Natural history of primary sclerosing cholangitis and prognostic value of cholangiography in a Dutch population


Published in:
Gut

DOI:
10.1136/gut.51.4.562

Link to publication

Citation for published version (APA):

General rights
It is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), other than for strictly personal, individual use, unless the work is under an open content license (like Creative Commons).

Disclaimer/Complaints regulations
If you believe that digital publication of certain material infringes any of your rights or (privacy) interests, please let the Library know, stating your reasons. In case of a legitimate complaint, the Library will make the material inaccessible and/or remove it from the website. Please Ask the Library: http://uba.uva.nl/en/contact, or a letter to: Library of the University of Amsterdam, Secretariat, Singel 425, 1012 WP Amsterdam, The Netherlands. You will be contacted as soon as possible.

*Gut* 2002;51:562-566
doi:10.1136/gut.51.4.562

Updated information and services can be found at:
http://gut.bmj.com/cgi/content/full/51/4/562

These include:

**References**

This article cites 13 articles, 5 of which can be accessed free at:
http://gut.bmj.com/cgi/content/full/51/4/562#BIBL

1 online articles that cite this article can be accessed at:
http://gut.bmj.com/cgi/content/full/51/4/562#otherarticles

**Email alerting service**

Receive free email alerts when new articles cite this article - sign up in the box at the top right corner of the article

**Topic collections**

Articles on similar topics can be found in the following collections

- Transplantation (284 articles)
- Other Epidemiology (1671 articles)
- Cancer: gastroenterological (1220 articles)
- Liver, including hepatitis (954 articles)

To order reprints of this article go to:
http://www.bmjjournals.com/cgi/reprintform

To subscribe to *Gut* go to:
http://www.bmjjournals.com/subscriptions/
Natural history of primary sclerosing cholangitis and prognostic value of cholangiography in a Dutch population


Background: Median survival of patients with primary sclerosing cholangitis (PSC) has been estimated to be 12 years. Cholangiography is the gold standard for diagnosis but is rarely used in estimating prognosis.

Aims: To assess the natural history of Dutch PSC patients and to evaluate the prognostic value of a cholangiographic classification system.

Patients: A total of 174 patients with established PSC attending a university hospital and three teaching hospitals from 1970 to 1999.

Methods: Charts were reviewed for validity and time of diagnosis, concurrent inflammatory bowel disease, interventions, liver transplantation, occurrence of cholangiocarcinoma, and death. Follow up data were obtained from the charts and from the attending clinician or family physician. Median follow up was 76 months (range 1–300). The earliest available cholangiography was scored using a radiological classification system for the severity of sclerosis, developed in our institution. Survival curves were computed by the Kaplan-Meier method. Cholangiographic staging was used to construct a prognostic model, applying Cox proportional hazards analysis.

Results: The estimated median survival from time of diagnosis to death from liver disease or liver transplantation was 18 years. Cholangiocarcinoma was found in 18 (10%) patients. Fourteen patients (8%) underwent liver transplantation. Cholangiographic scoring was inversely correlated with survival. A combination of intrahepatic and extrahepatic scoring, together with age at endoscopic retrograde cholangiopancreatography, proved strongly predictive of survival.

Conclusions: The observed survival was considerably better than reported in earlier series from Sweden, the UK, and the USA. Classification and staging of cholangiographic abnormalities has prognostic value.

PATIENTS AND METHODS

Patients

Records from all 181 patients, who were filed with a diagnosis of PSC between 1970–1999 from one tertiary referral centre and three teaching hospitals, were retrieved.

The diagnosis was reviewed together with cholangiograms and histology results when available. Seven patients were excluded because they lacked sufficient evidence for a diagnosis of PSC. Three of these patients had a Klatskin tumour, and four had complicated gall stone disease. From the remaining 174 patients the following data were recorded: age at diagnosis, sex, concurrent inflammatory bowel disease (IBD), cholecystectomy, endoscopic retrograde cholangiopancreatography (ERCP) findings, age at index ERCP (AGEERCP), disease duration until index ERCP, liver biopsies, total follow up time, follow up from index ERCP, clinical status at end of follow up, occurrence of CCA, and orthotopic liver transplantation (OLT). Follow up data were extracted from the charts and obtained from attending clinicians, primary care physicians, patients, or from one of the liver transplantation centres in the Netherlands.

Abbreviations: PSC, primary sclerosing cholangitis; CCA, cholangiocarcinoma; ERCP, endoscopic retrograde cholangiopancreatography; AGEERCP, age at index ERCP; IBD, inflammatory bowel disease; OLT, orthotopic liver transplantation; EHD, extrahepatic disease; PI, prognostic index.
Natural history assessment
Cumulative survival from the time of diagnosis to death from liver disease or OLT was estimated by Kaplan-Meier analysis. The same method was applied to assess the survival of patients with CCA from the time of diagnosis of this complication to death.

Cholangiographic scoring
All available cholangiograms were retrieved and scored by two investigators who were unaware of the patients’ clinical information, applying the modified scoring system as recently described (see table 1). Survival curves were computed to assess whether increasing intra- and extrahepatic scorings were correlated with a poorer prognosis.

Prognostic model computation
A variable based on the cholangiographic scoring system was defined using Kaplan-Meier and univariate Cox proportional hazards analysis. Subsequently, using backward regression, age at index ERCP, duration of disease until index ERCP, sex, concurrent IBD, and cholecystectomy were introduced into the regression analysis as possible covariates.

The assumption of proportional hazards for a Cox regression model fit was checked as described previously.

For statistical calculations the SPSS package version 8.0 (SPSS inc., Chicago, Illinois, USA) and S-plus version 4.5 (MathSoft, Seattle, Washington, USA) were used.

RESULTS
General characteristics
Among the 174 patients, 105 (60%) were male. Mean age at diagnosis was 40.4 (14.8) years. A total of 114 (66%) patients were known to have concurrent IBD (ulcerative colitis, n=83; Crohn’s disease, n=28; indeterminate colitis, n=3). Thirty nine patients (22%) underwent cholecystectomy, which not infrequently revealed a diagnosis of PSC.

Cholangiographic scoring
ERCPs were available from 133 patients, either initially (n=78) or within 2–257 months after diagnosis. Figures 2 and 3 show two typical examples of the cholangiographic spectrum of PSC.
The range of possible values for SUMIHDEHD′ scoring (IHD and the modified extrahepatic scoring (EHD) predicting survival (p=0.0026). This categorical parameter analysis yielded a single parameter that performed best in determining prognosis. Extensive Cox proportional hazards and extrahepatic scoring would be expected to play a role in abnormalities would reflect disease severity. Hence both intra-

The assumption was made that both intra- and extrahepatic disease types were combined into stage I, and EHD stages III and IV were negligible and not significant (p=0.91 and p=0.58, respectively). Therefore, stages 0 and I were combined into stage I′, and stages III and IV were grouped into stage III′. This resulted in a new parameter EHD′.

Computing a prognostic model

The assumption was made that both intra- and extrahepatic abnormalities would reflect disease severity. Hence both intra-

Figure 3 Cholangiography showing type II intrahepatic and type III extrahepatic disease.

Cumulative survival from the time of the index ERCP for the various intra- and extrahepatic disease types is displayed in figs 4 and 5, respectively. As can be seen in fig 5, differences in survival between extrahepatic disease (EHD) stages 0 and I, and EHD stages III and IV were negligible and not significant (p=0.91 and p=0.58, respectively). Therefore, stages 0 and I were combined into stage I′, and stages III and IV were grouped into stage III′. This resulted in a new parameter EHD′.

Table 2 Resultant SUMIHDEHD′ score from intrahepatic (IHD) and extrahepatic (EHD) disease. The score can be read from the intersection of the pertaining EHD row and IHD column (the combination 0–0 is non-existent because it would preclude a diagnosis of primary sclerosing cholangitis).
DISCUSSION

The results of this cohort study, which is the second largest reported (together with the Mayo Clinic series), containing the longest follow up so far, showed that median survival appears to be considerably better than reported previously. Three previously published studies involving more than 100 patients were reported on cohorts mainly from the 1970s and 1980s from the USA, the UK, and Sweden. All three recorded a median survival of approximately 12 years. In contrast, we found a 50% better median survival (18 years). The reasons for this discrepancy may be multifactorial. Selection bias may play a role. However, the general patient characteristics were similar in all four series. In the study from the Mayo Clinic, the date of referral was taken as the time of diagnosis which may have given rise to considerable underestimation of total disease duration. Moreover, death from any cause was regarded as an end point. However, when the six non-liver related deaths in our series were included as an end point in the survival analysis, the estimated median survival was 17.5 years, still much longer than that in the Mayo cohort. The King's College and Swedish studies used the same end points as in the present study. The Swedish study defined baseline time as the date of diagnosis, as we did. The British study defined baseline as the time of the first symptom consistent with—in retrospect—a diagnosis of PSC. Compared with our series, as well as with the Swedish cohort, this could give rise to a substantial increase in median survival time inasmuch as parameters such as elevated liver function tests may precede a diagnosis of PSC by many years. Yet median survival in our series was 50% longer than that in the series from King's College.

Alternatively, the time frame studied may be important. The series from the USA, the UK, and Sweden contained cohorts mainly from the 1970s–1980s while our cohort also spanned 1990–1999. Substantial therapeutic advances aimed at slowing down disease progression have not been made in the last decade but perhaps the diagnosis is now made earlier and hence ascertainment bias may well play a role.

The prevalence of CCA in the above mentioned studies was 6–8%. In our series, the prevalence of secondary CCA was somewhat worse (10%). This may be a result of the longer follow-up period in our series. The actual prevalence may be somewhat higher as not all patients who died were autopsied.

Several prognostic models have been proposed to predict the course of PSC and to tailor the timing of OLT. Independent variables for these models are age, histological staging, and serum bilirubin. However, hepatic histological biopsy findings can be non-specific, with a considerable degree of sampling variability, because the disease may not be evenly spread throughout the liver, and the pathognomonic onion skin lesion can easily be missed. Thus for diagnosis, liver biopsy is usually not done and histological staging may be variable. Likewise, serum bilirubin levels at presentation may be an unreliable indicator of disease stage. For instance, when a patient presents with supplicative cholangitis caused by a bile plugged distal dominant stricture, but otherwise patent biliary tree, the elevated serum bilirubin may soon revert to normal when the obstruction is relieved.

The mainstay of diagnosis is cholangiography and therefore it seems obvious to develop predictors based on cholangiographic abnormalities. Two studies have assessed the prognostic value of cholangiography. Both concluded that high grade intrahepatic strictures indicated poor prognosis. However, grading of intrahepatic strictures in these studies was rather detailed and may be difficult to reproduce because the degree of narrowing of strictures may vary considerably, depending on the amount of contrast injected into the biliary tree. On the other hand, the Amsterdam cholangiographic classification system uses a more qualitative appreciation of the abnormalities in the biliary tree in PSC patients. To date, this classification has not been evaluated clinically in terms of predicting prognosis. On the assumption that both intrahepatic scoring and extrahepatic scoring influence disease severity and consequently survival, a variable was defined representing both scorings. The resulting variable SUMIHDEHD showed that there was a clear inverse correlation between combined morphological stage and survival. Moreover, a PI was computed on the basis of SUMIHDEHD. The predicted survival plots for the various categories of SUMIHDEHD closely matched the actual estimated cumulative survival curves for these categories. However, the definitive value of our prognostic model can only be determined after cross validation.

In conclusion, survival of Dutch PSC patients is considerably better than reported previously. The Amsterdam cholangiographic classification system reflects disease stage and has the potential to serve as a predictor in determining prognosis. This may prove useful in patient selection and timing of liver transplantation, patient counselling, and stratifying patients for therapeutic trials.
ACKNOWLEDGEMENTS

The authors are indebted to Dr AWM van Milligen de Wit Red Cross Hospital, the Hague, and Dr W Bruins Slot, Spaarne Hospital, Haarlem, for providing some patient data.

Authors’ affiliations

C Y Ponsioen, S M E Vrouenraets, W Prawirodirdjo, R Rajaram, E A J Rauws, G N J Tytgat, Department of Gastroenterology and Hepatology, Academic Medical Centre, Amsterdam, the Netherlands

CJJ Mulder, Department of Internal Medicine, Rijnstate Hospital, Arnhem, the Netherlands

J B Reitsma, S Heisterkamp, Department of Epidemiology and Biostatistics, Academic Medical Centre, Amsterdam, the Netherlands

REFERENCES


