Significance and fate of septal fibrosis of the liver

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Received 15 September 2005; received in revised form 24 January 2006; accepted 30 January 2006

Available online 24 March 2006

Abstract

Septal fibrosis commonly occurs during chronic diseases of the liver. It is experimentally reproduced in a proportion of rats treated with pig-serum, and in 100% of rats infected with *Capillaria hepatica*. These models have only been used in relatively short-term studies. To contribute to the natural history and significance of hepatic septal fibrosis it is important to disclose its fate after prolonged observation, and following partial or total withdrawal of its inciting cause.

Adult Wistar rats were sacrificed 3, 6, 9 and 12 months following inoculation of 800 embryonated eggs of *C. hepatica*. Besides routine histology, liver sections were submitted to immunohistochemical, immunofluorescence and ultrastructural techniques for the identification of cells and extracellular matrix components present in the fibrous septa. Septal blood vessels were studied after portal vein perfusion with India-ink, while the hepatic functional profile and levels of anti-*C. hepatica* antibodies were determined in collected sera.

Results revealed that all parasites were already dead 2 months from inoculation, and the accumulated eggs in the liver lost their capacity to embryonate around the 4th–6th month, when progressive reduction in the number of cells and in the amount of collagen occurred in the septa. Septal fibrosis persisted throughout the time of experimentation (12 months). This fibrosis was seen as a supporting stroma for septal vessels that conducted portal blood directly to the sinusoids. Thus, persistence of fibrosis was probably related to its morphological and functional association with blood vessels.

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Keywords: *Capillaria hepatica*; Septal hepatic fibrosis; Fibrosis reversibility

1. Introduction

A peculiar type of septal fibrosis of the liver regularly occurs in *Capillaria hepatica*-infected rats [1]. During its life cycle, this helminth dies out inside the liver, around the 20th day of infection, soon after sexual maturity and egg lying. Necro-inflammatory lesions are then formed around the disintegrating parasites and their eggs. When such focal parasitic lesions start exhibiting signs of encapsulation and progressive resorption, which usually takes place around the 25th–30th day following inoculation, thin and long fibrous septa begin to sprout out from portal spaces throughout the liver, dividing the hepatic parenchyma into several polyhedral portions of different sizes. Earlier studies have noted that end-stage cirrhosis can be the outcome of severe *C. hepatica* infection in rodents [2]. Significantly, Lammler et al. [3] pointed out that these late lesions are difficult to be explained, since at this time egg production is ceasing and acute focal lesions are no longer observed. As a matter of fact, when *C. hepatica*-induced lesions were followed through different intervals, from 1 to 3 months, the process of septal fibrosis was seen to increase progressively with time [1]. However, the outcome of *C. hepatica*-induced septal fibrosis is not known, since sufficiently prolonged studies have not been made on this regard.

Also, in a much more studied model of hepatic septal fibrosis, the one induced in rats by repeated intraperitoneal
injections of whole pig-serum or its albumin fraction, only relatively short-term studies have been attempted [4–8]. Whether septal fibrosis, as seen in both experimental models just mentioned, will continuously evolve toward cirrhosis or will eventually stop and disappear, are questions of considerable interest to the study, not only for the understanding of the respective models, but for the general pathology of hepatic fibrosis.

The present investigation followed the changes of septal fibrosis in the liver of rats infected with C. hepatica, at several time intervals, from 3 months to over a year. While the amount of collagen considerably decreased with the extinction of parasitism, a residual septal fibrosis persisted. The reason why such fibrosis became irreversible was then investigated.

2. Materials and methods

2.1. Animals

A total of 54 healthy, outbred Wistar rats, weighing 200–500 g, of both sexes, were used. They were maintained in good housing conditions, with free access to a commercial balanced diet and water. All the animals were submitted to a surgical biopsy, with removal of a liver lobe at different times after inoculation, as follows: 3 months (n = 15), 6 months (n = 15), 9 months (n = 9) and 12 months (n = 15). At each time-point, blood was collected from a tail vein, for serum samples. At this same time, two animals were sacrificed soon after their portal vein system had been perfuncted with India-ink. The injection was made manually until the liver surface appeared uniformly black. India-ink was diluted (50–50) in 15% gelatin. Normal intact control rats were used for the obtaining of serum samples and for the studies of the portal vessels after India-ink perfusion.

2.2. Histopathology

Fragments of the liver were fixed for at least 48 h in buffered (pH 7.2) 10% formalin. Paraffin-embedded sections were stained with hematoxylin and eosin, sirius-red for collagen, the PAS method with and without diastase digestion, Perls’ method for iron, toluidine-blue for elastic fibers, peroxidase-antiperoxidase method for auto-hemopoietic cells and immunoperoxidase method for factor VIII and desmin. Normal intact control rats were used for the obtaining of serum samples and for the studies of the portal vessels after India-ink perfusion.

2.3. Transmission electron microscopy

Tiny fragments of liver tissue (about 1 mm3) were immediately fixed by immersion into 4% glutaraldehyde in 0.2 M cacodylate buffer, pH 7.4, for 1 h at 4 °C, washed in buffer and postfixed with 1% osmium tetroxide, dehydrated in graded concentrations of acetone and embedded in Poly-bed 812 (Embedding Media Polysciences, IVC). Selected ultrathin sections (50–70 nm) were made with a Reicbert (Ultramite Super Nova Leica) ultramicrotome and mounted on uncoated copper grids, contrasted with uranyl acetate and a lead citrate. Specimens were examined in a Zeiss EM-9 electron microscope, which was operated at an acceleration voltage of 50 kV.

2.4. Immunofluorescence

Fragments of liver tissue were immediately placed in liquid nitrogen for a few minutes and then kept frozen at −70 °C in airtight boxes, until the moment they were sectioned in a cryostat at −20 °C. The sections were submitted to an indirect immunofluorescence technique for the demonstration of collagen isotypes (I, II and IV), laminin and fibronectin. The specific anti-sera were polyclonal, obtained in rabbits (Institute Pasteur, France). They were used in dilution varying from 1:40 to 1:100. Details concerning their preparation and tests of specificity appeared elsewhere [9,10]. Secondary fluoresceinated anti-rabbit IgG was commercially obtained from Sigma, USA.

2.5. Immunohistochemistry

For the demonstration of factor VIII, dendritic cells and ED3, fragments of the liver were immediately embedded in Tissue-teck (OCT Compound-Miles Inc., Diagnostic Division, Elkhart, USA), frozen into liquid nitrogen, and cryopreserved in a freezer at −70 °C until the moment of sectioning. Sections of 6 μm were obtained in a cryostat at −20 °C on slides previously treated with poly-l-lysine at 10% (Sigma, St. Louis, MO, USA), fixed in dehydrated acetone, treated with PBS. For the blocking of non-specific ligations, sections were treated with 10% skimmed milk in PBS for 20 min at room temperature. Besides, blocking of endogenous avidin and biotin was performed with DAKO Biotin Blocking System (Code No. X0590, DAKO Corporation, USA). For the demonstration of SM-α actin, TGF-β1, TGF-β1-R, desmin, vimentin and PDGF, paraffin sections of formalin-fixed tissue were used. Antigen retrieval was accomplished through microwave treatment in citrate buffer at pH 6.0.

Sections were incubated with the primary antibodies overnight, at 4 °C in a humidified chamber. Primary antibodies were diluted in 2% BSA in PBS (pH 7.4). After washing in PBS, sections were incubated in 10% skimmed milk during 20 min for blocking non-specific ligations. The slides were then incubated with the secondary antibody: a sheep-anti-rat IgG conjugated to peroxidase (Dako envision system-labelled polymer (Dako, USA)), in the dilution of 1:1000 for 30 min at 37 °C in a humidified chamber. Blockade of the endogenous peroxidase was done with 0.3% H2O2 for 30 min, at room temperature. The color was devel-
oped with 0.06% 3,3′-diaminobenzidine tetrahydrochloride (DAB) (Sigma) and 0.06% H2O2 plus 1% dimethylsulphox-
ide (Sigma). Sections were counterstained with 1% methyl-
green for 2 min dehydrated and mounted with Permount.
Control sections in which primary antibody was either omit-
ted or replaced by normal rat serum, were used.

2.6. Functional liver tests
The serum concentrations of aminotransferases (ALT and AST), alkaline phosphatase (ALK P), albumin, biliru-
bins, and gamma-glutamyl transpeptidase (γ-GT) were deter-
mined in a Hitachi 917 automatic analyzer.

2.6.1. Hydroxyproline measurements
Biochemical determination of hydroxyproline concen-
tration was made according to the colorimetric method of
Bergman and Loxley.[11]

2.6.2. Serum antibodies
The levels of serum antibodies against C. hepatica
were determined by an enzyme-linked immunosorbant assay
(ELISA), to test for detection of total Ig antibodies, by using
a goat anti-rat IgG conjugated to peroxidase (Sigma). The
plates were sensitized overnight with 10 H9262 g/ml/per well of
C. hepatica-egg antigen, diluted in carbonate buffer, pH 9.9.

Lecture was made on a microplate reader “Molecular devices-
Thermomax” spectrophotometer, under wavelength 450 mm,
connected to a computer with MDS-Soft Max (Version 3.0).

3. Results
Multifocal parasitic lesions, containing worm debris and
immature eggs were already well encapsulated by 3 months
after inoculation (Fig. 1A). These lesions almost disappeared
by the 6th month of infection. Only clusters of eggs packed
inside scattered, dense, small fibrous scars, with or with-
out a few calcified foci, remained by the 6th month onward,
sometimes in the vicinity of fibrous bands and hepatic nod-
ules (Fig. 1B). Three months after infection, septal fibrosis
appeared throughout the liver. It was represented by thin
and long fibrous septa crisscrossing the hepatic parenchyma
(Fig. 1C). These thin septa connected portal spaces to por-
tal spaces. Few septa appeared projected from the capsule
of old parasitic scars, and from the external capsule of the
liver. The portions of hepatic tissue circumscribed by sep-
tal fibrosis, maintained the one-cell thick normal pattern of
the liver parenchyma. Six months after infection, the micro-
scopic appearance of septal fibrosis had changed. In H & E
preparations fibrous septa were hardly discernible, since
the place occupied by fibrosis appeared as a thin empty cleft, with

Fig. 1. (A) Dead and dying Capillaria hepatica adult worms appear packed within an encapsulated lesion present in the liver of a rat. 30 days following
infection. (B) Six-month-old C. hepatica infection discloses a “cirrhotic” appearance of the liver, besides two encapsulated and calcified lesions containing
worm debris and eggs. H & E, ×100. Pictures from C to I show early fibrous-cellular septa (C and D) contrasting with the later appearance they assume, when a
few, thin, vascular, wavy and interrupted septa, containing elastic fibers, are observed (E-I). (A-C and E) Hematoxylin & Eosin, (D and F-H) picric-sirius-red
staining for collagen, (I) orcein stain for elastic tissue. Magnification ×100, except (E) and (G) ×200.
a few fusiform cells here and there (Fig. 1E). Picro-sirius stained slides easily disclosed that septal fibrosis was still present as delicate, thin threads of red-stained fibers. The septa appeared wavy and interrupted, containing blood vessels and elastic fibers (Fig. 1F–I). These overall aspects tended to persist when materials taken from 9- and 12-month-old infections were examined. In short, besides the diminution in the amount of septal collagen from the 3rd to the 6th month following infection, all the other parameters considered, tended to remain the same, 9 and 12 months after infection. Isolation and cultivation of eggs taken from 3rd-month-old infection yielded embryonation, which did not occur when eggs from 6-, 9-, or 12-month-old infections were tested. Determination of hydroxyproline confirmed the impression from histological examination, showing a statistically significant difference in collagen concentration as compared to specimens taken from more prolonged infections. However, from the 6th month onward hydroxyproline measurements were about the same, indicating a stabilization of the process of collagen degradation (Fig. 2).

Collagen isotypes I and III were present within the fibrous septa at all observational stages. Collagen type IV and laminin marked basement membranes and thus disclosed the rich vascularization of the fibrous septa and also their diminution and persistence throughout the time of observation (Fig. 3A and B). Fibronectin in the fibrous septa presented decreased concentrations along the time of examination, as a matter of fact in a way comparable to what happened to the other components of the extracellular matrix evidenced by immunofluorescence.

The cellular elements in the fibrous septa were scarce even at the 3rd month of infection, and this was even more evident during the subsequent periods. No special aspects concerning the number and distribution of the cellular components of the septa, as identified by immunohistochemistry, were apparent. Cells positive for desmin, vimentin, PDGF, TGF-β, TGF-β-R, ED3 were few and appeared occasionally within the thin fibrous septa. Actin was positive for a few cells in the septa, but appeared prominent by revealing the muscular walls of blood vessels (Fig. 3C), the endothelial cells of which appeared positive for factor VIII (Fig. 3D). Late septa, from the 6th to the 12th months after inoculation, practically contained only a few collagen fibers and small thin-walled, blood vessels.
Ultrastructurally, the septa seen by the 3rd month exhibited a mixture of several cell types, with predominance of fibroblasts and a few myofibroblasts. Other cells eventually identified in the septa included fat-storing cells (HSC), mast cells, eosinophils and lymphocytes (Fig. 4A). From the 6th month on, the septa contained densely packed collagen fibers and one or other of the cells previously mentioned (Fig. 4B).

Sections of the liver previously injected with India-ink revealed that the fibrous septa contained vessels that terminated directly into sinusoids. Some of these vessels appeared straight, thin, without collaterals, running for some distance within the septum, before dividing into several terminal ramifications and going towards the sinusoids (Fig. 4C and D). These aspects were observed in all specimens taken from infected rats. Normal controls showed that portal vessels tended to form an arcing structure near the perportal parenchymal cords before giving off short communications to the sinusoids.

The exploration of liver function failed to reveal major alterations. The data obtained in several periods were within normal limits. The only exception was a slight increase in alkaline phosphatase, of 367.7 ± 5.0 IU/l (controls 139.4 ± 43.2) and aminotransferases, ALT 162.2 ± 3.5 IU/l (controls 140.8 ± 7.8) and AST 92.2 ± 22.4 IU/l (controls 68.4 ± 14.7) levels by the 3rd month of infection, a moment when focal necro-inflammatory parasitic lesions were prominent.

Levels of serum anti- *C. hepatica* antibodies are depicted in Fig. 5.

4. Discussion

The interesting fact about *C. hepatica*-induced fibrosis in rats is that its course is progressive during the first 3 months of infection [1], but it soon stops and starts revealing signs of progressive regression by the 6th month onward. This change of course correlates well with the extinction of parasitism, as demonstrated in the present study. These observations are in keeping with the well-established notion that not only hepatic fibrosis is reversible, but that the best way to produce such regression is to remove its cause [12–14]. To stay in the field of hepatic fibrosis due to parasitic diseases, a good example of this principle can be seen in both human and experimental schistosomiasis, when not only hepatic fibrosis, but its associated manifestations of portal hypertension, can com-
pletely disappear following curative treatment [9,10,14,15]. However, there are hepatic fibroses with different morphology and significance. Shibayama and Nakata [15] demonstrated that hepatic fibrosis induced by pig-serum injections into rats did not have a functional expression. By measuring oxygen consumption, the velocity of blood perfusion, and other parameters, they concluded that fibrosis by itself does not disturb liver function. Lottat-Jacob et al. [17] made the interesting observation that hepatic fibrosis in experimental schistosomiasis treated with interferon presented a differential sequence of disappearance, with perisinusoidal fibrosis disappearing first and more rapidly than that of peri-ovular granulomas. There are several morphological types of hepatic fibrosis and they do not have the same behavior as far as reversibility is concerned. There is a long controversy regarding cirrhosis reversibility, meaning that in some circumstances fibrosis may indeed be irreversible [18–20]. However, the debate is still lingering regarding human and experimental cirrhosis [21,22]. Although Wanless et al. [23] have listed a series of histological findings, collectively called the “hepatic repair complex”, supposed to represent parameters of cirrhosis regression, no one has ever documented the morphologic transition of a human cirrhotic liver to a normal one. When the present study was made 6 months after C. hepatica infection, the data showing extinction of parasitism and decrease of septal fibrosis left us under the impression that the whole structure of the liver would subsequently return to normal. However, septal fibrosis became stabilized for the next 6 months. Fibrosis then appeared to be merely a supportive stroma of a new vascular disposition. Adaptation of the liver on face of a continuous aggression may result in fibrosis with vascular changes. During the process of excess fibrous tissue removal, that tissue serving as a supporting stroma for physiologically functioning blood vessels tends to be spared. The expectation that such septal fibrosis can be removed is similar to expecting removal of the normal portal fibrous tissue by anti-fibrotic treatment or other means. Present observations support the conclusion that associated vascular changes may be a decisive factor regarding the reversibility of fibrosis. This would also explain why cirrhosis is clinically reversible, as recently discussed by Friedman [21], although probably morphologically irreversible.

References


