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CD44-Mediated Adhesion to Hyaluronic Acid Contributes to Mechanosensing and Invasive Motility

Yushan Kim and Sanjay Kumar

Abstract

The high-molecular-weight glycosaminoglycan, hyaluronic acid (HA), makes up a significant portion of the brain extracellular matrix. Glioblastoma multiforme (GBM), a highly invasive brain tumor, is associated with aberrant HA secretion, tissue stiffening, and overexpression of the HA receptor CD44. Here, transcriptomic analysis, engineered materials, and measurements of adhesion, migration, and invasion were used to investigate how HA/CD44 ligation contributes to the mechanosensing and invasive motility of GBM tumor cells, both intrinsically and in the context of Arg-Gly-Asp (RGD) peptide/integrin adhesion. Analysis of transcriptomic data from The Cancer Genome Atlas reveals upregulation of transcripts associated with HA/CD44 adhesion. CD44 suppression in culture reduces cell adhesion to HA on short time scales (0.5-hour postincubation) even if RGD is present, whereas maximal adhesion on longer time scales (3 hours) requires both CD44 and integrins. Moreover, time-lapse imaging demonstrates that cell adhesive structures formed during migration on bare HA matrices are more short lived than cellular protrusions formed on surfaces containing RGD. Interestingly, adhesion and migration speed were dependent on HA hydrogel stiffness, implying that CD44-based signaling is intrinsically mechanosensitive. Finally, CD44 expression paired with an HA-rich microenvironment maximized three-dimensional invasion, whereas CD44 suppression or abundant integrin-based adhesion limited it. These findings demonstrate that CD44 transduces HA-based stiffness cues, temporally precedes integrin-based adhesion maturation, and facilitates invasion.

Implications: This study reveals that the CD44 receptor, which is commonly overexpressed in GBM tumors, is critical for cell adhesion, invasion, and mechanosensing of an HA-based matrix. *Mol Cancer Res*; 12(10); 1416–29. ©2014 AACR.

Introduction

The prognosis of glioblastoma multiforme (GBM), a highly invasive and rapidly lethal brain tumor, has improved only incrementally in the past several decades. This dire prognosis is in large part attributed to the aggressively invasive nature of glioma cells, which, in turn, has fueled interest in exploring new strategies for slowing invasion, including identifying and limiting interactions between tumor cells and proinvasive components of the tumor extracellular matrix (ECM; ref. 1). In addition to invading brain tissue along vascular structures, GBM tumors are characterized by a diffuse single-cell invasion pattern of glioma cells into brain parenchyma (2, 3), which contains ECM characteristically devoid of the fibrillar adhesive proteins found in connective tissue, but rich in hyaluronic acid (HA; refs. 3 and 4). This

chemical signature of brain ECM is significant because an emerging body of literature has revealed that the HA/CD44 interaction can act as a powerful regulator of cell proliferation, survival, and antiapoptotic pathways (5–7). In the ECM of gliomas, HA is much more abundant than in normal brain (8), suggesting that its oversecretion may contribute to the aggressive invasion pattern of glioma cells in brain parenchyma. Finally, CD44 expression directly contributes to the survival of glioma stem-like cells hypothesized to be the driver of tumor recurrence (9).

Although changes in ECM biochemistry are key features of GBM, we and others have shown that ECM biophysical properties also strongly influence glioma cell invasion in an *in vitro* setting (10, 11). This is consistent with the observation that GBM tumors are stiffer than normal brain tissue, to the extent that ultrasound imaging can be used to delineate tumor margins intraoperatively (12, 13). This has led to the hypothesis that part of the aggressive nature of GBM may be regulated by biophysical interactions between glioma cells and the brain ECM. Matrix stiffness cues encoded in the ECM are traditionally thought to be transduced by integrins, and this signaling is altered in cells derived from a variety of cancer cell types. Although the importance of integrin-mediated signaling in these scenarios has been well characterized, the significance of non-integrin ECM adhesion receptors to

Department of Bioengineering, University of California, Berkeley, California.

Note: Supplementary data for this article are available at Molecular Cancer Research Online (<http://mcr.aacrjournals.org/>).

Corresponding Author: Sanjay Kumar, 274A Stanley Hall #1762, University of California, Berkeley, CA 94720-1762. Phone: 510-643-0787; Fax: 510-642-5835; E-mail: skumar@berkeley.edu

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tumor cell mechanobiology remains largely unexplored. The abundant presence of HA in brain ECM and the established role of CD44-mediated signaling in tumor progression beg the question of how HA/CD44 interactions contribute to glioma adhesion and invasion in ECMs composed of HA, integrin-adhesive domains, or both.

Despite the acknowledged association of GBM with altered HA deposition, CD44-based signaling, and tissue mechanics, little is known about the causal relationships between these phenomena in the pathogenesis of the disease, particularly with respect to tumor invasion. We therefore sought to investigate these connections by combining transcriptomic analysis of human GBM tumors to explore correlations in the expression of HA/CD44-related genes, engineered hydrogel materials to recapitulate the compliant, HA-rich nature of brain ECM (14–16), and biophysical studies of tumor cell adhesion, migration, and invasion. We find that GBM tumors preferentially express certain HA/CD44-related genes relative to normal brain tissue and that HA/CD44 interactions strongly contribute to tumor cell adhesion, mechanosensing, and invasive motility. These effects are both experimentally separable and functionally distinct from contributions of integrin-based adhesion.

Materials and Methods

The Cancer Genome Atlas gene expression analysis

Data from the publicly available data browser were queried for expression of *CD44* and related genes. The cBioPortal analysis tool from Memorial Sloan–Kettering was used to retrieve mRNA expression data for genes of interest from all available GBM tumor samples. Correlations with CD44 expression were analyzed by Pearson product-moment correlation coefficient.

HA hydrogel synthesis

HA hydrogels were synthesized as previously described (14, 17). Briefly, methacrylic anhydride was used to functionalize HA with methacrylate groups (Me-HA). The degree of methacrylation was characterized by ¹H NMR as detailed previously (14), and the Me-HA used for the experiments discussed here was characterized to have 50% of disaccharides methacrylated. Me-HA could then be conjugated via Michael Addition reactions with molecules containing free thiol groups. In some cases, Me-HA was conjugated with the cysteine-containing RGD peptide (Ac-GCGYGRGDSPG-NH₂, Anaspec) to add integrin-adhesive functionality at a concentration of 0.5 mmol/L. Finally, hydrogels were formed by crosslinking 5 wt% Me-HA in DMEM (Invitrogen) with varying concentrations of the bifunctional thiol dithiothreitol (DTT, Sigma-Aldrich), ranging from 2.79 mmol/L (to yield 0.15 kPa) to 22.3 mmol/L (to yield 6.9 kPa). After 1 hour crosslinking time, the hydrogels were rinsed thoroughly with PBS before cell seeding.

Rheological measurements

The shear modulus of various hydrogel formulations was measured using oscillatory rheometry as described

previously (14). Briefly, hydrogels were first crosslinked by incubation for 1 hour in a humidified 37°C chamber. Rheological testing consisted of frequency sweeps ranging from 100 to 0.1 Hz at 0.5% amplitude, also in a humidified 37°C chamber. Shear modulus was reported as the storage modulus at an oscillation frequency of 0.1 Hz.

Functionalization of HA hydrogels with full-length proteins

After crosslinking, some HA hydrogels were functionalized with adhesive proteins in a method adapted from a previous study (18). Because the hydrogels are resistant to passive protein adsorption, crosslinked hydrogels were conjugated with poly-L-lysine by carbodiimide chemistry using 0.5 M EDC (Pierce) and 0.5 M NHS (Sigma) in 0.1 M MES buffer at pH 5.8. After rinsing, a solution of 0.5% poly-L-lysine (Sigma) in PBS was added for 1.5 hours, then a 0.1 mg/mL solution of human plasma fibronectin (Millipore) or rat laminin (Life Technologies) was adsorbed for 1 hour at room temperature. Before cell seeding, the hydrogels were rinsed with PBS.

Cell culture

U373-MG and U87-MG human glioblastoma cells were obtained from the University of California, Berkeley Tissue Culture Facility, which sources its cultures directly from the ATCC. We note that ATCC U373-MG cells have been discovered to share common origins with SNB-19 and U251-MG cell lines, although these 3 lines seem to have since evolved to exhibit distinct karyotypes and drug sensitivities (19). Whatever their origins, all of these lines are understood to be fully distinct from U87-MG cells and thus serve as an appropriate culture model for comparison. Cells were cultured as previously described (10) in DMEM (Invitrogen) supplemented with 10% calf serum (JR Scientific), 1% penicillin–streptomycin, MEM nonessential amino acids, and sodium pyruvate (Invitrogen). Accutase (Innovative Cell Technologies, Inc.) was used to harvest cell cultures from tissue culture polystyrene.

Lentiviral shRNA transfection

To create CD44 knockdown cells, 2 distinct CD44-specific shRNA constructs (V2LHS_111682 and V2LHS_111684; Thermo Scientific) were screened in addition to a nontargeting scramble sequence (plasmid 1864; Addgene deposited by D. Sabatini) as a negative control. Viral particles were packaged using human embryonic kidney 293T cells, then transfected into U373-MG human glioblastoma cells with a multiplicity of infection of 1 IU/cell. Cells were selected using 1 μg/mL puromycin, and expression of the GFP-containing vectors was confirmed using flow cytometry.

Western blots

The efficiency of CD44 knockdown was confirmed using Western blots. RIPA buffer with protease and phosphatase inhibitors were used to lyse cells. Lysates were heated to 70°C, run through a 4% to 12% bis-tris gel, and then blotted onto a PVDF membrane. Membranes were probed with rat

anti-pan-CD44 primary antibody (Hermes-1; Pierce) or goat anti-RHAMM primary antibody (E-19; Santa Cruz Biotechnology), and mouse anti-GAPDH (Sigma-Aldrich). HRP-conjugated secondary antibodies against rat, mouse (Invitrogen), or goat (Zymed), and SuperSignal West Dura reagent (Thermo Scientific) were used for chemiluminescent detection. Membranes were imaged using the ChemiDoc XRS+ system (Bio-Rad), and bands were quantified using ImageLab software (Bio-Rad) and normalized to GAPDH content.

Centrifugal adhesion assay

In some cases, cells were pretreated by incubation in serum-free media with blockers of adhesion for 15 minutes, including pan-CD44-neutralizing antibody (Hermes-1; Pierce), rat IgG2a control antibody (eBioscience), and soluble RGD peptide (Ac-GCGYGRGDSPG-NH₂; Anaspec). Cells were seeded either on cross-linked HA hydrogels with or without RGD functionality. After a specified adhesion time, wells were completely filled with fresh serum-free medium, and the cell culture plate was sealed with an adhesive plate sealer. The plate was then inverted and centrifuged for 5 minutes at 100g. Cells remaining on the hydrogels were then fixed and stained with 4',6-diamidino-2-phenylindole (DAPI; Invitrogen). Automated Otsu thresholding analysis of the DAPI images were performed on ImageJ (NIH) to determine a total count of the number of cells on each hydrogel. Each condition was tested in at least 3 wells, over at least 3 experiments.

Integrin characterization assay

To characterize the integrin expression pattern of control and CD44 knockdown cells, an α/β Integrin-Mediated Cell Adhesion Array Combo Kit (ECM532; EMD Millipore) was used, in which antibodies specific for a particular integrin subunit or heterodimer are immobilized onto microtiter wells. Cells were nonenzymatically harvested by incubation in 5 mmol/L EDTA for 20 minutes, then incubated on microtiter plates for 2 hours in serum-free medium. Nonadherent cells were gently washed off, and the number of cells was quantified with a colorimetric readout per the manufacturer's instructions. Each integrin subunit or heterodimer antibody was assayed in duplicate wells, and the experiment was repeated 3 times. The specificity of each antibody was verified, and published accounts of the antibody clones (20, 21) used differed slightly from the kit description; these literature-validated antibody specificities are reported here.

Live cell time-lapse microscopy

U373-MG cells were seeded at least 4 hours before the beginning of each time-lapse experiment. Every 15 minutes, phase contrast images were obtained with a 10 \times objective. Migration speed was measured using manual tracking plugin in ImageJ. For each cell, the displacement of the nucleus for each 15 minute time interval was averaged over 6 hours to obtain the mean cell speed. At least 50 cells were analyzed per

condition, over at least three separate experiments. Persistence parameter was calculated as the end-to-end path distance divided by the total distance travelled during a 6-hour time span.

Immunofluorescence staining and cell area measurements

Mouse anti-vinculin primary antibody (Sigma) and Alexa Fluor 546 goat anti-mouse secondary antibody (Molecular Probes) were used to visualize vinculin. Rat anti-CD44 primary antibody (Hermes-1; Pierce) and Alexa Fluor 647 chicken anti-rat secondary antibody (Molecular Probes) were used to visualize CD44. Filamentous actin was stained using Alexa Fluor 488 phalloidin (Invitrogen), and nuclei were labeled with DAPI. Confocal images were obtained with a swept-field confocal microscope (Prairie Technologies). Cell area was measured by manually tracing the cell edges of live-cell phase-contrast images taken 24 hours after seeding using ImageJ software.

Transwell invasion assay

Transwell membrane inserts with 8- μ m pores, the smallest pore size we found to be permissible for cell invasion in U373-MG cells, were coated overnight with 2 mg/mL high-molecular-weight HA from *Streptococcus equi* (Sigma), 2 mg/mL low-molecular-weight HA (66-90 kDa; Lifecore Biomedical), 3.7 or 9.3 μ g/mL human plasma fibronectin (Millipore), or a combination of both HA and fibronectin. Membranes not adsorbed with fibronectin were blocked with BSA (Sigma). Membranes were then thoroughly rinsed with PBS, and cells were seeded at a density of 28,000 cells/cm² in cell medium supplemented with 1% calf serum. In some cases, soluble HA was added to the cell medium in both the upper and lower chambers at a concentration of 0.125 or 0.625 mg/mL. Phase contrast microscopy was used to visualize cells on either side of clear membranes. To visualize only cells on the underside of the membrane, which had therefore invaded through pores, GFP-expressing cells were seeded on Fluoroblok membranes and imaged with fluorescence microscopy. Cells were fixed 4 hours after seeding, and cells remaining on tops of membranes were removed by scraping with a pipette tip. To quantify cells that had invaded through the membrane, fluorescence images of DAPI-stained cells were thresholded, and the total number of nuclei in each membrane was counted with particle analysis software (ImageJ). To confirm specificity of CD44 with adsorbed HA, control cells were preincubated with CD44-neutralizing antibody and seeded on HA-coated transwells. To confirm specificity of RGD-ligating integrins with adsorbed fibronectin, cells were preincubated with soluble RGD peptide and seeded on fibronectin-coated transwells.

Statistical analysis

Analysis of percentage data for adhesion and invasion assays was preceded by an arcsin transformation. For normally distributed data, statistical significance was tested using ANOVA followed by Tukey-Kramer multiple

the mRNA level, correlation analysis of transcriptomic data from The Cancer Genome Atlas (TCGA) reveals that *CD44* is indeed frequently overexpressed in GBM tumors, with a median gene expression 4.25-fold higher than that of normal brain tissue (Fig. 1B), in agreement with many previous smaller-scale histologic studies (8, 23–26). Further analysis reveals that many other *CD44*-related genes that are also commonly overexpressed in GBM, particularly those involved in HA synthesis and degradation. The genes *HAS1*, *HAS2*, and *HAS3*, which encode the HA synthases, are aberrantly expressed. Importantly, induced *HAS2* expression has been shown to promote anchorage-independent growth and increases tumorigenicity (27). However, the level of HA ultimately found in the matrix is affected both by rates of HA synthesis and several post-transcriptional levels of regulation, so the median decrease in HA synthase transcription is not necessarily inconsistent with the elevated levels of HA in the tumor micro-environment. For instance, the expression of hyaluronidases or hyals, which are responsible for HA matrix catabolism, are also misregulated, with hyaluronidase-1 and hyaluronidase-3 generally poorly expressed relative to normal brain tissue. On the other hand, hyaluronidase-2 is expressed at high levels during development but at low levels in the adult brain (28), and its induced overexpression promotes tumor invasion in a mouse model (29).

In addition to proteins that regulate turnover of the HA network, other proteins that physically crosslink linear HA chains together also exhibit altered transcription patterns (Fig. 1A). Among these include the family of lecticans and tenascins, which, with HA, form the basic structural components of brain ECM (30). The expression of genes that encode proteins key to *CD44*-mediated intracellular signaling are also misregulated. A notable example is the ERM (ezrin, radixin, moesin) family of proteins, which mediates interactions between the cytoplasmic domain of *CD44* and the actin cytoskeleton, and have known pro-oncogenic function (31). All 3 ERM proteins are overexpressed in tumor tissue, with expression of *MSN* correlating strongly and positively with *CD44* expression. Interestingly, although receptor for HA-mediated motility (RHAMM) is generally described as a proinvasive HA receptor and has been found to be overexpressed in smaller GBM studies (32), TCGA analysis revealed that the median mRNA expression of the encoding gene *HMMR* in tumors is only 0.53-fold the expression level of normal brain tissues. This suggests that posttranscriptional regulation may have significant effects on *HMMR*, or that RHAMM overexpression may be a feature of a specific subset of GBM tumors. Finally, since HA functions as a key organizational scaffold for brain ECM proteins, we examined the gene expression of other ECM proteins (Fig. 1C). Indeed, GBM tumors overexpress several proteins that ligate integrins through RGD peptides, such as fibronectin. Together, these findings motivated us to more deeply investigate contributions of *CD44* and integrin binding to GBM tumor cell adhesion, migration, and invasion.

CD44-mediated adhesion is important even when RGD is abundant

Given the wealth of data supporting the importance of integrins in glioma cell–ECM adhesion, we first compared the biophysical contributions of *CD44* and integrin engagement to total cell–ECM adhesion. We initially chose to work with HA hydrogels with stiffnesses similar to that of brain tissue, which we fabricated according to our previous description (14). Using a centrifugal detachment assay, we measured cell adhesion strength to bare HA hydrogels and HA hydrogels functionalized with RGD peptide, both with a shear modulus of 4.6 kPa. Our use of the RGD sequence was motivated by the fact that this is the key integrin-binding sequence in fibronectin, which is enriched in the ECM of GBM tumors when compared with normal brain tissue. RGD is also ubiquitously used within synthetic biomaterials as a modular integrin-binding sequence (33–35), which allowed us to more readily compare our findings with work in these other materials systems. We seeded U373-MG human glioblastoma cells transduced virally with either a *CD44*-targeting shRNA (yielding 64% protein expression knockdown; Supplementary Fig. S1A and S1B) or a non-targeting control sequence, and allowed the cells to adhere for 0.5 or 3 hours in serum-free culture medium. We then measured adhesion strength by quantifying the percentage of cells that remained adherent after exerting a detachment force of 100 g to remove weakly adhered cells.

When cells were centrifuged 0.5 hours after initial seeding, 21% of control cells remained adherent to the bare HA hydrogel (Fig. 2A). Incorporation of an RGD peptide into the hydrogel yielded a modest but statistically significant increase in cell adhesion. When the same surfaces were presented to the *CD44* knockdown cells, bare HA hydrogels yielded negligible adhesion, whereas adhesion to HA-RGD hydrogels was reduced twofold relative to control cells. To confirm the molecular specificity of this effect with an alternative approach, we incubated control cells with a *CD44*-neutralizing antibody, which yielded qualitatively identical results. We also confirmed that *CD44* knockdown altered neither integrin subtype expression nor cell viability (Supplementary Fig. S2). Notably, and as reported previously (36), *CD44* suppression increased expression of the HA receptor RHAMM (Supplementary Fig. S1C and S1D), indicating that the effects of depleting *CD44* could not be fully rescued by enhancement of other HA adhesion proteins.

As expected, preincubation of both cell types with an isotype-matched IgG antibody led to similar attachment as the untreated control condition. Pretreatment of cells with a soluble RGD peptide reduced adhesion to HA-RGD but not to bare HA hydrogels, confirming that the cells adhered to the cross-linked HA through an RGD receptor-independent mechanism. When we repeated this assay for a longer adhesion time of 3 hours, the only conditions that resulted in increased adhesion were those with both *CD44*- and integrin-based adhesions (Fig. 2B). Together, these data suggest that *CD44* is largely responsible for adhesion on early time scales (0.5 hours), and that these initial adhesions

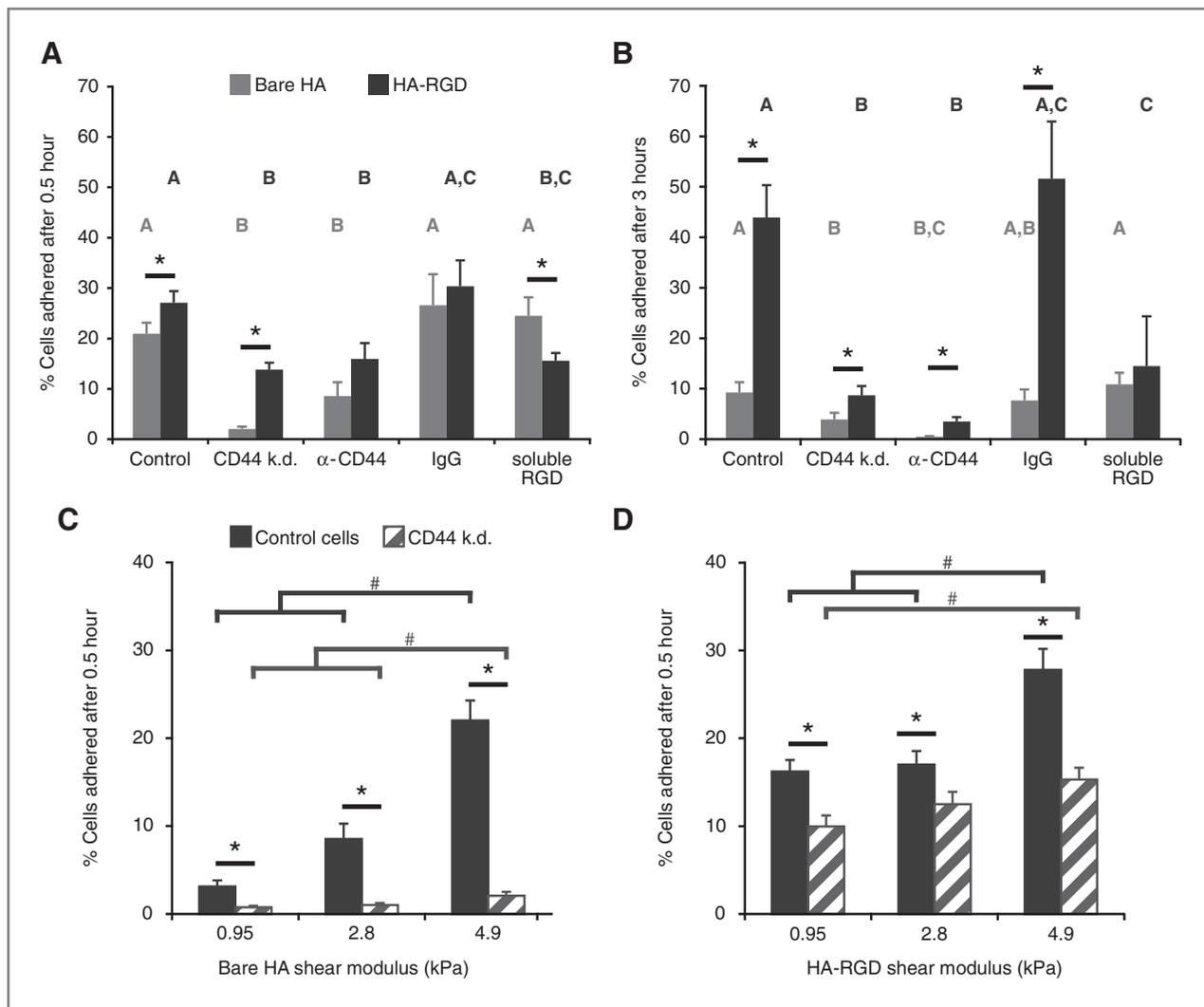


Figure 2. U373-MG cell adhesion to HA hydrogels is CD44 and stiffness dependent. A, adhesion of cells on 4.6 kPa hydrogels at short (0.5 hours) time scales. Control cells or CD44 knockdown (k.d.) cells were allowed to attach to substrates in the absence or presence of adhesion blockers, including α -CD44 neutralizing antibody, IgG isotype control antibody, or soluble RGD peptide; then centrifuged to induce detachment of weakly adhered cells. B, adhesion of cells on 4.6 kPa hydrogels at longer (3 hours) time scales. C, adhesion of cells on bare HA hydrogels as a function of stiffness after 0.5-hour adhesion time. D, adhesion of cells on RGD-functionalized HA hydrogels as a function of stiffness after 0.5-hour adhesion time. #, $P < 0.05$ differences between stiffnesses by Tukey-Kramer; *, $P < 0.05$ by Student *t* test.

are later reinforced by integrin-mediated adhesions, if available, on longer time scales (3 hours).

CD44-mediated adhesion, spreading, and motility is dependent on HA stiffness

It is well established that integrin-based adhesion and signaling are strongly sensitive to ECM stiffness (37–40). Given that GBM invasion is accompanied by HA deposition (8) and tissue stiffening (12), and that glioma invasion requires the ability to sense and transduce mechanical force, we wondered whether this integrin-independent HA/CD44-mediated adhesion might also be stiffness sensitive. We therefore systematically varied the stiffness of our bare HA hydrogels while keeping HA concentration constant at 5 wt%, and used the same centrifugation assay to ask whether

HA matrix stiffness altered cell adhesion (Fig. 2C and D). Our studies revealed that on bare HA hydrogels, adhesion of CD44-positive cells increased with stiffness, whereas adhesion of CD44 knockdown cells, while still sensitive to stiffness, was vastly reduced (Fig. 3D). Thus, CD44–HA adhesion is intrinsically mechanosensitive to matrix stiffness in a manner that does not require integrin ligation. Incorporation of RGD into the hydrogels increased adhesion at all stiffnesses for both cell types, but still resulted in a stiffness-dependent trend. Even with integrin ligation present, CD44 knockdown reduced cell adhesion at all stiffnesses tested (Fig. 2D).

To determine whether this CD44-mediated mechanosensing had functional consequences for cell behaviors relevant to tumor invasion, we next explored the relative roles of

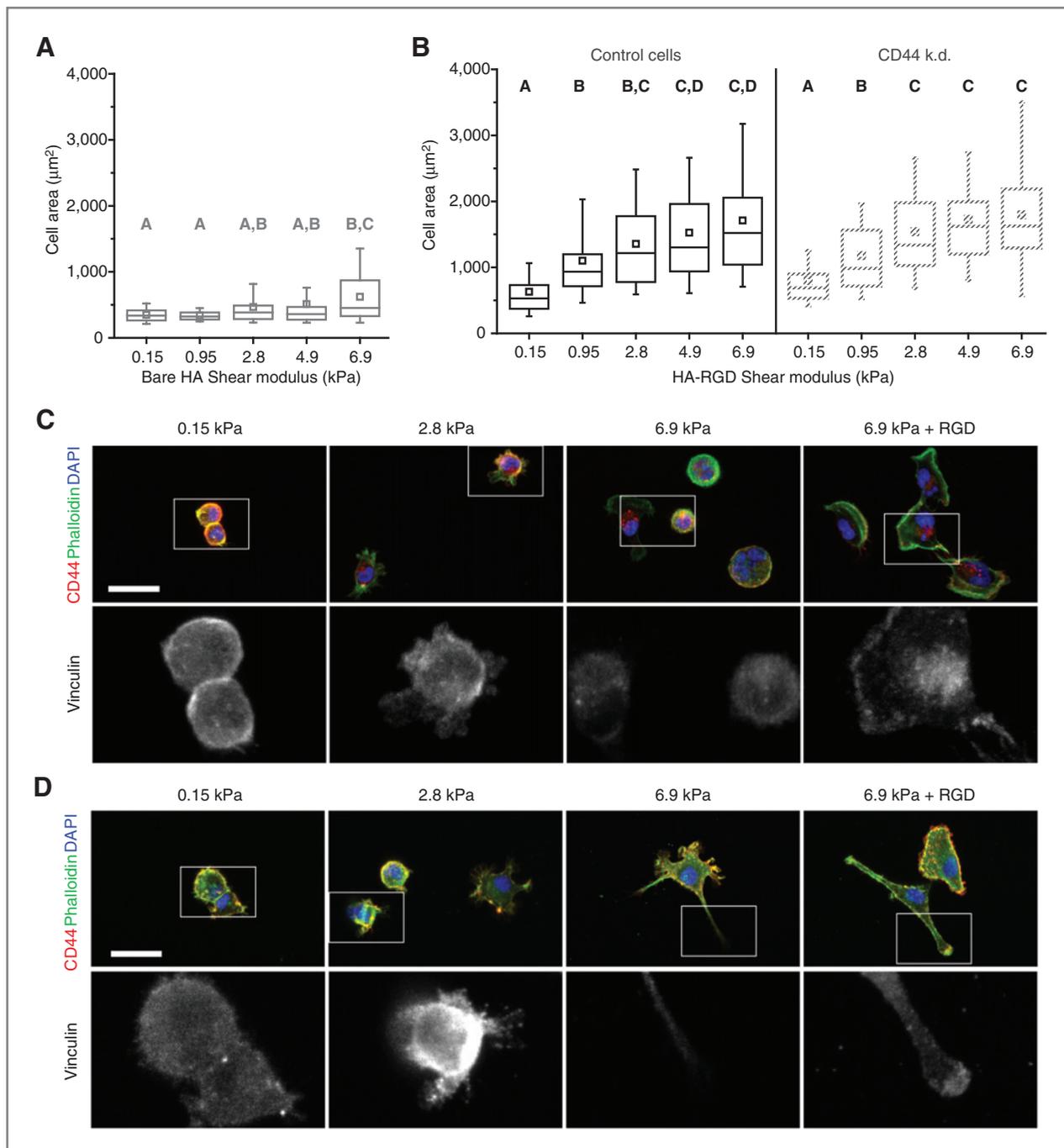


Figure 3. U373-MG glioma cell spreading and morphology on HA-based hydrogel is stiffness dependent. A and B, cell spread area after 24 hours of adhesion on bare HA hydrogels (A), or HA hydrogels functionalized with RGD (B). Hydrogel shear modulus is displayed on the x-axis. A, B, and C statistical families show $P < 0.05$ from Dunn test for multiple comparison of nonnormally distributed data. Boxes represent 25th and 75th percentiles, whiskers represent 10th and 90th percentiles. C, cell morphology of U373-MG human glioblastoma cells on HA hydrogel in the absence of integrin-based binding differs from cells spread on HA functionalized with RGD. Colors in top row represent localization of CD44 (red), F-actin (phalloidin, green), and nuclear DNA (DAPI, blue). Bottom row depicts localization of vinculin. Scale bar, 50 μm . D, cell morphology of U87-MG human glioblastoma cells with same conditions described in C.

CD44 and integrins in stiffness-dependent spreading and motility on HA-based hydrogels. When allowed to adhere over a period of 24 hours in medium containing 10% serum,

control U373-MG cells exhibited differing morphologies depending on stiffness and presence of RGD (Fig. 3). Cell area analysis performed on images obtained 24 hours after

seeding shows that, consistent with our past observations (14), HA hydrogels functionalized with RGD induced stiffness-dependent spreading (Fig. 3B). Surprisingly, cells seeded on HA substrates devoid of RGD still exhibited stiffness-dependent spreading, although the stiffness dependence was less steep than that of cells on HA-RGD hydrogels (Fig. 3A). On compliant bare HA hydrogels of 0.95 kPa, all cells remained rounded and formed sparse multicellular aggregates, as expected for a low-adhesion environment (41). On stiffer 4.9 and 6.9 kPa bare HA hydrogels, cells predominantly adhered as single cells, with CD44-positive processes and without vinculin-positive adhesive plaques. Cell spreading was dramatically reduced compared with cells on RGD-functionalized surfaces (Fig. 3C). In contrast, cells on 6.9 kPa HA-RGD hydrogels exhibited stress fibers, which were absent in bare HA hydrogels of all stiffnesses. Similar results were seen with U87-MG human glioblastoma cells, albeit with expected variations in morphology between cell lines (Fig. 3D). In both cases, addition of RGD to the substrate resulted in localization of vinculin staining to the tips of protrusions, whereas vinculin staining remained diffuse on bare HA hydrogels.

Consistent with the CD44 specificity of this adhesion, cell spreading measurements on bare HA hydrogels could not be obtained for CD44 knockdown cells because of insufficient adhesion to the hydrogel. With the addition of RGD to the matrix, however, these cells exhibited similar spreading as control cells on the same matrix (Fig. 3B). This implies that over timescales of roughly 24 hours and with the addition of 10% serum to the culture medium, integrin-based adhesions are able to compensate for the lack of CD44 adhesions on 2-dimensional (2D) cell spreading and motility.

To gain further insight into CD44-dependent cell motility, we used time lapse microscopy to record cell migration on HA and HA-RGD matrices 12 to 18 hours after seeding. Cells were able to productively migrate at comparable speeds with or without RGD functionalization. The speed of this migration was sensitive to HA matrix stiffness, both in the absence and presence of RGD peptide (Movie 1; Fig. 4A). Interestingly, the speed–stiffness relationships for the 2 materials crossed over, such that bare HA hydrogels supported faster migration for the lowest stiffnesses (<1 kPa), whereas HA-RGD hydrogels supported faster migration for the higher stiffnesses (>4 kPa). On all but the softest substrates tested, cells on bare HA surfaces at times exhibited protrusions that permitted productive motility, but were generally shorter lived than the larger lamellipodial structures seen on cells with RGD-integrin binding (Fig. 4D). Bare HA substrates supported less directionally persistent migration compared with hydrogels with RGD at all stiffnesses (Fig. 4B). Thus, despite the lack of classically "spread" 2D morphology, glioma cells are still able to productively migrate on 2D bare HA hydrogels using only CD44-based adhesion and without integrin-based adhesion. As with spread area and morphology, CD44 knockdown cells exhibited RGD concentration-dependent migration equivalent to that of control cells (Fig. 4C). In other words, knockdown of CD44 does not affect cell spreading or motility when

abundant integrin-based signaling is available. To demonstrate that these results generally apply to a range of integrin ligands and not just RGD peptides, we functionalized HA hydrogels with full-length fibronectin or laminin and obtained similar cell spread area and cell speed dependence on hydrogel stiffness (Supplementary Fig. S3).

CD44-mediated adhesion promotes glioma invasiveness

The above studies reveal that CD44 can mediate mechanotransductive signaling and support matrix adhesion and 2D motility without giving rise to mature, vinculin-positive focal adhesions. To test whether this mode of adhesion might also support the highly aggressive invasion of glioma cells through the narrow interstices in brain parenchyma, we used a 3D transwell invasion model.

To confirm that adsorbed HA and fibronectin would still engage CD44 and integrins in a manner similar to that of the cross-linked HA hydrogels (as seen in Fig. 2), we first performed a series of adhesion control experiments with full-length fibronectin, which revealed similar ligand density-dependent spreading behavior as with RGD (Supplementary Fig. S3). To confirm that CD44- and integrin-specific adhesions are capable of inducing transwell invasion, we next performed receptor-blocking experiments (Fig. 5A and B). First, control cells were preincubated with CD44-neutralizing antibody in suspension, then seeded on transwell membranes coated with HA and blocked with BSA. The resulting drop in cell invasion compared with cells not blocked with CD44-neutralizing antibody indicated that an appreciable component of the adhesion to adsorbed HA could be attributed to CD44 (Fig. 5A). Second, CD44 knockdown cells were preincubated with soluble RGD peptide, and the resulting abrogation of cell invasion on fibronectin-coated surfaces compared with unblocked cells indicated that the same receptors, presumably integrins, engage both RGD and fibronectin (Fig. 5B).

To test how CD44 and integrin ligation contribute to cell invasion when both cognate matrix elements were present, we then coated transwell membrane inserts containing 8- μ m pores with HA, various concentrations of fibronectin, or both HA and fibronectin. The transwell invasion model has previously been shown to be predictive of glioma cell invasion through tissue (42). Membranes coated only with HA were also blocked with BSA, and membranes coated only with BSA were used as a negative control to measure baseline invasion with minimal receptor-mediated adhesion. Interestingly, CD44-expressing control cells were most invasive across membranes coated with HA with BSA, but not fibronectin (Fig. 5C). This phenomenon is CD44-specific, because the invasive motility of CD44 knockdown cells across these same membranes was significantly lower and indistinguishable from results obtained with BSA-coated membranes. Moreover, inclusion of fibronectin on HA-coated membranes reduced invasive motility in a concentration-dependent fashion; adsorption from a solution of 9.3 μ g/mL fibronectin with HA reduced invasion of control cells 2.2-fold relative to HA-coated membranes, whereas 3.7 μ g/mL fibronectin with HA yielded intermediate levels of

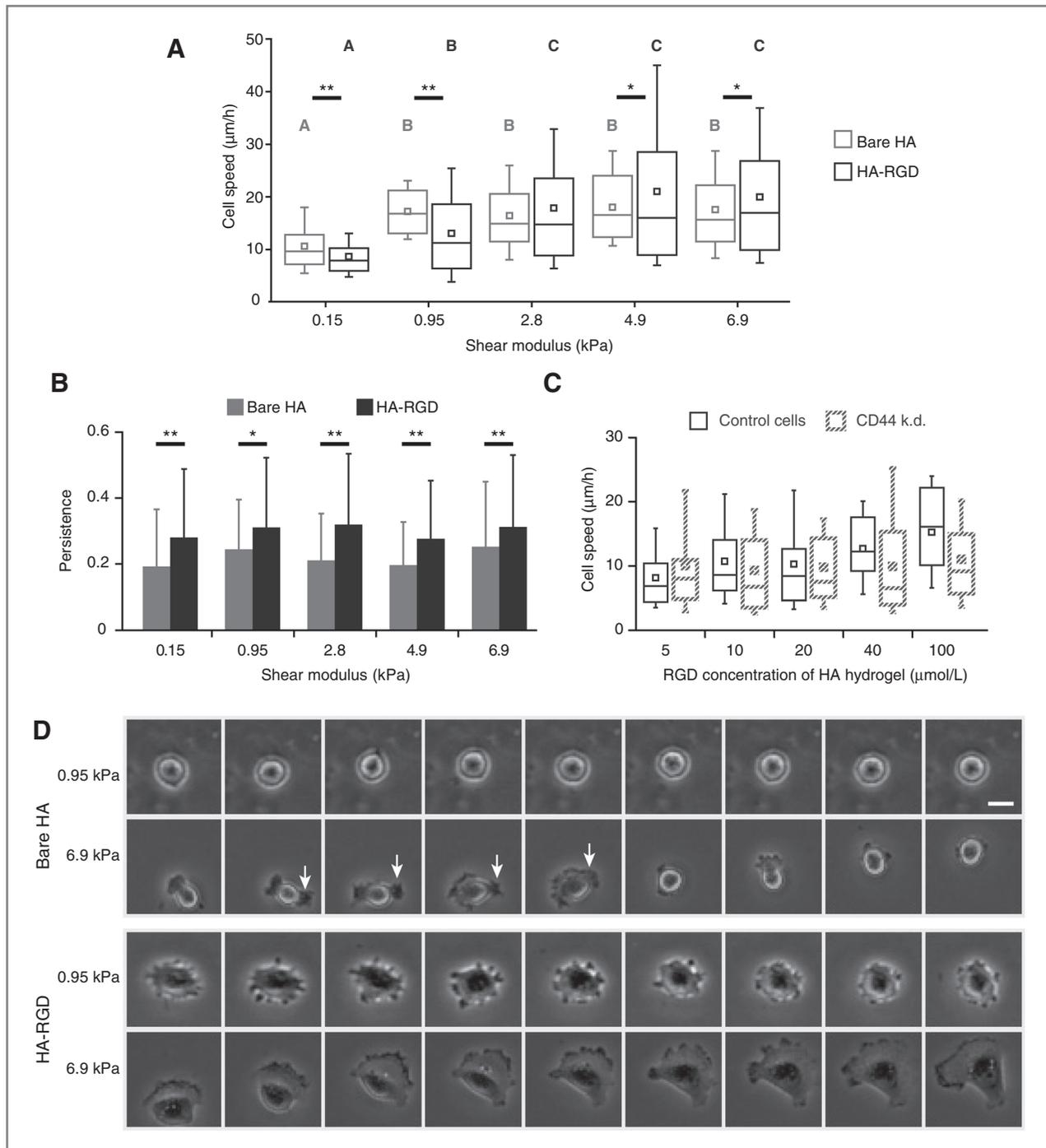


Figure 4. U373-MG glioma cell motility on HA-based hydrogel is CD44- and stiffness-dependent. **A**, cell migration speed 12 to 18 hours after seeding on bare HA hydrogels and on RGD-functionalized HA hydrogels. Hydrogel shear modulus is displayed on the x-axis. A, B, and C statistical families show $P < 0.05$ Dunn test for multiple comparison of nonnormally distributed data. Boxes represent 25th and 75th percentiles, whiskers represent 10th and 90th percentiles. For pairwise comparisons between cell speeds on HA and HA-RGD hydrogels at a given shear modulus, *, $P < 0.1$, **, $P < 0.05$ by Student *t* test. **B**, comparison of cell migration speed of control U373-MG cells with CD44 knockdown counterparts on HA hydrogels of varying RGD concentration. Comparison could not be made on bare HA hydrogels because of insufficient adhesion of CD44 knockdown cells in the absence of RGD functionalization. With HA functionalization of 1 $\mu\text{mol/L}$ RGD, very few CD44 knockdown cells attached. At all other RGD concentrations tested, no statistical difference in the migration speed between the 2 cell lines was found. Under all conditions tested, differences between control and CD44 knockdown cell speeds were not statistically significant. **C**, persistence parameter of random cell motility paths on bare HA- and RGD-functionalized hydrogels. *, $P < 0.1$, **, $P < 0.05$ by Student *t* test. **D**, still images taken from time lapse microscopy. Each column represents a 0.5-hour interval. On bare HA of 0.95 kPa, cells consistently remain rounded. On bare HA of 6.9 kPa, cells extend small protrusions that rupture quickly, such as the one indicated by white arrows, which lead to productive migration. In contrast, cells on stiff HA-RGD hydrogels exhibit broad, stable lamellipodia. Scale bar, 25 μm .

invasion. To compare the effect of soluble HA to adsorbed HA, we added soluble HA in the media of both the top and bottom chambers of the transwell membrane. This did not lead to a significant increase in cell invasion compared with the BSA control.

To gain mechanistic insight into these results, we obtained both phase contrast images of cells adhered to either side of each membrane and fluorescence images of GFP-expressing cells that had invaded through the membrane to the undersides of Fluoroblok membranes (Fig. 5D). Consistent with adhesion results on HA and HA-RGD gels (Fig. 2A), control cells attached readily to membranes coated with HA and/or fibronectin. However, control cells invaded membranes coated in HA much more than membranes coated with either fibronectin alone or both HA and fibronectin. This implies that fibronectin enhanced ECM engagement and cell spreading, but hindered invasion. When neither type of adhesion was available (CD44 knockdown cells on HA + BSA matrix, or any cell population on BSA matrix), cells rarely attached or spread, which also precluded invasion. Only in an intermediate range of adhesion strength, in which HA-CD44 adhesion was available but integrin adhesion was not, did invasion proceed appreciably higher than baseline levels.

Discussion

Given the biochemical and biophysical changes in brain tissue associated with GBM progression, we were motivated to investigate how glioma invasion is affected by the nature of cell-matrix adhesions and the mechanical properties of the ECM. In addition to exposing previously underappreciated correlations between expression of CD44, pro-oncogenic signals, and HA-synthesizing proteins, we have also found evidence that CD44-HA adhesion is intrinsically mechanosensitive and occurs on a faster time scale than integrin-based binding. Most importantly, we find that although both adhesion systems support robust 2D motility, CD44-based adhesion is accompanied by extension of small, short-lived processes, whereas integrin-based adhesion is associated with broad lamellipodia and more directionally persistent migration. These differences extend to 3D migration through constricted pores, as CD44-HA adhesion drives optimal invasion in a transwell paradigm.

To explain our transwell results, we assembled a model based on previously described models of glioma invasion through constricted spaces (42) and adhesion strength on cell migration (43). Based on these and our observations, we propose a model in which optimal cell invasion requires a balance between cell-matrix adhesion and turnover (Fig. 6). In other words, we postulate that although CD44 alone can support cell adhesion, these adhesions are less mechanically reinforced than those formed by integrins. Consistent with this idea, integrin ligation is accompanied by formation of actomyosin bundles and focal adhesions (Fig. 3C and D), which slows migration through narrow pores. This model is supported by our measurements of adhesion, spreading, and

migration on 2D bare HA surfaces, where the finding that CD44 can support adhesion without lamellipodial spreading suggests that the latter is not required for robust adhesion or migration. Thus, invasion through 3D pores is not directly predictable from 2D motility alone, which is broadly consistent with the recent observation that 2D protrusive propensity predicts 3D migration speed much more accurately than does 2D migration speed (44). Therefore, previously described causal relationships between CD44 expression, tumor size, and survival time *in vivo* (7) may not only be due to CD44-mediated stimulation of canonical pro-oncogenic signaling, but also through promotion of glioma invasion by biophysical mechanisms.

Using transwell assays and engineered HA hydrogels, we have demonstrated that in a range of timescales from 0.5 to 24 hours, CD44 is capable of supporting glioma cell adhesion in the absence of integrin engagement. Ultimately, our results indicate that CD44 may serve a directly proinvasive function through the formation of short-lived matrix adhesions. Previous studies support the first observation that HA promotes glioma cell invasion, either by addition of HA into a Matrigel coating (45), addition of soluble HA in the cell culture medium (46), or overexpression of the HA synthases HAS1 (47) or HAS2 (48). Consistent with these earlier studies, we have directly shown that adsorption of HA onto transwell membranes strongly enhances invasion in a CD44-dependent manner. Although integrins have been extensively demonstrated to contribute to GBM tumor growth and invasion (49, 50), our data suggest a complex interplay between integrins and CD44 that remains to be fully elucidated.

Remarkably, glioma cell adhesion, spreading, and 2D motility are all sensitive to HA stiffness via CD44. We note that cell spreading is less sensitive to stiffness when CD44, rather than integrins, mediate adhesion; whether this is because of differences in the receptors themselves, the signals they transduce, or integrin-CD44 interactions remains to be investigated. This CD44-mediated mechanosensing has functional consequences for cell behaviors relevant to tumor invasion. It has long been appreciated that the molecular weight of HA can vary widely from oligomers of 4 disaccharides to macromolecules composed of up to 30,000 disaccharides, and these can have strikingly different effects on the activation of downstream signals (51). Full-length HA is anti-angiogenic, whereas oligosaccharides of HA trigger angiogenesis (52) by a variety of mechanisms, including activation of PKC α and Src (53). However, the biophysical basis of HA-molecular-weight effects remains unexplored. In particular, the recent discovery that exceptionally high-molecular-weight HA strongly promotes cancer resistance and longevity in the naked mole rat (54) adds further relevance to our observations. Our data suggest a mechanism in which cells are able to differentiate HA molecular weight in part by detecting the mechanical rigidity of CD44-bound ligands. To date, one previous study has provided evidence that CD44 may play a role in sensing these rigidity differences in fibroblast cells (55), and another recent study indicates that CD44 can support or alter integrin-mediated

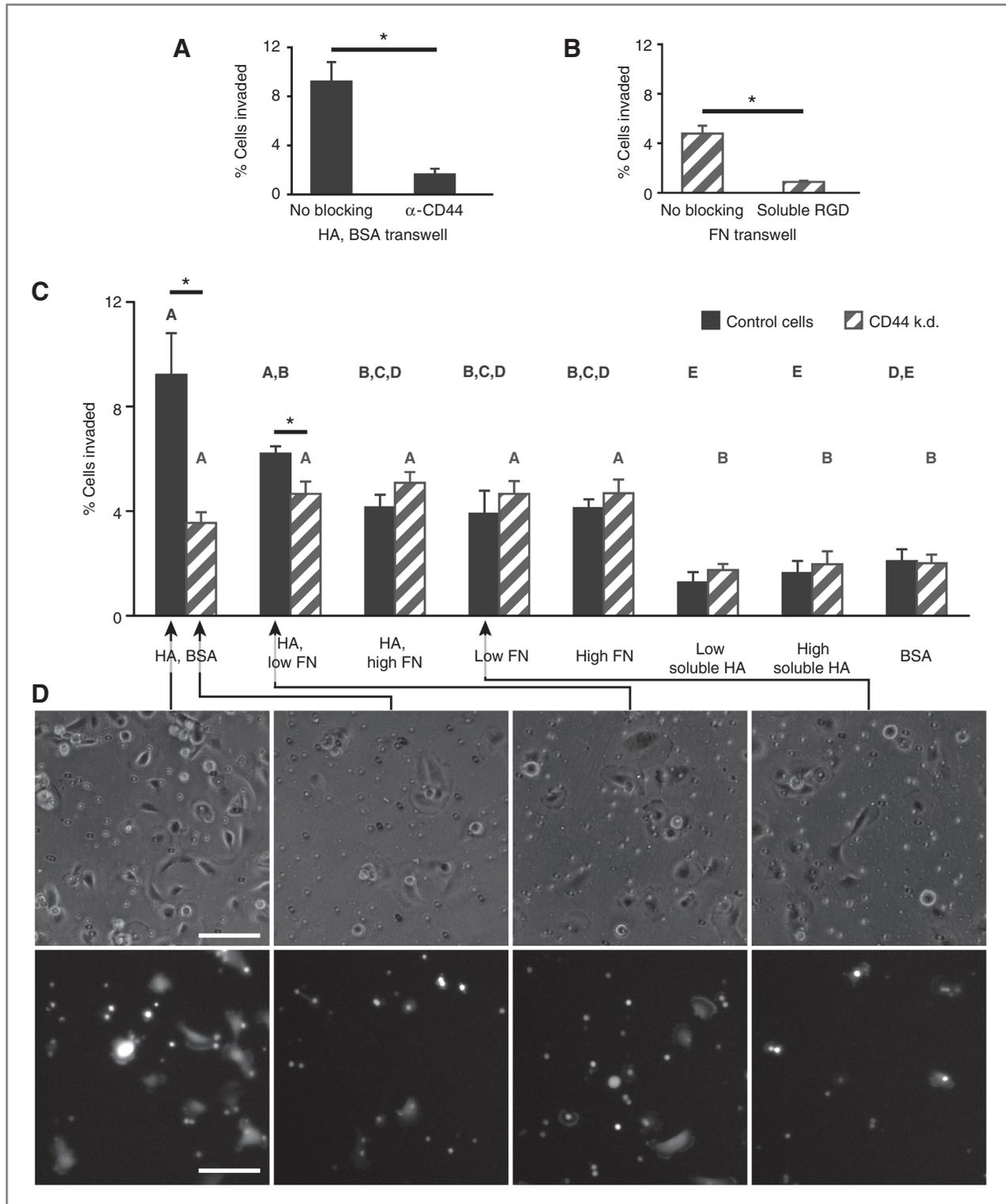
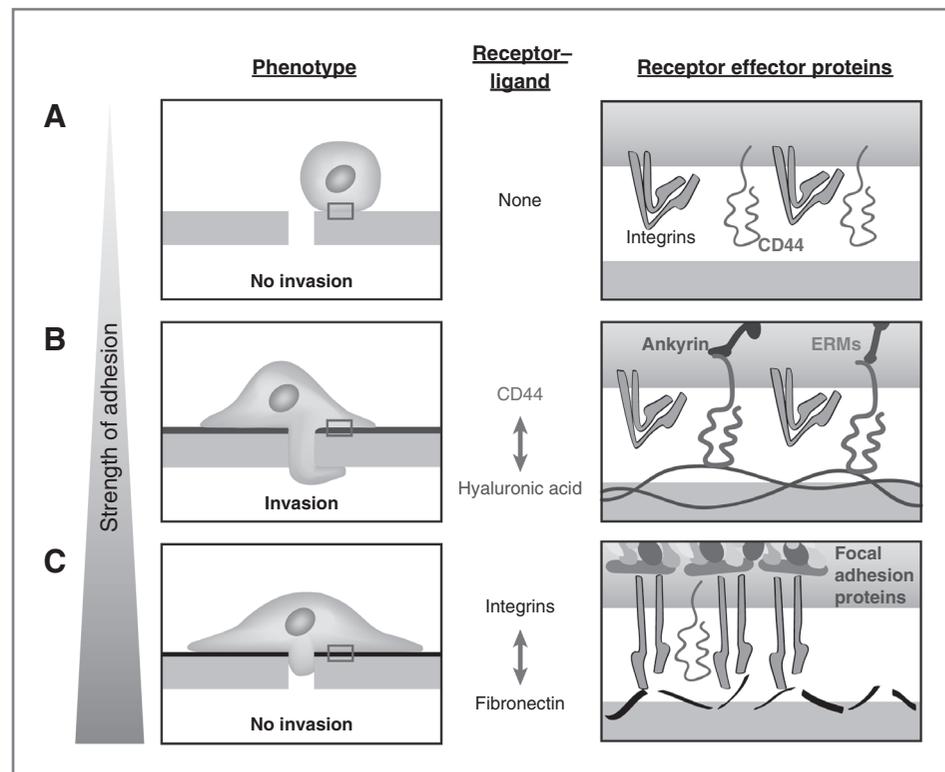


Figure 5. U373-MG glioma cell invasion through transwell membranes is promoted by HA-CD44 adhesion, but not by fibronectin-integrin adhesion. **A**, incubating control cells plated on HA + BSA coated transwells with CD44-neutralizing antibody drastically reduces invasion, demonstrating that interaction with HA is CD44 dependent. **B**, incubating CD44 knockdown cells with soluble RGD peptide on membranes coated with 3.7 μ g/mL fibronectin strongly attenuates invasion, demonstrating that adhesion to fibronectin is integrin dependent. #, $P < 0.05$ multiple comparison by Tukey-Kramer; *, $P < 0.05$ by Student t test. (Continued on the following page.)

Figure 6. Proposed model for influence of adhesion strength on cell invasion. We propose a model for cell invasion that involves a balance between formation and turnover of cell adhesions. A, with little to no cell adhesion to the substrate, extrusion of the cell body through a confined space is not possible. B, with an optimal balance between adhesion and detachment, as is the case with the relatively labile CD44–HA interaction, cells have the highest invasion potential. C, when abundant integrin-based adhesion is present, the cells spread extensively and adhere robustly on the substrate but do not invade.



mechanical signals in cardiac myocytes (56). In GBM, the extreme overexpression of *HYAL-2* (Fig. 1A), typically seen only in developing brain, suggests that tumor cells may resurrect the promigratory and angiogenic processes associated with development (28) by generating low molecular weight HA fragments.

An important innovation of our study is the use of solid-state HA substrates, which are a more physiologically mimetic form of the high-molecular weight HA that composes much of brain ECM. This also added another level of control by enabling us to vary HA stiffness. HA must be chemically methacrylated before crosslinking, which raises the potential concern that this functionalization may alter the binding affinity of HA for CD44. However, our adhesion assay results (Fig. 2A and B) demonstrate that HA methacrylation preserves CD44 binding function. Furthermore, the main binding site of CD44 for HA does not include the hydroxyl group of the N-acetyl glucosamine monomer, which is most likely to become methacrylated, and this hydroxyl group points away from the N-acetyl group that is essential for CD44 binding (57). Ultimately, additional layers of complexity are needed to capture key

aspects of perivascular invasion, in which GBM cells infiltrate tissue by following vascular structures.

According to the "motor-clutch" model for cellular mechanosensing (58, 59), the comparatively weak CD44–HA bond would shift stiffness-sensitivity to lower stiffness values than the stronger integrin-based adhesions. This is supported by previously reported binding affinities for CD44 to HA₁₀, 50 μmol/L (57), which are much lower than that of activated α₅β₁ for fibronectin, reported as 90 nmol/L (60). Given that brain matrix is a relatively compliant tissue that presumably limits cell-induced tissue stresses, it stands to reason that under these conditions, the weaker CD44–HA adhesion pair might serve as an important mechanotransducer. Indeed, our 2D motility data agree with this model; addition of RGD to the HA matrix expands the range over which cell speed is sensitive to matrix stiffness, whereas cell speed on bare HA hydrogels plateaus at lower stiffness values. Finally, our data also show that there is a time-dependent cooperation between the 2 adhesion receptors, in which CD44–HA adhesions form relatively quickly within 0.5 hours, and integrin–RGD adhesions take longer to mature. If CD44 is not available, then over a period of 0.5 to 3 hours,

(Continued.) C, percent of control or CD44 knockdown U373-MG glioma cells that invaded through 8-μm pores in membranes coated with combinations of HA, varying concentrations of fibronectin (FN) and BSA, or with HA dissolved in the cell media (soluble HA). Statistically significant differences of $P < 0.05$ from multiple comparisons within cell line groups by Tukey–Kramer are grouped by A, B, and C statistical families. *, $P < 0.05$ *t* test pairwise comparisons of cell lines on the same substrate. D, representative phase contrast images of cells either above or below clear transwell membranes (top row), and GFP-expressing cells below Fluoroblok membranes (bottom row). Both types of membranes have 8-μm pores, and images were taken 2.5 hours after cell seeding. Scale bars, 100 μm.

adhesion is drastically decreased if serum is not present, but over longer timescales of 24 hours in the presence of serum, integrin-based adhesion is able to compensate. This lag time between the maturation of the 2 types of adhesions may be because of the large ensemble of proteins that are required to form mature focal adhesions, whereas CD44 adhesions require the association of relatively few components. The initiation of adhesion by cell membrane-bound HA to substrates temporally followed by integrin-based interaction has previously been observed with chondrocytes on protein-coated glass (61), but our results demonstrate definitively that CD44 is responsible for this early adhesion, and that the ECM can be engineered to modulate these interactions.

Our study raises a number of new questions in both cellular mechanobiology and the pathophysiology of GBM: What is the molecular basis of CD44-based mechanosensing, and to what extent do these mechanisms crosstalk with integrin-based mechanosensing? How do these CD44-based mechanotransductive signals ultimately influence gene expression? And, finally, can interruption of these mechanotransductive signals limit tumor growth and invasion *in vivo*? Orthogonal control of these 2 adhesive systems with the use of engineered materials, systems biological tools, and animal models will help clarify these important issues, and potentially yield unexpected new insights into the biophysical basis of tumor invasion.

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Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

Authors' Contributions

Conception and design: Y. Kim, S. Kumar

Development of methodology: Y. Kim, S. Kumar

Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): Y. Kim

Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): Y. Kim, S. Kumar

Writing, review, and/or revision of the manuscript: Y. Kim, S. Kumar

Study supervision: S. Kumar

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