Erythropoietin Production by Fetal Mouse Liver Cells in Response to Hypoxia and Adenylate Cyclase Stimulation*

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ABSTRACT. This study was done to investigate aspects of control of extrarenal erythropoietin (Ep) production. To this end we studied the effects of three stimuli of renal Ep production in the adult, i.e. hypoxia, cobalt, and activation of adenylate cyclase on Ep generation by cultured fetal mouse liver cells. The fetal liver was taken as a model for extrarenal Ep production because this organ is considered the predominant site of extrarenal Ep production. We found that Ep production by the cells increased as the oxygen concentration was decreased in the incubation atmosphere from 20% to 1%. Cobalt (10⁻⁴-10⁻⁵ M) had no effect on Ep production. Activation of adenylate cyclase by forskolin (10⁻⁵ M) or isoproterenol (10⁻⁶ M) greatly enhanced Ep production. These findings indicate that the Ep-stimulating effect of cobalt is specific for the kidney. However, oxygen depletion and activation of adenylate cyclase seem to be more general stimuli in Ep-producing cells. Furthermore we found that Ep production in hypoxia correlated with lactate formation in the cultured liver cells. This finding suggests that Ep production in fetal livers under hypoxic conditions parallels the shift from aerobic to anaerobic cellular energy metabolism.

Erythropoietin (Ep) is a growth factor known to stimulate erythropoiesis. Ep has been demonstrated in a variety of tissues including kidney (1), liver (2), submandibulary glands (3), and macrophages (4). Although the existence of Ep in an organ does not imply production by the organ the liver is considered as the primary site of Ep production during fetal life (5, 6). After birth Ep production shifts to the kidney (7, 8), and in the adult extrarenal Ep production accounts for less than 15% of the total (9). There is considerable information regarding the control of Ep production by the adult kidney. In contrast, information regarding the regulation of extrarenal Ep production is limited, and it is not clear whether extrarenal Ep production is activated by the same stimuli as renal Ep production. Recently, Zucali and Mirand (10) reported in a series of investigations that cultures of fetal mouse liver cells are capable of producing Ep, and we have shown that renal cell cultures are suitable for study of the mechanisms of Ep production in the kidney (11). Thus, in the present studies, we employed cultures of fetal mouse livers to investigate the effects of hypoxia, cobalt, and activation of the adenylate cyclase by isoproterenol and forskolin on hepatic Ep production. We found that hypoxia and activation of the adenylate cyclase enhanced Ep formation whereas cobalt had no effect. Furthermore, we obtained evidence that Ep production stimulated by hypoxia tends to parallel the increase in lactate production which marks the shift of cellular energy metabolism from aerobic to anaerobic pathways.

Materials and Methods

Cell cultures of fetal mouse livers

Fetal mouse liver cell cultures were established basically as described by Zucali and Mirand (12). In brief, fetal livers were obtained from NMRI-strain mice (Dr. Ivanovas, Kisslegg, West Germany) at the 13th to 14th day of gestation. Livers were aseptically removed from the fetuses and subsequently disaggregated in a single cell suspension by vigorous pipetting with a Pasteur pipette. Cell debris was removed by washing the cell suspension twice with culture medium (RPMI 1640, 100 U/ml penicillin, 100 μg/ml streptomycin, 10% fetal bovine serum; all agents were from Boehringer, Mannheim, West Germany). Cells were seeded at a density of 1 × 10⁶ nucleated cells/cm² into 75 cm² culture flasks (Greiner, West Germany) containing 10 ml culture medium. Cells were cultivated in incubators.
maintaining a humidified gas atmosphere 1:4:15 of carbon dioxide-oxygen-nitrogen at 37 C, respectively. The first renewal of the culture medium was done 1 day after seeding of the cells. Most nonadherent cells were removed by this procedure. The next refeeding of the cells was done 3 days later (day 4 of culture).

Studies on the Ep production rate

Ep production rate studies of the cultures were done between the fourth and seventh day of culture, using a culture medium that contained 5% instead of 10% fetal bovine serum. For determination of the effect of oxygen tension on the Ep production rate, cultures were kept at either 20%, 10%, 5%, 2%, or 1% oxygen for 3 days between the fourth and seventh day of culture using O2 controlled incubators (Heraeus, Hanau, West Germany). The effects of cobalt, forskolin, and isoproterenol on Ep production were examined by adding aliquots of freshly prepared stock solutions of cobaltous chloride (10^-2 M), forskolin (10^-5 M), and isoproterenol (10^-5 M) to the cultures for 3 days and at 5% oxygen. PO2 was determined in the gas atmosphere in the incubator using a PO2 electrode.

Ep activity

Ep activity in the culture medium was assayed with the posthypoxic polycythemic mice assay for Ep exactly as described (13). In brief, NMRI-strain female mice (25 g) were made polycythemic by exposure to 0.42 atmosphere for 20 h/day for 2 weeks. On the sixth and seventh posthypoxic days, five to six mice per sample were injected sc with standard doses of human urinary Ep (calibrated against the International Standard Preparation B) or with the unknown samples. On the eighth day, each mouse received 0.5 μCi 59Fe-citrate (Amersham Buchler, Braunschweig, West Germany). Forty-eight hours later, the mice were bled by heart puncture and the percentage of radioiron incorporation into the blood was determined. Ep titers in the unknowns were calculated from log dose-log response curves of the Ep standards. Figure 1 shows an averaged dose response curve of seven different assays.

If not otherwise indicated, culture media from 3-day incubations were assayed. Heretofore the supernatants of 30 (75 cm^2) flasks (300 ml) were pooled and concentrated 60-fold by ultrafiltration on Amicon YM-10 membranes. Each assay mouse received 2 × 0.5 ml 60-fold concentrated medium. Ep activities of culture media shown in Results were corrected for the Ep activity of pure culture medium which was determined in each assay as a control. Sixty-fold concentrated control medium caused an iron incorporation of 1.02 ± 0.12% in five different assays. From the double-logarithmic plotting of the dose response curve for human standard Ep (Fig. 1) the mean Ep activity of 60-fold concentrated control medium was calculated as 15 mU/ml.

Lactate concentrations

Lactate concentrations in the culture media were determined with a commercial kit for l-lactate (Boehringer).

Determination of intracellular cAMP levels

To determine intracellular cAMP levels, the culture medium was substituted by prewarmed L-15 medium (Boehringer) with or without forskolin (10^-5 M) or isoproterenol (10^-6 M), and the culture dishes (7 cm^2) were placed on a heater maintaining a temperature of 37 C. Five minutes after addition the L-15 medium was withdrawn and the dishes were placed on an ice block. After addition of 0.4 ml ice-cold buffer (containing 5 mM potassium phosphate, 0.2 mM EDTA, 0.5 mM 3-isobutyl-methylxanthine, and 150 mM KCl, pH 6.8) the cells were scraped off with a teflon policeman. The obtained suspension was sonicated, boiled for 5 min, and centrifugated. The supernatants were assayed for cAMP using a cAMP binding assay for cAMP exactly as described (14).

Isoproterenol concentrations

Isoproterenol concentrations in the samples assayed for Ep in the polycythemic mice were determined by HPLC. Protein was precipitated by addition of perchloric acid (final concentration, 0.1 M) to each sample. After centrifugation, aliquots of the supernatants were analyzed on a prepacked column (Hyperchrome C 18, 5 μm, Bischoff, Dören, West Germany). The solvent system for the isocratic elution was buffer-methanol = 93:7 (buffer: L-octylsulfonic acid, sodium salt, 12 mg/liter; citric acid-monohydrate, 8 g/liter; Na2EDTA, 50 mg/liter; pH 4.0 in water). The column was calibrated with isoproterenol dissolved in 0.1 M perchloric acid.

Results

Effect of hypoxia on Ep formation of cultured fetal livers

Figure 2a shows the Ep production rate of the cultured livers at different oxygen tensions in the incubation
It is evident that the Ep production rate increased as the pO$_2$ decreased in the incubation atmosphere from 120 to 7 torr. The Ep production rate at 7 torr was about 6-fold higher than that at 120 torr. The oxygen dependent Ep production rate ($y$) can be described as an inverse function of the pO$_2$ ($x$) as follows: 

$$y = \frac{156}{x} + 4.2; \ r = 0.97.$$ 

Figure 2b shows the lactate formation rate of the cultured livers at different oxygen tensions. It is obvious that lactate production by the cultures also increased with decreasing pO$_2$. The oxygen dependence of lactate formation ($y$) also can be described by the inverse function of pO$_2$ ($y = \frac{5424}{x} + 72; \ r = 0.94$).

The effect of cobaltous chloride (10$^{-4}$ M and 10$^{-5}$ M) on Ep formation at 20% and 5% oxygen was examined three times, but in no case could a stimulation of Ep formation by cobalt be observed.

**Effect of adenylate cyclase stimulation on Ep formation by cultured fetal livers**

Isoproterenol and forskolin are well documented stimulators of adenylate cyclase (15). To ensure that these agents also stimulate adenylate cyclase of cultured fetal livers cells, we determined intracellular cAMP levels at 5 and 30 min after addition of isoproterenol and forskolin to the cells. Table 1 shows that both isoproterenol (10$^{-5}$ M) and forskolin (10$^{-5}$ M) caused significant increases in intracellular cAMP levels. The effects of these agents on Ep production by the cultures are shown in Fig. 3a. Both agents enhanced Ep production by a factor of 7-8. As a control we determined the effect of culture medium that had been incubated with isoproterenol (10$^{-5}$ M) or forskolin (10$^{-5}$ M) for 3 days. These controls failed to stimulate iron incorporation into the assay mice. Since high concentrations of isoproterenol are known to enhance iron incorporation in mice (16), we determined the isoproterenol concentrations in the medium samples by HPLC. It turned out that the concentration of isoproterenol was lowered from 10$^{-5}$-10$^{-7}$ M after a 3-day incubation period.

Figure 3b shows the lactate formation of the cultured livers in the presence of isoproterenol or forskolin. It is evident that the production of lactate was not stimulated by isoproterenol or by forskolin.

**Discussion**

To determine whether the same stimuli of Ep production in the adult kidney also are active in the fetal liver, we investigated the effects of hypoxia, cobalt, and activation of adenylate cyclase on Ep production by cultured fetal mouse liver cells. We found that the production of Ep by the cultured fetal livers was increased by lowering the oxygen tension in the incubation atmosphere. This finding is in agreement with the results of a study by Zucali (17) who reported a 2-fold increase in Ep production by fetal mouse liver cultures, when the oxygen concentration in the incubator was lowered from 19% to 13%. Note that we found a more significant stimulation of Ep production at much lower oxygen concentrations...
Our results also are in agreement with earlier in vivo studies of Zanjani et al. (18) showing that the plasma Ep concentration in fetal blood is elevated during hypoxia. Philipps et al. (19) also found a dependence of plasma Ep concentrations on the oxygen content in the blood in in vivo experiments. It can be concluded, therefore, that the oxygen sensing mechanism which leads to an increased Ep production in fetal liver is not disturbed by cultivation of the cells. Our results furthermore show that the lactate production rate of the cultured liver had a very similar oxygen dependence as the Ep production rate (Fig. 2). If one accepts lactate production as an index for the shift of the cellular energy metabolism from the aerobic to the anaerobic pathway, one can conclude from this result that Ep production in the fetal liver tends to parallel the shift of the energy metabolism into the anaerobic pathway. Interestingly we found the same relationship between lactate and Ep production in cultures of renal mesangial cells (unpublished results). A comparison of the oxygen dependencies of Ep production in cultured fetal liver and cultured mesangial cells suggests that the oxygen sensors of both tissues have similar sensitivity under cell culture conditions (Fig. 4).

In contrast to its effect on the kidney, cobalt had no effect on Ep production by cultured liver cells. Since the mechanism by which cobalt enhances Ep production in the kidney is not well understood, we cannot conclude from our finding which specific pathway of stimulating Ep formation is lacking in the fetal liver.

Prostaglandins (20) and β-adrenergic agonists (16) are potent stimulators of renal Ep production. Since these agents are considered to act mainly by activating adenylate cyclase (23), we investigated the effects of the adenylate cyclase activators isoproterenol and forskolin on Ep production in the cultured fetal liver cells. Both agents enhanced cAMP levels and Ep production by the cells. It might be argued that erythropoietic activity of these culture media could have resulted from the agents...
being carried over into the assay mice rather than by Ep itself; isoproterenol can indeed enhance radioiron incorporation in the mouse (16). However, control media containing isoproterenol or forskolin failed to stimulate iron incorporation. Furthermore we determined the isoproterenol concentrations in the samples by HPLC before assay of Ep and found that the samples contained less than \(10^{-7} \text{M} \) isoproterenol. We calculated that one assay mouse received less than 0.5 \(\mu\)g isoproterenol/day and kg body weight. Thus a small dose of isoproterenol cannot account for the measured Ep activity of about 360 mU/ml (16). The lowered concentration of isoproterenol after 3 days of culture is not unexpected in view of the short half-life of isoproterenol in the presence of oxygen and nonacid pH values (22). Our results, therefore, further support the idea that activation of adenylate cyclase enhances Ep production in Ep-generating fetal liver cells. It is unlikely, also, that isoproterenol exerted its effect on Ep production by increasing the cellular oxygen consumption causing intracellular hypoxia (23). We found that isoproterenol did not stimulate lactate formation by the cultured livers.

The question arises whether the mechanism(s) by which hypoxia and activation of adenylate cyclase stimulate Ep synthesis are related. In cultures of mesangial cells we had obtained evidence that activation of adenylate cyclase mediates the effect of hypoxia on Ep formation (24). Adenylate cyclase in these cells probably was activated by prostaglandin \(E_2\), for which an increased synthesis during hypoxia was found (24). Preliminary studies suggest that prostaglandins may be involved in the stimulation by hypoxia of Ep production in fetal liver cells. When the prostaglandin synthase inhibitor indomethacin \((10^{-5} \text{M})\) was added to cultured fetal livers, Ep formation was abolished even in presence of an incubator oxygen tension of 7 torr (unpublished observation). We reported a similar observation in cultured mesangial cells (24). It seems likely, therefore, that hypoxia stimulates Ep production in the kidney and in the fetal liver by enhancing the production of certain prostaglandins.

Finally we may consider the type of cell that could be responsible for the production of erythropoietin in liver cultures. Carbon particle ingestion studies (25, 26) and immunofluorescent studies (8, 27) have localized the site of hepatic Ep production to the Kupffer cells. More recently Paul et al. (27) clearly demonstrated that isolated rat Kupffer cells are capable of secreting Ep. Therefore, we determined the percentage of Kupffer cells in the present cultures, using their capability of phagocytosing latex beads. We found that primary cultures of fetal mouse liver cells regularly contained about 30% of latex phagocytosing cells between 4 and 7 days of culture. By a simple trypsinization procedure we obtained subcultures containing about 90% Kupffer cells. The culture media of these cells were repeatedly assayed for Ep activity in the polycythemic mouse assay, and in no instance was significant Ep activity detected. The culture density of Kupffer cells in these subcultures, however, was only one-fifth of that of the primary cultures. Considering the rather high detection limit for Ep of the polycythemic mouse assay it was not clear whether fetal Kupffer cells in cultures were incapable of Ep production, or whether Ep activity in the culture media of the Kupffer cells was low because of the low cellular density. The results of the study by Paul et al. (28), clearly support the second explanation.

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