Emergence of cancer stem cells in hepatocellular carcinoma

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Liver cancer represents the second most deadly human malignancy. The major histological subtype called hepatocellular carcinoma (HCC) arises by chronic inflammation-triggered regenerative responses of normally quiescent hepatocytes and progenitors, respectively. Such regenerative stress accelerates the accumulation of genetic and epigenetic changes (Yamashita & Wang, 2013), while detailed mechanisms remain uncertain. In this issue of The EMBO Journal, Nikolaou et al present a novel HCC model that facilitates both isolation and molecular characterization of self-renewing, HCC-propagating cancer stem cells that could instruct future interventions (Nikolaou et al, 2014).

See also: KC Nikolaou et al (February 2015)

Tumor cells show heterogeneity in terms of their expression profiles and their epigenetic landscapes. In the recent years, the notion has emerged that some tumor cells possess stemness properties, meaning that they have the ability to self-renew and to generate a whole tumor after transplantation. These so-called cancer stem cells (CSCs) are especially important targets of cancer therapy, since they are thought to orchestrate tumor growth. CSCs are highly resistant against therapeutic measures and have high metastatic and invasive capacity. Therefore, continuous efforts are undertaken to isolate CSCs and to trace their developmental origin(s) (Yamashita & Wang, 2013; Kreso & Dick, 2014). It is currently assumed that HCC is induced and/or maintained by CSCs, either arising from hepatocyte de-differentiation or HPC transformation. Systematic studies to address these hypotheses are currently lacking (Yamashita & Wang, 2013; Oishi et al, 2014).

Cancer stem cells emergence from de-differentiating hepatocytes is supported by experiments that validated their reprogramming capacity. For instance, inactivation of the Hippo pathway, a regulator of cell proliferation, is sufficient to de-differentiate adult hepatocytes into cells with progenitor characteristics (Yimlamai et al, 2014). Furthermore, the activation of the NOTCH pathway in hepatocytes results in their transdifferentiation to biliary epithelial cells (Yang et al, 2013). Finally, the simultaneous activation of the NOTCH and AKT pathways leads to the onset of intrahepatic cholangiocarcinoma originating exclusively from hepatocytes (Fan et al, 2012).

Support for the HPC-origin of CSCs arises from their phenotypic similarities as well as their common ability to self-renew and to differentiate. Moreover, CSCs found in HCC have been shown to express similar oncofetal marker as HPCs (Yamashita & Wang, 2013). Those similarities strongly suggest that HPCs could develop into CSCs, while definitive experimental proof was so far missing.

The experiments conducted by Nikolaou et al established a new model for HCC addressing this gap and facilitating direct isolation of CSCs. They functionally analyzed the histone methyltransferase PR-SET7 in liver development. PR-SET7 is the sole enzyme that catalyzes histone H4K20 monomethylation (H4K20me1). This methylation plays an important role in many cellular processes like DNA replication, DNA damage response and mitotic condensation (Jørgensen et al, 2013). PR-SET7 knock-out (ko) is lethal in mice and flies and leads to cell cycle arrest and apoptosis (Driskell et al, 2012; Jørgensen et al, 2013). Using a cre-lox system, Nikolaou et al knocked out PR-SET7 in embryonic mouse livers. Consistent with the essential role of PR-SET7, prenatal lethality was observed. E18.5 embryos were anemic and had a severely reduced liver volume. In contrast, mice with PR-SET7 deletion in adult liver survived and only displayed areas of necrotic cell death. Necrosis was found in regenerating areas of the liver and was caused by the attempted proliferation of hepatocytes which die due to the lack of PR-SET7. To stimulate liver regeneration, the authors decided to perform partial hepatectomy (PHx), which leads to compensatory growth of the remaining tissue by proliferation of adult hepatocytes (Fig 1). PR-SET7-deficient cells die upon proliferation, and enhanced necrotic areas in PR-SET7-deficient PHx livers coincided with chronic inflammation. Interestingly, all PR-SET7 ko mice with PHx developed HCC between postnatal days 240–300. Thus, PHx in PR-SET7 ko livers reflects the inflammatory aspect of human HCC, although in humans hepatocyte proliferation is not as severely impaired.

Remarkably, the tumor cells isolated by Nikolaou et al showed CSC properties: they could be cultured in vitro over several passages and they could give rise to tumors with characteristics of the parental tumor when engrafted into immunodeficient mice. These properties distinguish them from a previously described HCC model which was only transplantable into already damaged livers (He et al, 2013). Why the tumor progenitor cells isolated by He et al and the CSCs isolated by Nikolaou et al differ, is currently unclear. One possibility is that they might stem from different origins. The inability of PR-SET7-deficient hepatocytes to...
CSCs in hepatocellular carcinoma

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Hepatocellular carcinoma (HCC) is a highly lethal cancer with about 300,000 deaths per year. CSCs in HCC have been uncovered primarily through transplantation-based experiments in non-human primates, demonstrating that CSCs drive tumor growth. Immunodeficient mice can host xenografts of human HCC cells, which can be transplanted into immunodeficient mice. In these cell models, CSCs self-renew and can be transplanted into immunodeficient mice, where they give rise to new tumors.

- Remarkable feature of these tumors is the presence of CSCs with stemness properties: they can self-renew and proliferate and can be transplanted into immunodeficient mice, where they give rise to new tumors.

- Additional experiments like lineage tracing are required to clarify the origin of these cells.

- In summary, Nikolaou et al created a new and very informative HCC model. Arising HCCs present with very high penetrance and surprisingly little tumor heterogeneity. Notably, the tumor cell population includes CSCs that have the ability to self-renew and to initiate tumors when transplanted into immunodeficient mice. Although these cells have been phenotypically and functionally well characterized, we still know very little about their genetic and epigenetic aberrations. Further analyses should reveal CSC-specific oncogenes and tumor suppressor genes. Furthermore, the high penetrance of CSCs in this tumor model will allow for a better understanding of their biological features such as the regulation of proliferation and drivers for their metastatic capacity. Even more intriguing will be potential translation of these animal studies for human HCCs.

References


