The Novel Anti-tumor Therapy Targeting the “Functional” Cancer Stem Cell Markers

Go J Yoshida* and Hideyuki Saya
Division of Gene Regulation, Institute for Advanced Medical Research, School of Medicine, Keio University, 35 Shinonomachi, Shinjuku-ku, Tokyo 160-8582, Japan

Abstract
It has recently been reported that molecular targeting drugs fail to overcome the “oncogene/oncogenic signal-addiction” of cancer cells. Tumor tissue is composed of heterogeneous cancer cells, so that the therapeutic response is difficult to predict. Recent advance in cancer research has strongly suggests that cancer stem cells contribute to the formation and maintenance of the heterogeneous cellular society in the tumor tissue. After all, this heterogeneity is the major cause of the acquired resistance to anti-tumor therapies. In this commentary, we would like to briefly explain the promising therapeutic targets of CD44 variant isoform and EpCAM, the “functional” cancer stem cell markers.

Keywords: Cancer stem cell; Heterogeneity; Stemness; CD44; xCT transporter; Redox stress; EpCAM; Nutrient microenvironment

Tumor Heterogeneity and Cancer Stem Cells
Tumor tissue is composed not only cancer cells but also various types of cells in the stroma; Cancer-associated fibroblasts (CAF), tumor-associated macrophages (TAM) etc. Cancer cells show invasion and metastasis in collaboration with CAF and TAM. Accumulating evidence suggests that tumor cells educate stroma cells as they enhance the malignant potential of tumor cells [1]. Given that cancer cells frequently have mutations in the cell-cycle regulatory proteins, tumor cells are mistaken as homogeneous. However, there exists a cellular heterogeneity in tumor tissues [2], which is why therapeutic response to the molecular targeting drugs is different depending on the genetic or epigenetic background of each tumor cell. Thus, the high degree of heterogeneity renders tumor tissue more easily to acquire the resistance to the molecular-targeting therapies focusing on “oncogene/oncogenic signal-addiction” [3]. Only the resistant clones survive, so that the level of heterogeneity is transiently diminished by the anti-cancer treatment, which is called “bottleneck effect”. Generally speaking, this “bottleneck effect” is lost and the heterogeneity is again recognized in the recurrent tumor tissue [2].

An emerging concept of the cancer stem cells (CSCs) simply explains how the hierarchy of tumor cells is formed. CSCs have the self-renewal ability and multi-lineage differentiation potential [4,5], both of which are necessary to architect the secondary heterogeneous tumor lesion from a single cell (Figure 1). This diversity is responsible for the formation of minimal residual disease (MRD) [2], which leads to the latent relapse and the distant metastasis. MRD is frequently enriched in therapy-resistant clones with the “stemness” [6-8].

The Significance of CSC in Terms of Therapeutic Strategy
CSCs have been well-defined and intensively researched, but much remains to be elucidated about the function of CSCs in the tumor development and metastasis. We would like to focus on the biological characteristics of CSCs, which contributed to the therapeutic resistance. Stemness is composed of the various phenotypes of CSCs; resistance to redox stress via the synthesis of glutathione (GSH) and G0 cell cycle arrest under hypo-nutrient or growth factor conditions (Figure 1). Better understanding of the regulatory system of “stemness” is critical to develop the novel anti-tumor therapy focusing on CSCs.

Epithelial-mesenchymal transition (EMT) has widely been recognized as a crucial step in the invasion and metastasis as well as normal tissue development and wound healing [9]. EMT has long been believed to increase the number of CSCs at the invasive front and metastatic foci [10,11]. CD44 is an adhesion molecule to extracellular matrix such as osteopontin and hyaluronic acids [12]. CD44 has a numerous isoforms because of alternative splicing machinery mainly regulated by RNA binding protein epithelial splicing regulatory protein 1 (ESRP1) and epigenetic modulation of the Histone H3 lysine 9 trimethylolation (H3K9me3) [13-15]. While CD44 variant isoform (CD44v) is...

Keywords: Cancer stem cell; Heterogeneity; Stemness; CD44; xCT transporter; Redox stress; EpCAM; Nutrient microenvironment

Tumor Heterogeneity and Cancer Stem Cells
Tumor tissue is composed not only cancer cells but also various types of cells in the stroma; Cancer-associated fibroblasts (CAF), tumor-associated macrophages (TAM) etc. Cancer cells show invasion and metastasis in collaboration with CAF and TAM. Accumulating evidence suggests that tumor cells educate stroma cells as they enhance the malignant potential of tumor cells [1]. Given that cancer cells frequently have mutations in the cell-cycle regulatory proteins, tumor cells are mistaken as homogeneous. However, there exists a cellular heterogeneity in tumor tissues [2], which is why therapeutic response to the molecular targeting drugs is different depending on the genetic or epigenetic background of each tumor cell. Thus, the high degree of heterogeneity renders tumor tissue more easily to acquire the resistance to the molecular-targeting therapies focusing on “oncogene/oncogenic signal-addiction” [3]. Only the resistant clones survive, so that the level of heterogeneity is transiently diminished by the anti-cancer treatment, which is called “bottleneck effect”. Generally speaking, this “bottleneck effect” is lost and the heterogeneity is again recognized in the recurrent tumor tissue [2].

An emerging concept of the cancer stem cells (CSCs) simply explains how the hierarchy of tumor cells is formed. CSCs have the self-renewal ability and multi-lineage differentiation potential [4,5], both of which are necessary to architect the secondary heterogeneous tumor lesion from a single cell (Figure 1). This diversity is responsible for the formation of minimal residual disease (MRD) [2], which leads to the latent relapse and the distant metastasis. MRD is frequently enriched in therapy-resistant clones with the “stemness” [6-8].

The Significance of CSC in Terms of Therapeutic Strategy
CSCs have been well-defined and intensively researched, but much remains to be elucidated about the function of CSCs in the tumor development and metastasis. We would like to focus on the biological characteristics of CSCs, which contributed to the therapeutic resistance. Stemness is composed of the various phenotypes of CSCs; resistance to redox stress via the synthesis of glutathione (GSH) and G0 cell cycle arrest under hypo-nutrient or growth factor conditions (Figure 1). Better understanding of the regulatory system of “stemness” is critical to develop the novel anti-tumor therapy focusing on CSCs.

Epithelial-mesenchymal transition (EMT) has widely been recognized as a crucial step in the invasion and metastasis as well as normal tissue development and wound healing [9]. EMT has long been believed to increase the number of CSCs at the invasive front and metastatic foci [10,11]. CD44 is an adhesion molecule to extracellular matrix such as osteopontin and hyaluronic acids [12]. CD44 has a numerous isoforms because of alternative splicing machinery mainly regulated by RNA binding protein epithelial splicing regulatory protein 1 (ESRP1) and epigenetic modulation of the Histone H3 lysine 9 trimethylolation (H3K9me3) [13-15]. While CD44 variant isoform (CD44v) is...
transcriptional factor Prrx1 induces EMT but decreases the number of CSGs. mesenchymal-epithelial transition (MET) is an important step for cancer cells to colonize and proliferate at the pre-metastatic niche. That is why the knockdown of Prrx1 induces MET and promotes the lung metastasis of breast cancer cells [18].

Furthermore, it has been demonstrated that tumor cells maintain epithelial phenotype with the high expression of CD44v at the invasive front [10]. This seems paradoxical to the current concept, but some researcher insist on the presence of CAFs with CSCs at the invasive front [19]. Indeed, CD44v is strongly expressed at the invasive front and negatively correlated with the expression of oncoprotein c-Myc. Redox stress-induced Wnt signal activation is responsible for the inversed relationship between CD44v and c-Myc [10]. The irreversible quiescence of CSCs is finely regulated by the modulation of several kinds of stress derived from tumor microenvironment.

**Novel Therapy Attacking the “Functional” CSC Markers**

There are many molecules exclusively expressed in CSCs. The specific isoform of CD44 and EpCAM are positively correlated with each other even in normal epithelial cells [20]. However, both the CD44v and Epithelial Cell Adhesion Molecule (EpCAM) are not just the surface markers of CSCs. We would like to briefly introduce the recent findings on this therapeutic approach.

**Novel therapy attacking the robustness against oxidative stress of CSCs**

The heterogeneous expression pattern of CD44 is caused by the heterogeneous expression of ESRP1, which is mainly responsible for the alternative splicing of CD44 [13,15]. Epigenetic modulation of the histone at the site of ESRP1 is dynamic depending on the tumor microenvironment, which strongly supports the idea of "dynamic stemness model" [21], which is contrary to the widely-accepted idea of CSC hierarchy model. CD44v-positive cells are not necessarily express CD44v. After all, the H3K4me3 of the ESRP1 promoter lesions promote ESRP1-inducing CD44v expression, whereas H3K27me3 suppresses it [15]. Intra-tumor heterogeneity in terms of CD44v expression is considered to be determined by this dynamic epigenetic change (Figure 2).

There are several isoforms of CD44v, depending on which variable exons are inserted. Furthermore, different isoforms of CD44v exhibit different biological function and it has been suggested that CD44 variant is a promising therapeutic-target in the cancer therapy [22]. In particular, CD44v8-10, CD44 variant isoform including the variable exon 8-10, stabilizes xCT transporter at the cellular membrane. xCT forms a heterodimer with CD98 heavy chain (CD98hc), also referred to as 4F2. xCT amino acid transporter help CSCs exchange glutamate and cystine [23]. Cysteine, which is converted from cysteine, is a scarce substrate of GSH, major anti-oxidant molecule. Cancer cells are exposed to excessive amount of reactive oxygen species (ROS), which induces apoptosis, autophagy, cellular senescence, and differentiation [24-26] (Figure 3). GSH works as a gatekeeper for CSCs to prevent the ROS accumulation and maintain the "stemness". This CD44v-xCT-GSH axis enables CSCs to survive and proliferate under the redox stress conditions.

Importantly enough, xCT-GSH axis is not limited to tumors of epithelial tissues; glioma cells are reported to express xCT without CD44v and the secreted glutamate is the cause of glioma-associated brain edema [27]. Besides, triple negative breast cancer tissues, pathologically classified into basa-type and medul (low)-type
developed predominantly expressed in epithelial cancer cells, the CD44 standard isoform (CD44s) is mainly expressed in mesenchymal cancer cells. Epithelial tumor cells no longer express CD44v after EMT, and instead, express CD44s with high migratory and invasive phenotypes [13] (Figure 2). Mesenchymal tumor cells with CD44s expression after EMT tend to show the activation of phosphatidylinositol-3 kinase (PI3K)/Akt signaling pathway in the metastatic foci in the lungs of the breast cancer cells [13].

Several transcription factors have been identified to induce EMT; FOXC2 in the basal-like breast cancer [16], SIP1/ZEB2 in ovarian, breast, and hepatic tumors [17], and Snail, Slug, Twist in several kinds of cancers [9]. Contrary to the accepted wisdom, the homeobox
mesenchymal tumors, highly express xCT and show the "glutamine addiction" independently of CD44v expression [28]. Sulfasalazine (SSZ), an anti-inflammatory drug long used for patients with ulcerative colitis, has been shown to prevent the function of xCT transporters [15,23,28,29] and expected to break down the robustness of CSCs against redox stress (Figure 3).

Novel therapy attacking the sensitivity to nutrient microenvironment of CSCs

EpCAM is expressed in the circulating tumor cells (CTCs) in the blood and several neutralizing antibodies have been developed for the prevention and the treatment for tumor metastasis [30,31]. For instance, catumaxomab is the antibody preparation against human EpCAM and CD3 and clinically effective for the patients with advanced ovarian tumors with recurrent symptomatic malignant ascites [32]. However, it still remains unknown how EpCAM contributes to the survival potential of CSCs under oppressive conditions.

In the perspective of membrane-molecular function, EpCAM has recently been identified as a sensor of growth factor microenvironment by the formation of super-complex with amino-acid transporters [33,34]. For example, EpCAM is positively correlated with the expression and localization of amino acid transporter L-type amino acid transporter (LAT) I promotes the uptake of leucine, thereby enhancing malignant target of rapamycin (mTOR) signaling pathway [33,35]. There exists a crossstalk between mTOR and S6-Adenosine monophosphate-activated protein kinase (AMPK) signal pathways. In the steady state with enough amount of energy, EpCAM-low or negative cancer cells tend to be activated as compared with EpCAM-high CSCs. Serum starvation leads to AMPK signal activation exclusively in CSCs in the specific type of prostate cancers [33]. In other words, EpCAM exists as an upstream molecule for AMPK signal pathway to sense the change of the nutrient microenvironment by the formation of super-complex with amino acid transporters in cooperation with CD147 (EMMPRIN) and CD98hc (the chaperon referred to as 4F2) [34].

Thus, it is highly likely that EpCAM expression makes CSCs sensitive to the change in glucose and growth factors by the regulation of the localization and stabilization of several monocarboxylate and amino acid transporters. Prostate cancer cells respond to lack of growth factors differently depending on the EpCAM expression amount [33]. EpCAM-positive CSCs may rapidly adapt to the change in nutrient microenvironment. Long-term serum starvation promotes cell cycle arrest and the acquisition of a quiescent phenotype, often observed in CTCs. EpCAM-high cancer cells entered the G0 phase via the amino acid transporters in cooperation with CD147 (EMMPRIN) and CD98hc (the chaperon referred to as 4F2) [34].

References


