Liver Cancer Stem Cells

Sameh Mikhail and Aiwu Ruth He

Lombardi Comprehensive Cancer Center, Georgetown University Hospital, 3800 Reservoir Road Northwest, Washington, DC 20007, USA

Correspondence should be addressed to Aiwu Ruth He, aiwu.r.he@gunet.georgetown.edu

Received 27 February 2011; Accepted 30 March 2011

Academic Editor: Pierce Chow

Copyright © 2011 S. Mikhail and A. R. He. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Hepatocellular carcinoma is the most common primary malignancy of the liver in adults. It is also the fifth most common solid cancer worldwide and the third leading cause of cancer-related death. Recent research supports that liver cancer is a disease of adult stem cells. From the models of experimental hepatocarcinogenesis, there may be at least three distinct cell lineages with progenitor properties susceptible to neoplastic transformation. Identification of specific cell surface markers for each of the liver cell types, production of corresponding monoclonal antibodies and cell sorting techniques have together revolutionized the characteristics of normal stem cells. In hepatocarcinogenesis, multiple signaling transduction pathways, important for stem cell proliferation and differentiations, are deregulated. Strategies are being developed to identify and characterize the liver cancer stem cells. Targeting liver cancer stem cells may bring hope to curing hepatocellular carcinoma.

1. Introduction

Hepatocellular carcinoma (HCC) is the most common primary malignancy of the liver in adults. It is also the fifth most common solid cancer worldwide and the third leading cause of cancer-related death [1, 2]. Moreover, HCC incidence and death rate are rising in the United States and demonstrate the highest annual percent increase of the top 15 cancers by incidence [3]. The worldwide incidence of HCC varies according to the prevalence of hepatitis B (HBV) and hepatitis C (HCV) infection and the age-standardized incidence rates vary from 4.9 per 100,000 population in North America to 80 per 100,000 population in China [4]. It is worth noting that chronic HCV infection is the leading cause of HCC in Europe, Japan, and the United States whereas HBV infection is the leading cause in the majority of Asian and African countries [2].

Recent research supports that cancer is a disease of adult stem cells (SC). Adult stem cells are the only cells that persist in the tissue for a sufficient length of time to acquire the requisite number of genetic changes for neoplastic development. In contrast to intestinal mucosa and epidermis where a steady flux of cells occurs from the stem cell zone to the terminally differentiated cells that are imminently to be lost, liver normally exhibits a very low level of cell turnover. However, when abnormal hepatocyte loss occurs, such as after partial hepatectomy (PH) or toxic injury, the liver demonstrates an enormous regenerative capacity. The clonality of HCC is now well established based upon the studies examining viral integration sites of HBV in tumor samples [5], as well as on the determination of restriction fragment length polymorphisms of X-linked genes in tumor cells [6]. However, the cell type that has given rise to HCC has not been universally accepted. From the models of experimental hepatocarcinogenesis, there may be at least three distinct cell lineages with progenitor properties susceptible to neoplastic transformation [7].

2. The Stem Cell Origin of Liver Cancer

2.1. Hepatocytes Have “Stem Cell” Properties. Hepatocytes in normal adult liver have a lifespan of over a year. However, in response to parenchymal cell loss, the hepatocytes restore the liver mass by self-replication. In rodents, the liver can restore its original volume after two-thirds partial hepatectomy (PH) in approximately 10 days [8, 9]. Serial transplantation experiments have shown that hepatocyte can divide at least 69 times, demonstrating the clonogenic potential of hepatocytes—one of the crucial properties of an SC [10]. In HCV infected liver, the hepatocyte proliferation
rate increases with increasing cellular damage [11]. Many models of liver cancer utilize a brief exposure to a genotoxic carcinogen at a time when the liver is in a proliferative state, either during the period after a PH or necrogenic insult [12]. Hepatocytes have been found to be directly involved in carcinogenesis of HCC in 2-acetylaminofluorene and DEN-treated rats where hepatocytes were labeled with β-galactosidase [13]. Hepatocytes in proliferation appear to be the origin of cancer.

2.2. Oval/Liver Progenitor Cells as Targets for Malignant Transformation. When hepatocyte and/or cholangiocytes are damaged or inhibited in their proliferation, a potential SC compartment located within the smallest branches of the intrahepatic biliary tree, the ductules, and canals of Hering gets activated [14]. The “oval” cells in rodent or “the liver progenitor cells” in human liver involve a population of cells that are bipotential and capable of differentiating into hepatocytes or cholangiocytes. The oval/progenitor cells are labeled by over 30 surface markers including biliary-type cytokeratin (CK), CK7, CK19, oval cell markers OV6 and OV1, neuroendocrine marker chromogranin A, neural cell adhesion molecule and parathyroid hormone-related peptide, and connexin 43. The origin of HCC from hepatic progenitor cells (HPC) is often suggested from the fact that tumors contain an admixture of mature cells and cells phenotypically similar to HPCs [15, 16]. Oval/HPC proliferations and activations are observed after severe liver parenchyma injury, viral hepatitis, alcoholic hepatitis, and nonalcoholic fatty liver disease. HPC/oval cell activation accompanies many instances of liver damage, irrespective of etiology, suggesting such cells are carcinogen targets during hepatocarcinogenesis. Oval cells from p53-null mice formed HCC when transplanted into athymic nude mice [17]. A origin from oval cells is suggested by the fact that if oval cell expansion is blocked in the CDE diet mouse modeling by targeting c-Kit with imatinib mesylate, the HCC formation is reduced [18]. Furthermore, the gene expression profile from a selected group of HCC is consistent with the profile of HPCs.

2.3. Bone Marrow-Derived Stem Cells. Petersen et al. demonstrated that hepatocytes could be derived from circulating bone marrow cells [19]. Hematopoietic Stem Cells (HSC) from wild-type mice were able to repopulate the liver of FAH-deficient (fah−−) mice [20]. In the setting of sex-mismatched bone marrow transplantation, bone marrow-derived hepatocyte are found in the recipient liver with a large variation in their frequency ranging from less than 1% to >40%. However, in a chimerical mouse model with genetically labeled bone marrow, there was no malignant transformation of the bone marrow-derived liver SC during hepatocarcinogenesis induced by chemical carcinogen [20]. These results suggest that bone marrow-derived liver SC may not be targets for malignant transformation in HCC.

2.4. Isolation of Liver Cancer Stem Cells. In the last decade, identification of specific cell surface markers for each of the liver cell type, production of corresponding monoclonal antibodies, and cell sorting techniques have together revolutionized the characteristics of normal stem cells. It has been show that cancer SCs in HCC can be identified by several cell surface antigen CD133, CD90, CD44, OV6, and epithelial cell adhesive molecule (EpCAM), or by selecting the side population (SP) cells in Hoechst dye-staining [21–25]. Table 1 shows markers that are associated with liver cancer SC. The surface markers enrich HCC cells with greater tumorigenicity in immunodeficient mice, higher colon-forming efficiency, and proliferation ability in vitro. In addition, most of the markers are found to be expressed in only a minute proportion of HCC cells, and the expression of the markers correlate with poor prognosis and tumor recurrence. However, it remains to be seen how much overlap there is between these various markers, or whether there is a “one-fits-all” marker for cancer SCs in HCC. Most of the markers which are used for isolating cancer SCs from primary tumor samples were established and adapted from established cancer cell lines. It is not clear whether the cancer SCs that are derived from established cancer cell lines and cultured in vitro reflect the SCs from primary tumor in the gene expression of these surface marker. It remains a challenge to isolate enough clonally derived cancer SCs from primary tumor without in vitro propagation for lineage tracking and differentiation experiments, identification of deregulated signaling pathways that lead to the malignant transformation of normal adult SC to cancer SCs.

2.5. Pathways Important for Stem Cell Function Are Deregulated in Hepatocarcinogenesis. From HCC animal model and gene array analysis, a growing body of research suggests that many signaling pathways known to be involved in SC maintenance, self-renewal, and pluripotency, are altered in HCC. This alteration may result in the malignant transformation of liver SC [26]. These observations support the hypothesis that molecular changes in HCC originate in cancer SC [27]. Moreover, these pathways could serve as diagnostic markers and targets for therapeutic interventions [27].

3. WNT/β-Catenin

Disrupted Wnt signaling is observed in approximately one-third of all HCC which underscores its importance in carcinogenesis [28]. The Wnt pathway has a fundamental role in embryogenesis with signaling effects on proliferation and apoptosis in developing cells [29]. Wnt pathway activation is essential for maintenance of SC compartment and regulates cellular differentiation [30]. The “canonical Wnt pathway” describes a cascade of events beginning with the translation of β-catenin from the cell membrane into nucleus, where β-catenin then acts as a coactivator of the TCF/LEF family of transcription factors, these in turn regulate specific target genes including c-myc, cyclin D1, and survivin [31]. The signaling cascade is normally initiated when Wnt ligand binds to Frizzled (FZD), a transmembrane receptor [32]. FZD then signals to β-catenin to escape its association with E-cadherin. The cytoplasmic elements of the activated Wnt pathway prevent β-catenin from being phosphorylated by a
of lung cancer SCs [34].

A wide range of secreted factors regulate SC proliferation and fate. TGF-β shows remarkable functional conservation between species and between tissues that self-renew through asymmetric divisions or populational asymmetry. TGF-β signaling is important for embryonic hepatocyte proliferation, as well as in the formation of gastrointestinal cancer [37–39]. Tang et al. demonstrated that lack of responsiveness to TGF-β pathway in liver SC led to carcinogenesis [40]. Subsequently, it has been shown that targeting this pathway using indirect modulation of IL6/STAT3 appeared to be effective in eradication of cancer SC [26, 40].

5. Hedgehog

The Hedgehog signaling pathway consists of a complex suite of molecules which regulate cell differentiation, regeneration, and stem cell biology. The pathway plays important roles in the development and homeostasis of the gut tissue [41]. Studies have identified a possible role for this pathway in HCC with expression of Sonic, the predominant ligand of the Hedgehog pathway in liver, that is present in up to 60% of human HCC samples [42, 43]. The Hedgehog pathway is deregulated in hepatocarcinogenesis [42]. Genes involved in the Hedgehog pathway are highly expressed in tumorigenic CD133⁺ liver cancer SC [21]. Suppression of Hedgehog pathway not only decreased HCC cell proliferation but also chemosensitized HCC cells to 5-fluorouracil and to the induction of cell apoptosis [44].

6. Target CSCs in the Treatment of HCC

Cancer SCs are predicted to mediate tumor recurrence after chemo- and radiation-therapy due to the relative inability of these modalities to effectively target cancer SCs. Eradicating Cancer SCs brings the hope for cure. Interesting results have been demonstrated in inhibiting breast cancer SC by targeting TGF-β and Notch pathways [45]. Similarities between normal and malignant SC, at the levels of cell-surface proteins, molecular pathways, cell cycle quiescence, and microRNA signaling present challenges in developing cancer SC-specific therapeutics. Treatment against cancer SCs should be developed targeting known stem cell regulatory pathways that are deregulated in cancer SCs compared to normal SC, as well as through unbiased high-throughput siRNA or small molecule screening. Both experimental approaches require identification and characterization of the putative liver cancer SC in order to target liver cancer SC specifically to decrease the toxicities. The current strategies of identifying cancer SCs are based on the expression of extracellular markers, the growth of cancer SCs, and greater tumorigenicity in immunedeficient mice compared cancer cells that are not cancer stem cells. Despite the amount of literature on liver cancer SCs, it is still not clear as to what constitutes a universal liver cancer SC-specific profile. Given the fact that the diverse etiology for hepatocarcinogenesis and multiple types of progenitor cells are involved in malignant transformation, it is unlikely that a universal liver cancer SC-specific profile will be used for therapeutics development. Ultimately, targeting liver cancer

| Table 1: Markers that have aided in the identification of stem cells. |
|-----------------|-----------------|
| **Markers**     |                  |
| Cluster of differentiation (CD) 133+ |                  |
| CD44+           |                  |
| CD45-           |                  |
| CD90⁺           |                  |
| CD34            |                  |
| OV6             |                  |
| Side population (SP) |          |
| Epithelial cell adhesion molecule (EpCAM) |          |
| OC.2, OC.3, OC.4, OC.5, OC.10 |          |
| BDS7            |                  |
| Thy-1           |                  |
| c-kit           |                  |
| ABCG2/BCRP1 (breast cancer resistance protein) |          |
| Connexin 43     |                  |
| Tumor rejection antigen 1-81 (TRA-1-81) |          |
| TRA-1-60        |                  |
| Sry-box containing gene 2 (SOX2) |          |
| Surface antigen stage-specific embryonic antigen 3 (SSEA-3) |          |
| CK7, CK19, CK14 |                  |
| α-fetoprotein (AFP) |          |
| γ-glutamyltranspeptidase |          |
| Placental form of glutathione-S-transferase |          |
| Flt-3 ligand    |                  |
| DMBT1 (deleted in malignant brain tumor 1) |          |
| Neural cell adhesion molecule 1 (NCAM)/CD56 |          |
| Chromogranin A  |                  |
| Parathyroid hormone related peptide (PTHrP) |          |
SCs in treating HCC will be in the context of personalized medicine.

7. Concluding Remark

Current research supports that HCC derived from malignant transformation of HPC. There may be at least three distinct cell lineages with progenitor cell properties susceptible to neoplastic transformation: hepatocyte, oval/hepatic progenitor cells, and bone marrow-derived stem cells. Multiple signaling transduction pathways important for stem cell proliferation and differentiations are found deregulated during hepatocarcinogenesis. Strategies are being developed to identify and characterize the liver cancer SCs. Targeting liver cancer SCs may bring hope in curing HCC.

References

[1] A. Jemal, R. Siegel, J. Xu, and E. Ward, “Cancer statistics, 2010,” CA: A Cancer Journal for Clinicians, vol. 60, no. 5, pp. 277–300, 2010.
[2] F. X. Bosch, J. Ribes, M. Diaz, and R. Cleries, “Primary liver cancer: worldwide incidence and trends,” Gastroenterology, vol. 127, no. 5, supplement 1, pp. S5–S16, 2004.
[3] B. K. Edwards, E. Ward, B. A. Kohler et al., “Annual report to the nation on the status of cancer, 1975–2006, featuring colorectal cancer trends and impact of interventions (risk factors, screening, and treatment) to reduce future rates,” Cancer, vol. 116, no. 3, pp. 544–573, 2010.
[4] S. F. Altekruse, K. A. McGlynn, and M. E. Reichman, “Hepatocellular carcinoma incidence, mortality, and survival trends in the United States from 1975 to 2005,” Journal of Clinical Oncology, vol. 27, no. 9, pp. 1485–1491, 2009.
[5] M. Esumi, T. Aritaka, M. Ari et al., “Clonal origin of human hepatoma determined by integration of hepatitis B virus DNA,” Cancer Research, vol. 46, no. 11, pp. 5767–5771, 1986.
[6] S. H. Zhang, W. M. Cong, and M. C. Wu, “Focal nodular hyperplasia with concomitant hepatocellular carcinoma: a case report and clonal analysis,” Journal of Clinical Pathology, vol. 57, no. 5, pp. 556–559, 2004.
[7] S. Sell, “Mouse models to study the interaction of risk factors for human liver cancer,” Cancer Research, vol. 63, no. 22, pp. 7553–7562, 2003.
[8] N. Fausto, “Liver regeneration and repair: hepatocytes, progenitor cells, and stem cells,” Hepatology, vol. 39, no. 6, pp. 1477–1487, 2004.
[9] G. K. Michalopoulos and M. C. DeFrances, “Liver regeneration,” Science, vol. 276, no. 5309, pp. 60–66, 1997.
[10] K. Overturf, M. al-Dhalimi, C. N. Ou, M. Finegold, and M. Grompe, “Serial transplantation reveals the stem-cell-like regenerative potential of adult mouse hepatocytes,” American Journal of Pathology, vol. 151, no. 5, pp. 1273–1280, 1997.
[11] O. Falkowski, H. J. An, I. A. Ianus et al., “Regeneration of hepatocyte ‘buds’ in cirrhosis from intrabiliary stem cells,” Journal of Hepatology, vol. 39, no. 3, pp. 357–364, 2003.
[12] V. M. Craddock, “Effect of a single treatment with the alkylating carcinogens dimethylnitrosamine, diethylnitrosamine and methyl methanesulphonate, on liver regenerating after partial hepatectomy. II. Alkylation of DNA and inhibition of DNA replication,” Chemico-Biological Interactions, vol. 10, no. 5, pp. 323–332, 1975.
[13] G. M. Williams, R. Gebhardt, H. Sirma, and F. Stenback, “Non-linearity of neoplastic conversion induced in rat liver by low exposures to diethylnitrosamine,” Carcinogenesis, vol. 14, no. 10, pp. 2149–2156, 1993.
[14] S. Yang, A. Koteish, H. Lin et al., “Oval cells compensate for damage and replicative senescence of mature hepatocytes in mice with fatty liver disease,” Hepatology, vol. 39, no. 2, pp. 403–411, 2004.
[15] L. Libbrecht and T. Roskams, “Hepatic progenitor cells in human liver diseases,” Seminars in Cell and Developmental Biology, vol. 13, no. 6, pp. 389–396, 2002.
[16] N. D. Theise, J. L. Yao, K. Harada et al., “Hepatic ‘stem cell’ malignancies in adults: four cases,” Histopathology, vol. 43, no. 3, pp. 263–271, 2003.
[17] M. L. Dumble, E. J. Croager, G. C. T. Yeoh, and E. A. Quail, “Generation and characterization of p53 null transformed hepatic progenitor cells: oval cells give rise to hepatocellular carcinoma,” Carcinogenesis, vol. 23, no. 3, pp. 435–445, 2002.
[18] B. Knight, J. E. Tirinzzi-Parker, and J. K. Olynyk, “C-kit inhibition by imatinib mesylate attenuates progenitor cell expansion and inhibits liver tumor formation in mice,” Gastroenterology, vol. 135, no. 3, pp. 969.e1–979.e1, 2008.
[19] B. E. Petersen, W. C. Bowen, K. D. Patrene et al., “Bone marrow as a potential source of hepatic oval cells,” Science, vol. 284, no. 5417, pp. 1168–1170, 1999.
[20] H. Ishikawa, K. Nakao, K. Matsumoto et al., “Bone marrow engraftment in a rodent model of chemical carcinogenesis but no role in the histogenesis of hepatocellular carcinoma,” Gut, vol. 53, no. 6, pp. 884–889, 2004.
[21] S. Ma, K. W. Chan, L. Hu et al., “Identification and characterization of tumorigenic liver cancer stem/progenitor cells,” Gastroenterology, vol. 132, no. 7, pp. 2542–2556, 2007.
[22] Z. F. Yang, D. W. Ho, M. N. Ng et al., “Significance of CD90+ cancer stem cells in human liver cancer,” Cancer Cell, vol. 13, no. 2, pp. 153–166, 2008.
[23] Z. F. Yang, P. Ngai, D. W. Ho et al., “Identification of local and circulating cancer stem cells in human liver cancer,” Hepatology, vol. 47, no. 3, pp. 919–928, 2008.
[24] T. Yamashita, M. Forgues, W. Wang et al., “EpCAM and alpha-fetoprotein expression defines novel prognostic subtypes of hepatocellular carcinoma,” Cancer Research, vol. 68, no. 5, pp. 1451–1461, 2008.
[25] T. Chiba, K. Kita, Y. W. Zheng et al., “Side population purified from hepatocellular carcinoma cells harbors cancer stem cell-like properties,” Hepatology, vol. 44, no. 1, pp. 240–251, 2006.
[26] L. Mishra, T. Banker, J. Murray et al., “Liver stem cells and hepatocellular carcinoma,” Hepatology, vol. 49, no. 1, pp. 318–329, 2009.
[27] J. U. Marquardt, V. M. Factor, and S. S. Thorpeairson, “Epigenetic regulation of cancer stem cells in liver cancer: current concepts and clinical implications,” Journal of Hepatology, vol. 53, no. 3, pp. 568–577, 2010.
[28] S. Boyault, D. S. Rickman, A. de Reynies et al., “Transcriptome classification of HCC is related to gene alterations and to new therapeutic targets,” Hepatology, vol. 45, no. 1, pp. 42–52, 2007.
[29] M. Branda and J. R. Wands, “Signal transduction cascades and hepatitis B and C related hepatocellular carcinoma,” Hepatology, vol. 43, no. 5, pp. 981–992, 2006.
[30] Y. Li, B. Welm, K. Podosyspania et al., “Evidence that transgenes encoding components of the Wnt signaling pathway preferentially induce mammary cancers from progenitor cells,” Proceedings of the National Academy of Sciences of the United States of America, vol. 100, no. 26, pp. 15853–15858, 2003.
[31] M. Le Pierre et al., “Small-molecule antagonists of the oncogenic Tcf/beta-catenin protein complex,” Cancer Cell, vol. 5, no. 1, pp. 91–102, 2004.

[32] A. H. Huber and W. I. Weis, “The structure of the beta-catenin/E-cadherin complex and the molecular basis of diverse ligand recognition by beta-catenin,” Cell, vol. 105, no. 3, pp. 391–402, 2001.

[33] R. H. Giles, J. H. van Es, and H. Clevers, “Caught up in a Wnt storm: Wnt signaling in cancer,” Biochimica et Biophysica Acta, vol. 1653, no. 1, pp. 1–24, 2003.

[34] Y. Teng, X. Wang, Y. Wang, and D. Ma, “Wnt/beta-catenin signaling regulates cancer stem cells in lung cancer A549 cells,” Biochemical and Biophysical Research Communications, vol. 392, no. 3, pp. 373–379, 2010.

[35] S. Inagawa, M. Itabashi, S. Adachi et al., “Expression and prognostic roles of beta-catenin in hepatocellular carcinoma: correlation with tumor progression and postoperative survival,” Clinical Cancer Research, vol. 8, no. 2, pp. 450–456, 2002.

[36] T. Yamashita, J. Ji, A. Budhu et al., “EpCAM-positive hepatocellular carcinoma cells are tumor-initiating cells with stem/progenitor cell features,” Gastroenterology, vol. 136, no. 3, pp. 1012–1024, 2009.

[37] M. Weinstein, X. Yang, and C. X. Deng, “Functions of mammalian Smad genes as revealed by targeted gene disruption in mice,” Cytokine and Growth Factor Reviews, vol. 11, no. 1-2, pp. 49–58, 2000.

[38] H. Chang, C. W. Brown, and M. M. Matzuk, “Genetic analysis of the mammalian transforming growth factor-beta superfamily,” Endocrine Reviews, vol. 23, no. 6, pp. 787–823, 2002.

[39] J. Massague, S. W. Blain, and R. S. Lo, “TGFbeta signaling in growth control, cancer, and heritable disorders,” Cell, vol. 103, no. 2, pp. 295–309, 2000.

[40] Y. Tang, K. Kitisin, W. Jogunoori et al., “Progenitor/stem cells give rise to liver cancer due to aberrant TGF-beta and IL-6 signaling,” Proceedings of the National Academy of Sciences of the United States of America, vol. 105, no. 7, pp. 2445–2450, 2008.

[41] J. Taipale and P. A. Beachy, “The Hedgehog and Wnt signalling pathways in cancer,” Nature, vol. 411, no. 6835, pp. 349–354, 2001.

[42] J. K. Sicklick, Y. X. Li, A. Jayaraman et al., “Dysregulation of the Hedgehog pathway in human hepatocarcinogenesis,” Carcinogenesis, vol. 27, no. 4, pp. 748–757, 2006.

[43] S. Huang, J. He, X. Zhang et al., “Activation of the hedgehog pathway in human hepatocellular carcinomas,” Carcinogenesis, vol. 27, no. 7, pp. 1334–1340, 2006.

[44] Q. Wang, S. Huang, L. Yang et al., “Down-regulation of Sonic hedgehog signaling pathway activity is involved in 5-fluorouracil-induced apoptosis and motility inhibition in Hep3B cells,” Acta Biochimica et Biophysica Sinica, vol. 40, no. 9, pp. 819–829, 2008.

[45] A. Pannuti, K. Foreman, P. Rizzo et al., “Targeting Notch to target cancer stem cells,” Clinical Cancer Research, vol. 16, no. 12, pp. 3141–3152, 2010.