Adhesion rings surround invadopodia and promote maturation

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Summary

Invasion and metastasis are aggressive cancer phenotypes that are highly related to the ability of cancer cells to degrade extracellular matrix (ECM). At the cellular level, specialized actin-rich structures called invadopodia mediate focal matrix degradation by serving as exocytic sites for ECM-degrading proteinases. Adhesion signaling is likely to be a critical regulatory input to invadopodia, but the mechanism and location of such adhesion signaling events are poorly understood. Here, we report that adhesion rings surround invadopodia shortly after formation and correlate strongly with invadopodial activity on a cell-by-cell basis. By contrast, there was little correlation of focal adhesion number or size with cellular invadopodium activity. Prevention of adhesion ring formation by inhibition of RGD-binding integrins or knockdown (KD) of integrin-linked kinase (ILK) reduced the number of ECM-degrading invadopodia and reduced recruitment of IQGAP to invadopodium actin puncta. Furthermore, live cell imaging revealed that the rate of extracellular MT1-MMP accumulation at invadopodia was greatly reduced in both integrin-inhibited and ILK-KD cells. Conversely, KD of MT1-MMP reduced invadopodium activity and dynamics but not the number of adhesion-ringed invadopodia. These results suggest a model in which adhesion rings are recruited to invadopodia shortly after formation and promote invadopodium maturation by enhancing proteinase secretion. Since adhesion rings are a defining characteristic of podosomes, similar structures formed by normal cells, our data also suggest further similarities between invadopodia and podosomes.

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Introduction

An increasing number of studies have shown that deregulated ECM-integrin signaling is a critical driver of cancer aggressiveness, both in human tumors and in model systems (Lahlou and Muller, 2011; Lehmann et al., 2011; Parekh and Weaver, 2009; Provenzano et al., 2008a; Provenzano et al., 2008b; Stewart et al., 2004). One of the most feared and lethal consequences of deregulated adhesion signaling that can lead to metastatic spread is invasive tumor cell behavior; however, the underlying cellular mechanisms are complex and include cross-regulation of cell-cell adhesion, migration and extracellular matrix (ECM) degradation (Guo and Giancotti, 2004). ECM degradation is likely to be particularly critical for invasion of epithelial tumors surrounded by dense crosslinked basement membranes, since they cannot be traversed without proteolytic activity (Parekh and Weaver, 2009).

A number of studies have shown that actin-rich protrusions known as invadopodia are key cancer cell structures for ECM degradation due to the focused concentration of matrix-degrading proteases (Linder et al., 2011; Weaver, 2006). Similar structures have been identified in 3-dimensional systems and in mouse models, and are dependent for their activity on key invadopodium molecules (Gligorijevic et al., 2012; Quintavalle et al., 2010; Rottiers et al., 2009; Wolf et al., 2007). In addition, normal cells such as macrophages and osteoclasts that need to remodel matrix form similar structures called podosomes (Linder et al., 2011; Murphy and Courtneidge, 2011). Matrix metalloproteinases (MMPs) are particularly important for the ECM-degrading activity of invadopodia and recent papers have shown that vesicle trafficking is important for delivery of MMPs to initial invadopodium actin puncta. The transmembrane proteinase MT1-MMP in particular is trafficked from a late endocytic compartment to invadopodia, dependent on VAMP7, IQGAP, and the exocyst complex (Sakurai-Yageta et al., 2008; Steffen et al., 2008). MT1-MMP is essential in many systems for invadopodium activity, traversal of basement membranes and in vivo tissue invasion (Artym et al., 2006; Hotary et al., 2006; Sabe et al., 2004).

We recently found that both the chemical composition and rigidity of ECM could regulate the activity of cancer cell invadopodia (Alexander et al., 2008; Liu et al., 2010; Parekh et al., 2011). Similarly, integrins and downstream adhesion adaptor proteins have also been found to regulate invadopodia (Chan et al., 2009; Lucas et al., 2010; Nakahara et al., 1996) and invadopodium-like structures formed in src-transformed cells (Badowski et al., 2008; Destaing et al., 2010). Although focal
Adhesions are the cancer cell structures most highly associated with ECM-integrin signaling, we and others have identified Myosin IIa- (Alexander et al., 2008), integrin α5 (Mueller et al., 1999), and RhoC- (Bravo-Cordero et al., 2011) positive structures surrounding invadopodia that resemble the adhesion rings of podosomes (Gimona et al., 2008; Linder and Aepfelbacher, 2003), suggesting direct on-site signaling. In addition, Spinardi et al. described a “podosome-like” structure formed in bladder carcinoma cells, with adhesion rings surrounding actin cores within hemidesmosome plaques (Spinardi et al., 2004). However, the presence of adhesion rings surrounding podosomes has been suggested to be a defining difference between podosomes and invadopodia (Gimona et al., 2008; Linder, 2009; Linder et al., 2011). Furthermore, unlike many podosome-forming cells, for which podosomes are the only adhesion structures, cancer cells usually contain focal adhesions (FA) which may compete with invadopodia for adhesion signaling proteins (Chan et al., 2009) or induce necessary signals from a distance (Wang et al., 2005). Therefore, whether cancer cell invadopodia constitute bona fide adhesion structures is unclear, as is the exact role of adhesion proteins in the invadopodium lifecycle.

In this study, we followed up on the intriguing possibility that invadopodia form adhesion rings and investigated how adhesion signaling regulates invadopodium activity. We find that virtually all invadopodia do form adhesion rings and that the adhesion rings are formed shortly after actin assembly. Inhibition of adhesion signaling by targeting RGD-binding integrins and integrin-linked kinase (ILK) led to a selective reduction in adhesion-ringed and ECM-degrading invadopodia, and reduced the rate of MT1-MMP accumulation at invadopodia. In addition, integrin- and ILK-inhibited cells exhibited a reduction in the recruitment of the vesicular adaptor and ILK-binding protein IQGAP to invadopodia. These data suggest a model in which integrin-ILK signaling at invadopodium precursor structures leads to adhesion ring formation, IQGAP recruitment and capture of MT1-MMP-containing vesicles to produce fully mature ECM-degrading invadopodia.

**Results**

Podosome-like adhesion rings form around invadopodia and correlate with invadopodium activity

To understand how adhesion signaling associated with either FA or invadopodia regulates invadopodium activity, we began by immunostaining invadopodium-forming cancer cells for the FA markers paxillin or vinculin as well as the invadopodium markers cortactin or actin. CA1d breast cancer and SCC61 head and neck squamous cell carcinoma (HNSCC) cells have been previously described and were plated on invadopodium substrates coated with crosslinked 1% gelatin overlaid with FITC-fibronectin (FN) (Alexander et al., 2008; Clark et al., 2007). Interestingly, visualization by confocal microscopy revealed that paxillin and vinculin not only localized to FA but also to podosome-like adhesion rings surrounding actively degrading cortactin- or actin-containing invadopodia (Fig. 1A,B; supplementary material Fig. S1). Due to the high cytoplasmic levels of these adhesion proteins, these rings were not visible in wide-field epifluorescent images. Additional adhesion markers were also localized to either the rings (activated β1 integrin, α5 integrin), or invadopodium puncta (activated Src) (supplementary material Fig. S1).

In order to better understand the relationship between adhesion rings, FA, and invadopodium activity, we performed a correlative analysis of invadopodium-associated ECM degradation with the number of adhesion-ringed invadopodia or FA. To provide a richer dataset for analysis and because we previously found that cells modulate invadopodium activity according to substrate rigidity, we chose 3 rigidity conditions for the analysis and cultured CA1d or SCC61 cells on FITC-FN/1% gelatin overlaying polyacrylamide (PA) or glass invadopodium substrates of defined rigidity (Soft PA = 1 kPa, Hard PA = 10 kPa, glass = 1 GPa) as previously described (Alexander et al., 2008; Parekh et al., 2011). After 18 hours, the cultures were fixed and immunostained for cortactin and paxillin. Cortactin-positive invadopodium puncta have a characteristic morphology and size ≥1 μm (Clark et al., 2007), and were analyzed for association with ECM degradation and number per cell. FA number and size were analyzed from paxillin-positive structures at the periphery of the cell, and invadopodium-associated adhesion rings were identified manually (see Materials and Methods for more details, and Fig. 1 and supplementary material Fig. S1 for ringed invadopodium images). Overall, analysis across all conditions revealed that a higher percentage of adhesion-ringed invadopodium puncta were associated with degraded ECM than were non-ringed invadopodia (Fig. 1C; supplementary material Fig. S1C), suggesting that adhesion rings may promote invadopodium-associated degradation. We also correlated the number of FA and invadopodium ring structures per cell with the amount of degradation per cell. Interestingly, the number of paxillin-ringed invadopodia peaked on the same rigidity that elicited the highest invadopodium activity per cell, on the hard PA gel for the CA1d cells and on the glass surface for the SCC61 cells (supplementary material Fig. S1D,E). Using a Spearman’s rank correlation to test the statistical dependence between two variables, we found that the number of adhesion-ringed invadopodia per cell significantly and positively correlated with ECM degradation area per cell on a cell-by-cell basis (Fig. 1D). The number of non-adhesion-ringed invadopodia also significantly correlated with ECM degradation per cell but to a lesser extent than adhesion-ringed invadopodia. By contrast, except for SCC61 cells plated on glass substrates, neither the number nor size of FA correlated with the degradation area per cell on a cell-by-cell basis (Fig. 1D; supplementary material Fig. S1D,E). We also found no relationship between FA dynamics and ECM degradation (live cell imaging, data not shown). Finally, we did not observe ECM degradation colocalized with FA or in streak-like FA-type distribution (Wang and McNiven, 2012).

Altogether, these correlative data indicate that adhesion ringed-invadopodia are highly associated with ECM degradation.

Adhesion rings form shortly after invadopodium formation

Since SCC61 cells cultured on the glass surface degraded the most ECM and formed the most adhesion-ringed invadopodia (supplementary material Fig. S1D,E), this condition was used for subsequent mechanistic experiments to identify the role of adhesion rings in invadopodium function. Current models of invadopodium stages include early initiation of actin polymerization, followed shortly by recruitment of matrix metalloproteinases (MMPs) such as MT1-MMP and subsequent ECM degradation (Aytym et al., 2006; Oser et al., 2009). Adhesion rings might be recruited early, before actin polymerization to trigger initiation of invadopodia, or alternatively, later, to promote persistence of invadopodia and/or recruitment of MMPs. To determine the real-time dynamics of
adhesion rings forming at invadopodia, SCC61 cells stably expressing GFP-paxillin were transiently transfected with td-Tomato-F-Tractin (Tom-Tractin) to mark actin-positive invadopodia (Johnson and Schell, 2009; Laukaitis et al., 2001). These cells were then cultured overnight in medium without serum on MatTek dishes coated with 1% crosslinked gelatin/FITC-FN (“Matrix”, green in triple merge) and immunostained for vinculin (Vinc) or paxillin (Pax) to mark adhesion structures (red in merges) and actin or cortactin (Cort) to mark invadopodia (blue in merges). Example focal adhesions (black arrows) and adhesion rings surrounding invadopodia (white arrows) are indicated. Dark areas in the matrix are where the FITC-FN/gelatin was degraded. Scale bars = 5 μm. (B) Zooms of boxed areas in A. Scale bars = 5 μm. 

Percentage of adhesion ringed- and non-adhesion ringed-invadopodia colocalized with degradation of the underlying matrix in CA1d or SCC61 cells stained for paxillin and cortactin. Data are compiled from cells plated on soft PA, hard PA and glass conditions. Data separated by condition are shown in supplementary material Fig. S1. n > 180 cells per condition from 3 independent experiments. (D) Spearman correlation of the invadopodium ring or focal adhesion characteristics of an individual cell with the degradation area of FITC-FN underneath the same cell for CA1d or SCC61 cells plated on the noted underlying surfaces and stained for paxillin and cortactin. n = 60 cells per condition from 3 independent experiments. *P < 0.05; **P < 0.01; ***P < 0.001 vs. no correlation.

1. In general, adhesion rings formed quickly after the formation of the Tom-Tractin-positive actin puncta, with respective mean and median times of 2.6 and 2 minutes after formation of the invadopodia (Fig. 2B). In addition, ~90% of Tom-Tractin-positive invadopodium puncta recruited an adhesion ring over the lifetimes observed in the 1 hour movies (Fig. 2C). Finally, as has been previously noted (Bravo-Cordero et al., 2011), we frequently saw an oscillation of actin intensity at invadopodium puncta. These actin oscillations were paralleled by ring oscillations and the recovery of GFP-paxillin intensity was generally slightly delayed in onset with respect to the Tom-Tractin intensity.
recovery (Fig. 2D kymograph). Occasionally, adhesion rings were lost from the invadopodia and did not recover (Fig. 2E kymograph) or arrived late (Fig. 2B), which we speculate may explain why many invadopodia in fixed analyses did not have surrounding rings. These data indicate that adhesion rings are typically recruited shortly after invadopodum initiation and suggest that they may contribute to the stability of invadopodia and/or recruitment of proteinases.

Integrin activity is critical for adhesion ring formation and invadopodium activity

To directly test the role of adhesion proteins in invadopodium formation, adhesion ring formation, and invadopodium-associated ECM degradation, we blocked ligand binding of multiple integrin classes by incubating SCC61 cells with a peptide that blocks the binding of RGD-binding integrins to their ligands (gRGDsp) or with a combination of binding of RGD-binding integrins to their ligands (gRGDsp) and integrin blocking antibodies (AIIB2+LM609) (Cheresh, 1987; Hall et al., 1990) (Fig. 3; supplementary material Fig. S2). These integrin targets were chosen based on the ECM substrate used in our assay (FITC-FN) as well as their known contribution to podosome formation and activity (Destaining et al., 2010; Nakamura et al., 2007). We began with a standard immunofluorescence invadopodium assay and quantitated the ECM degradation area per cell. In addition, we quantitated the presence of various invadopodium features, including the total number of cortactin-positive characteristica invadopodium puncta, which blocks the binding of RGD-binding integrins to their ligands (gRGDsp). We speculated that cortactin puncta had associated ECM degradation (inactive) vs. “Active” invadopodia). Although it is possible that some of the “Inactive” invadopodia may represent other cortactin-positive structures, the size criteria of ≥1 μm diameter minimizes this possibility and use of this metric allows evaluation of invadopodium maturation. Furthermore, we complement such analyses with live cell imaging (next paragraph). Compared to gRGESP or IgG controls, integrin inhibition reduced the median area of ECM degradation per cell from 4 to 1.4 μm² and 6.4 to 3.1 μm², respectively (Fig. 3A,B; supplementary material Fig. S2). Integrin inhibition also significantly reduced the median number of adhesion-ringed invadopodia from 1 to 0 per cell, but had little effect on total invadopodium numbers (Fig. 3C; supplementary material Fig. S2). In addition, integrin inhibition with these reagents reduced the median number of active invadopodia from 3 to 1 per cell, while increasing or not affecting the number of inactive invadopodia (Fig. 3C; supplementary material Fig. S2). Of note, incubating cells with either β1 or αvβ3 blocking antibodies alone respectively inhibited or had no significant effect on the activity of invadopodia, suggesting that β1 may be the main integrin involved in our system (supplementary material Fig. S2C,D). Overall, these data indicate that integrins significantly contribute to adhesion ring formation and invadopodium-associated ECM degradation.

To directly determine if inhibition of integrins leads to a loss of proteinase recruitment, or conversely to alterations in invadopodium dynamics, we performed live cell imaging. To allow exclusive detection of extracellular MT1-MMP, SCC61 cells were engineered to stably express low levels of the critical invadopodium proteinase MT1-MMP tagged with superspectral-pHLuorin-GFP (MT1-MMP-pHLuorin). The pHLuorin tag has greatly enhanced fluorescence at the extracellular pH of 7.4 and is quenched in the more acidic intracellular vesicles (Lizarraga et al., 2009; Miesenböck et al., 1998). After transient transfection with Tom-Actin to visualize invadopodia (supplementary material Movie 2), dual color widefield images were captured on a DeltaVision microscope at a frame rate of 1 per 15 seconds and computationally deconvolved to remove out of focus light. From these movies, the times of invadopodium formation and disassembly were determined as the IC50 points of the background-subtracted Tom-Actin fluorescence intensity fit to a sigmoid curve at the initiation or disappearance of the invadopodia. Invadopodium lifetime was calculated as the time between formation and disassembly. Invadopodium formation rate was manually calculated as the number of invadopodia per cell per hour. MT1-MMP recruitment to invadopodia over time...
was measured as the background-subtracted fluorescence intensity of MT1-MMP-pHLuorin at each time point subtracted from the initial fluorescence intensity at the time of invadopodium formation. Using this approach, we found that inhibition of RGD-binding integrins with gRGDsp had no significant effect on the rate of invadopodium formation or on invadopodium lifetime compared to treatment with control gRGEsp peptide (Fig. 3D,F). However, there was a significant reduction in the accumulation of MT1-MMP at invadopodia. Thus, in control gRGEsp-treated cells, the MT1-MMP signal at Tom-Tractin-positive invadopodia increased steadily over the lifetime of individual invadopodia (Fig. 3E,G; supplementary material Movie 2). By contrast, in gRGDsp-treated cells, the overall MT1-MMP accumulation rate was significantly decreased, suggesting that the major effect of integrin inhibition was on invadopodium maturation (Fig. 3E,G; supplementary material Movie 3).

Integrin-linked kinase controls adhesion ring formation, invadopodium dynamics and activity

Our live and fixed imaging results with RGD blocking peptide indicated that integrins are involved in both adhesion ring formation and the accumulation of MT1-MMP at invadopodia. Since the majority of invadopodial MT1-MMP is derived from exocytosis of late endocytic vesicles (Steffen et al., 2008), we hypothesized that downstream signaling from integrins might facilitate this process. Although little is known about the relationship between integrin signaling and exocytosis (Balasubramanian et al., 2010; Gupton and Gertler, 2010; Wickström and Fässler, 2011; Wickström et al., 2010a), a potential downstream candidate is integrin-linked kinase (ILK), based on its reported function in linking caveolar vesicles to IQGAP at adhesions (Wickström et al., 2010a). Since cholesterol-rich caveolin-positive membranes and IQGAP have been separately shown to mediate proteinase trafficking to invadopodia (Caldieri et al., 2009; Sakurai-Yageta et al., 2008; Yamaguchi et al., 2009), ILK seemed a likely linker between integrins and MT1-MMP containing vesicles that might promote invadopodium degradative activity. To test that hypothesis, we knocked down ILK in SCC61 cells. Using two separate shRNA constructs in SCC61 cells, ILK expression was decreased by 58% and 82% in ILK-KD1 and -KD2 cells compared with a non-targeting control (Fig. 4B). When assessed by fixed cell immunofluorescence, ILK-KD cells exhibited a significant reduction in invadopodium maturation (Fig. 4E,G; supplementary material Movie 3).
KD had no further effect on invadopodium-associated ECM degradation compared with either treatment alone, suggesting that they are likely to function in the same pathway (Fig. 4E).

We also performed live cell imaging in GFP-paxillin, Tom-Tractin-expressing control and ILK-KD cells to analyze the effect of ILK-KD on MT1-MMP accumulation and invadopodium dynamics. Unlike the effect of RGD peptide (Fig. 3), ILK-KD led to a decrease in invadopodium formation, from a median of 28 to 12 invadopodia per cell per hour, and in invadopodium lifetime from a median of 35 to 21 minutes. (Fig. 4F,G). Similar to the effect of integrin inhibition with gRGDsp peptide, KD of ILK1 led to a large reduction in the rate of MT1-MMP recruitment to invadopodia. We note that unlike with RGD-inhibited cells, the decrease in extracellular MT1-MMP accumulation only occurred after the first 20 minutes (Fig. 4H,I; supplementary material Movies 4, 5). However, overall these data indicate that integrin and ILK-mediated adhesion signaling is critical for accumulation of extracellular MT1-MMP at invadopodia. Since the accumulation was quantitated at already-formed invadopodia, the decreases in protease accumulation for both integrin- and ILK-inhibited cells were independent of any effects on invadopodium formation or lifetime.

Integrins and ILK recruit IQGAP to invadopodia

Based on our finding that integrins and ILK affect invadopodium activity and MT1-MMP recruitment and the known role of the ILK-binding partner IQGAP in capturing caveolar vesicles (Sakurai-Yageta et al., 2008; Wickström et al., 2010a), we decided to test whether IQGAP localization at invadopodia was affected by integrin-ILK signaling. 3-D confocal imaging of IQGAP localization to actin- and vinculin-stained invadopodia demonstrated that IQGAP is localized in an overlapping distribution with both the actin puncta and the vinculin ring.
(Fig. 5A, horizontal line scans). Vertical line scans through the center of invadopodia also show that IQGAP is more concentrated at the top of the actin-positive area (Fig. 5A). To determine whether IQGAP is recruited to invadopodia dependent on integrins and ILK, RGD-inhibited or ILK-KD SCC61 cells were cultured overnight on invadopodium substrates and fixed and stained for actin and IQGAP. IQGAP and actin intensities at invadopodia were quantitated from confocal images taken under identical laser and camera conditions for all samples and the ratio of IQGAP to actin intensity at each invadopodium was plotted. Consistent with a model in which integrins and ILK facilitate IQGAP recruitment to invadopodia, inhibition of RGD-binding integrins or ILK-KD led to a decrease in the intensity of IQGAP at invadopodia (Fig. 5B–E). The median ratio of IQGAP:actin intensity decreased from 0.57 to 0.49 with RGD treatment and from 0.49 to 0.34 and 0.37 with ILK shRNA #1 and #2, respectively, compared to controls.

MT1-MMP regulates ECM degradation and invadopodium dynamics, but not adhesion ring formation

Recruitment of MT1-MMP and other proteinases is thought to be a late stage in invadopodium formation; however, several studies have reported reduced numbers of invadopodia per cell in proteinase-inhibited cells (Artym et al., 2006; Clark et al., 2007; Steffen et al., 2008), suggesting feedback from proteolytic activity.
To test whether MT1-MMP acts upstream or downstream of adhesion ring formation, SCC61 cells transduced with shRNA targeting MT1-MMP were analyzed for invadopodia and adhesion ring formation and ECM degrading ability in fixed cell assays (Fig. 6). Consistent with previous reports (Artym et al., 2006; Steffen et al., 2008), ECM degradation area per cell was virtually abolished with MT1-MMP knockdown (Fig. 6C). Furthermore, there was a 4-fold decrease in the number of active invadopodia per cell and a corresponding increase in inactive invadopodium numbers (Fig. 6D). There was no significant change in either the number of adhesion-ringed invadopodia or the number of total invadopodia. Live cell imaging further revealed that the rate of invadopodium formation was reduced in MT1-MMP-KD cells (Fig. 6E). There was also a small but significant decrease in the median lifetime of invadopodia in MT1-MMP-KD cells from 45.5 to 35.5 minutes (Fig. 6F), suggesting that there is indeed a positive feedback loop that results from MT1-MMP activity at invadopodia (Artym et al., 2006; Clark et al., 2007; Steffen et al., 2008). These data are consistent with a model in which adhesion rings form upstream of MT1-MMP accumulation to promote ECM degradation. MT1-MMP activity itself also has significant effects on invadopodium dynamics, both in the formation and stability phases.

**Discussion**

In this study, we examined the role of adhesion signaling in the regulation of invadopodium formation and activity. We find that podosome-like adhesion rings surround invadopodia shortly after their formation and their presence is highly correlated with invadopodium activity. Integrins and ILK are critical for adhesion ring formation, localization of the vesicular adaptor protein IQGAP and the transmembrane protease MT1-MMP to invadopodia, and ECM degradation. Overall these data indicate that adhesion signaling is critical for invadopodium maturation.

**Adhesion ring formation is not a distinguishing feature of podosomes**

Invadopodia and podosomes are both actin-rich structures that can degrade underlying matrix. Despite many similarities, including common molecular constituents and pathways of activation, invadopodia are currently classified as distinct structures that form in cancer cells and have separate properties.
from podosomes such as longer lifetimes, protrusive behavior, and morphological classifications (Gimona et al., 2008; Murphy and Courtneidge, 2011). One major distinction has been the presence of adhesion rings surrounding actin puncta in podosomes (Gimona et al., 2008; Linder, 2009; Linder et al., 2011). Here, we showed that two separate invadopodium-producing cancer cell lines exhibit adhesion ring formation around invadopodial protrusions. These rings consist of multiple adhesion components such as paxillin, vinculin, and activated β1 integrin and occur at the basal cell surface by confocal microscopy, suggesting that the ring structures represent bona fide adherent structures. However, they were less prominent with respect to staining intensity and morphology than typical podosome structures formed in cells such as macrophages, perhaps due to competition with focal adhesions for components (Chan et al., 2009) or due to greater distribution of adhesion components into a soluble cytoplasmic pool in cancer cells. Interestingly, in src-transformed fibroblasts, β1 integrin was shown to be important for not only the organization but also the formation of “invadosome” structures, which have similarities to both podosomes and invadopodia and are directly induced by mutated src kinase (Destaing et al., 2010). Although in our cancer cell invadopodium system we do find some role for ILK in invadopodium formation, the major role that we observed for both integrins and ILK was a very specific effect on invadopodium adhesion ring formation and maturation. Similarly, in osteoclasts, despite a small effect on actin podosome puncta diameter, knockout of integrin β1, β2, and/or αv or kindlin3 led to severe defects in podosome organization and bone resorption (Schmidt et al., 2011). In addition, Badowski et al. examined the role of paxillin in invadosome formation by src-transformed cells and podosome formation by osteoclasts. In those systems, they found that loss or mutation of paxillin primarily affected podosome organization and ECM degradation (Badowski et al., 2008). Overall, our data suggest that adhesion ring formation is a common feature of both podosomes and invadopodia and indicate that a major function of those rings is to promote recruitment of proteinases and ECM degradation.

A new model of invadopodium stages

Altogether, our data are consistent with a model in which adhesion rings are assembled shortly after invadopodium actin puncta assembly and promote recruitment of proteinases to allow ECM degradation (Fig. 7). RGD-binding integrins and ILK are crucial components of this process as inhibition or knock-down reduces adhesion ring formation, MT1-MMP recruitment to invadopodia and ECM degradation. Furthermore, inhibiting integrins in ILK-KD cells had no further effect on ECM degradation, suggesting that integrins and ILK reside in the same pathway with respect to invadopodium-associated ECM degradation. We place MT1-MMP at a later stage than adhesion formation since KD of MT1-MMP had no effect on the number of adhesion-ringed invadopodia but inhibition of adhesion signaling led to decreased recruitment of MT1-MMP. After exocytosis, MT1-MMP and other proteases might also interact with integrins via a direct docking mechanism (Gálvez et al., 2002; Mueller et al., 1999) that could provide additional positive feedback to enhance invadopodium lifetime and/or ECM degradation.

Whereas integrin inhibition with gRGDsp did not affect invadopodium formation or lifetime, ILK-KD did, indicating that ILK may not function solely downstream of integrins for those activities. Although ILK is thought to primarily function as a downstream effector of integrins in complex with PINCH and parvin (Sakai et al., 2003; Stanchi et al., 2009; Wickström et al., 2010b), ILK can also be regulated by growth factor signaling (Ho and Dagnino, 2012; Serrano et al., 2012) and PI3K/PTEN (Wu and Dedhar, 2001). Since invadopodium formation can be activated by both growth factor and PI3K signaling (Yamaguchi et al., 2005; Yamaguchi et al., 2011), one possibility is that the regulation of invadopodium dynamics by ILK may occur downstream of growth factor rather than integrin signaling. Another possibility is that the inhibition of RGD-binding integrins with gRGDsp was less effective at blocking the integrin-ILK-IQGAP-MT1-MMP pathway than ILK-KD. Since knockdown of MT1-MMP itself can affect invadopodium dynamics (Fig. 6), a more complete block of the pathway with ILK-KD might lead to the difference that we noted for invadopodium dynamics.

Regulation of exocytosis by integrins and ILK

Integrins and adhesion signaling have recently been shown to affect diverse exocytic processes. Assembly and localization of the exocyst complex were shown to be regulated by integrin adhesion and paxillin through activation of the RalA GTPase (Balasubramanian et al., 2010; Spiczka and Yeaman, 2008). β1...
Integrin-ECM interaction was also recently shown to promote exocytosis of lipid rafts by a mechanism that links ILK, the scaffold protein IQGAP1 and the formin mDia1 to capture of microtubules and subsequent vesicle delivery to the plasma membrane (Wickström and Fässler, 2011; Wickström et al., 2010a). Furthermore, integrin adhesion to collagen in 3D was shown to promote polarized trafficking of MT1-MMP (Bravo-Cordero et al., 2007). Of note, several microtubule motors have been shown to regulate exocytosis of MT1-MMP at podosomes (Cornfine et al., 2011; Wiesner et al., 2010). Furthermore, the exocyst complex localizes to invadopodia through IQGAP1 and is important for MT1-MMP localization to invadopodia (Sakurai-Yageta et al., 2008). In light of these data and our finding that IQGAP localization to invadopodia is reduced in integrin- and ILK-KD cells (Fig. 5), we speculate that adhesion rings surrounding invadopodia may promote targeted MT1-MMP recruitment to invadopodia through capture of vesicles via a similar ILK-IQGAP-mediated mechanism (Fig. 7).

Deregulation of β1 integrin and ILK signaling has been implicated in a wide range of aggressive tumor behaviors, including cancer invasion (Dai et al., 2003; Edwards et al., 2008; Guo and Giancotti, 2004; Lu et al., 2008; Poincloux et al., 2011; Sawai et al., 2006). With regard to ECM degradation and cancer invasion, our data suggest that at least one key mechanism may be via integrin-ILK-mediated regulation of protease secretion at invadopodia. Interestingly, ILK-regulated secretion has also been implicated in tumor angiogenesis (Edwards et al., 2008) and caveolar formation (Wickström et al., 2010a). Future studies should address how general is the secretory role of integrin-ILK signaling and to what extent invadopodia represent “hotspot” secretion sites for lipid raft-carried molecules.

Materials and Methods

Reagents

Antibodies: Cortactin, 4F11 (Upstate Biotechnology, Lake Placid, NY), Paxillin, Y113 (Abcam, Cambridge, MA), Vinculin, bvn-m-1 (Sigma, St. Louis, MO), IQGAP, H-109 (Santa Cruz, Santa Cruz, CA), Alexa Fluor 546 or 633 phalloidin (Invitrogen, Grand Island, NY). Peptides: Gly-Arg-Gly-Asp-Ser-Pro (GRGDSP, Sigma), and Gly-Arg-Gly-Glu-Ser-Pro (GRGESP, American Peptide Co., Sunnyvale, CA). tTomato-F-tractin (Tom-Tract) was a gift of Dr. Robert Fischer at NHLBI and was created by cloning the 9-52 stretch of the F-actin binding protein ITPKA (Johnson and Schell, 2009) into pCMV-tdTomato (Clontech, Mountain View, CA).

Cell culture and stable cell lines

MCF10A-Ca1d breast cancer cells were obtained from Dr. Fred Miller (Karnovsky Institute, MI) and have previously been described (Santner et al., 2001). Cells were cultured in DMEM:F12 supplemented with 5% horse serum (HyClone, Thermo Scientific, Lafayette, CO), 0.1 μg/ml cholera toxin (Calbiochem, Merck KGaA, Darmstadt, Germany), 10 μg/ml insulin (Gibco, Invitrogen), 0.5 μl ml hydrocortisone (Sigma), and 100 ng/ml EGF (Invitrogen) at 37°C with constant humidity. SCC61 HNSCC cells were previously described (Clark et al., 2007; Wechselbaum et al., 1986) and were cultured in DMEM supplemented with 20% fetal bovine serum (HyClone) and 0.4 mg/ml hydrocortisone at 37°C.

Tom-Tract was transiently transfected into SCC61 cells at a 1:1 μg DNA:μL Lipofectamine 2000 (Invitrogen) ratio for 6 hours. GFP-paxillin was cloned from pEGFP-N3-Paxillin, a gift of Dr. Donna Webb (Vanderbilt), into the pENTR-TOPO vector and recombined into the pCMV-Tomato vector using the Gateway recombination system (Invitrogen). MT1-MMP-pHLuorin (MT1-pHLuor) was a gift from Dr. Philippe Chavrier (Institut Curie, Paris). MT1-pHLuor was PCR-cloned into the pLentiv lentiviral vector (Invitrogen). Control (non-targeting shRNA (NTC), Addgene) and two shRNA constructs targeting ILK (sh1: 5'-CGGAACAACACUUCCGGAU-3' (Thermo Scientific), sh2-5'-CAGCAGATGAACTTCTTCA-3', sh2-5'-CAAGCTCCTACTCATCCTTCA-3') or shILC2 as a control were subcloned into the lentiviral vector pLenti-BlockIt (Invitrogen). Retrovirus and lentivirus were produced using Phoenix (Dr. Gary Nolan, Stanford) or 293FT cells, respectively, and SCC61 cells were incubated with viral supernatant overnight and later selected with the appropriate markers.

Fixed cell microscopy and invadopodium analysis

The ECM degradation assay has been previously described (Bowden et al., 2001; Clark et al., 2007). Briefly, MatTek culture dishes (MatTek Corp., Ashland, MA) were coated with a thin layer of 1% gelatin that was crosslinked with 0.5% glutaraldehyde prior to addition of 50 μg/ml FITC-conjugated fibronectin (FITC-FN). Cells were cultured in a 1:1 ratio of DMEM:RPMI-1640 (or L–15 for live cell imaging) with 5% NuSerum (Gibco), 10% FBS, and 100 ng/ml EGF for 20 hours before fixation with 4% paraformaldehyde and immunostaining. Polyacrylamide substrate preparation was also previously described (Alexander et al., 2008; Parekh et al., 2011).

Widefield fluorescent images were captured on a Nikon Eclipse TE2000-E microscope with a 40× Plan Fluor oil immersion objective lens. Degradation area per cell was determined using MetaMorph software (Molecular Devices, Sunnyvale, CA) by tracing a region around the cellular actin footprint and measuring the thresholded area of dark spots in the FITC-FN in that region.

Fixed cell confocal images were taken using a Zeiss LSM 510 microscope with a Plan Apo 63× oil immersion objective lens with Argon-488 nm, HeNe-543 nm, and HeNe-633 nm lasers. Images were taken at a scan speed of 8 and 4 scans were averaged per acquisition. Degradation area per cell was determined as above. Invadopodia were identified as morphologically characteristic round actin or cortactin puncta >1 μm in diameter found at the bottom of the cell (Clark et al., 2007) and quantitated from confocal images. If ECM degradation was associated with an invadopodium, it was classified as “active”; otherwise it was classified as “inactive”. The presence of an adhesion ring was defined as paxillin or vinculin fluorescence above background surrounding the invadopodia. Invadopodium metrics (adhesion ringed, non-ringed, total, active and inactive invadopodia) were manually counted. For focal adhesion analyses on paxillin-immunostained cells, the central region of the cell with high background cytoplasmic fluorescence was excluded and the remaining periphery was thresholded. The number and area of peripheral focal adhesions for each cell were then determined with the “Analyze Particles” function in ImageJ (NIH).

For IQGAP localization analyses, SCC61 cells were plated on 1% gelatin/FITC-FN and fixed and stained for actin and IQGAP. Images were obtained with the Zeiss LSM 510 confocal as described above. The pinhole was opened slightly (1.4 μm optical slice) to account for the slightly higher Z-axis localization of IQGAP at invadopodia. Equal laser power, digital gain and offset settings were used in each experiment and actin and IQGAP signals were within the dynamic range. Invadopodia were traced using the actin immunostaining and average intensity of the actin and IQGAP signals within the traced regions was obtained using ImageJ.

Live cell imaging

Adhesion ring imaging

For adhesion ring imaging, live cell confocal images were taken using a Zeiss LSM 710 microscope with a Plan Apo 63× oil immersion objective lens with Argon-488 nm and HeNe-561 nm lasers. Time-lapse images were obtained every 15 seconds and were processed with 10 iterations of constrained iterative deconvolution using Softworx 5.0 (Applied Precision, Issaquah, WA). For live-cell invadopodium analyses, the actin-containing invadopodia were traced at their largest area and fluorescence intensity of Tom-TRACT and MT1-pHLuorin was determined at each time point using Metamorph. The Tom-TRACT and MT1-pHLuorin signal at each time point was background corrected at each invadopodium with an identical traced region placed in a similar area of the cell but lacking an invadopodium. With these data, invadopodium maturation was determined.

Imaging of Tom-TRACT alone or with MT1-pHluorin

Imaging of Tom-TRACT alone or with MT1-pHluorin was performed using an Applied Precision DeltaVision Core microscope with a Plan Apo 60× oil immersion objective lens. SCC61 cells expressing Tom-TRACT and/or MT1-pHluorin were seeded in the ECM degradation assay on 1% gelatin/unlabeled-FN and placed in a heated microscope chamber at 37°C for 2 hours prior to imaging. When used, RGE or RGD peptides were added at the time of cell seeding. Images were then obtained every 15 seconds and were processed with 10 iterations of constrained iterative deconvolution using Softworx 5.0 (Applied Precision, Issaquah, WA). For live-cell invadopodium analyses, the actin-containing invadopodia were traced at their largest area and fluorescence intensity of Tom-TRACT and MT1-pHluorin was determined at each time point using Metamorph. The Tom-TRACT and MT1-pHluorin signal at each time point was background corrected at each invadopodium with an identical traced region placed in a similar area of the cell but lacking an invadopodium. With these data, invadopodium formation rate, lifetime, and MT1-pHluorin accumulation at invadopodia were determined as described in the manuscript.

Statistics

Data sets were tested for normality using the Kolmogorov-Smirnov normality test in GraphPad Instat3 (GraphPad Software, La Jolla, CA). Groups of non-normal data were compared using the Mann-Whitney rank sum test, while parametric data were compared with an unpaired t-test. Non-normal data were plotted as box and whiskers, which show respectively the 25–75th and 5–95th percentiles with the dotted red line indicating the mean and the black line indicating the median. For
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the Spearman rank analysis performed in Fig. 1, comparisons between degradation area per cell and each other data set were performed in GraphPad InStat3. A perfect positive correlation = 1, no correlation = 0 and a perfect negative correlation = −1, and p values represent a correlation significantly different than no correlation.

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Competing Interests

The authors declare that there are no competing interests.

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