Relevance of epithelial-to-pericyte transition in cancer

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To cite this article: Anitha K. Shenoy & Jianrong Lu (2016): Relevance of epithelial-to-pericyte transition in cancer, Molecular & Cellular Oncology, DOI: 10.1080/23723556.2016.1260672

To link to this article: http://dx.doi.org/10.1080/23723556.2016.1260672

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Accepted author version posted online: 22 Nov 2016.
Published online: 22 Nov 2016.

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Epithelial-to-mesenchymal transition (EMT) is a cellular reprogramming process by which well-knit epithelial cells lose intercellular adhesion and acquire the increased migratory capacity of mesenchymal cells. The cardinal features of EMT have led to the popular hypothesis that EMT is a prerequisite for carcinoma metastasis. However, recent studies have shown that EMT is dispensable for spontaneous metastasis in various genetically engineered mouse models of breast and pancreatic cancers. Therefore, the significance of EMT in cancer progression remains to be determined, and the fates and roles of epithelial tumor cells naturally transitioning to a mesenchymal state are largely elusive.

In our study we tracked epithelial cancer cells that underwent inducible or spontaneous EMT in tumor transplantation models. Unlike epithelial cells, the majority of EMT cancer cells preferentially occupy the perivascular space and are closely associated with blood vessels, thus simulating pericytes. This intriguing observation prompted us to examine EMT cancer cells in tumor vascularization. EMT markedly upregulates various pericyte markers in carcinoma cells, including platelet-derived growth factor receptor (PDGFR)-β and N-cadherin. In tumor xenografts, induction of EMT in cancer cells increases pericyte coverage of tumor vasculature. In tumors that are generated from EMT-prone carcinoma cells, a substantial fraction of tumor pericytes are derived from naturally occurring EMT cancer cells. Depletion of these EMT cells reduces the number of pericytes, destabilizes blood vessels, and attenuates tumor growth. Therefore, EMT cancer cells phenotypically and functionally resemble pericytes and are indispensible for vascular stabilization and sustained tumor growth. We propose that in the primary tumor, a small subset of epithelial cancer cells undergo EMT and the resulting enhanced mobility enables EMT cancer cells to migrate within the tumor mass. Moreover, acquisition of PDGFR-β expression by EMT cells allows their chemotaxis toward the endothelium, and homodimerization of N-cadherin on the plasma surface of EMT cells and endothelial cells (ECs) establishes their intercellular adhesion. Like pericytes, EMT cancer cells improve vascular support for growth of the bulk tumor. These findings suggest that EMT confers key pericyte attributes on cancer cells and may often represent epithelial-to-pericyte transition (EPT) (Fig. 1). EMT consists of a broad spectrum of intermediate phenotypes between the completely epithelial state and the completely mesenchymal state. Given the mesenchymal nature of pericytes, it is likely that EPT cells exhibit a complete EMT phenotype. Our study also reinforces the importance of tumor vascularization. Avascular tumors are severely restricted in their growth due to the lack of a blood supply. Cancer cells are well known to be able to induce angiogenesis, the formation of new blood vessels, for expansion of the tumor mass. Furthermore, certain cancer cells may mimic ECs to form de novo perfusable vascular-like networks by themselves, a phenomenon termed vascular mimicry. Pericyte coverage is critical for the maturation of nascent vasculature. Glioblastoma stem cells can indeed differentiate into functional pericytes. Our study further suggests that EPT may be a general mechanism by which cancer cells perform pericyte functions. Our findings uncover a key role of EPT cancer cells in tumor growth, but the potential implication of EPT in metastasis remains to be elucidated. In our study we did not observe evident distant metastasis by labeled EMT cancer cells. This might be attributed to the metastatic incompetence of the chosen cancer cells and/or the relatively short experimental duration that may be insufficient for the development of metastasis. It has been suggested that deficient pericyte coverage of tumor vasculature increases interstitial fluid pressure and facilitates cancer cell intra-vasation; thus, pericytes may stabilize the vasculature to limit metastasis. As EPT cancer cells function like pericytes to stabilize...
blood vessels, this special EMT program may potentially suppress blood-borne metastasis. On the other hand, migration of cancer cells toward blood vessels in the primary tumor is a natural part of the intravasation process. Association of EPT cancer cells with ECs may expedite their entry into the circulation. It is possible that the EMT process may generally enable cancer cells to be chemo-attracted to and associated with blood vessels, but whether such cells stabilize the vasculature or intravasate for metastasis might be determined by their intrinsic malignant properties.

A key contribution of EMT to malignancy is therapy resistance. Recent studies showed that, although it is dispensable for metastasis, EMT promotes chemoresistance in vivo. While the EMT process per se confers resistance to cell death induced by various cancer therapies, the vascular association of EPT cancer cells observed in our study may further contribute to therapy resistance. It has been recognized that capillary ECs are not just passive conduits for delivering blood but also form vascular niches that produce a variety of growth factors and cytokines (e.g., platelet-derived growth factor [PDGF], hepatocyte growth factor [HGF], stromal cell-derived factor 1 [SDF1]), Wnts, Notch ligands, and adhesion molecules. These factors are defined collectively as “angiocrine factors,” and act in a paracrine or juxtacrine manner to promote the survival of cancer cells in the vicinity. It is well established that vascular niches in the bone marrow provide a sanctuary for subpopulations of leukemic cells to resist chemotherapy-induced death. EMT rewires signaling pathways in carcinoma cells; for instance, through the EMT process mouse mammary epithelial tumor cells downregulate epidermal growth factor receptor (Egfr), Her2, and Her3, but upregulate Pdgfrs, Axl, nerve growth factor receptor (Ngfr), HGF receptor Met, and C-X-C chemokine receptor type 4 (Cxcr4). Many of these newly acquired receptors can recognize EC-derived angiocrine factors. EPT cancer cells that reside in proximity to capillary ECs and express cognate receptors for angiocrine factors are primed to respond to perivascularly enriched angiocrine signals. In contrast, non-EMT carcinoma cells do not share the same receptor repertoire and/or close distance to blood vessels. We thus envision that under chemotherapy EPT cancer cells display a selective survival advantage, in part due to protection by vascular niches, and are able to withstand the cytotoxic effects of the treatment. Therefore, the functional interactions of EPT cancer cells with capillary ECs may promote therapy resistance.

Disclosure of potential conflicts of interest
No potential conflicts of interest were disclosed.

Acknowledgment
We thank Joseph Garcia for assistance with figure preparation.

References