Decision to treat chronic hepatitis B in France in 2008-2009

C. Larsen¹, C. Pioche¹, P. Couzigou², E. Delarocque-Astagneau³, C. Brouard¹, O. Goria⁴, D. Guyader⁵, P. Hillon⁶, P. Marcellin⁷, F. Roudot-Thoraval⁸, D. Roulot⁹, C. Silvain¹⁰, J.-P. Zarski¹¹, F. Denis¹², S. Chevaliez⁸, C. Semaille¹ and the hepatology reference centers and laboratories network for chronic hepatitis B surveillance

1/ French Institute for Public Health Surveillance (InVS), Saint-Maurice, France – 2/ University Hospital, Rouen, France – 5/ University Hospital, Rennes, France – 6/ University Hospital, Dijon, France – 7/ APHP-Bichat Hospital, Clichy, France 8/ APHP-Henri Mondor Hospital, Créteil, France – 9/ APHP-Avicenne Hospital, Bobigny, France – 10/ University Hospital, Poitiers, France – 11/ University Hospital, Grenoble, France – 12/ University Hospital, Limoges, France

Background

- France is a low endemic European country for chronic hepatitis B virus (HBV) infection with an HBsAg prevalence estimates of 0.65% in the general population.
- Annual mortality related to HBV is estimated to 2.2/100 000 inhabitants.
- As part of the national hepatitis B control program, a surveillance network was set up in 2008 including hepatology reference centers and laboratories of university hospitals throughout France.

Objectives

To describe the epidemiologic, clinical and virological characteristics of patients with chronic hepatitis B at first referral in the hepatology reference centers.

Methods

- Chronic hepatitis B is defined as persistent HBsAg \geq six months.
- Data collected: country of birth, ALT and HBV DNA levels, co-morbidities, viral co-infections, antiviral therapy initiation.
- Excessive alcohol consumption is defined as >210g/week of ethanol for women and >280g/week for men.
- Liver fibrosis (Metavir scoring) was assessed either by biopsy and/or noninvasive methods (serum markers, transient elastography).
- For the analysis, moderate to severe liver fibrosis was defined using:
- scores \geq F2 at biopsy and if not available;
- scores \geq F2 at elastography or serum markers if performed solely;
- or scores \geq F2 at elastography and serum markers when both methods matched.
- Indications of treatment are analysed according to EASL guidelines:
- "Patients with HBV DNA levels>2000 IU/ml and/or the ALT levels>upper limit of normal for the laboratory, and liver biopsy (or non-invasive markers when validated in HBV-infected patients) shows moderate to severe active necroinflammation and/or fibrosis using a standardised scoring system should be considered for treatment".





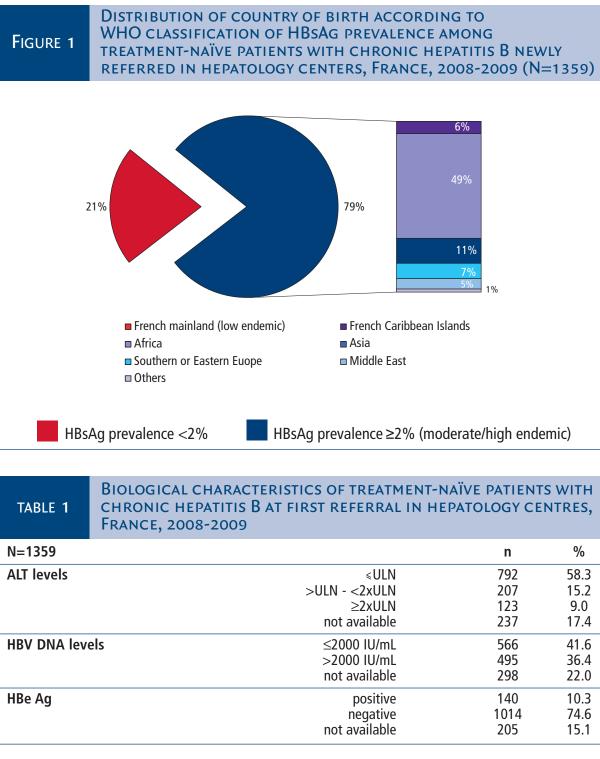
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FRENCH INSTITUTE FOR PUBLIC HEALTH SURVEILLANCE

References

Meffre C *et al.* J Med Virol 2010;82:546-55. Marcellin P et al. J Hepatol 2008;48:200-7.

- 1 359 treatment naïve HBsAg-positive patients included in 33 wards.
- 59% male.
- 79% born in moderate to high endemic country for HBV (figure 1).
- Median age: 36 years (interquartile range: 17 years).
- 633 (47%) patients with ALT>upper limit of normal or HBVDNA>2000 IU/mL.
- Characteristics of patients are presented in tables 1-3.



Results

n	%
792	58.3
207	15.2
123	9.0
237	17.4
566	41.6
495	36.4
298	22.0
140	10.3
1014	74.6
205	15.1
	792 207 123 237 566 495 298 140 1014

	CO-MORBIDITIES AND VIRAL CO-INFECTIONS AMONG
E 2	TREATMENT-NAÏVE PATIENTS WITH CHRONIC HEPATITIS B AT FIRST
	REFERRAL IN HEPATOLOGY CENTRES, FRANCE, 2008-2009

TABL

N=1359		n	%
Excessive alcohol consumption ^a	no	1040	76.5
	yes	95	7.0
	not available	224	16.5
BMI	<25 kg/m ²	511	37.6
	≥25 kg/m²	386	28.4
	not available	462	34.0
Anti-HIV	positive	19	1.4
	negative	937	69.0
	not available	403	29.6
Anti-HCV	positive	37	2.7
	negative	1029	75.7
	not available	293	21.6
Anti-HDV	positive	33	2.4
	negative	911	67.0
	not available	415	30.6

^a>210g/week of ethanol for women and >280g/week for men; BMI, body mass index; overweight /obesity when BMI ≥25Kg/m².

DECISION TO TREAT ACCORDING TO EASL GUIDELINES

- 429 patients with increased ALT (>upper limit of normal) or HBV DNA >2000 IU/mL and an available assessment of liver fibrosis.
- 157 (36.6%) patients with liver fibrosis \geq F2 including: - 58 patients with liver fibrosis F4;
- 93 patients treated (out of 114 with available information on treatment).

TABLE 3INDEPENDENT FACTORS ASSOCIATED WITH DECISION TO TREAT CHRONIC HEPATITIS B IN HEPATOLOGY CENTERS, FRANCE, 2008-2009			
		aOR	CI95%
Liver fibrosis ≥F2 31.7 9.8-103.0		9.8-103.0	
ALT level >ULN 7.4 3.2-17.1		3.2-17.1	
HBV DNA>2000 UI/ml 3.8 1.4-10.1		1.4-10.1	
Age >36 years		3.5	1.8- 6.9

aOR, adjusted Odds Ratio; CI, confidence interval; ULN, upper limit of normal;

Other factors associated with decision to treat in univariate analysis (sex, country of birth, HBeAg status, history of excessive alcohol consumption) did not remain significant in the multivariate analysis; significant interactions are taken into account

Conclusion

- In France, most of the patients with chronic hepatitis B were originating from a moderate/high endemic country for HBV and were HBeAg negative.
- Decision to treat chronic hepatitis B in 2008-2009 is taken in respect of the EASL guidelines.

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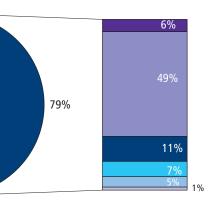
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	WHO CLASSIFICATION OF H TREATMENT-NAÏVE PATIENT REFERRED IN HEPATOLOGY
	21%
	 French mainland (low endemic) Africa Southern or Eastern Euope Others
HBs	Ag prevalence <2%
TABLE 1	BIOLOGICAL CHARACTERIST CHRONIC HEPATITIS B AT F FRANCE, 2008-2009
ALT levels	
IDV DIVA IEV	els
HBe Ag	els
	CO-MORBIDITIES AND VIRA TREATMENT-NAÏVE PATIENT REFERRAL IN HEPATOLOGY
HBe Ag TABLE 2 N=1359	Co-morbidities and vira treatment-naïve patient referral in hepatology
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HBe Ag TABLE 2 N=1359 Excessive alc BMI Anti-HIV	Co-morbidities and vira treatment-naïve patient referral in hepatology

OF BIRTH ACCORDING TO BSAG PREVALENCE AMONG S WITH CHRONIC HEPATITIS B NEWLY CENTERS, FRANCE, 2008-2009 (N=1359)



French Caribbean Islands 🗖 Asia

Middle East

sAq prevalence $\geq 2\%$ (moderate/high endemic)

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