Overexpression of TGF-α and EGFR, a key event in liver carcinogenesis, is induced by hypoxia specifically in hepatocytes

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Abstract

Transforming growth factor alpha (TGF-α) is a mitogenic factor for hepatocyte and a ligand of the epithelial growth factor receptor (EGFR). TGF-α promotes liver carcinogenesis. TGF-α is also overexpressed in regenerative nodules of the cirrhotic liver but the mechanism of this expression is poorly known. Because hypoxia is a feature of cirrhotic livers and hypoxia may induce TGF-α and EGFR expressions, the aim of this study was to determine whether the TGF-α/EGFR pathway is affected by hypoxia in liver cells. Cell isolates were prepared from normal Wistar rats. Liver myofibroblasts were obtained in culture by activation of hepatic stellate cells (HSC), and by outgrowth of portal myofibroblasts from bile duct segments. Hepatocytes, Kupffer cells and liver myofibroblasts in culture were submitted to hypoxia for 4-24 hours. Hypoxia was achieved using a catalytic system, which reduces oxygen concentration to less than 1% within 30 minutes. The absence of toxicity was verified by lactate dehydrogenase dosing in cell supernatant. Vascular endothelial growth factor (VEGF) served as a hypoxia-inducible control gene. Gene expression was assessed by real-time reverse transcription polymerase chain reaction (RT-PCR). Under normoxia, the expression of TGF-α was significantly higher in hepatocytes than in non-parenchymal liver cells (~1.7-fold). EGFR transcripts were also more abundant in Hepatocytes than in myofibroblasts (~3-fold) or in Kupffer cells (~22-fold). Hypoxia induced an increase in VEGF mRNA to a similar extent in all cell types. By contrast, hypoxia caused an increase in TGF-α transcripts mainly in Hepatocytes (112 ± 7 vs 32 ± 2 under normoxia), also but to a lesser extent in portal myofibroblasts (35 ± 5 vs 17 ± 4), but not in HSC-derived myofibroblasts nor in Kupffer cells. An increase in EGFR expression was induced by hypoxia also predominantly in Hepatocytes (125 ± 12 vs 44 ± 6), and to a much lesser extent in other cell types. These results demonstrate that hypoxia induces TGF-α and EGFR overexpression in hepatocytes and, thereby, might act as a promoting event in liver carcinogenesis upon cirrhotic liver.

Introduction

TGF-α is a mitogenic factor for hepatocytes and a ligand of the EGF receptor (EGFR). TGF-α can promote liver carcinogenesis, as illustrated in TGF-α transgenic mice, which constantly develop hepatocellular carcinoma [1]. TGF-α is also overexpressed in regenerative nodules of the cirrhotic liver but the reason for this expression is unknown. Because local hypoxia is a constant feature of cirrhotic livers and hypoxia may induce TGF-α and EGFR expressions, the aim of this study was to determine whether the TGF-α/EGFR pathway may be affected by hypoxia in liver cells.

Materials and methods

Cell isolation and culture

Hepatocytes were isolated from normal Wistar rats by a method derived from Seglen [2]. Experiments were conducted in compliance with the national ethical guidelines for the care and use of laboratory animals. All experiments were performed after the approval of Institutional Animal Use and Care Committee. The animals were anesthetized with a subcutaneous injection of chlorpromazine (2 mg/kg) and ketamine (20 mg/kg). The liver was perfused in situ, for 15 min with a Ca²⁺-free 10 mM Hepes buffer at a flow rate of 30 ml/min⁻¹, and for 15 min with 0.025 % collagenase B (Boehringer Mannheim, Meylan, France) in Ca²⁺-containing 10 mM Hepes buffer at a flow rate of 20 ml.min⁻¹. The hepatocytes were then separated from the connective tissue by gentle agitation in Leibovitz-15 (L15) medium (Sigma) containing 0.2 % bovine serum albumin (BSA, Sigma). The resulting cell suspension was filtered through gauze, allowed to sediment for 20 min at 4°C, and washed in BSA-containing L15. Hepatocytes were 80-85% pure, as assessed by characteristic cell size under phase-contrast microscopy, and cell viability exceeded 90%. Hepatocytes were plated onto 100 mm collagen I-coated culture dishes at a density of 1.0-1.2 x 10⁶ cells/cm² in 10 ml William’s medium E (GIBCO BRL, Life Technologies, Cergy-Pontoise) containing 10% fetal bovine serum (GIBCO BRL), 5 mM Hepes buffer, 5 µg/mL insulin (Novo Nordisk, Boulogne Billancourt, France) containing 10% fetal bovine serum (GIBCO BRL), 5 mM Hepes buffer, 5 µg/mL insulin (Novo Nordisk, Boulogne Billancourt, France), and 100,000 IU/100 mg/L penicillin-streptomycin (GIBCO BRL). After 3 h, the medium was replaced by a serum-free medium supplemented with 1 µM hydrocortisone 21-hemisuccinate (Sigma). Medium was changed 24 h after plating, and then hepatocytes were subjected to hypoxic treatment. Liver myofibroblasts were obtained in culture by activation.
of hepatic stellate cells HSC (HSC), and by outgrowth of portal myofibroblasts from bile duct segments. Hepatic stellate cells (HSC) were isolated by an established method [3] with modifications [4]. The purity of HSC isolates was higher than 99%, as assessed by fluorescence of retinoid-containing vacuoles under ultraviolet excitation, and cell viability exceeded 90%, as tested by erythrosine exclusion. Portal myofibroblasts were obtained as previously described [5]. HSC and portal myofibroblasts were grown on uncoated plastic in 75 cm² flasks until confluence in DMEM (Sigma), containing 1% penicillin/streptomycin, 1% Hepes and 10% fetal bovine serum. The cells were maintained at 37°C, under air/CO₂ 95 %/5 %. Medium was changed 24 hours after plating, and every 48 hours thereafter until confluence. Liver myofibroblasts were used within passages 3 and 5. Cultures were serum-starved just before incubation under hypoxic conditions. Kupffer cells were isolated from liver cell suspension and separated by density gradient centrifugation through 8.2 % Nycodenz (Sigma) interface. Cells were washed in 25 mL MEM-E at 500g for 7 min, suspended in Medium 199 supplemented with 20% serum, and plated in 60-mm uncoated plastic tissue culture dishes at a concentration of 4-5 × 10⁵ cells/mL. [3]. After 20 min, nonadherent cells and residual serum were washed off the monolayer with L-15 salts and then replaced with fresh medium containing 20% serum [6]. Twenty-four hours later the culture was incubated for 3 min at 37°C with L-15 salts containing 0.5% trypsin and 0.02% EDTA. EDTA/trypsinized cells were removed by gently aspirating the monolayer with a sterile pipet and the culture was continued by addition of fresh medium containing 20% serum and then Kupffer cells were subjected to hypoxic treatment.

Cell hypoxia

Hypoxic condition was achieved using AnaeroGen system (Oxoid, Dardilly, France), which catalytically reduces oxygen concentration to less than 1% within 30 minutes. Culture dishes were placed into a 2.5 l airtight jar with an AnaeroGen sachet, and the lid was closed immediately. The jar was then incubated at 37°C until opening. After a designated time period, the jar was opened, and the cells were immediately lysed in order to extract RNAs and proteins. Controls included parallel cultures in which cells were subjected to normoxia (21% oxygen tension). In preliminary experiments, we defined durations involved in hypoxia (125 ± 12 vs 44 ± 6, figure 1, middle panel), and to a much lesser extent in portal myofibroblasts (35 ± 5 vs 17 ± 4, Figure 1, upper panel), but not in HSC-derived myofibroblasts nor in Kupffer cells. An increase in EGFR expression was induced by hypoxia also predominantly in hepatocytes (125 ± 12 vs 44 ± 6, figure 1, middle panel), and to a much lesser extent in other cell types (Figure 1, middle panel).

Reverse-transcription and real time polymerase chain reaction

Total RNA was extracted from cells, using RNeasy Mini kit (Qiagen). Complementary DNA was synthesized from total RNA (5 pg), using the Moloney Murine Leukemia Virus (MMLV) reverse transcriptase (Life Technologies) and pd(N), primers (Pharmacia Biotech). Quantitative real-time polymerase chain reaction was performed using the Sybr Green PCR Core Reagents Kit (PE Applied Biosystems) on a PE Applied Biosystems 7700 Sequence Detector [4]. The primers were designed according to published rat cDNA sequences in Genbank database, using the Primer Express software v1.5 (PE Applied Biosystems) 

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Discussion and conclusion

Cellular responses to low oxygen tension are primarily mediated by the hypoxia inducible factors (HIFs) [7,8]. Two oxygen labile subunits (HIF-1α and HIF-2α) have been identified as the sensors that are co-opted to mediate adaptive transcription changes involved in tumour growth pathways [9,10]. Well-characterized tumour promoting functions of both HIF-1α and HIF-2α include expression of growth factors promoting proliferation (TGF-α) [11] as well as angiogenesis (VEGF) [10,12]. Furthermore, studies have shown that lifting of hypoxia would decrease the expression of HIF-α thus inhibiting angiogenesis and tumour growth [10].

TGF-α is a known ligand to EGFR, which is overexpressed as result of HIF-2α activation [12]. The EGFR pathway is basically implicated in hepatocarcinoma emergence [13] and represents a target for antitumoral therapy (tyrosine kinase inhibitors) [14,15].

Our results show that hypoxia stimulates to a similar extent, the expression of EGFR, TGF-α and VEGF, in hepatocytes. By contrast, in HSC-derived myofibroblasts, hypoxia induced the expression of VEGF, but not that of TGF-α or EGFR. The central role of liver myofibroblasts in hypoxia-induced angiogenesis during hepatic wound healing has already been shown by Corpechot et al. [16]. Hepatic angiogenesis can be triggered not only by hypoxia, but also by a chronic pseudohypoxic state mediated by the accumulation of HIF-1α in HSC [17].

Kupffer cells responded to hypoxia by overexpressing TGF-α expression whereas portal myofibroblasts showed an increase in the expression of all three genes, TGF-α, EGFR and VEGF, but in lesser proportion compared to hepatocytes.
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Figure 1. Response of the different liver cell types to hypoxia. Hepatocytes, Kupffer cells (KF), hepatic stellate cell-derived myofibroblasts (HSC-MF), and portal myofibroblasts (PMF), in culture, were maintained in normoxia (□) or exposed to hypoxia (■), and then subjected to TGF-α, EGFR and VEGF mRNA quantification by reverse-transcription real-time polymerase chain reaction. *P value< 0.01, between hypoxia and normoxia.
In conclusion, the present study provides further evidence to indicate that hepatocellular hypoxia enhances both angiogenesis and cell proliferation and that different liver cell types are implicated in growth factors pathways which are implicated in hypoxia-related cell proliferation. We suggest that hypoxia-induced TGF-α and EGFR overexpression in hepatocytes might act as a promoting event in liver carcinogenesis in cirrhosis and could be a potential target for liver cancer prevention and therapy.

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Statement of ethics

Experiments were conducted in compliance with the national ethical guidelines for the care and use of laboratory animals. All experiments were performed after the approval of Institutional Animal Use and Care Committee. No human subjects were submitted to any experimental testing.

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