

# Evidence for Persistent Hepatitis C Virus (HCV) Infection in Hemophiliacs

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## Abstract

Hepatitis C virus (HCV) is the major etiologic agent associated with non-A, non-B hepatitis. This study was designed to assess virologic and serologic markers in hemophiliacs exposed to non-heat-treated and/or virus-inactivated plasma derivatives. Serial bleeds from 48 hemophilic patients were analyzed for the presence of HCV viral RNA sequences as detected by polymerase chain reaction (PCR) and antibodies to structural (core) and nonstructural (C-100 and 33C) proteins by specific dot immunoblot assay. All patients exposed to non-heat-treated products, and four of six patients exposed only to virus inactivated products, had evidence of HCV infection. However, over the 5-yr study period, six exposed patients (13%) consistently lacked detectable anti-C-100 and seven (15%) lost this antibody. HCV viremia (PCR positive) was found in 91% of exposed patients, and was significantly more frequent in HIV seropositive hemophiliacs ( $P < 0.05$ ). Six patients had high antibody level to HCV and elevated ALT, but appeared to clear viremia. Four hemophiliacs were HCV seropositive but lacked detectable viremia. These data indicate that hemophiliacs remain persistently infected by HCV and that antibody to the core antigen of HCV is a reliable marker of this transfusion transmissible agent. (*J. Clin. Invest.* 1991; 88:1672-1679.)  
**Key words:** Non-A hepatitis • non-B hepatitis • liver disease • Factor VIII concentrate • Factor IX concentrate • virus inactivation

## Introduction

Since the early 1980s, viral inactivation of plasma derivatives used for the treatment of hemophiliacs has been the object of intensive research. A large number of techniques for inactivation of infectious agents in plasma derivatives ranging from heat treatment at 60–80°C in dry, humidified, or liquid state (1–6), to detergents or chemicals (7), have been developed and evaluated in chimpanzees as well as in previously untreated hemophiliacs. Clinical trials showed that most of these methods effectively inactivated HIV but were only partially effective with the agent of non-A, non-B hepatitis. Unequivocal evidence of non-A, non-B hepatitis prevention was obtained only with high temperature ( $\geq 75^\circ\text{C}$ ), pasteurization in liquid

phase, or the solvent-detergent (SD)<sup>1</sup> method (6–8). These clinical results were later confirmed serologically with the discovery of hepatitis C virus (HCV), the virus implicated in post-transfusion non-A, non-B hepatitis, and the development of screening assays for antibody to HCV. Several studies indicated that the prevalence of antibody to the HCV nonstructural C-100 antigen ranged from 61 to 89% in patients with hemophilia A receiving non-heat-treated Factor VIII concentrates, and 22 to 56% in patients receiving exclusively virus-inactivated material (9–13). Investigators raised questions as to why all potentially exposed patients were not seropositive to HCV, and why, over time, between 1985–88, approximately 10% of seropositive hemophiliacs lost reactivity with the C-100-based screening ELISA (9, 14). One hypothesis suggested that some patients may not develop antibody to this HCV nonstructural protein. Alternatively, the use of virus-inactivated products may limit or eliminate HCV antigenic stimulation and patients who no longer produce virus may return to a seronegative state (seroreversion) (14).

In order to test these hypotheses, serial serum samples collected 1985–89 from a group of 48 hemophiliacs were tested to assess the presence of circulating HCV by polymerase chain reaction (PCR) and the immune response to structural and nonstructural antigens of HCV by dot immunoblot. Results indicate that most HCV-infected hemophiliacs remain chronically infected while receiving virus-inactivated plasma derivatives and that antibody to the core antigen of HCV was a very reliable marker of past and present HCV infection.

## Methods

**Patients.** A group of 48 persons with hemophilia A (40) or B (8) ranging in age from 6 to 63 yr (median 23, mean 24 y) was clinically monitored one to four times a year between 1985 and 1989 at the International Hemophilia Training Center, Kremlin-Bicêtre, France. Standard check-up included clinical examination, measurement of alanine amino transferase (ALT), HIV antibody detection, testing for hepatitis surface antigen (HBsAg), antibody to hepatitis B surface (anti-HBs) and core (anti-HBc) antigens. In addition, residual sera were kept frozen at  $-30^\circ\text{C}$  in 0.5-ml aliquots. One aliquot of serum for each year was sent frozen in dry ice to the USA, divided into five, 100- $\mu\text{l}$  aliquots on arrival, and kept frozen below  $-80^\circ\text{C}$  until used. Each patient was classified into one of four liver disease evaluation categories on the basis of clinical examination and liver enzyme profile. Category 1 included patients with or without hepatomegaly who had persistently elevated ALT levels ( $\geq 45$  IU/liter). Category 2 included patients who developed acute non-A, non-B hepatitis during the course of the study. Category 3 included patients with minor (below twice the upper nor-

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1. Abbreviations used in this paper: ALT, alanine amino transferase; anti-HBc, antibody to hepatitis B core; anti-HBs, antibody to hepatitis B surface; Dr, density of reflectance; HBsAg, hepatitis surface antigen; HBV, hepatitis B virus; HCV, hepatitis C virus; PCR, polymerase chain reaction; SD, solvent-detergent.

mal level), erratic elevation of ALT, and no hepatomegaly. Category 4 included hemophiliacs with no clinical or biological evidence of liver disease. Patients with hemophilia A received a variety of Factor VIII-rich plasma derivatives that were non-heat-treated before October 1985, heat-treated between October 1985 and 1987, and, since 1988, treated by a SD method (7). Before 1985, most patients were treated with freeze-dried cryoprecipitate or intermediate purity Factor VIII concentrate prepared by the Centre National de Transfusion Sanguine (CNTS), Paris, France. However, some patients also received Factor VIII concentrates of American origin. In 1985–87, patients were treated with intermediate purity concentrate heat-treated as humidified powder under high pressure (CNTS) as described (5). Since 1988, all patients were treated exclusively with a high purity SD-treated Factor VIII (Bio-Transfusion, Les Ulis, France). Patients with hemophilia B received Factor IX concentrate (PPSB, CNTS) non-heat-treated, heat-treated, and SD as for patients with hemophilia A. There were six patients (five with severe and one with moderate hemophilia) who received exclusively a combination of heat-treated and SD-treated products. These six patients and nine others were HIV seronegative, whereas 33 patients were HIV seropositive in 1985 and remained positive. 12 patients age 15 or younger had been vaccinated with HEVAC B (Pasteur vaccine) and all had anti-HBs as only marker of hepatitis B. In addition, 2 of the 36 patients who were HBV infected by transfusion were HBsAg positive; 34 had anti-HBs, 25 anti-HBc, and one had none of these HBV markers although he had received whole blood and plasma derivatives for more than 30 yr.

**Detection of HCV genomic sequences by PCR.** The genomic RNA isolation, cDNA synthesis, and PCR analysis were performed as described by Weiner et al. (15) with PCR primers described by Kubo et al. (16). Briefly, 50  $\mu$ l of patient serum were incubated in lysis buffer (50 mM Tris pH 8, 1 mM EDTA, 0.1 M NaCl, 0.5% SDS, 100  $\mu$ g/ml proteinase K) for 1 h at 37°C. RNA was purified by phenol/chloroform extractions. The cDNA synthesis was performed according to manufacturer's protocols (Bethesda Research Laboratories, Gaithersburg, MD). PCR conditions were: denaturation at 95°C for 1 min, annealing at 42°C for 2 min, and extension at 72°C for 3 min for 35 cycles. After first round amplification, the PCR-amplified sample was reamplified with a second pair of primers which were internal to the original primer pair. The nested-PCR conditions were the same as the first round except only 30 cycles were performed. The PCR products were analyzed on a 1.2% agarose gel, blotted to Hybond-N (Amersham Corp., Arlington Heights, IL), and hybridized to a random primed <sup>32</sup>P-labeled HCV DNA insert. Samples positive by PCR were considered viremic.

**Dot blot immunoassay.** Antibodies to HCV gene products were assayed with a semiautomated dot blot immunoassay (MATRIX; Abbott Laboratories, North Chicago, IL) which employs an array of purified recombinant antigens spotted on nitrocellulose. This technology has been described in detail (17, 18). The test panels consisted of the C-100 antigen, which was purchased from Chiron Corporation (Emeryville, CA), plus polypeptides corresponding to putative HCV NS-3 (33C), and 5'-structural sequences encoding for the putative core of HCV (core) which were expressed in *Escherichia coli* as fusions with cytosine monophosphate keto deoxy octulosonate (CMP-KDO) synthetase (19). Each array also contained procedural controls (anti-human IgG and human IgG) to verify sample and reagent addition, as well as a negative control (casein) to adjust for nonspecific binding to the nitrocellulose surface.

The prepared solid phase was incubated in a reaction cell with the test sample (diluted 1:100) for 1 h at 35°C followed by sequential 30-min incubations at 35°C with biotin-labeled goat anti-human IgG (heavy and light chains), alkaline phosphatase-labeled rabbit anti-biotin, and bromochloroindolyl phosphate. Upon completion of the incubation with the chromogen, the nitrocellulose surface was dried and reflectance at defined locations within the array was determined to provide an objective measure of the extent of the individual reactions. Reactivity cutoff values were calculated for each antigen based on a 50-member antibody-negative population; sample to cutoff (S/CO) or density of reflectance (Dr) ratio values > 1 were considered positive.

For typical HCV-positive donor specimens, antigen reactivities varied linearly with antibody concentration for S/CO values < 100 (data not shown).

**Statistical analysis.** The two tailed Fischer's exact tests were used to assess statistical differences between comparison groups.

## Results

**HCV infection.** By serological criteria, 46 out of 48 multitransfused hemophiliacs (95.8%) had antibody to HCV. Two patients who had received exclusively heat-treated and SD Factor VIII concentrates had no convincing clinical, serologic, or virologic evidence of HCV infection. All 46 HCV seropositive patients had antibodies to at least two of the tested antigens. However, among these, six were consistently negative for anti-C-100 and one was consistently negative for anti-33C, but all were positive for antibody to the core protein. The mean Dr ratio calculated on the highest ratio for each patient was 48.5 $\pm$ 48.2 for anti-C-100, 234.0 $\pm$ 92.1 for anti-33C, and 206.9 $\pm$ 54.1 for anticore. Immunoreactivity to C-100 often appears transient and markedly lower than to the 33C or core antigens, although the Dr ratios cannot be directly compared due to the presence of an unknown number of available epitopes for each protein on the reactive areas.

**HCV viremia.** HCV viremia was assessed by PCR using primer pairs from the nonstructural NS3/NS4 region of the viral genome. Overall, 42 out of 46 patients (91.3%) with serological evidence of HCV infection were found viremic in at least one of the three to five samples collected over a period of 4 to 5 y. The data collected were grouped according to four different patterns. Patterns 1, 3, and 4 are presented in detail in Table I and summarized in Table II.

Pattern 1 included 10 patients whose samples all tested positive for HCV genome in serum. In addition to apparent, persistent HCV viremia based on PCR data, it is significant that all patients were also seropositive for HIV. Nine patients had either clear (seven) or moderate (two) signs of liver disease and only patient 39 had consistently normal ALT level. HCV antibodies to the three antigens tested were present in all patients on entry. Over time, antibody level to C-100 tended to decline in seven patients and became undetectable in four of them: The mean maximum Dr ratio was 77.5 $\pm$ 62.7 (Table II). Antibody to 33C was initially at high or moderate level, and decreased significantly during the course of the study in five patients. Except for patient 38, antibody to core was at high level in all patients and remained rather stable during the study period.

Pattern 2 included 26 patients (56.5%) whose HCV viremia was intermittently detectable (Table II). Due to the large number of patients in this group and the absence of particular features, serial sample results are not presented in Table I. Acute (two patients) or chronic liver disease (11 patients) was frequent, while four patients showed no sign of liver abnormality. HIV seropositivity was found in 18 patients (69%) from this group. Four patients had no detectable antibody response to C-100 (three patients) or 33C (one patient); all responded to the core protein. The mean Dr ratios of antibody to C-100, 33C, and core were similar to group 1 (Table II). During the study period, antibody to C-100 ratio declined in 13 patients, becoming undetectable in three of them. Antibody to 33C also declined in 13 patients and as did antibody to core in four patients.

Table 1. HCV Viremia, Antibody to HCV, Liver Disease, and HIV Status in Hemophiliacs with Patterns 1, 3, and 4

Patient	Date	HCV RNA	Dot immunoblot (S/CO)			Alt* level	HIV antibody status	Liver† disease
			C-100	33-C	Core			
a. Pattern 1								
5	11-84	+	46	324	205	46	Pos	3
	06-86	+	30	272	201	51		
	08-87	+	10	297	252	33		
	08-88	+	4	250	234	20		
	10-89	+	1	74	149	21		
13	04-85	+	41	145	182	80	Pos	1
	02-86	+	39	110	222	46		
	08-87	+	20	19	238	23		
	09-88	+				44		
	06-89	+	7	4	193	39		
20	06-85	+	13	72	171	179	Pos	1
	06-86	+	6	42	126	158		
	11-87	+	0	1	260	156		
	07-88	+	0	1	158	99		
	08-89	+	0	0	120	117		
28	01-87	+				32	Pos	1
	04-88	+	145	267	78	121		
	09-89	+	177	298	83	138		
37	06-86	+	60	258	160	34	Pos	1
	04-87	+	89	304	102	45		
	10-88	+	83	284	105	17		
38	09-89	+				60	Pos	1
	01-85	+	29	290	3	138		
	02-86	+	31	220	9	87		
	04-87	+	ND			85		
	04-88	+	25	76	6	95		
39	10-89	+	14	48	5	120	Pos	4
	05-85	+	83	294	193			
	01-86	+	155	291	147	28		
	01-87	+	62	206	177	25		
	02-88	+	86	231	98	14		
40	01-89	+	ND <sup>‡</sup>			25	Pos	3
	03-85	+	22	227	130	40		
	04-86	+	5	159	211	42		
	10-87	+	3	110	161	23		
	10-88	+	2	10	171	79		
45	09-89	+	0	26	119	43	Pos	1
	05-85	+	173	283	126	77		
	05-86	+	101	319	230	75		
	04-87	+	93	268	138	126		
	08-88	+	54	367	217	165		
48	08-89	+	17	269	178	152	Pos	1
	01-85	+	2	301	129			
	08-86	+	28	247	222	152		
	03-87	+	19	267	136	148		
	05-88	+	11	340	196	110		
	06-89	+	0	303	182	80		
b. Pattern 3								
9	02-85	+	1	68	196	60	Pos	1
	10-86	+	1	39	170	61		
	09-87	+	0	0	177	57		
	12-88	—	0	0	177	57		

Table I. (Continued)

Patient	Date	HCV RNA	Dot immunoblot (S/CO)			Alt* level	HIV antibody status	Liver <sup>‡</sup> disease
			C-100	33-C	Core			
11	01-85	+	26	220	183	19	Pos	3
	04-86	+	19	239	206	16		
	04-87	+	37	289	204	10		
	09-88	—	7	321	292	50		
	12-89	—	0	359	306	44		
12	01-85	+	27	202	188	34	Neg	4
	04-86	+	14	180	186	33		
	04-87	—	27	237	150	23		
	03-88	—	14	249	279	19		
	03-89	—	9	205	304	43		
15	05-85	+	0	228	180	40	Neg	3
	86	+	3	283	128	45		
	04-88	—	3	299	255	28		
	06-89	—	4	301	227	66		
18	02-85	+	0	17	227	30	Pos	3
	01-86	+	0	3	217	28		
	07-87	—	0	3	275	16		
	07-88	—	0	3	283	40		
	02-89	—	0	0	138	47		
32	06-85	+	1	218	82	42	Pos	1
	05-86	+	1	255	79	31		
	09-87	+	0	286	149	23		
	09-88	+	0	264	164	30		
	06-89	—	0	271	84	58		
c. Pattern 4								
19	03-85	—	39	234	67	56	Neg	4
	11-86	—	3	54	5	14		
	05-87	—	4	46	9	18		
	05-88	—	2	31	6	13		
	02-89	—	2	31	6	19		
26	09-85	—	ND			51	Pos	1
	02-86	—	37	134	191	57		
	02-87	—	9	79	187	39		
	10-88	—	28	256	115	39		
	03-89	—	0	36	151	59		
29	07-85	—	7	257	237	135	Neg	1
	07-86	—	19	247	204	186		
	07-87	—	10	204	234	36		
	07-88	—	1	121	219	15		
	07-89	—	3	313	132	223		
42	03-85	—	59	210	178	81	Pos	1
	03-86	—	70	321	228	85		
	04-87	—	64	292	220	89		
	04-88	—	66	299	164	52		
	06-89	—	81	290	163	195		
27	09-85	—	0	0	0	6	Neg	4
	10-86	—	0	0	0	14		
	10-87	—	0	0	0	13		
	09-88	—	0	0	0	50		
	09-89	—	0	0	0	18		
33	08-86	+ <sup>  </sup>	0	0	0	20	Neg	4
	07-88	—	0	0	0	10		
	07-89	—	0	0	0	13		

\* In IU/liters, upper normal level is 46 IU/liters.

<sup>‡</sup> 1 corresponds to chronic liver disease; 2 corresponds to acute non-A, non-B hepatitis becoming chronic; 3 indicates minor, erratic elevation of ALT level; 4 indicates consistently normal ALT level and normal clinical examination.<sup>§</sup> ND, Not done.<sup>||</sup> This patient came to the Center for check-up once a year from Morocco. No additional sample was available to verify this surprising result obtained in two separate experiments.

Table II. Characterization of HCV-Infected Hemophiliacs

Patient group	HCV cDNA PCR		Mean HCV antibody* DR ratio (±SD)			Liver disease†				HIV antibody	
	Pattern‡	Patient No.	C-100	C-33	Core	1	2	3	4	POS	NEG
1	+/+	10	77.5 (48.5)	266.1 (87.6)	185.8 (76.8)	7		2	1	10	
2	+/-	26	46.6 (42.7)	223.2 (87.2)	195.5 (55.1)	11	2	9	4	18	8
3	+/-	6	11.3 (15.0)	213.3 (125.9)	251.3 (54.0)	2		3	1	4	2
4	-/-	4	44.0 (22.7)	281.0 (36.9)	180.7 (67.9)	3			1	2	2

\* Calculated as mean of maximum reflectance to background ratio observed in each patient.

† See Table I.

‡ +/+ positive in all tested samples; +/- intermittently positive; +/- initially positive becoming negative; -/- consistently negative.

Pattern 3 included six patients who were initially viremic and subsequently became negative by PCR. As shown in Table Ib, these patients had chronic (two patients) or minor liver disease (three patients), and one patient had no sign of liver disease. ALT levels were mostly normal or less than twice the upper normal level. Three patients had no detectable antibody to C-100; the other three had either low or declining levels of antibody. In contrast, four had consistently high anti-33C and two had low antibody titers that became undetectable. All patients had consistently high anticore immune response. When compared with patients in pattern 1, the mean maximum of anti-C-100 was significantly lower in patients with pattern 3 ( $P < 0.05$ ) and the level of anticore was higher although the difference was not significant.

Pattern 4 included four patients with no evidence of HCV viremia, although two of them (patients 29 and 42, Table Ic) had severe chronic hepatitis. Patient 19 (Fig. 1) was HIV antibody negative, had no sign of liver disease, and a decreasing level of all three antibodies to HCV. Patient 26 had fluctuating levels of antibody to 33C and core, and had no detectable anti-C-100 in the final sample. He had a slightly elevated ALT level. Patient 29 also had fluctuating antibody levels, whereas patient 42 had persistently high levels of each antibody. Patients 27

and 33 had no evidence of HCV infection and normal ALT levels, and are presented at the end of Table Ic.

*HCV infection in hemophiliacs receiving virus inactivated plasma derivatives.* Four out of six patients who received exclusively heat-treated and SD-treated products (Table III) had evidence of HCV infection. Patients 1, 25, and 30 were 6–8 yr old and had severe hemophilia A. In 1985, they already had antibody to the three HCV antigens tested and viremia was found in two to three samples of each patient over the study period. The fourth patient (number 16, Fig. 2), a 22-yr-old with mild hemophilia A, as well as patient 24 (data not shown) who had received both heat-treated and non-heat-treated products, were HCV seronegative in 1985 but seroconverted to HCV. The immune response to C-100 was apparently lower than to 33C or core. Over time, the antibody level to C-100 declined rapidly but levels of anti-33C and core remained high. HCV viremia was detected one (patient 16) and two years (patient 24) after seroconversion, respectively. Interestingly, patient 24 maintained normal ALT levels during the entire study period.

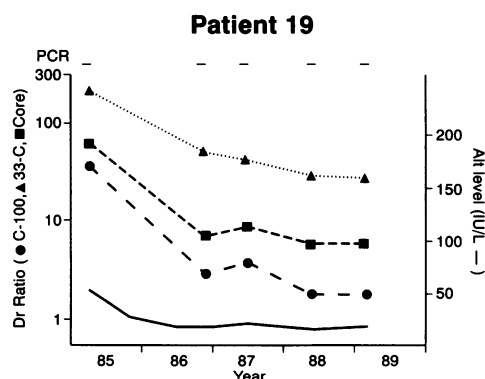


Figure 1. HCV markers in a nonviremic hemophiliac. ALT levels, HCV virologic and serologic markers of patient 19 followed up from 1985 to 1989.

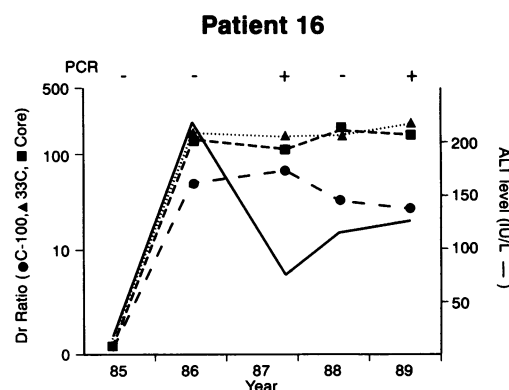


Figure 2. Acute HCV infection in a hemophiliac. ALT levels, HCV virologic and serologic markers of patient 16, a 22-yr-old patient with moderate hemophilia A who received exclusively heat or SD-treated Factor VIII concentrate. Acute hepatitis C is indicated by a peak of ALT, seroconversion to all three markers of HCV and delayed HCV viremia.

Table III. Treatment and Serological Pattern of Six Hemophiliacs Who Received Only Heat-treated Concentrates

Patient No.	Type of product used*			Maximum ALT level	HBV†		HCV‡		
					anti-S	anti-C	anti-C-100	33-C	Core
	(Total number of units)								
				<i>IU/liter</i>					
1	Hemofil T	(34,450)	1985–87	45	+	–	+	+	+
	CNTS/SD	(55,000)	1988–89						
16	Hemofil T	(4,000)	1985	284	+	+	+	+	+
25	Hemofil T	(30,560)	1984–85	36	+	–	+	+	+
	Kryobulin	(10,000)	1986						
27	CNTS/SD	(14,000)	1988–89	50	+	–	–	–	–
	Kryobulin	(18,500)	1985–87						
30	CNTS/SD	(41,500)	1988–89	22	+	–	–	–	+
	CNTS	(8,000)	1985–87						
33	CNTS/SD	(46,000)	1988–89	20	+	–	–	–	–
	CNTS/PPSB	(4,000)	1985–89						

\* Hemofil T was from Hyland, Glendale, CA, USA; CNTS heat-treated, Centre National de Transfusion Sanguine, Les Ulis, France; SD, inactivated with solvent detergent method; Kryobulin (TIM 3) was from Immuno AG, Vienna, Austria.

† Only patient 16 was naturally infected with hepatitis B; all others were vaccinated.

‡ Antibody pattern in 1989.

Data compatible with HCV superinfection during the course of the study were found in three additional cases. Patients 44 (Fig. 3), 41, and 48 (data not shown) had a very low level of antibody to C-100 (range 2–3 Dr ratio) and high levels of antibody to 33C and core in 1985. Levels of antibody to C-100 subsequently increased to a mean reflectance ratio of  $82.2 \pm 40.1$  (range 28–141) and then decreased over 2–3 yr to a mean level of  $36 \pm 29.7$  (range 0–77). In two of the three cases (patients 41 and 44), HCV RNA was not detectable initially but viremia became evident in these patients simultaneously or within 2 yr of the increased C-100 level. All patients had a concomitant elevation of ALT level. Patient 41 was a chronic carrier of hepatitis B surface antigen and had a previously reported delta hepatitis (20). He was receiving French heat-treated Factor VIII concentrate approximately three times a month during the period of anti-C-100 response. Patient 44

was under a prophylactic regimen of 20 U/kg of heat-treated intermediate purity Factor VIII every other day in 1986–87 (Fig. 3). He had developed immunity against HBV. Patient 48 had hemophilia B, received non-heat-treated and subsequently heat-treated PPSB during the period of his response to C-100 (30 U/kg per mo). He had serologic evidence of immunity to HBV.

**Role of HIV in HCV infection.** The various results collected in this study were analyzed according to the HIV immune status of patients. Among 46 HCV infected hemophiliacs, 33 (72%) were HIV seropositive and 13 seronegative. The mean levels of antibody to C-100, 33C or core, and the variations in levels of antibody to C-100 (undetectable, declining, or consistently high over time), were similar between the two groups. In contrast, the percentage of viremic samples in HIV seropositive patients (105/161 or 65.2%) was significantly higher than in seronegative patients (21/60 or 35%,  $P < 0.01$ ). A majority of HIV seropositive patients had chronic liver disease. 20 of the 33 seropositive patients (67%) were classified as code 1 (See Table I), compared to 5 of the 13 patients (38%) in the HIV seronegative group. However, the mean levels of ALT were similar between the two groups ( $63.6 \pm 51.0$  in HIV seropositive vs.  $49.0 \pm 60.6$  in HIV seronegative patients).

## Discussion

The data collected in this study differ from previously reported results on HCV in hemophilia. All exposed patients had serological evidence of HCV infection with the presence of antibody to the core protein of HCV and, in most cases, to the nonstructural 33C antigen. These two proteins were apparently more immunogenic than C-100 since antibody levels were higher and persisted longer. Thus, our data demonstrate that current seronegativity in 20% of the patients was related to a decline of antibody to C-100. However, this apparent seroreversion took place despite continued infection as shown in some patients with patterns 1 or 3 (Table I). In other anti-C-

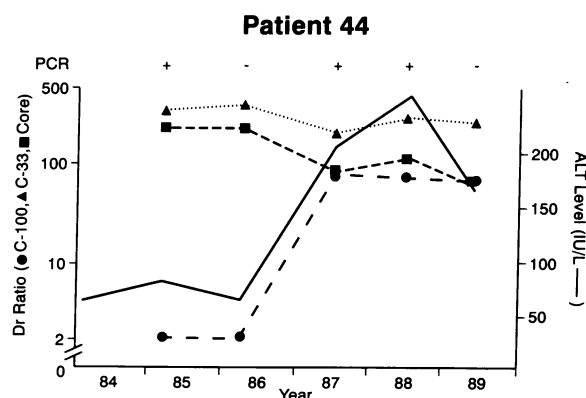


Figure 3. HCV markers suggestive of superinfection in a hemophiliac. ALT levels, HCV virologic and serologic markers of patient 44 who had chronic hepatitis before 1984. While antibody to HCV 33C and core remained at a high level, an immune response to C-100 was observed concomitant with a peak of ALT and transitory HCV viremia.

100 negative patients (15%), the lack of detection by current screening assays (C-100 based) may be attributed in part to the absence of a significant immune response to C-100. These results are consistent with previous observations in similar populations (10, 14, 21). In contrast, seroreversion for antibody to 33C was infrequent (6.5%) and not observed with anticore. There was no evidence that HIV infection played a significant role in this mechanism since it was observed with similar frequency irrespective of HIV status.

One of the features of this study was the frequency of viremia found in over 90% of patients over a 5-yr period during which they received successively heat-treated and SD-treated products. In a limited number of cases, the viremia patterns we described had been identified previously by Garson et al. (21). However, contrary to what was proposed by these investigators, several lines of evidence suggest that most HCV-infected hemophiliacs in this study have a persistent hepatitis C virus infection. First, HCV viremia was found in 42/46 (91.3%) exposed patients and, during the study period, there was no evidence for a decline in viremia. 29 hemophiliacs were viremic in the 1985 sample and 27 in the 1989 sample. The fact that viremia was detected intermittently in more than 50% of exposed patients is not surprising since the PCR method is known to have limitations (15, 22, 23). It has been reported that in persistent viral diseases like HIV or herpes, viral production may fluctuate over time (24, 25). Second, although we, like others, have found a clear influence of HIV infection in HCV viral production (Table Ia), apparently persistent HCV infection indicated by viremia was found in 11 out of 13 HIV seronegative hemophiliacs (85%). This frequency of HCV infection was not different from the HIV antibody positive group. Third, the cause of viral production does not appear related to superinfection to any large extent. The heat-treated concentrates used from 1985 to 1987 had a lower level of HCV infectivity than non-heat-treated concentrate, and there is no evidence that the SD-treated products might transmit hepatitis C (7). However, as exemplified in Fig. 3, we have found three cases whose antibody profile and viremia pattern suggested superinfection.

Some evidence of virus clearance and disease recovery was found only in a few patients with viremia pattern 3 or 4 (Table Ib and c). Patient 19 (Fig. 1) associated a consistent absence of detectable viremia, a progressive decline of antibody level to all three HCV antigens and a prolonged normalization of ALT level. Patients with pattern 3 may correspond to a stage preceding the features of patient 19. In four out of six patients, PCR was repeatedly negative and antibody level to C-100 was undetectable, low, or declining. However, antibody to core and, in most cases, to 33C remained elevated, and, except in patient 12, ALT levels were still moderately or occasionally elevated. In view of the limitations of the PCR method (14) and the potential fluctuations of viremia, one cannot exclude that some, if not all, of these patients might in fact belong to pattern 2. The use of more efficient cDNA synthesis and primer pairs for detection of viral RNA (26), availability of liver biopsies as a source of viral RNA, and testing of prospective samples from these patients may allow us to better categorize them. Unfortunately, liver biopsies are not easily obtained in hemophiliacs.

Patients 26, 29, and 42 appear immune to HBV, are clearly HCV seropositive, and have clinical and biological evidence of chronic liver disease, yet no detectable HCV RNA (Table Ic).

The liver condition is probably not related to HBV since HBsAg is not detected and antibodies to surface and core antigens are present. In addition, patients 26 and 42 are HIV infected. Four main hypotheses can be proposed to explain these findings. One is a false negativity of the HCV PCR due to inadequate precautions during sample handling or storage, or, alternatively, significant sequence heterogeneity at the primer or probe levels. Such hypothesis could be confirmed by using a primer pair from a more conserved region of the HCV genome such as the 5' noncoding region (26). A recent report has emphasized the relative diversity of nucleotide sequences in the NS3/NS4 regions that we used in our detection system (14). The second hypothesis is a variant of HBV or, as previously described, a level of HBV production below the HBsAg detection level (27). However, the presence of both anticore and antisurface antibody to HBV does not support this explanation. The third hypothesis is that the virus may be replicating in the liver but not released in the blood stream. Finally, a still unidentified hepatotropic virus may be involved. This hypothesis is unlikely since similar concentrates have been used for this and other groups of patients.

This study indicates that in hemophilia all exposed subjects have serological markers of HCV and suggest that HCV is a persistent infection. This feature may not be the rule in other groups of HCV infected patients since the interaction of HIV, HCV superinfection and potentially HBV might interfere with the natural history of HCV infection. The role of HBV, however, is likely to be minimal since eight of nine HBV vaccinated hemophiliacs had persistent HCV viremia. Prospective studies of patients receiving noninfectious blood products and examination of liver biopsies for the presence of viral RNA in hepatocytes should provide a final answer. The fact that, irrespective of HIV, hemophiliacs can be persistently infected and possibly superinfected with HCV suggest a limited efficacy of their immune system against HCV.

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