MicroRNA-141 suppresses prostate cancer stem cells and metastasis by targeting a cohort of pro-metastasis genes

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MicroRNAs play important roles in regulating tumour development, progression and metastasis. Here we show that one of the miR-200 family members, miR-141, is under-expressed in several prostate cancer (PCa) stem/progenitor cell populations in both xenograft and primary patient tumours. Enforced expression of miR-141 in CD44+ and bulk PCa cells inhibits cancer stem cell properties including holoclone and sphere formation, as well as invasion, and suppresses tumour regeneration and metastasis. Moreover, miR-141 expression enforces a strong epithelial phenotype with a partial loss of mesenchymal phenotype. Whole-genome RNA sequencing uncovers novel miR-141-regulated molecular targets in PCa cells including the Rho GTPase family members (for example, CDC42, CDC42EP3, RAC1 and ARPC5) and stem cell molecules CD44 and EZH2, all of which are validated as direct and functionally relevant targets of miR-141. Our results suggest that miR-141 employs multiple mechanisms to obstruct tumour growth and metastasis.
Human cancers are heterogeneous containing cancer stem cells (CSCs) that possess high capacities for tumour propagation and metastasis. Metastasis causes >90% of cancer-related deaths but our understanding of the molecular mechanisms that regulate metastasis remains limited. The invasion–metastasis cascade is a multistep cellular process that involves dissemination of cancer cells through the surrounding extracellular matrix, survival in the circulation and initial seeding followed by subsequent expansion (colonization) in the foreign microenvironment. Recent evidence suggests that microRNAs (miRNAs), small (~20–22 nt) non-coding RNAs that modulate multiple biological processes, play important roles in regulating multiple biological processes, play important roles in regulating transition (EMT), an essential developmental process important negative regulators of epithelial to mesenchymal progression and metastasis.

miR-141 in regulating PCSCs, especially in the context of PCa tumourigenic cells that fulfill the CSC definition. In a previous miRNA library screening for PCSC-regulating miRNAs, we identified miR-141, one of the miR-200 family members, to be a potent tumour suppressor, is prominently under-expressed in CD44+ PCa cells. As the CD44+ PCa cells are prominently devoid of miR-141, we first restored its expression in CD44+ PCa cells and performed CSC assays including stringent clonal (hocolone) and single cell-derived sphere-formation assays. Freshly purified CD44+ DU145 cells were transfected with the miR-141 mimics for 48 h. Importantly, miR-141 oligo transfection inhibited both tumour incidence and weight in vivo compared with the same cells transfected with NC oligos.

**Results**

miR-141 inhibits PCSC properties and tumour regeneration. As the CD44+ PCa cells are prominently devoid of miR-141, we first restored its expression in CD44+ PCa cells and performed CSC assays including stringent clonal (hocolone) and single cell-derived sphere-formation assays. Freshly purified CD44+ DU145 cells were transfected with the miR-141 mimics (miR-141) or control (NC) oligos for 48 h and then used in clonal and sphere assays. Transfection of 30 nM miR-141 mimics for 48 h resulted in significantly increased, that is, ~128-fold in PPC-1 cells to >67,000-fold in PC3 cells, miR-141 levels compared with the same cells transfected with NC oligos. Direct measurement of miR-141 copy numbers in DU145 cells indicated that cultured DU145 cells expressed a wide range of endogenous miR-141 levels from barely detectable to ~6,000 copies per cell (n = 11 independent measurements).

Herein, we report that miR-141 is under-expressed in CD44+ PCa cells and PCSCs from both xenograft and patient tumours, and miR-141 exhibits tumour and metastasis-suppressing effects in PCa. Whole-genome RNA sequencing (RNA-Seq) analysis identified multiple pro-metastasis genes including CD44, Rho GTPase and EZH2 as direct and functionally relevant targets of miR-141.

**miR-141 is under-expressed in CD44+ PCa cells in patient tumours.** Systematic studies from our lab have established that the CD44+ PCa cell population is enriched in clonogenic and tumourigenic cells that fulfill the CSC definition. In a previous miRNA expression profiling of a library of ~310 sequence-validated human miRNAs, we observed that miR-141 was significantly under-expressed in CD44+ and several other PCa stem/progenitor cell populations. To further explore this observation, we purified CD44+ PCa cells from LAPC9, LAPC4 and DU145 xenografts, VCaP cultures and, for comparisons, CD133+ cells from LAPC4 xenografts and integrin α2β1+ cells from DU145 xenografts, and performed quantitative reverse transcription–PCR (qRT–PCR) analysis (Supplementary Table 1) of mature miR-141 levels relative to the corresponding marker-negative populations. We found that miR-141 was commonly under-expressed in these PCa stem/progenitor populations, including all CD44+ subpopulations (Fig. 1a). Furthermore, correlation analysis in eight PCa xenograft/culture-derived cell types revealed that the miR-141 messenger RNA levels overall inversely correlated with the abundance of CD44+ cells (Fig. 1b). Importantly, the miR-141 levels were much lower in the CD44+ cells freshly purified from 21 primary human PCa (HPCa) patient samples (Supplementary Table 2) compared with the corresponding CD44− HPCa cells (Fig. 1c). We further adopted a qRT–PCR approach to estimate the copy numbers of miR-141 in PCa cells and subpopulations (Supplementary Fig. 1 and see Methods). We observed that on normalizing to the standard curve (Supplementary Fig. 1a,b), the endogenous miR-141 levels in five PCa cell types varied widely (Supplementary Fig. 1c). Significantly, miR-141 levels in bulk HPCa cells from 20 patient tumour samples also varied widely ranging from barely detectable to ~100,000 copies per cell (Supplementary Fig. 1d). Notably, in the majority of the 20 samples analysed, miR-141 levels were lower in CD44+ cells than in the corresponding CD44− cells (Supplementary Fig. 1d,e).
Figure 1 | Underexpression of miR-141 in CD44+ PCa cells and its PCa-suppressive functions. (a) Relative expression levels of miR-141 in CSC populations (compared with the corresponding marker-negative cells) in xenografts including CD44+ populations from LAPC4, LAPC9, VCaP and DU145, α2β1+ from DU145 and CD133+ from LAPC4. (b) Relative expression levels of miR-141 in four AR+ and four AR– PCa cell types. Average % of CD44+ cells (as determined by flow analysis; n = 3–5) is indicated for each cell type (UD, undetectable). (c) Relative expression levels of miR-141 in CD44+ cells (as % of the corresponding CD44– cells) from 21 primary HPCa samples. Samples marked by * indicate early generation PDX tumours. (d-f) miR-141 inhibits clonal and clonogenic properties of CD44+ DU145 cells as shown in clonal (d,e) and sphere (f) assays. (g-j) miR-141 inhibits prostate tumour regeneration and growth. Shown are tumour images, cell numbers injected (100,000 cells per injection for all models), site of injection (s.c., subcutaneously) and days (d) when tumours were collected. Tumour weight and incidences with corresponding P-values were calculated using upaired student’s t-test for panels e and f, tumor weight comparisons in panels g-j, and sphere numbers for panel l. P values were calculated using upaired student’s t-test for panels e and f, tumor weight comparisons in panels g-j, and sphere numbers for panel l. P values for tumour incidences in panels g-j were calculated using Chi-square test. NC, not significant. *P < 0.05 and **P < 0.01. Scale bar, 1 cm (d,k).}

 significanly suppressed xenograft tumour growth in CD44+ LAPC9 (Fig. 1h) and in bulk DU145 and PPC-1 (Fig. 1j) cells. A similar trend of tumour-inhibitory effect of miR-141 on CD44+ VCaP cells was also observed (Supplementary Fig. 2b). miR-141 suppression of PCa clonal and tumour growth was associated with inhibition of cell proliferation as evidenced by reduced 5-ethyl-2′,3′-deoxyuridine (EdU) incorporation in miR-141-overexpressing cells (Supplementary Fig. 2c–e) and reduced Ki-67+ cells in miR-141-overexpressing tumours (Supplementary Fig. 2f). miR-141 expression showed minimal effects on cell death and senescence (not shown). These results, together, suggest that miR-141 inhibits the regeneration and growth of both AR+ (LAPC9 and VCaP) and AR– (DU145 and PPC-1) PCa, and that its PCa-suppressive functions are associated with inhibition of proliferation.

Of clinical relevance, lentiviral-mediated miR-141 overexpression inhibited clonal growth (Fig. 1k) and serial sphere formation in CD44+ (Fig. 1l,m) and bulk (Supplementary Fig. 2g) HPCa cells from patient tumours. miR-141 inhibits metastasis and exhibits therapeutic efficacy. The CD44+ PCa cell population acutely purified from xenograft models possesses high metastatic potential13,20,25; thus, we determined whether manipulating miR-141 levels in CD44+ or bulk PCa cells would affect their metastatic capabilities. To this end, we introduced miR-141 via lentiviral infection into luciferase
miR-141 inhibits PCa cell invasion and directly targets CD44.

To determine molecular mechanisms underlying the metastasis-suppressing effects of miR-141 in PCa cells, we performed Matrigel invasion assays, which revealed that miR-141 significantly inhibited invasive capabilities of several different types of PCa cells including AR\textsuperscript{−} (DU145 and PC3; Supplementary Fig. 4a,b, primary patient HPCa (Supplementary Fig. 4c,d) and AR\textsuperscript{+} (VCaP, LAPC4 and LNCaP C4-2; Supplementary Fig. 4e) PCa cells. The miR-200 family members, including miR-141, are well-established negative regulators of EMT\textsuperscript{7–10}. As miR-141 inhibits PCa cell invasion and metastasis, we wondered whether miR-141 might be exerting its effects by suppressing EMT.

Collectively, these results indicate that miR-141 inhibits the metastasis in both AR\textsuperscript{−} (PC3 and DU145) and AR\textsuperscript{+} (LAPC9) PCa models.

Figure 2 | miR-141 inhibits lung metastasis of CD44\textsuperscript{+} PCa cells. (a–d) miR-141 inhibits lung metastasis of CD44\textsuperscript{hi} PC3 cells. Tumour growth and metastasis were monitored using bioluminescence imaging (IVIS). (a,b) Representative images (a) and quantification (b, in total photon flux) of primary tumours before termination of the experiment. (c,d) Representative images (c) and quantification (d) of lung metastasis with the primary tumours covered. In a, four animals showed lung metastasis without their primary tumours covered (in red box). Inserts in d are the representative GFP IHC images in the lungs (note significantly reduced GFP\textsuperscript{+} metastatic foci in the Lenti-141 lung). (e) miR-141 inhibits lung metastasis of CD44\textsuperscript{+} DU145 cells. Shown were representative IHC images of human-specific Ki-67 staining in mouse lung. (f–i) miR-141 inhibits LAPC9 metastasis. Tumour volumes were measured (f). Mice were terminated at day 40 and endpoint tumours were harvested and weighted (g; incidence indicated in bars). Lungs (and LN and some other organs) were harvested (h) to analyse metastasis by assessing GFP\textsuperscript{+} metastasis foci (i). Values represent mean ± s.d., for b,d (n = 9 for each group). For g, the animal # for each group was indicated by the denominators (that is, 5, 8, 5 and 6, respectively), For i, n = 3 for each group. For images in d (insets), the original scale bars are indicated. P values were calculated using unpaired student’s t-test.
Fig. 3a). Consistently, miR-141 increased CDH1 and reduced ZEB1 protein levels in bulk DU145 and PC3 cells (Fig. 3c).

Interestingly, however, quantitative PCR (qPCR) analysis showed that in bulk DU145 cells, miR-141 did not reduce the mRNA levels of most other EMT markers including FN, ZEB2, SNAI1, SNAI2 and TWIST (Fig. 3a). These latter observations suggest that miR-141 might be inducing a 'partial' MET phenotype in PCa cells in that it induces a strong epithelial phenotype (evidenced by increased CDH1) but only a partial loss of mesenchymal genes. Recent studies have suggested that CSCs may not always be associated with a complete mesenchymal phenotype—rather, cancer cells with...
a partial EMT phenotype that allows a greater proliferative capability of epithelial cells and the morphological plasticity of mesenchymal cells will have the best chance to survive and establish a tumour or metastatic colony. Consistent with this notion, CD44+ PCa/HPCA cells, although possessing high clonogenicity, tumour-regenerating and metastatic potentials, did not manifest a ‘pure’ mesenchymal gene expression profile (Supplementary Fig. 4f–j).

In our previous study, miR-34a, which was under-expressed in CD44+ PCa/HPCA cells, was found to directly target CD44 mRNA at two sites of the 3'-untranslated region (3'-UTR) (Fig. 3d). As miR-141 was also under-expressed in PCa (Fig. 1a) and HPCA (Fig. 1c) cells, and there was a good inverse correlation between % CD44+ PCa cells and relative miR-141 levels (Fig. 1b), we suspected that miR-141 might also target CD44. Indeed, several miRNA target prediction programmes identified a putative miR-141 binding site in the 3'-UTR of CD44. As miR-141-overexpressing PCa cells expressed luc activity higher than cells co-transfected with the reporter construct plus NC oligos (Fig. 3e), we suspected that miR-141 might also target CD44. Indeed, several miRNA target prediction programmes identified a putative miR-141 binding site in the 3'-UTR of CD44. Partial overexpression of miR-141 in bulk DU145 and PC3 cells or in CD44+ DU145 cells significantly reduced CD44 mRNA and protein levels (Fig. 3f,g). Notably, transfection in DU145 and PPC-1 cells with miR-141 mimics at 3 nM, which only minimally increased miR-141 levels (Supplementary Fig. 1g), significantly reduced CD44 mRNA levels in these cells (Supplementary Fig. 5a;b; left). Infection of cells with low MOI of Lenti-141 (that is, 2.5 and 5) similarly reduced CD44 mRNA levels (Supplementary Fig. 5a,b right). Immunofluorescence staining for CD44 in PC3 cells that had been transduced with Lenti-141 revealed a stunning contrasting pattern such that the miR-141-expressing (that is, GFP+) cells were largely devoid of CD44 expression (Fig. 3h). We also found that the endpoint tumours derived from miR-141-overexpressing PCa cells expressed lower levels of CD44 mRNA (Fig. 3f) and protein (Fig. 3i) than the control tumours. Finally, we performed functional ‘rescue’ invasion assays by overexpressing a CD44 complementary DNA lacking the miR-141 target site. As shown in Fig. 3j, miR-141 oligo transfection inhibited DU145 cell invasion, which was partially overcome by expressing exogenous CD44. Altogether, these studies (Fig. 3d–j) identify CD44 as a direct target of miR-141 in PCa cells.

RNA-Seq analysis reveals novel pathways regulated by miR-141. To identify novel targets of miR-141, we performed genome-wide RNA-Seq in DU145 (Fig. 4) and LAPC9 (Supplementary Fig. 6) cells transfected with miR-141 and NC oligos. Of the 13,456 mapped genes in DU145 cells, we identified 2,253 differentially expressed genes (DEGs; false discovery rate (FDR) < 0.05) including 949 down- and 1,304 upregulated genes (Fig. 4a and Supplementary Data 1). Ingenuity Pathway Analysis (IPA) in the 2,253 DEGs revealed ‘prostate cancer signalling’ and several growth factor signalling pathways including PI3K/AKT, ERK/MAPK, IGF-1-insulin receptor and epidermal growth factor/ERBB, and growth hormone signalling among the pathways most significantly affected by miR-141 (Fig. 4b). These latter findings suggest that miR-141 may dampen mitogenic signalling to inhibit cell cycle progression, which would be consistent with our earlier observations that miR-141 suppresses cell proliferation (Supplementary Fig. 2c–f). In further support, Gene Set Enrichment Analysis (GSEA) revealed that the miR-141 gene expression profile was negatively associated with the KEGG REACTOME ‘CELL CYCLE’ term (Fig. 4c). Moreover, the miR-141 DEGs were enriched in signal transducer and activator of transcription 3 (STAT3) and JAK/Stat3 signalling (Fig. 4b), a well-known regulator of stem cells, suggesting that miR-141 may negatively have an impact on (cancer) stem cell signalling in DU145 cells. In support, GSEA demonstrated that the miR-141-induced gene expression changes negatively correlated with embryonic stem cells (ESC) core genes (Fig. 4d) and positively correlated with ‘mammary stem cell downregulated’ (Fig. 4e). Together, these analyses indicate that miR-141 targets genes that normally promote cell cycle progression and stem cell activities, consistent with our preceding biological studies.

RNA-Seq data also provided strong support to our earlier observations that miR-141 inhibits PCa cell invasion and metastasis (Fig. 4f–k). For instance, functional annotations of the miR-141 downregulated 949 genes in DU145 cells revealed ‘invasion/metastasis’ and ‘cell cycle’ to be the two major classes of the affected genes (Fig. 4h). Furthermore, consistent with well-established regulation of EMT by miR-200 family7–10, miR-141 caused changes in genes involved in EMT, transforming growth factor-β, cell–cell junction and cytoskeleton, and other related signalling processes (Fig. 4f). Many epithelial markers including E-cadherin, Claudins (for example, CLDN7 and CLDN3) and cytokeratins (KRT) were strongly induced by miR-141 (Fig. 4g). GSEA also showed that the miR-141 gene profile was strongly enriched in normal prostatic luminal epithelial cells we recently reported (normalized enrichment score (NES) = 1.87; P = 0.000; FDR = 0.000). Meanwhile, miR-141 downregulated TGFb2, Zeb1, MAP2K4 and VIM but caused inconsistent changes in ZEB2, TWIST2 and SNAI2 (Fig. 4g). These results indicate, again, that miR-141 induces a strong epithelial phenotype but only a partial loss of mesenchymal gene expression (Fig. 4i–k).

Strikingly, we observed, among the miR-141 downregulated genes, a strong over-representation of genes associated with cytoskeleton remodelling and cell motility including Rho GTPases, as well as its upstream regulators and downstream effectors (Fig. 4f, shaded). A recent study using Ago-HITS-CLIP approach identified a role of miR-200 family targets in coordinated control of actin cytoskeleton dynamics. As the miR-200 members share largely overlapping downstream targets7–10, we took advantage of this argonate high throughput sequencing-crosslinking immunoprecipitation (CLIP-Seq) data that provided direct target association information of miR-200a, which shares identical seed sequence with miR-141, and compared it with our RNA-seq data to identify potential targets of miR-141 in PCa cells. From the 2,253 DEGs, 532 transcripts (263 down- and 269 upregulated) were found to have binding sites to miR-200a/miR-141 (Fig. 4a, Supplementary Fig. 6 and Supplementary Data 2). Among the top 35 downregulated ‘direct target’ genes were 4 known miR-141 targets (that is, TGFb2, CDK6, SEC23A and MAP2K4), 5 genes involved in Rho GTPase signalling (MAP2K4, ARPC5, CDC42EP3, CDC42 and RAC1) and CD44 (Fig. 4i and Supplementary Fig. 6).

An RNA-Seq experiment in xenograft LAPC9 cells over-expressing miR-141 also revealed that: (1) miR-141 significantly downregulated metastasis-associated genes; (2) miR-141 impeded actin cytoskeleton and Rho GTPase signalling; (3) miR-141 upregulated CDH1 and many epithelial genes, whereas downregulated numerous mesenchymal genes; and (4) miR-141 directly targeted ~150 genes, many of which were shared with the predicted targets in DU145 cells (Supplementary Figs 6 and 7a–h, Supplementary Data 3 and 4, and Supplementary Note 1).
Rho GTPase signalling components as direct targets of miR-141. The Rho family of small GTPases plays critical roles in regulating actin dynamics, organelle development, cell movement and other cellular functions. Three major Rho GTPases, RhoA, RAC1 and CDC42, regulate different yet intertwined aspects of cell movement: RhoA mainly modulates (actin) stress fibres and focal adhesion formation, and RAC1 mainly regulates lamellipodia via ARP2/3 complex, whereas CDC42 is not only a major inducer of filopodia but also can activate ARP2/3 complex to form invadosomes. ARPC5 encodes one of the seven subunits of human ARP2/3 complex and is a major regulator of actin cytoskeleton. CDC42EP3 encodes the effector protein downstream of CDC42, which is involved in actin cytoskeleton re-organization during cell shape changes including pseudopod formation.

As our RNA-Seq analysis implicated RAC1, CDC42, CDC42EP3 and ARPC5 as direct miR-141 targets in PCa cells (Fig. 4l and Supplementary Fig. 7a–h), we performed western blotting (WB) analysis to determine the impact of miR-141 on the protein levels of these four molecules. miR-141 downregulated, to varying degrees, the four proteins (Fig. 5a,b). Notably, miR-141 mimics at low concentrations (that is, 3 and 15 nM) and Lentivirus containing miR-141 at low MOI (for example, 5) also downregulated the mRNA levels of RAC1 and CDC42 in PCa cells (Supplementary Fig. 7i). Luciferase reporter assay indicated that miR-141 inhibited the luc activities of the constructs containing the 3’-UTR of RAC1, CDC42, CDC42EP3 or ARPC5 (Fig. 5c). Importantly, site-specific mutagenesis of the miR-141-binding sites abrogated the miR-141 suppression of the reporter activities (Fig. 5d), confirming direct miR-141 binding to these Rho GTPases. To further validate that these molecules represent the functional mediators of miR-141 effects, we performed ‘rescue’ invasion experiments, in which we overexpressed RAC1 and CDC42 coding constructs that lacked the miR-141 binding site(s) at the 3’-UTRs. As shown in Fig. 5e,
overexpression of miR-141 inhibited DU145 cell invasion, but the inhibitory effect was overcome by the expression of non-targetable RAC1 or CDC42 cDNA. Furthermore, miR-141 overexpression suppressed the GTPase activities of RAC1 and CDC42 but not RhoA (Fig. 5f) and, interestingly, RhoA was not predicted to be a direct miR-141 target. Of note, infection of PCa cells with Lenti-141 at low MOI (that is, 2.5 and 5) also decreased RAC1, CDC42, ARPC5 or CDC42EP3 3’-UTRs (polycomb repressive complex 2) that is frequently overexpressed in aggressive forms of prostate and other cancers, and involved in CSC maintenance, invasion and metastasis43–46, as a miR-141 ‘upstream regulator’ (P = 2.46 × 10^−7), suggesting that EZH2 might be one of the mediators in miR-141-elicited global gene changes. In support, miR-141-induced gene expression profile was associated with ‘EZH2 Targets DN’ in several data sets (Supplementary Fig. 6f). We observed a putative miR-141-binding site in the 3’-UTR of EZH2 mRNA (Fig. 6a) and luc reporter assays showed that miR-141 suppressed the luc activities of the wild type but not mutated EZH2 3’-UTR (Fig. 6b). In addition, the CD44^+ PCa cells generally expressed higher levels of EZH2 mRNA (Fig. 6c) and miR-141 overexpression

EZH2 represents yet another novel target of miR-141. Interestingly, IPA upstream regulator analysis implicated EZH2, a histone methyltransferase and a critical component of the PRC2 (polycomb repressive complex 2) that is frequently overexpressed in aggressive forms of prostate and other cancers, and involved in CSC maintenance, invasion and metastasis43–46, as a miR-141 ‘upstream regulator’ (P = 2.46 × 10^−7), suggesting that EZH2 might be one of the mediators in miR-141-elicited global gene changes. In support, miR-141-induced gene expression profile was associated with ‘EZH2 Targets DN’ in several data sets (Supplementary Fig. 6f). We observed a putative miR-141-binding site in the 3’-UTR of EZH2 mRNA (Fig. 6a) and luc reporter assays showed that miR-141 suppressed the luc activities of the wild type but not mutated EZH2 3’-UTR (Fig. 6b). In addition, the CD44^+ PCa cells generally expressed higher levels of EZH2 mRNA (Fig. 6c) and miR-141 overexpression

Figure 5 | Rho GTPase signalling pathway members are direct targets of miR-141. (a) WB of RAC1, CDC42, ARPC5 and CDC42EP3 in DU145 cells transfected with NC or miR-141 oligos at the two doses and time intervals. Indicated below are relative expression levels (normalized to respective GAPDH and compared to NC) of each protein determined by densitometric scanning of the individual bands. (b) Bar graph of relative protein level of RAC1. Values represent mean ± s.d. from four independent experiments. *P<0.05 and **P<0.01. (c) Luc reporter assays in DU145 cells co-transfected with wild-type RAC1, CDC42, ARPC5 or CDC42EP3 3’-UTRs and 30 nM miR-141 or NC oligos. Values represent mean ± s.d. (n = 4). *P<0.05 and **P<0.01. (d) Luc reporter assays in DU145 cells co-transfected with wild-type or mutant RAC1, CDC42, ARPC5 or CDC42EP3 3’-UTRs together with NC or miR-141 oligos (30 nM) for 72 h. Values represent mean ± s.d. (n = 4). *P<0.05. (e) Invasion assays in DU145 cells co-transfected with NC or miR-141 oligos together with pcDNA3.1 empty vector or pcDNA-RAC1 or pcDNA-CDC42 vector. Values represent mean ± s.d. from triplicates in three independent experiments, P-values are indicated above. (f) Rho GTPase activities were measured in DU145 cells infected with Lenti-141 or control lentivirus. Values represent mean ± s.d. of duplicates from three independent experiments. *P<0.05. (g–i) Invasion assays in DU145 cells transfected with the antisense oligos to NC (anti-141) (g), or transfected with 10 nM of scramble siRNA control (siCon) or siRNAs targeting RAC1 (h), or CDC42 (i), either alone or in combination of anti-141. Values represent mean ± s.d. of triplicates from three independent experiments. *P<0.05 and **P<0.01. P-values were determined using unpaired student’s t-test.
Figure 6 | EZH2 is a direct target of miR-141. (a) Putative binding site of miR-141 in EZH2 3′-UTR. (b) Luc reporter assays in DU145 cells co-transfected with either wild-type EZH2 3′-UTR or the 3′-UTR with the predicted miR-141-binding site mutated and miR-141 or control oligos. (c) qPCR analysis of EZH2 mRNA levels in CD44+ PCa cells. (d) qPCR of EZH2 in DU145 cells overexpressing miR-141 using either oligo transfection or lentiviral infection. (e) WB of EZH2 and SUZ12 in PC3 and DU145 cells overexpressing miR-141. Relative protein levels were measured by densitometry and normalized to GAPDH levels (indicated below). (f) Invasion assay in DU145 cells treated with the EZH2 inhibitor DZNep. Inset, WB showing reduced EZH2 proteins by DZNep. (g) Invasion assays in DU145 cells co-transfected with 30 nM NC or miR-141 oligos together with empty vector or pCMV-EZH2 construct. (h) qPCR of EZH2 target genes including FBN1, RAD51C, KIAA0101 and CDKN2A in DU145 cells infected with lenti-Ctrl or lenti-141. (i) WB analysis of H3K27me3 in DU145 cells overexpressing miR-141 using either lentiviral vectors or oligo transfection. Total H3 and actin were included as endogenous controls. (j) miR-141 exhibits pleiotropic anti-tumour and anti-metastasis effects in PCa by targeting multiple molecules associated with CSCs, cell cycle and proliferation, and cytoskeleton, cell motility/invasion and metastasis. Values represent mean ± s.d. of triplicates from three to five independent experiments. *P < 0.05 and **P < 0.01. P values were determined using unpaired student’s t-test.

EZH2 ORF

EZH2 3′-UTR

3′ ggauAGAAAUUGUC - - - - - - UGUCACAAu 5′ hsa-miR-141

UGCCUUCUCACAGCCAGAAGUUGUUUGUACCA TATCACAA

(mutations)

significantly reduced both RNA and protein levels of EZH2 in PCa cells (Fig. 6d,e). Interestingly, miR-141 expression also reduced the protein levels of SUZ12, another component of the PRC2 complex (Fig. 6e); SUZ12 (repressed) targets were enriched in our miR-141 downregulated genes (Supplementary Fig. 8a) and we observed one putative miR-141-binding site at the 3′-UTR of SUZ12 using miRNA target prediction database (MicroRNA.org). These observations, overall, suggest that EZH2 (and perhaps SUZ12) is a direct target of miR-141. Consistent with this suggestion, 3-Deazaneplanocin A hydrochloride, a small-molecule EZH2 inhibitor, reduced EZH2 protein and blocked DU145 cell invasion (Fig. 6f). EZH2 knockdown (Supplementary Fig. 8b) also inhibited PCa cell invasion, which was alleviated by anti-141 (Supplementary Fig. 8c). Significantly, miR-141 inhibition of PCa cell invasion could be partially relieved by overexpressing an exogenous EZH2 cDNA that lacked the miR-141-targeting sequence (Fig. 6g).

Importantly, we interrogated the functional relationship between miR-141 and EZH2. As expected, EZH2 knockdown led to derepression of four EZH2-repressed genes (FBN1, KIAA0101, RAD51C and CDKN2A)44,45 (Supplementary Fig. 8d). Remarkably, miR-141 similarly derepressed the expression of 4 EZH2 targets (Fig. 6h) and decreased levels of H3K27me3 (Fig. 6i). These results, altogether, provide convincing evidence that EZH2 represents a functionally relevant target of miR-141 in PCa cells.
Similar to miR-34a, exhibits PCa metastasis-suppressing effects, similar inhibitory effects (Supplementary Fig. 8e).

Overexpressing miR-34a, Let-7b and miR-106a, both miR-34a and Let-7b possess strong tumour- and/or metastasis-suppressive functions. By first confirming the under-expression of miR-141 in PCa stem/progenitor populations, here we show that miR-141 is most highly under-expressed (compared with the other three miRNAs) in patient tumour-derived CD44+ HPCa cells (Fig. 1c, 14). miR-141 re-expression inhibits clonal and clonogenic properties in CD44+ and bulk PCa/HPCa cells in vitro, as well as tumour regeneration in four xenograft models. The tumour-suppressive functions of miR-141 are associated with inhibition of cell proliferation, as evidenced by EdU and Ki-67 staining, and supported by RNA-Seq data (Fig. 6j). Of interest, a side-by-side comparison in DU145 cells by analysing the impact of overexpressing miR-34a, Let-7b and miR-141 on cell growth and invasion revealed that miR-34a demonstrated the strongest inhibitory effects, whereas miR-141 and Let-7b showed overall similar inhibitory effects (Supplementary Fig. 8e).

One of the most important findings here is that miR-141, similar to miR-34a, [exhibits PCa metastasis-suppressing effects, as observed in three orthotopic models (that is, PC3, DU145 and LAPC9) in which miR-141 overexpressing PCa cells produced less metastasis. Interestingly, pilot studies with tail vein-injected cells revealed that mice with the prostate-implanted miR-141-expressing PCa cells showed less circulating GFP+ HPCa cells in the peripheral blood than the animals that received control PCa cells (Supplementary Fig. 8f), suggesting that miR-141 probably suppresses the metastatic cascade at an early stage. Consistently, miR-141 enforced a strong epithelial phenotype by upregulating E-cadherin, cytokeratins and many other epithelial markers as demonstrated by both qPCR and RNA-Seq analyses. Consequently, miR-141 inhibits cell motility and invasion. Interestingly, although miR-200 family members are well-known suppressors of the actin cytoskeleton, cell motility and invasion, these novel findings further suggest that miR-141 negatively regulates early metastatic events. EZH2 has been shown to be oncogenic in PCa via regulating cell invasion and CSC activities. Of interest, PRC2 epigenetically represses the expression of miRNAs including miR-200, miR-181 and miR-203 (ref. 49). Of these observations, together with miR-141 directly targeting EZH2, suggest a regulatory circuitry between miRNAs and the PRC2 epigenetic silencing machinery.

Some functional studies herein were conducted by transfecting PCa cells with 30 nM synthetic miR-141 oligos and miRNA mimic transfection has been shown to cause ‘supraphysiological’ levels of intracellular miRNA and accumulation of artificial RNA species. Indeed, transfection of 30 nM of miR-141 in PCa cells can, in a cell type-dependent manner, result in >100–67,000-fold increase in miR-141 levels compared with NC transfected cells. On the other hand, measurement of copy numbers in 20 freshly purified patient tumour-derived HPCa cells, surprisingly, reveals a wide range of ‘physiological’ levels of endogenous miR-141 (from undetectable to 103,800 copies per cell (Supplementary Fig. 1d,e)). Even cultured and xenograft-derived PCa cells exhibit a wide variation in endogenous miR-141 copy numbers. These findings suggest that PCa cells in vitro and in vivo express a range of, rather than a set, ‘physiological’ levels of miR-141. Notably, 3 nM of miR-141 mimics, which did not lead to a dramatic increase in intracellular miR-141, also significantly reduced the mRNA levels of CD44 (Supplementary Fig. 5a,b) and RAC1 (Supplementary Fig. 7i). Importantly, we performed and validated many of our biological studies using lentiviral-mediated overexpression, which does not cause supraphysiological levels of miRNA or accumulation of artefactual RNA species. Thus, constitutive lentiviral-mediated miR-141 expression, which led to a slight increase in miR-141 copy numbers, impressively inhibited clonal and clonogenic capacities of primary HPCa cells (Fig. 1k–m), xenograft tumour regeneration and growth (Fig. 1h–j and Supplementary Fig. 2b), and metastasis (Fig. 2a–e and Supplementary Fig. 3a,b). Furthermore, Dox-inducible lentiviral miR-141 expression caused only 11–13-fold increases in miR-141 levels but greatly inhibited metastasis in several PCa models (Fig. 2f–i and Supplementary Fig. 3c–g). Finally, Lenti-141, at low MOI (that is, 2.5 and 5), inhibited the mRNA levels of CD44 (Supplementary Fig. 5a,b) and the Rho GTPase activities of RAC1 and CDC42 (Supplementary Fig. 7k). Collectively, numerous complimentary and cross-validating approaches we took in our studies deliver a clear message that miR-141 possesses prostate tumour and metastasis suppressive functions via targeting critical molecules including CD44, Rho GTPases and EZH2. One caveat to note is that the original CLIP-Seq experiment was performed in breast cancer cells. Although miR-141 shares identical target seed sequence with miR-200a and we have experimentally and functionally validated a few deduced targets (for example, CD44, RAC1, ARPC5, CDC42 and CDC42EP3), ideally, miR-141-specific CLIP-Seq should be conducted in PCa cells.

Previous miRNA expression profiling studies in prostate tumour tissues and (unfractionated) bulk PCa cells have revealed conflicting results, that is, no change, downregulation or upregulation according to miR-141 levels in PCa compared with normal/benign (N/B) tissues. Serum levels of miR-141 have been reported to be elevated in advanced PCa patients. We find that prostate tumours in the TCGA PRAD database express significantly higher levels of miR-141 mRNA than neighboring N/B tissues (Supplementary Fig. 8g). Our own copy number measurements in 11 HPCa and 7 N/B samples (Supplementary Table 4) also revealed higher miR-141 levels in tumours (Supplementary Fig. 8h). Pilot in situ
hybridization studies in three pairs of PCa and matching N/B tissues similarly revealed higher levels of miR-141 in tumours (Supplementary Fig. 8i). These results may appear seemingly contradictory to our observations here that miR-141 is under-expressed in tumorigenic PCa cells and miR-141 possesses tumour- and metastasis-suppressive functions. There are several possible explanations. First, PCSCs in untreated tumours always represent a minor cell population24–26,61. As miR-141 is predominantly expressed in the non-CSCs, which represent the bulk of the tumour, the miR-141 expression levels are thus expected to be higher in tumours. Second, PCa cells frequently have increased AR expression and/or AR activity, and there has been some evidence for cross-regulations of miR-141 and androgen signalling58,60. Indeed, we find that ARþ PCa cell lines generally express higher levels of miR-141 than AR− PCa lines (Fig. 1b) and there exist several putative AR-binding sites in mir-141 upstream genomic region (Supplementary Fig. 8j). Therefore, higher levels of miR-141 in prostate tumours could be partially related to increased AR expression/activity, consistent with the findings that PCSCs generally have lower AR expression22–26,61 (thus lower miR-141). Finally, higher miR-141 levels in the sera/plasma of advanced PCa patients30 are probably related to the disintegration of the prostate organ integrity, leading to the leaking of PCa cells to the blood stream. This is analogous to reduced prostate-specific antigen (PSA) mRNA/protein expression in PCSCs25,25 but significantly increased serum PSA levels in PCa patients.

In summary, the present study demonstrates that miR-141, devoid in PCSCs, suppresses prostate tumour growth and metastasis by targeting a cohort of protumour genes including CD44, EZH2 and Rho GTPases (Fig. 6j). Coupled with our earlier studies on miRNA regulation of PCSCs13,14, the results herein reinforce the concept that lack of expression of several key tumour/metastasis suppressive miRNAs concertedly and coordinately confers CSC properties4. Our observations provide further rationale for developing this CSC-targeting miRNAs into novel, combinatorial anti-tumour and anti-metastasis replacement therapeutic61, as illustrated by miR-34a that is now in clinical trial62,63.

Methods

Cell lines and animals. DU145, LNCaP, PC3, PPC-1 and VCaP cells were obtained from American Type Cell Culture and cultured in RPMI-1640 (Life Technologies, Carlsbad, CA) plus 7% fetal bovine serum (FBS) with the exception of VCaP, which were cultured in DMEM medium (Life Technologies) supplemented with 15% FBS. Human xenograft prostate tumours, LAPC9 (bone metastasis; positive for androgen receptor (AR) and PSA), LAPC4 (lymph node metastasis; AR− and PSA−) and DU145 (brain metastasis; AR− and PSA−) were maintained in NOD/SCID mice. These cell and xenograft lines have been routinely used in our lab13,14,18,20,21,23,25,33, regularly authenticated by our institutional CCSC Cell Line Characterization Core using short tandem repeat analysis and checked to be free of mycoplasma contamination using the Agilent (Santa Clara, CA) MycoSensor QPCR Assay Kit (catalogue number 302107).

NOD/SCID mice were produced mostly from our own breeding colonies with occasional purchases from the Jackson Laboratories and maintained in standard conditions according to the Institutional Guidelines. All animal experiments were approved by our Institutional Animal Care and Use Committee.

RNA isolation and qPCR analysis. Total RNA was extracted using miRNA isolation kit (Life Technologies) according to the manufacturer’s instructions. miRNA qPCRs were performed using TaqMan miRNA assays (Life Technologies). qPCR data for miRNAs was normalized to RNU48, whereas qPCR data for mRNAs was normalized to GAPDH. Primer information was listed in Supplementary Table 1.

Determination of mir-141 copy numbers. Copy numbers of mir-141 per cell in PCa cells or tissues were estimated using a qRT–PCR method, which contained a stem-loop RT followed by real-time PCR. We first synthesized miR-141 over several orders to construct a standard curve, in which the stem-loop RT primers were annealed to miR-141 and extended in the presence of reverse transcriptase (TaqMan MicroRNA Reverse Transcription Kit, catalogue number 4366596, Ambion). Next, miR141-specific forward primer, TaqMan probe and reverse primer for stem-loop were designed with miR-141 (Primer Express version 2.0; Applied Biosystems). Transgene expression levels were normalized with four housekeeping controls (SiRNA1 (17673), SiRNA2 (17379), Taqman Human Universal Master Mix II, catalog number 4440004). Quantification of miRNAs was estimated based on measured Ct values. Finally, copy numbers per cell was estimated based on standard curve of miR-141 and miRNA assuming 15 pg total RNA per cell.

Primary HPCa processing. All HPCA samples used in this study (Supplementary Table 2) were obtained from PCa patients undergoing Da Vinci based radical prostatectomy with the written informed patient consent in accordance with federal and institutional guidelines and with the approved Institutional review Board (IRB) protocol (MDACC LAB04-0498). The protocol for processing HPCA samples to obtain high purity epithelial cancer cells was previously described13,14.

WB analysis. WB analysis was routinely performed using primary antibodies listed in Supplementary Table 3; ECL mouse IgG, horseradish peroxidase-linked whole antibody (NA931V, GE Healthcare Life Sciences, Pittsburgh, PA) or ECL rabbit IgG and horseradish peroxidase-linked whole antibody (NA934V, GE Healthcare Life Sciences). Several representative WB full films are presented in Supplementary Fig. 9.

Transient transfection with oligonucleotides. PCa cells were transfected with 30 nmol-1 of mir-141 miRNA mimics or non-targeting negative control (NC) oligos, or the Anti-miR-141 and Anti-NC antisense oligos (Life Technologies) using Lipofectamine RNAiMax (Life Technologies) per the manufacturer’s instructions. After culturing for 48 h, transfected cells were harvested for in vitro and in vivo studies.

Lentiviral-mediated overexpression of mir-141. Three different lentiviral vectors over-expressing miR-141 were generated in our lab, that is, Lenti-miR-141 (based on Lenti-miR over-expression system from System Biosciences, Mountain View, CA; Supplementary Fig. 1b), pGIPZ-miR-141 (based on pGIPZ lentiviral backbone from Open Biosystems, pGIPZ-NS as the control; Supplementary Fig. 1c) and pTRIPZ-141 (based on pTRIPZ-inducible lentiviral backbone from Open Biosystems, pTRIPZ-NS as the control; Supplementary Fig. 2d). pGIPZ and pTRIPZ lentiviral vectors were produced in HEK293T packaging cells with Trans-Lentiviral packaging system (Open Biosystems). Lenti-miR-141 and Lenti-ctl viruses were produced in 293T packaging cells with the packaging plasmid mix as previously described13. Titres were determined by infecting and counting the GFPþ 293T cells. PCa cells (from either cultures or xenografts) were infected with the lentiviral supernatant at various MOI in the presence of 8 μg ml−1 polybrene and harvested 48–72 h after infection for experiments.

Clonal and sphere formation assays. For clonal assays, cultured PCa or HPCA cells freshly purified from patient primary tumours were plated at a clonal density (that is, 100 cells per well) in a six-well plate. The number of holooclones was counted several days later. For Matrigel-based sphere formation assays, cells in medium were plated (generally 1,000 cells per well) in Matrigel at 1:1 ratio in a total of 100 μl in 96-well plate. Spheres were enumerated 1–2 weeks after plating. For floating sphere formation assays, cells were plated in serum-free epithelial basal medium supplemented with B27 (Invitrogen) and 20 ng ml−1 epidermal growth factor and basic fibroblast growth factor in ultra-low attachment plate. Floating spheres that arose in 1–2 weeks were counted. For all these experiments, a minimum of triplicate wells was run for each condition and repeat experiments were performed when necessary and feasible.

Migration and invasion assays. For Boyden Chamber invasion assays, we employed Bicor Matrigel Invasion Chamber (BD, Franklin Lakes, NJ) following the manufacturer’s instructions. Briefly, 100,000–200,000 of PCa cells after transfection with NC/miR-141 oligos or infection with miR-141 overexpressing lentivirus were seeded into each well. Medium with 20% FBS was used in the lower chamber as chemo-attractant. After 20 h, cells were fixed and stained using Hema 3 staining kit (Fisher Scientific). Representative images were taken for each membrane and cells were counted. Migration assays were simultaneously performed in control wells using identical protocol with Transwell (Costar; 8 μm PET) without Matrigel. In some experiments, DU145 cells were plated 1 day before and treated with 1, 2 and 5 μM of 3-Deazaneplanocin A for 96 h after which cells were harvested and used in invasion assays and WB analysis.

EdU flow cytometry. EdU is a nucleoside analogue to thymidine and incorporated into DNA during active DNA synthesis. EdU incorporation assays were performed using Click-IT EdU Flow Cytometry Assays kit (C10418, Invitrogen) as per the manufacturer’s instructions. Briefly, PCC-1 bulk cells or CD44+ DU145 cells transfected with NC or miR-141 oligos (30 nM) for 48 h were pulsed with 10 μM EdU for 2.5h and harvested. After fixation and permeabilization, cells were
processed for EdU immunostaining and also labelled with Pacific Blue azide and 5-
Luc reporter assays

The human CD44 3′-UTR was amplified from DU145 genomic DNA using primers 5′-GAGCTC-CATCGTCACTCCTC-CAACAC-3′, as previously described. For site-specific mutagenesis, the region in the CD44 3′-UTR complementary to the seed sequence of miR-141 was mutated (5′-TAGTGT-3′ to 5′-GGCCGGG-3′; see Fig. 3d). The human EZH2 3′-UTR was amplified from DU145 genomic DNA using primers 5′-GAGCTC-CATCGTCACTCCTC-CAACAC-3′ and 5′-TAAGCTTGAAGTCTCAGGAGACAC-3′, as previously described.31

For site-specific mutagenesis, the region complementary to the seed sequence of miR-141 was mutated (5′-AAAGTTGT-3′ to 5′-TATCACAA-3′; see Fig. 6a). The PCR fragments were cloned into pGEM-T vector (Promega, Madison, WI) and sequence confirmed. Cells were plated in reporter plasmid and 5 ng complementary to the seed sequence of miR-141 was mutated (5′-TAGTGT-3′ to 5′-GGCCGGG-3′; see Supplementary Fig. 5f). CD42 (5′-GTTGT-3′ to 5′-CAACAA-3′; see Supplementary Fig. S3) and ARPC5 (5′-GTTGT-3′ to 5′-CAACAA-3′; see Supplementary Fig. S3) 3′-UTR luc reporter constructs were obtained from SwitchGear Genomics (Menlo Park, CA) and luc activities were measured as per the manufacturer’s instructions using LightSwitch luc assay kit.

Rho family GTPase activity assay

The activity of Rho family of small GTPases including RhoA, RAC1 and CDC42 were detected using G-MLA small G-protein activation assays as per the manufacturer’s instructions (Cytoskeleton, Inc., Denver, CO; catalogue number BK135)

RNA-Seq and bioinformatics

DU145 and LAPC9 cells were transfected with 30 nM of miR-141 or NC oligos for 48 h. Total RNA was purified using RNeasy mini kit (Qiagen, Hilden, Germany). We first validated the overexpression of miR-141 by qPCR. One hundred nanograms of total RNA samples was then converted to cDNA using a NuGEN Ovation RNA-Seq System v2 according to the manufacturer’s protocol (NuGEN, San Carlos, CA). NuGEN-amplified double-stranded cDNAs were fragmented into ~180 bp using a Covaris system (Covaris, Woburn, MA). Fragmented cDNAs were run on a SPIRE-TE library construction system (Beckman Coulter, Fullerton, CA) and during the adapter ligation step, uniquely indexed NEXTflex adapters (Boo Scientific, Austin, TX) were used for each of the samples to allow for multiplexing. Adapter-ligated libraries were enriched by PCR using a KAPA library amplification kit (KAPA Biosystems, Wilmington, MA) (1 cycle at 98 °C for 45 s and 7 cycles at 98 °C for 15 s, 65 °C for 30 s and 72 °C for 30 s; 1 cycle at 72 °C for 1 min) and purified with AMPureXP beads (Beckman Coulter, Pasadena, CA). The purified libraries were quantified using a KAPA library quantification kit. The libraries were loaded on a cBot (Illumina, San Diego, CA) at a final concentration of 10 pM to perform cluster generation, followed by 2 × 76 bp sequencing on HiSeq 2000 (Illumina). Sequencing reads were mapped to reference human genome sequence (NCBI 36.1 [hg19] assembly by TopHat (Version 2.0.6). The number of fragments per kilobase of transcript per million mapped reads (FPKM) was calculated (as per the Genome Browser on March 9, 2012) was enumerated using htseq-count from HTSeq package (version 0.5.49). Genes with less than ten fragments in all the samples were removed before differential expression analysis. The differential expression between conditions was statistically assessed by R/Bioconductor package edgeR (version 3.0.8). Genes with FDR of ≤0.05 and > 200 bp were called as differentially expressed.

For Gene Ontology analysis, IPA (Qiagen, Valencia, CA) and GSEA (Broad Institute) were performed. We followed the standard procedure (http://www.broadinstitute.org/gsea/doc/GSEAUserGuideFrame.html) as described by GSEA guide and curated gene sets C2 of the Molecular Signature Database version 4.0 to compute overlaps between our gene set and genes in Molecular Signature Database. Some of the data sets presented in the text are annotated here. For example, Wong et al.44 identified a core ESC-like gene module containing genes coordinately upregulated in mouse ESC that are shared with the human ESC signature (Fig. 4e). In another study, Lim et al.45 identified a gene signature (Fig. 4e) that was consistently downregulated in mammary stem cells in both mouse and human. Through whole-genome DNA microarrays in 31 breast cancer cell lines, Charrafer-Joutrel et al.69 found a list of genes (Fig. 4i) that was upregulated in luminal-like breast cancer cell lines compared with the mesenchymal-like breast cancer lines. Onder et al.66 performed gene expression analysis in HMLE cells (immortalized non-transformed mammary epithelium) after E-cadherin (CDH1) knockdown by RNA interference and identified genes that were downregulated after loss of E-cadherin (Fig. 4j). To analyse EMT-causing pathways in tumorigenesis, Agrner et al.67 identified a gene signature that was upregulated in MDA-MB-231 breast cancer cells after knockdown of ZEB1 (Supplementary Fig. 4a) by RNA interference (Fig.4k) representing transcriptional targets of E-cadherin repressor ZEB1 in invasive human cancer cells. The miR-141 downregulated genes in PCa cells were found to be greatly enriched in EZH2-repressed genes reported by Lu et al.68 and Nuyten et al.75 (Supplementary Fig. 6f). Finally, the predicted miR-141 targets in PCa cells significantly overlapped with a cohort of SUZ12 targets in embryonic stem cells21 (Supplementary Fig. 8a).

Merging our RNA-Seq data with the miR-200a Ago-HITS-CLIP-Seq data

miR-200 Ago-HITS-CLIP results12 were available at https://bitbucket.org/sagcl/bracken-hits-clip. 2013. We used the authors’ in-house scripts (also available through the above link) for sequence alignment and peak calling. The CLIP-Seq results containing the binding information were further extracted and merged with our DU145 or LAPC9 RNA-Seq data by gene symbol. Genes that were shared on both lists were called out to generate Supplementary Data 2 and 4.

siRNA-mediated knockdown

siRNAs targeting RAC1, CDC42 and EZH2 were purchased from Origene (Rockville, MD). Three unique 27mer siRNA duplexes for each target were provided. siRNAs were transfected at 10 nM for 48 h using Lipofectamine RNAiMax (Life Technologies). Knockdown efficiency was determined by qPCR. The sequences for all siRNAs are listed in Supplementary Table 3.

Statistical analyses

In general, experiments were done in triplicates for each condition when feasible. Results are presented as mean ± s.d. calculated using Microsoft Excel. Statistical differences were determined using unpaired two-tailed Student’s t-test for most analyses, except for tumour incidence for which we used χ² test. P-values <0.05 are considered statistically significant. No statistical method was used to pre-determine sample size and no samples were excluded for any analysis.

Data availability

The RNA-Seq data discussed in this publication have been deposited in NCBI Gene Expression Omnibus with the accession number GSE1756. All other relevant data are available from the corresponding authors upon request.

References

1. Kreso, A. & Dick, J. E. Evolution of the cancer stem cell model. Cell Stem Cell 14, 275–291 (2014).
2. Medema, J. P. Cancer stem cells: the challenges ahead. Nat. Cell Biol. 15, 338–344 (2013).
3. Tang, D. G. Understanding cancer stem cell heterogeneity and plasticity. Cell Res. 22, 457–472 (2012).
4. Liu, C. & Tang, D. G. MicroRNA regulation of cancer stem cells. Cancer Res. 71, 5950–5954 (2011).
5. Adams, B. D., Kasinski, A. L. & Slack, F. J. Aberrant regulation and function of miRNAs in cancer. Curr. Biol. 24, R762–R776 (2014).
6. Feng, X. Y. & Gao, W. O. Roles of microRNAs in cancer metastasis and tumor progression. Oncogene 33, 135–147 (2014).
7. Pencheva, N. & Tavazoie, S. F. Control of metastatic progression by microRNA regulatory networks. Nat. Cell Biol. 15, 546–554 (2013).
8. Burk, U. et al. A reciprocal repression between ZEB1 and members of the miR-200 family promotes EMT and invasion in cancer cells. EMBO Rep. 9, 582–589 (2008).
9. Gregory, P. A. et al. The miR-200 family and miR-205 regulate epithelial to mesenchymal transition by targeting ZEB1 and SIP1. Nat. Cell Biol. 10, 593–601 (2008).
10. Park, S. M., Gaur, A. B., Lengyel, E. & Peter, M. E. The miR-200 family determines the epithelial phenotype of cancer cells by targeting the E-cadherin repressors ZEB1 and ZEB2. Genes Dev. 22, 894–907 (2008).
11. Korpal, M. et al. Direct targeting of Sec32a by miR-200s influences cancer cell secretion and promotes metastatic colonization. Nat. Med. 17, 1101–1108 (2011).
12. Bracken, C. P. et al. Genome-wide identification of miR-200 targets reveals a regulatory network controlling cell invasion. EMBO J. 33, 2050–2056 (2014).
13. Liu, C. et al. The microRNA miR-34a inhibits prostate cancer stem cells and metastasis by directly repressing CD44. Nat. Med. 17, 211–215 (2011).
14. Liu, C. et al. Distinct microRNA expression profiles in prostate cancer stem/progenitor cells and tumor-suppressive functions of let-7. Cancer Res. 72, 3393–3404 (2012).
15. Lin, Y. et al. The microRNA miR-34a regulates proliferation and differentiation of normal and malignant breast cells. PLoS Genet. 8, e1002751 (2012).
16. Patrawala, L. et al. Side population is enriched in tumorigenic, stem-like cancer cells, whereas ABCG2 and ABCG2− cancer cells are similarly tumorigenic. Cancer Res. 65, 6207–6219 (2005).
19. Collins, A. T. et al. Prospective identification of tumorigenic prostate cancer stem cell lines. *Cancer Res.*, 65, 10946–10951 (2005).

20. Patrawala, L. Highly purified CD44+ prostate cancer cells from xenograft human tumors are enriched in tumorigenic and metastatic progenitor cells. *Oncogene* 25, 1696–1708 (2006).

21. Patrawala, L., Calhoun-Davis, T., Schneider-Broussard, R. & Tang, D. G. Hierarchical organization of prostate cancer cells in xenograft tumors: the CD44+ CD29+ cell population is enriched in tumor-initiating cells. *Cancer Res.* 67, 6796–6805 (2007).

22. Rajasekhar, V. K., Studer, L., Gerald, W., Soci, N. D. & Scher, H. I. Tumour-initiating stem-like cells in human prostate cancer exhibit increased NF-kB signalling. *Nat. Commun.* 2, 162 (2011).

23. Qin, J. et al. The PSA−/− prostate cancer cell population harbors self-renewing long-term tumor-propagating cells that resist castration. *Cell Stem Cell* 10, 556–569 (2012).

24. Domingo-Domenech, J. et al. Suppression of acquired docetaxel resistance in prostate cancer through depletion of notch- and hedgehog-dependent tumor-initiating cells. *Cell Cancer Cell* 22, 373–382 (2012).

25. Liu, X. et al. Systematic dissection of phenotypic, functional, and tumorigenic heterogeneity of human prostate cancer cells. *Oncotarget* 6, 23595–23596 (2015).

26. Rybak, A. P., Bristow, R. G. & Kapoor, A. Prostate cancer stem cells: deciphering the origins and pathways involved in prostate tumorigenesis and aggression. *Oncotarget* 6, 1900–1919 (2015).

27. Kalluri, R. & Weinberg, R. A. The basics of epithelial-mesenchymal transition. *J. Clin. Invest.* 119, 1420–1428 (2009).

28. Thery, J. P. & Seaman, J. P. Complex networks orchestrate epithelial-mesenchymal transitions. *Nat. Rev. Mol. Cell Biol.* 7, 131–142 (2006).

29. Iorio, M. V. et al. MicroRNA signatures in human ovarian cancer. *Cancer Res.* 67, 8699–8707 (2007).

30. Mitchell, P. S. et al. Circulating microRNAs as stable blood-based markers for cancer detection. *Proc. Natl Acad. Sci. USA* 105, 10513–10518 (2008).

31. Cheng, H. et al. Circulating plasma miR-141 is a novel biomarker for metastatic colon cancer and predicts poor prognosis. *PLoS ONE* 6, e17745 (2011).

32. Madhavan, D. et al. Circulating miRNAs as surrogate markers for circulating tumor cells and prognostic markers in metastatic breast cancer. *Clin. Cancer Res.* 18, 1415–1426 (2012).

33. Li, H., Chen, X., Calhoun-Davis, T., Claypool, K. & Tang, D. G. PC3 human prostate carcinoma cell holoclines contain self-renewing tumor-initiating cells. *Cancer Res.* 68, 1820–1825 (2008).

34. Yu, M. et al. Circulating breast tumor cells exhibit dynamic changes in epithelial and mesenchymal composition. *Science* 339, 580–584 (2013).

35. Lovisa, S. et al. Epithelial-to-mesenchymal transition induces cell cycle arrest and parenchymal damage in renal fibrosis. *Nat. Med.* 21, 998–1009 (2015).

36. Jolly, M. K. et al. Coupling the modules of EMT and stemness: a tunable ‘stemness window’ model. *Oncotarget* 6, 6130–6135 (2017).

37. Li, S. et al. Breast cancer stem cells transition between epithelial and mesenchymal states that regulate their response to chemotherapeutic agents. *Cell Stem Cell* 2, 333–344 (2008).

38. Lim, E. et al. Transcriptome analyses of mouse and human mammary cell subpopulations reveal multiple conserved genes and pathways. *Breast Cancer Res.* 12, R21 (2010).

39. Charalae-Jaufret, E. et al. Gene expression profiling of breast cell lines identifies potential new basal markers. *Oncotarget* 25, 2273–2284 (2006).

40. Onder, T. T. et al. Loss of E-cadherin promotes metastasis via multiple downstream transcriptional pathways. *Cancer Res.* 68, 3645–3654 (2008).

41. Aigner, K. et al. The transcription factor ZEB1 (deltaEF1) promotes tumour cell dedifferentiation by repressing master regulators of epithelial polarity. *Oncotarget* 6, 6979–6988 (2017).

42. Lu, C. et al. Regulation of tumor angiogenesis by EZH2. *Cancer Cell* 18, 147–157 (2010).

43. Nuyten, M. et al. The transcriptional repressor NIP11 is an essential player in EZH2-mediated gene silencing. *Oncogene* 27, 1449–1460 (2008).

44. Ben-Porath, I. et al. An embryonic stem cell-like gene expression signature in poorly differentiated aggressive human tumours. *Nat. Genet.* 40, 499–507 (2008).

Acknowledgements
For the work conducted at the M.D. Anderson Cancer Center (MDACC), we thank P. Whitney for FACS, Histology core for help in IHC, D. Kusewitt and C. Perez for IVIS, P. Whitney for FACS, Histology core for help in IHC, D. Kusewitt and C. Perez for IVIS, C. Perez for Image Cytometry and Genomics. Finally, we thank all other Tang lab members.
Author contributions
C.L., R.L. and D.G.T. conceived and designed the study. C.L., H.-P.C., Y.T., K.L., Y.L., Y.Z. and J.S. performed the RNA-seq analysis. C.L., R.L. and Q.D. carried out experiments and data analysis. B.L., K.R. and J.K. provided technical help and helpful discussions. C.L., R.L. and D.G.T. wrote the manuscript with input from all other authors. D.G.T. supervised the project.

Additional information
Supplementary Information accompanies this paper at http://www.nature.com/naturecommunications

Competing financial interests: The authors declare no competing financial interests.

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How to cite this article: Liu, C. et al. MicroRNA-141 suppresses prostate cancer stem cells and metastasis by targeting a cohort of pro-metastasis genes. Nat. Commun. 8, 14270 doi: 10.1038/ncomms14270 (2017).

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