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Exosome-mediated Transfer of $\alpha_v\beta_3$ Integrin from Tumorigenic to Non-Tumorigenic Cells Promotes a Migratory Phenotype

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Abstract

The $\alpha_v\beta_3$ integrin is known to be highly up-regulated during cancer progression and promotes a migratory and metastatic phenotype in many types of tumors. We hypothesized that the $\alpha_v\beta_3$ integrin is transferred through exosomes and, upon transfer, has the ability to support functional aberrations in recipient cells. Here, for the first time, it is demonstrated that $\alpha_v\beta_3$ is present in exosomes released from metastatic PC3 and CWR22Pc prostate cancer cells. Exosomal $\alpha_v\beta_3$ is transferred as a protein from donor to non-tumorigenic and tumorigenic cells since the β_3 protein or mRNA levels remain unaffected upon transcription and translation inhibition in recipient cells. Furthermore, it is shown that upon exosome uptake, *de novo* expression of $\alpha_v\beta_3$ increases adhesion and migration of recipient cells on $\alpha_v\beta_3$ ligand, vitronectin. To evaluate the relevance of these findings, exosomes were purified from the blood of TRAMP mice carrying tumors where the expression of $\alpha_v\beta_3$ is found higher than in exosomes from wild-type mice. In addition, it is demonstrated that $\alpha_v\beta_3$ is co-expressed with synaptophysin, a biomarker for aggressive neuroendocrine prostate cancer.

Implications: Overall this study reveals that the $\alpha_v\beta_3$ integrin is transferred from tumorigenic to non-tumorigenic and cancer cells via exosomes, and its *de novo* expression in recipient cells promotes cell migration on its ligand. The increased expression of $\alpha_v\beta_3$ in exosomes from mice bearing tumors points to its clinical relevance and potential use as a biomarker.

Introduction

Prostate cancer is the second leading cause of cancer related deaths among men in the United States. According to the NCI, there will be 180,890 estimated new cases and 26,120 estimated deaths in 2016. Although, there have been continuous advances in the understanding of diagnosis and treatment of locally advanced and metastatic prostate cancer, the current therapies only provide partial disease stabilization. Androgen receptor (AR) targeted therapies are the standard of care for prostate cancer (1) but androgen-independent or AR-negative forms of cancer such as the neuroendocrine (NE) cancer are very challenging to treat (2-4).

Integrins are transmembrane receptors that comprise an α and a β subunit, known to be deregulated as prostate cancer progresses to advanced stages (5, 6). Compelling evidence indicates that signals originating from integrin ligand binding orchestrate key mechanisms of tumor progression, including cell survival, adhesion, proliferation, gene expression and modulation of the migratory/invasive phenotypes (5, 7-9). There is a strong interplay between integrins, extracellular cell matrix (ECM), cancer cells and their microenvironment as prostate cancer progresses towards a metastatic stage with bone being one of the major sites for metastasis (5, 10). Prostate tumors are highly heterogenous (11) making it challenging to generate novel treatment strategies and effective drugs. The $\alpha_v\beta_3$ integrin is present at very low levels in normal human prostate but is highly up-regulated in primary cultures of epithelial cells from human prostate adenocarcinoma and promotes adhesion and invasion of cancer cells to ECM proteins such as vitronectin (VN) (12, 13). The $\alpha_v\beta_3$ integrin promotes tumor growth within the bone, plays a role in the formation of osteoblastic lesions mediated by prostate cancer cells (14) and is often overexpressed in melanoma and metastatic colorectal cancer (15). Given its wide spread distribution in advanced cancer, many therapeutic approaches have been used to target $\alpha_v\beta_3$ such as inhibitory monoclonal antibodies (mAbs) and small

molecules (16, 17).

Exosomes (Exo) are small (30-150 nm) extracellular vesicles (EV) found in a number of biological fluids like blood, urine, amniotic fluid and cell culture media (18, 19). There has been debate about the nomenclature for EVs and recently, a new term “small EVs” has been introduced by Kowal et al (20) for EVs that are isolated by high-speed ultracentrifugation methods. However, the term Exo will be used for this study until further analysis is performed in our laboratory. Exo are enriched in tetraspanins such as CD63 and CD81. CD63 is mostly present in the compartments of the endosomal/lysosomal system and binds to many adaptor proteins (21). CD63 has been shown to bind to syntenin-1 which plays a role in Exo biogenesis (22). CD81 is another tetraspanin which not only has multiple binding partners that play a role in interaction with cytoskeletal proteins but it has also been shown to bind to the GTPase Rac, thus regulating tumor cell migration (21). More importantly, CD81 is known to regulate outside-in signaling for integrins (23). Exo and larger vesicles including oncosomes play a role in cancer progression by horizontally transferring bioactive molecules to recipient cells (24). Specifically, transfer of proteins and mRNAs to non-tumorigenic and tumorigenic cells has been shown. Regarding Exo transfer to non-tumorigenic cells, Mian He et al have demonstrated that hepatocellular carcinoma-derived Exo transfer pro-tumorigenic RNAs and proteins to immortalized hepatocytes, thereby, inducing motility of these cells (25). It has also been reported that prostate cancer Exo containing H-ras and K-ras transcripts cause neoplastic reprogramming of adipose stem cells *in vivo* (26) and that Exo purified from breast cancer patient sera are able to induce normal epithelial cells to form tumors in a dicer-dependent fashion (27). In contrast, for Exo transfer to tumorigenic cells, Tauro et al have shown that Exo obtained from H-ras transformed MDCK cells contain integrins which may induce EMT of recipient cells (28). Exo containing different tumor-derived integrins have also been shown to prepare a fertile microenvironment for organ-specific cancer metastasis (29). We have recently demonstrated that the $\alpha_v\beta_6$ integrin is expressed in Exo from prostate cancer cells and is transferred via Exo; however, only transfer of the $\alpha_v\beta_6$ integrin among cancer cells was shown (30).

It is evident that Exo play a major role in cell-cell communication and several studies have shown that

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Exo promote cancer progression (27, 29, 30). The protein content of Exo is of great interest and in this study, we investigated whether the $\alpha_v\beta_3$ integrin is expressed in Exo from prostate cancer cells and is transferred from tumorigenic to non-tumorigenic cells.

We demonstrate for the first time that exosomal $\alpha_v\beta_3$ integrin is transferred from tumorigenic to non-tumorigenic and cancer cells leading to functional changes in recipient cells such as increase in cell adhesion and migration. We also show higher $\alpha_v\beta_3$ expression in Exo from tumor-bearing mice indicating that $\alpha_v\beta_3$ integrin is a potential biomarker for prostate cancer.

Materials and Methods

Cell Lines

PC3, C4-2B, BPH-1 cell lines and culture conditions have been previously described (31). CWR22Pc cells were cultured as previously described (32).

Antibodies

The following antibodies (Abs) were used for immunoblotting (IB): mouse mAbs to CD63 (#ab 8219), CD81 (#ab 23505), or rabbit polyclonal Abs (pAbs) to FLOTILLIN-1 (FLOT-1) (#ab 41927) from Abcam; mouse mAb to ubiquitin (sc-8017) or rabbit pAbs to ERK (#sc 93), AKT (#sc 8312), CALNEXIN (CANX) (#sc 11397), rat mAb to CD9 (#sc 18869) from Santa Cruz; rabbit pAb to ACTIN (#A2066) from Sigma and rabbit pAb to SYNAPTOPHYSIN (SYN) (#180130) from Invitrogen; rabbit pAb serum against the cytoplasmic domain of human β_3 has been described (12). A rabbit mAb against β_3 (#ab 75872) from Abcam was used in immunofluorescence (IF). The AP3 mAb against β_3 (ATCC) was used for FACS analysis. A mouse anti-human $\alpha_v\beta_3$ integrin (VN receptor) mAb LM609 (#MAB1976) from Millipore and an isotype negative control Ab were used in adhesion and migration assays.

Exosome Isolation and Analysis

Exo were isolated from culture supernatant (SN) collected 48 hours after starvation by differential ultracentrifugation (19). Briefly, the SN was spun down at 10,000 x g at 4°C for 35 minutes. The SN was collected in a fresh ultracentrifuge tube without disturbing the pellet. The collected SN was then spun at 100,000 x g at 4°C for 1 hour, the pellet was washed in PBS followed by a second spin at 100,000 x g for 1 hour at 4°C. The final Exo pellet was resuspended in PBS. Proteins were extracted from Exo and lysates were prepared. Equal amounts of proteins were separated by SDS-PAGE and analyzed by IB as described

before (30). Chemiluminescence kits from Thermo scientific and Bioexpress were used for visualization. Most of the pelleting material at 10,000 x g has been shown to have a mean size of 200 nm by Nanoparticle Tracking Analysis (NTA) by Kowal et al (20). We also performed NTA analysis on the 10,000 x g pellet and saw similar results (data not shown) suggesting that microvesicles were removed.

Nanoparticle Tracking Analysis

NTA was used to determine the size distribution and concentration of Exo released from both PC3 and CWR22Pc cells. Exo from plasma of tumor-bearing TRAMP and wild-type Non-TRAMP mice were also analyzed. Exo were resuspended in PBS and diluted 1:1000. The samples were loaded in the instrument manually and analysis performed according to the manufacturer's instructions using the NTA software (NS300, Malvern Instruments, MA). The temperature for all experiments was 25°C.

Sucrose Gradient

Exo analysis was performed using a 0.25 M (top) – 2.0 M (bottom) continuous sucrose gradient as described before (19). A dual piston gradient maker (Jule Biotechnologies, Inc) was used. Briefly, the Exo pellet was resuspended in 2.5 M sucrose, loaded at the bottom of the ultracentrifuge tube and centrifuged overnight at 210,000 x g at 4°C (Sorvall, SW41, Swinging – bucket Ultracentrifuge rotor). One mL fractions were collected and resuspended in 20 mM HEPES solution. This was followed by centrifugation at 110,000 x g at 4°C for 1 hour (TLA-100.2 rotor). The final Exo pellet was resuspended in RIPA buffer. The density of each fraction was determined using refractive index with ABBE-3L refractometer (Fisher Scientific).

Analysis of Exosome-mediated $\alpha_v\beta_3$ transfer via FACS and Immunoblotting

BPH-1 and C4-2B cells were serum-starved for 24 hours and then incubated with 20 $\mu\text{g}/\text{mL}$ of PC3 Exo. After 24 hours, the cells were trypsinized, washed with PBS and subsequently stained with the AP-3 Ab specific to β_3 . Samples were then incubated with Alexa 488 rabbit anti-mouse Ab (Molecular Probes),

washed in PBS and the data were analyzed using the FACS Calibur flow cytometer (BD Biosciences). The β_3 subunit transfer to BPH-1 and C4-2B recipient cells was also analyzed through IB. In some experiments, the recipient cells that were incubated with PC3 Exo for 24 hours were also subjected to acid wash treatment before cell lysis as described previously (30).

Analysis of Exosome-mediated $\alpha_v\beta_3$ transfer upon Actinomycin D or Cycloheximide treatment

BPH-1 and C4-2B cells were serum starved for 24 hours. C4-2B cells were either treated with Actinomycin D (Act D) (10 $\mu\text{g}/\text{mL}$) (Acros Organics) or Cycloheximide (CHX) (10 $\mu\text{g}/\text{mL}$) (33) (Sigma) followed by incubation with or without PC3 Exo (15 $\mu\text{g}/\text{mL}$). BPH-1 cells were treated with CHX (1 $\mu\text{g}/\text{mL}$) (34) followed by incubation with or without PC3 Exo (18 $\mu\text{g}/\text{mL}$). After 24 hours, cells were lysed and analyzed by IB as described above.

Quantitative Real Time PCR

Quantitative Real Time PCR (qRT-PCR) analysis was performed as described earlier (35). The following primers were used: β_3 F 5'-ACTTCTCCTGTGTCCGCTACAAG-3' and β_3 R 5'-GGTGTCAGTACGCGTGGTACA-3' (36) or GAPDH F 5' - GGAAGGTGAAGGTCGGAGT-3' and GAPDH R 5'-GTTCTCAGCCTTGACGGTGC-3'. GAPDH was used as a housekeeping gene for normalization. Each reaction was carried out at least in triplicates and the fold differences in the β_3 expression were determined relative to GAPDH. Delta graph software was used to plot the data.

Immunofluorescence and Confocal Microscopy

Analysis of membrane protrusive events- BPH-1 cells were allowed to attach on VN coated glass coverslips for 1 hour in media without FBS. C4-2B cells were allowed to attach on VN for 1 hour in complete media followed by starvation for 1 hour. Cells were then subjected to incubation with or without PC3 Exo (20 $\mu\text{g}/\text{mL}$) for 2 hours. Cells were processed for IF as previously described (30). Cells were incubated with

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Alexa Fluor 488 Phalloidin (ThermoFisher Scientific) for 30 minutes to stain for F-Actin. The slides were analyzed using an inverted confocal microscope (LSM510, Carl Zeiss). A minimum of 50 cells was counted for both cell lines in each conditions to analyze membrane protrusions.

PKH26 labeled Exosome uptake and β_3 expression- PC3 Exo were labeled with PKH26 dye as previously described (30). BPH-1 and C4-2B cells were serum starved followed by incubation with or without labeled PC3 Exo (40 $\mu\text{g}/\text{mL}$). After 24 hours, cells were stained using the same IF protocol as described above except that the cells were not permeabilized. Cells were incubated with mouse mAb specific to β_3 for 1 hour followed by Alexa 488 rabbit anti-mouse Ab incubation for 1 hour at RT. The secondary Ab alone was used as a control for non-specific binding. Intensity analysis for β_3 expression was performed on at least 35 cells in each condition using Image J software, and the ratio of β_3 expression intensities was calculated.

Adhesion and Migration Assays

The top and bottom of Transwell chambers (12 μm pore diameter, Millipore) were coated with VN (10 $\mu\text{g}/\text{mL}$) overnight at 4°C. BPH-1 and C4-2B cells were incubated with or without PC3 Exo (20 $\mu\text{g}/\text{mL}$). After 24 hours, recipient cells were trypsinized and either treated with GRGDSPK (RGD) (1 mg/mL) or GRGESP (RGE) peptides (Gibco BRL) (1 mg/mL) for 30 minutes at 4°C. For experiments using $\alpha_v\beta_3$ integrin blocking Ab, cells were incubated with LM609 Ab (15 $\mu\text{g}/\text{mL}$) or isotype control Ab (15 $\mu\text{g}/\text{mL}$) for 1 hour at 4°C following a 24 hours incubation with Exo. Cells were then seeded on VN coated Transwell chambers for 16 hours. After fixation with 3.7% paraformaldehyde, the cells attached on top and bottom of the filter were stained with DAPI (Sigma). Cell adhered to the top and bottom of the filter were counted and then the cells that were attached on the top of the filter were removed using a cotton swab. Cells that had migrated to the bottom of the filter were then counted (37). Migration results were analyzed as described previously (30). VN was purified from plasma by affinity chromatography.

Generation of TRAMP mice

TRAMP mice, expressing SV40 large T antigen in the prostatic epithelium were generated and characterized as described before (38). All mice were maintained under specific pathogen-free conditions. Care and handling of animals was in compliance with IACUC experimental protocols.

Exosome Isolation from Mouse Plasma

TRAMP (n=5) and Non-TRAMP wild-type (n=5) mice (age range from 32.5 – 37.5 weeks) were subjected to intracardiac puncture immediately after euthanasia for blood withdrawal. The blood withdrawn from these mice was collected in 3.8% Na-citrate and spun down to obtain plasma. The plasma samples were processed for isolation of Exo using ExoQuick™ as per the manufacturer's instructions (Systems Biosciences). Quantitative analysis of plasma-derived Exo was performed by determining protein concentration using BCA assay followed by IB to detect expression of the $\alpha_v\beta_3$ integrin and exosomal markers such as CD9 and FLOT-1.

Immunofluorescence of TRAMP tissues

Prostates were dissected from TRAMP mice and the tissues were preserved in formalin and embedded in paraffin. Antigen retrieval was performed on these samples by incubation in 10 mM Na-citrate buffer (pH 6.0) for 23 minutes at 95°C. The sections were blocked for 1 hour at RT with PBS/5% BSA. Staining with β_3 Ab (Abcam), SYN Ab (Invitrogen) or non-immune rabbit IgG was performed by incubation of tissue samples with primary Abs (1:100) for 1 hour at RT, followed by incubation with Alexa Fluor 488-goat anti rabbit Ab (1:250) (Molecular Probes) for 20 minutes at RT. Nuclei were counterstained using DAPI. After three washes, coverslips were mounted on the sections and analyzed by confocal microscopy as described above.

Statistical Analysis

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Statistical significance between datasets was calculated using Excel (Microsoft) software. Chi square tests and Student's *t*-tests (2 sided) were performed. *P* value of < 0.05 was considered statistically significant.

Results

The $\alpha_v\beta_3$ integrin is expressed in exosomes released from prostate cancer cells

We investigated whether the $\alpha_v\beta_3$ integrin, an ECM receptor known to be up-regulated in prostate cancer (12, 13), is present in Exo secreted by different prostate cancer cells. In our experimental design, we cultured PC3 and CWR22Pc cells and isolated Exo from their supernatants by ultracentrifugation; our IB analysis shows that the $\alpha_v\beta_3$ integrin is expressed in both PC3- (Fig. 1A *left panel*) and CWR22Pc- (Fig. 1B *left panel*) derived Exo. Each Exo preparation was characterized by detection of CD63 and CD81, exosomal markers, that appear enriched in Exo lysates compared to total cell lysates (TCL). In contrast, CANX, an endoplasmic reticulum protein, is not detected in these Exo preparations (Figs. 1A and 1B *left panels*). The size distribution of both PC3- and CWR22Pc-derived Exo was determined using NTA. The majority of the Exo from both cell lines was in the accepted size range for Exo with peaks for particle size between 120-150 nm (18) for PC3 Exo (Fig. 1A *right panel*) and 70-120 nm (18) for CWR22Pc Exo (Fig. 1B *Right panel*). Since CWR22Pc are AR positive and PC3 are AR negative cells, we conclude that the expression of the $\alpha_v\beta_3$ integrin in Exo is not affected by the presence or absence of AR. To further characterize the fidelity and purity of Exo, a continuous sucrose gradient was performed to analyze the Exo released from CWR22Pc cells. The data show that the β_3 integrin subunit expression is enriched at the expected density range between 1.13 to 1.19 g/mL (18). Expression of CD63, CD81 and FLOT-1 but not CANX, is also enriched in the same fractions as the β_3 integrin subunit (Fig. 1C). This analysis was also performed using Exo from PC3 cells and similar results are observed; expression of the α_v subunit is enriched in the same fractions as the β_3 integrin subunit along with exosomal markers CD63 and CD81 (data not shown). Our results show that the $\alpha_v\beta_3$ integrin is expressed in Exo secreted by PC3 and CWR22Pc prostate cancer cells.

Exosomal transfer of $\alpha_v\beta_3$ integrin occurs from tumorigenic prostate cells to either non-tumorigenic or tumorigenic prostate epithelial cells

We focused our attention on whether Exo derived from prostate cancer cells that contain the $\alpha_v\beta_3$ integrin have the ability to transfer $\alpha_v\beta_3$ to non-tumorigenic BPH-1 cells or other tumorigenic cells such as C4-2B. After incubating PC3 Exo with BPH-1 cells for 24 hours, we observe using IB analysis that there is a significant increase in the $\alpha_v\beta_3$ integrin levels (Fig. 2A). In contrast, in cells incubated with PBS (vehicle) alone, there is no detectable $\alpha_v\beta_3$ expression. AKT and ACTIN are used as loading controls. Similarly, we observe an increase in $\alpha_v\beta_3$ integrin level in C4-2B cells treated with Exo (Fig. 2B). To exclude any possibility of Exo being bound externally on the surface of the recipient cells, we subjected the BPH-1 cells to acid wash treatment upon 24 hour incubation with PC3 Exo. No change in $\alpha_v\beta_3$ expression is observed after acid wash (Fig. 2C). Using FACS analysis, we showed that $\alpha_v\beta_3$ is found on the surface of C4-2B cells when treated with PC3 Exo in comparison to cells treated with vehicle (Fig. 2D). We next explored whether the $\alpha_v\beta_3$ integrin is transferred as a protein or mRNA. We performed qRT-PCR on C4-2B cells either treated with PC3 Exo or vehicle. The results reveal no change in mRNA levels of the β_3 integrin subunit upon PC3 Exo treatment (Fig. 2E) and suggest that the $\alpha_v\beta_3$ integrin is not transferred as mRNA. To further exclude the possibility that exosomal β_3 mRNA would transfer or cytokines may induce transcription and translation of $\alpha_v\beta_3$ in the recipient cells, C4-2B cells were treated with either a transcription inhibitor Act D or a protein biosynthesis inhibitor CHX before incubation with PC3 Exo. IB analysis shows that $\alpha_v\beta_3$ integrin levels remain unaffected by Act D or CHX treatment of C4-2B cells (Fig. 2F *Upper Panel*). Act D and CHX activity at the used concentrations was confirmed by evaluating their effect on ubiquitin expression levels (data not shown). Similarly, no change in $\alpha_v\beta_3$ expression is observed in BPH-1 cells upon CHX treatment (Fig. 2F *Lower Panel*). This shows that $\alpha_v\beta_3$ integrin is transferred as a protein via Exo. BPH-1 cells were not subjected to Act D since a concentration as low as 0.1 μ g/mL was toxic to these cells. Overall, these data

demonstrate that extracellular $\alpha_v\beta_3$ integrin is transferred from PC3 Exo to non-tumorigenic as well as tumorigenic cells and localizes at the surface of the recipient cells.

Higher $\alpha_v\beta_3$ integrin expression and increased membrane protrusions are observed in recipient cells upon exosome internalization

Based on the IB data shown in Figure 2 which confirmed exosomal transfer of $\alpha_v\beta_3$ integrin among cells, we performed IF experiments to show PC3 Exo internalization and changes in $\alpha_v\beta_3$ integrin expression in recipient cells. BPH-1 cells were seeded on VN coated coverslips followed by incubation with PKH26 labeled PC3 Exo for 24 hours after which the cells were processed for IF. We show that there is Exo internalization in the recipient cells (Fig. 3A *Upper middle and right panels*). Z-stack analysis was also performed to confirm Exo uptake (data not shown). We also show an increase in $\alpha_v\beta_3$ integrin level in BPH-1 cells upon Exo uptake (Fig. 3A *Lower middle and right panels*). $\alpha_v\beta_3$ integrin expression was below the level of detection in the Exo alone, however, it was increased 2.6 fold in cells that were incubated with PC3 Exo as evaluated by Image intensity analysis. Similar results were observed using C4-2B cells (data not shown). This further validates our results that the $\alpha_v\beta_3$ integrin is horizontally transferred between prostate cancer cells and non-tumorigenic cells. We then sought to investigate if Exo internalization led to any changes in recipient cells such as increased cell spreading and filopodia formation. Therefore, we incubated BPH-1 cells that were seeded on VN coated coverslips with PC3 Exo for 2 hours and processed samples for IF by staining for F-Actin filaments. We observe membrane protrusions in 81% of BPH-1 cells that were treated with Exo (Fig. 3B). About 31% of the cells that were treated with vehicle developed membrane protrusions. The experiment was also conducted in C4-2B cells and similar results were observed (data not shown). Overall, these data confirm that $\alpha_v\beta_3$ integrin is transferred to recipient cells via Exo. Our results also indicate that cells become more motile with increased membrane protrusions upon Exo internalization.

$\alpha_v\beta_3$ integrin transferred through exosomes supports $\alpha_v\beta_3$ -dependent adhesion and migration of non-tumorigenic cells

We next investigated whether the transferred $\alpha_v\beta_3$ integrin is functionally active. It is known that the $\alpha_v\beta_3$ integrin plays a role in prostate cancer cell migration (12) and modulates bone metastatic growth by mediating adhesion and migration of cancer cells to the bone (14). We asked whether non-tumorigenic cells are able to acquire an $\alpha_v\beta_3$ -specific phenotype by measuring cell adhesion and migration on VN which is a substrate for this integrin. To test this hypothesis, BPH-1 cells were incubated with either PC3 Exo or vehicle. After 24 hours, these cells were incubated with GRGDSPK (RGD) peptide or GRGESP (RGE) control peptide before seeding the cells onto VN coated filters. RGD peptides have been used extensively in other studies to block the $\alpha_v\beta_3$ integrin function (39) and have been shown to enhance therapeutic efficacy in treating prostate cancer bone metastasis (40). We observe that there is a significant increase in adhesion when BPH-1 cells (Fig. 4A) were treated with Exo compared to cells treated with vehicle. Recipient cells that were preincubated with Exo and then RGD, were affected at a higher extent than control RGE-treated cells. These results show that RGD is able to block binding of the $\alpha_v\beta_3$ integrin that has been transferred from Exo, to its ligand VN. Representative images of BPH-1 cells attached to VN in the Exo treatment conditions described above are shown in Fig. 4B. Migration assays were also performed using the same treatment conditions. We observe that a higher number of BPH-1 cells incubated with PC3 Exo migrate to the bottom of the filter compared to cells treated with vehicle alone (Fig. 4C). Representative images of migrated BPH-1 cells are shown in Fig. 4D. In order to further establish that the increase in cell adhesion and migration of recipient BPH-1 cells is $\alpha_v\beta_3$ integrin specific, we treated BPH-1 cells with LM609 Ab (an $\alpha_v\beta_3$ integrin inhibitory Ab) for 1 hour after a 24 hour incubation with Exo and then the cells were seeded onto VN coated filters. We observe that there is a 50% reduction in cell attachment (Fig. 5A) and a 50% reduction in cell migration (Fig. 5C) when cells that were pretreated with Exo are incubated with LM609 Ab. Adhesion and migration are not significantly affected when BPH-1 cells that were pretreated with Exo are incubated with isotype negative control Ab. Representative images of adhered and migrated BPH-1 cells in the treatment conditions described

above are shown in Fig 5B and 5D. Overall, these data show that the increase in cell adhesion and migration is dependent on the $\alpha_v\beta_3$ integrin that is transferred through Exo.

$\alpha_v\beta_3$ integrin transferred through exosomes leads to increase in migration of tumorigenic C4-2B cells

Migration assays were performed using C4-2B cells under the same treatment conditions described for BPH-1 cells. A similar trend in cell migration increase was observed when C4-2B cells were incubated with PC3 Exo (Fig. 6A). Likewise, there is a significant decrease in cell migration of C4-2B cells treated with PC3 exo and RGD versus cells treated with Exo and RGE. Migration assays were also performed using LM609 Ab as previously described. There is a significant decrease in cell migration when cells were treated with Exo and $\alpha_v\beta_3$ integrin blocking Ab versus cells treated with Exo and negative control Ab (Fig. 6B). These results demonstrate that PC3 cells are able to propagate their migratory phenotype by transferring $\alpha_v\beta_3$ integrin to recipient cells via Exo.

Increased $\alpha_v\beta_3$ integrin expression is observed in exosomes derived from plasma of tumor-bearing mice

Based on the observations made thus far, we next sought to investigate whether the $\alpha_v\beta_3$ integrin is expressed in Exo derived from plasma of mice carrying tumors and whether there are any differences in its expression pattern compared to wild-type mice. We first performed NTA analysis on Exo from TRAMP and Non-TRAMP mice to show that we could obtain Exo that were within the accepted size range from both groups of mice (Fig. 7A). We then analyzed these Exo via IB and demonstrated that $\alpha_v\beta_3$ integrin is present in Exo from murine plasma and that there is a significant increase in $\alpha_v\beta_3$ in Exo from tumor-bearing TRAMP (Fig. 7B) compared to Exo from non-tumor bearing mice. Increased expression of CD9, an exosomal marker (Fig. 7B) in Exo obtained from tumor-bearing TRAMP mice compared to wild-type mice is also shown, whereas a different marker FLOT-1 is not changed.

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We also observe co-expression of $\alpha_v\beta_3$ integrin and SYN in the prostate tissues from TRAMP mice (Fig. 7C). Based on this observation, we investigated if exosomal transfer of the $\alpha_v\beta_3$ integrin between prostate cancer cells leads to induction of NE phenotype. After a 24 hour incubation with PC3 Exo, SYN expression levels were unaffected compared to C4-2B cells treated with vehicle alone (Fig. 7D).

Discussion

In this study, our novel findings show that the exosomal $\alpha_v\beta_3$ integrin is transferred from tumorigenic cells to non-tumorigenic as well as cancer cells and is functionally active in recipient cells upon Exo uptake. We also show for the first time that $\alpha_v\beta_3$ expression is higher in Exo from plasma of tumor-bearing mice compared to Exo from wild-type mice. These results provide new insights into cancer cell communication with the surrounding environment.

The $\alpha_v\beta_3$ integrin is a key player in prostate cancer progression (12, 15). Given its unique functions, this integrin provides specificity in response to environmental cues and upon transfer, increases cell motility. Our study shows that the $\alpha_v\beta_3$ integrin is expressed in Exo released from prostate cancer cells. We have demonstrated that Exo uptake promotes membrane protrusions, transfers $\alpha_v\beta_3$ integrin which relocates to the cell surface of the recipient cells and is functionally active. Although we observe a robust increase in $\alpha_v\beta_3$ expression in our IB analysis in BPH-1 cells upon Exo incubation (Fig. 2A and C), a lower level of $\alpha_v\beta_3$ in the recipient cells was detected by FACS analysis. This suggests a possibility that either only a small amount of the integrin is loaded into the cells via Exo or even though a larger amount of exosomal $\alpha_v\beta_3$ integrin transfer may occur, only a portion of the $\alpha_v\beta_3$ receptor is exposed at the plasma membrane of the recipient cells.

It is well established now that Exo uptake by recipient cells may occur in different ways such as through filopodia clathrin-dependent endocytosis, macropinocytosis, caveolin-mediated and receptor-mediated internalization (41, 42). The $\alpha_v\beta_3$ integrin in dendritic cells has been shown to play a role in exosome uptake (43). This suggests a possibility that $\alpha_v\beta_3$ in Exo may also play a role in its uptake in recipient cells but further studies need to be performed to figure out the exact mechanism of Exo internalization and subcellular $\alpha_v\beta_3$ localization. It has also been recently shown that filopodia play a role in facilitating the Exo entry into the recipient cells by either filopodia grabbing, surfing or pulling (41). Since

we observe increased membrane protrusion formation in our recipient cells upon Exo incubation, it is possible that Exo are captured by the membrane protrusions that develop, facilitating Exo uptake. It also remains to be tested whether $\alpha_v\beta_3$ travels as a unit in the Exo or β_3 is transferred by itself through Exo, ultimately being able to heterodimerize with the α_v subunit in the recipient cells to become functional. Further studies beyond the scope of this report will need to be performed to understand these and other aspects of this process in greater detail.

Our study also shows that the $\alpha_v\beta_3$ integrin is transferred through Exo from prostate cancer cells to non-tumorigenic or cancer cells and leads to a functional increase in adhesion and migration of recipient cells on an $\alpha_v\beta_3$ integrin ligand, VN. Recently, it has been shown that Exo from sera of breast cancer patients have the ability to induce tumorigenic properties to normal epithelial cells and lead to formation of tumors (27). Our findings are in agreement with these studies suggesting that cancer cell-derived Exo not only have the ability to communicate with other cancer cells but are also able to transfer cargo proteins to non-tumorigenic cells inducing them to migrate and develop other cancer-related phenotypes.

We have shown in our study for the first time that the $\alpha_v\beta_3$ integrin is found in Exo from plasma of mice bearing prostate tumors and is expressed at a higher level in comparison to Exo from wild-type mice. We also provide the first evidence that the $\alpha_v\beta_3$ integrin is highly expressed in the tissues of TRAMP mice with NE phenotype. The $\alpha_v\beta_3$ integrin is known to play a role in the formation of osteoblastic lesions in prostate cancer (14), and inhibition of $\alpha_v\beta_3$ substantially reduces bone destruction, tumor burden (44), adhesion and migration on components of the bone matrix (45). Our results suggest that Exo may transfer $\alpha_v\beta_3$ with its adhesive and migratory abilities to ligand-rich metastatic sites such as the bone, thereby, may offer an alternative and synergistic manner to express $\alpha_v\beta_3$. Further studies to investigate the fate of lesions in the bones of mice carrying prostate tumors upon transfer of the $\alpha_v\beta_3$ integrin will be conducted. Higher expression of the $\alpha_v\beta_3$ integrin in our tumor models implicates it as a possible biomarker but patient studies need to be performed for further validation. As an example, an integrin, $\alpha_3\beta_1$ has been detected in urine Exo

of metastatic prostate cancer patients and found at higher levels in these patients as compared with benign prostate hyperplasia patients (46).

Tetraspanins are known to be enriched in Exo and play an important role in Exo biogenesis as well as Exo targeting and uptake by recipient cells (21). CD9 is considered an exosomal marker and has been shown to play a role in integrin internalization as well as protein sorting into EVs (21). We observe that there is a significant increase in the expression of CD9 in Exo from plasma and sera of tumor-bearing mice compared to Exo from wild-type mice. It is possible that formation of a subset of CD9 enriched Exo occurs that also have enriched $\alpha_v\beta_3$ integrin. CD9 has also been shown to incorporate junctional adhesion molecule-A (JAM-A) in a complex with the $\alpha_v\beta_3$ integrin in endothelial cells resulting in endothelial cell migration upon basic fibroblast growth factor stimulation (47). A possible interaction between JAM-A and the $\alpha_v\beta_3$ integrin in Exo may play a role in the regulation of cell migration of recipient cells upon Exo transfer of $\alpha_v\beta_3$. Further exploration of this possible interaction between these three molecules in prostate cancer epithelial cells and Exo released by these cells may give insights into the mechanism and properties of the $\alpha_v\beta_3$ integrin both in Exo and recipient cells.

Finally, integrins are known to interact with multiple cytokine receptors to activate various signaling pathways. The $\alpha_v\beta_3$ integrin has been shown to directly bind with IGFIR (48) and PDGF receptor (49) and it is involved in both signaling pathways. Our lab has recently shown that the $\alpha_v\beta_3$ integrin does not bind to TGF β receptor type II (T β RII) in prostate cancer cells (50). However, it does not rule out the possibility that this interaction may occur in Exo given that Exo cargo composition and interaction with other molecules may be different in the entire cell as compared to Exo. Further studies will be performed to investigate if T β RII and the $\alpha_v\beta_3$ integrin interact with each other in Exo and are able to activate downstream signaling pathways upon delivery to recipient cells.

Overall, we have shown that the $\alpha_v\beta_3$ integrin is transferred from tumorigenic to non-tumorigenic cells as well as cancer cells via Exo and its *de novo* expression in recipient cells promotes cell migration

through interaction with its ligands. The increased expression of the $\alpha_v\beta_3$ integrin in Exo from mice bearing tumors points to its clinical relevance and potential use as a biomarker for prostate cancer.

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Figure Legends

Figure 1. The $\alpha_v\beta_3$ integrin is expressed in exosomes released by prostate cancer cells. *A, Left Panel:* Exo from PC3 cells were purified via ultracentrifugation; 5 μg of Exo lysate and TCL were loaded on 12.5% SDS-PAGE gel. IB analysis shows expression of β_3 , CANX and exosomal markers CD63 and CD81. A representative preparation of Exo out of multiple Exo preparations is shown. *Right panel:* Nanoparticle size distribution analysis for PC3-derived Exo (n=5). *B, Left panel:* IB analysis of Exo released from CWR22Pc cells. β_3 , CANX and CD81 expression is shown. 10 μg protein were loaded. A representative Exo preparation out of multiple Exo preparations is shown. *Right panel:* Nanoparticle size distribution analysis for CWR22Pc-derived Exo (n=2). A representative preparation is shown for both A and B, *Right panels.* *C,* Sucrose gradient analysis of Exo secreted by CWR22Pc cells was performed as described in the Materials and Methods. Expression of β_3 , CD63, CD81, FLOT-1 and CANX is shown. The expected density range for Exo is 1.13-1.19 g/mL.

Figure 2. Exosome-mediated transfer of the $\alpha_v\beta_3$ integrin between non-tumorigenic and tumorigenic cells. *A,* Purified Exo (20 $\mu\text{g}/\text{mL}$) secreted from PC3 cells were incubated with BPH-1 cells. After 24 hours, expression levels of the $\alpha_v\beta_3$ integrin were analyzed through IB. BPH-1 cells treated with vehicle were used as a negative control. AKT and ACTIN were used as loading controls. A representative IB out of 3 experiments is shown. *B,* C4-2B cells were incubated with PC3 Exo and $\alpha_v\beta_3$ integrin expression was analyzed after 24 hours. AKT and ERK were used as loading controls. A representative IB out of 4 experiments is shown. *C,* BPH-1 cells were incubated with PC3 Exo (20 $\mu\text{g}/\text{mL}$) for 24 hours. The cells were washed with acid wash buffer twice followed by IB analysis. Expression of $\alpha_v\beta_3$ was analyzed (n=2). ERK was used as loading control. *D,* FACS analysis of C4-2B cells incubated with PC3 Exo for 24 hours.

Exosome-mediated Transfer of $\alpha_v\beta_3$ Integrin

C4-2B cells were incubated with AP-3, a mAb specific to β_3 or non-specific mIgG (negative control) (n=4). E, Quantification of mRNA levels of β_3 integrin by qRT-PCR; mRNA was isolated after incubation of C4-2B recipient cells with PC3 Exo for 24 hours. β_3 mRNA expression is normalized to GAPDH, $P \leq 0.02$ (n=3). F, C4-2B cells were treated with Act D (10 $\mu\text{g}/\text{mL}$) or CHX (10 $\mu\text{g}/\text{mL}$) followed by incubation with PC3 Exo (15 $\mu\text{g}/\text{mL}$) for 24 hours. BPH-1 cells were treated with CHX (1 $\mu\text{g}/\text{mL}$) and incubated with PC3 Exo (18 $\mu\text{g}/\text{mL}$) for 24 hours. $\alpha_v\beta_3$ integrin expression is shown (n=2). ERK was used as a loading control.

Figure 3. Increased $\alpha_v\beta_3$ and membrane protrusions are observed upon Exosome transfer. A, BPH-1 cells were allowed to attach on VN-coated coverslips. Cells were then incubated with PKH26 labeled PC3 Exo (40 $\mu\text{g}/\text{mL}$) for 24 hours (+ PC3 Exo) or vehicle alone (- PC3 Exo). Cells were then fixed and incubated with AP3, a mAb to β_3 , followed by FITC-tagged anti-mouse secondary Ab (lower panels “ β_3 integrin”) or with secondary Ab alone (upper panels “-”). PKH26-Exo signal was readily detected in confocal images taken through the interior of the recipient cells (upper panels, + PC3 Exo). A maximum projection image at higher enlargement (bottom right) revealed the presence of increased β_3 integrin and PKH26-Exo signal. B, BPH-1 cells were seeded on VN-coated coverslips and attached for one hour in FBS depleted media. Cells were then incubated with PC3 Exo (20 $\mu\text{g}/\text{mL}$) for 2 hours and processed for IF as described in the Materials and Methods. F-Actin staining is shown in green.

Figure 4. Transfer of $\alpha_v\beta_3$ positive exosomes increases adhesion and migration of non-tumorigenic BPH-1 cells. A, BPH-1 cells that were incubated with or without PC3 Exo (20 $\mu\text{g}/\text{mL}$) were treated with GRGDSPK (RGD) (1 mg/mL) or GRGESP (RGE) (1 mg/mL) peptide before being added onto the filter coated with $\alpha_v\beta_3$ ligand, VN (10 $\mu\text{g}/\text{mL}$). Cells were washed, fixed and stained with DAPI; pictures of 10-15 random fields were taken. BSA was used as a non-specific substrate for cell attachment. The average number of attached cells was determined. Error bars depict SEM for 2 independent experiments **, $P < 0.01$ compared

with controls. B, Representative images of control and PC3 Exo treated BPH-1 cells attached to VN were stained with DAPI are shown. C, Migration assay of BPH-1 cells either incubated with $\alpha_v\beta_3$ positive PC3 Exo or vehicle alone. Cells were seeded for 16 hours on BSA (1%) or VN (10 $\mu\text{g}/\text{mL}$) and cell migration was evaluated as described under Materials and Methods. Cells that migrated to the bottom of the filter were counted and the average number of cells per field was determined. Error bars depict SEM for 2 independent experiments. **, $P<0.01$. D, Representative images of control and PC3 Exo-treated BPH-1 cells stained with DAPI in various conditions are shown.

Figure 5. $\alpha_v\beta_3$ integrin blocking antibody inhibits adhesion and migration of recipient BPH-1 cells. A, Cell adhesion and migration assays were performed by incubating recipient cells with LM609 Ab for 1 hour after 24 hours incubation with PC3 Exo (20 $\mu\text{g}/\text{mL}$). Cells were incubated on VN-coated filters for 16 hours followed by washing, fixing and staining with DAPI. An isotype Ab was used as a negative control. A, Cell adhesion was evaluated by counting the average number of cells per field. Error bars depict SEM for 2 independent experiments. **, $P<0.01$ compared with controls. B, Representative images of control and PC3 Exo treated cells stained with DAPI are shown. C, The migrated cells were counted and the average cell number per field was determined. Error bars depict SEM for 2 independent experiments. **, $P<0.01$. D, Representative images of migrated cells.

Figure 6. Gain of function in prostate cancer C4-2B cells upon transfer of exosomal $\alpha_v\beta_3$. A, Cell migration assays were performed using C4-2B cells as shown in Figure 4. The average number of cells per field was counted. Error bars depict SEM for 2 independent experiments. **, $P<0.01$. B, Cells were treated with LM609 Ab for 1 hour after 24 hours of Exo incubation and analysis was performed after 16 hours. An isotype Ab was used as a negative control. The migrated cells were counted and the average cell number per field was determined. Error bars depict SEM for 2 independent experiments. *, $P<0.05$, **, $P<0.01$

Figure 7. Increased $\alpha_v\beta_3$ integrin expression in exosomes from plasma of mice carrying prostate tumors. A, NTA analysis for Exo from plasma of tumor bearing TRAMP mice (n=3) and wild-type Non-TRAMP mice (n=2). B, IB evaluation of expression of the $\alpha_v\beta_3$ integrin in Exo purified using ExoQuick™ from plasma of 5 TRAMP versus 5 Non-TRAMP wild-type mice; $\alpha_v\beta_3$ integrin levels in 3 out of 5 specimens are shown. The levels of $\alpha_v\beta_3$ and FLOT-1 were evaluated in reducing conditions while CD9 was evaluated in non-reducing conditions. CD9 results were obtained from a different gel using the same mouse Exo samples. C, IF was utilized to analyze the expression of SYN and $\alpha_v\beta_3$ in serial sections of TRAMP prostate tumor tissues (n=24 biological replicates and 1 technical replicate/biological replicate). Staining with non-immune rabbit IgG was used as a negative control. Nuclei were stained with DAPI. D, Expression levels of SYN after 24 hours incubation of C4-2B cells with PC3 Exo (n=4). CANX was used a loading control.

Authors' contributions

Conception and design: A. Singh, L.R. Languino

Development of Methodology: A. Singh

Acquisition of data (provided animals, provided facilities, etc): A. Singh, H. Lu

Analysis and interpretation of data (e.g, statistical analysis, biostatistics, computational analysis): A. Singh, C. Fedele, H. Lu, J.H. Keen, L.R. Languino

Writing, review, and/or revision of manuscript: A. Singh, J.H.Keen, L.R. Languino

Administrative, technical, or material support (i.e, reporting or organizing databases): A. Singh, M.T. Nevalainen, J.H. Keen, L.R. Languino

Study supervision: L.R. Languino

Abbreviations

The abbreviations used are: Exo, Exosomes; NE, Neuroendocrine; VN, Vitronectin; FLOT-1, Flotillin-1; CANX, Calnexin; TCL, Total Cell Lysate; T β RII, TGF β receptor type II; IB, Immunoblotting; NTA, Nanoparticle Tracking Analysis