation of ADF/cofilin abolishes EGF-induced actin nucleation at the leading edge and subsequent lamellipod extension. *J. Cell Biol.* 151:1119–28
- Zipfel WR, Williams RM, Christie R, Nikitin AY, Hyman BT, Webb WW. 2003a. Live tissue intrinsic emission microscopy using multiphoton-excited native fluorescence and second harmonic generation. *Proc. Natl. Acad. Sci. USA* 100:7075–80
- Zipfel WR, Williams RM, Webb WW. 2003b. Nonlinear magic: multiphoton microscopy in the biosciences. *Nat. Biotechnol.* 21:1369–77