

Seroprevalence of HbsAg, Anti-HBc and Anti-HCV in Southern Brazil, 1999-2001

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The prevalence of infection by the hepatitis B (HBV) and C (HCV) viruses varies among geographical regions. We evaluated 263,795 blood donor samples collected from 1999-2001 in various cities in the state of Santa Catarina to determine the prevalence of HbsAg, anti-HBc and anti-HCV markers. The markers were analyzed by immunoenzymatic tests, as determined by the Ministry of Health, and the data were obtained from blood banks and from ANVISA (the Brazilian National Agency for Sanitary Vigilance). There was a significant reduction in the mean frequency of HbsAg and anti-HBc during the study period, from 0.98% to 0.64% and from 8.83% to 5.35%, respectively, though they varied considerably among the different regions. There was also a decrease in the mean frequency of anti-HCV, although it was not significant, decreasing from 0.38% to 0.34%. Even with this reduction, the frequency of these markers was still high compared with that found in other countries, indicating high rates of infection by hepatitis B and C viruses. This emphasizes the urgency of vaccination programs against HBV, especially in some regions of Santa Catarina state, in order to reduce the prevalence of this infection and consequently reduce the risk of transmission through sexual relations or from the donation of blood and/or hemocomponents.

Key Words: Blood donors, hepatitis B, hepatitis C, HbsAg, anti-HBc, anti-HCV.

Infection by the hepatitis B (HBV) and hepatitis C (HCV) viruses is the most common cause of post-transfusion hepatitis [1,2]. However, with the emergence of HIV infection more emphasis has been given to the control of blood utilized in transfusions and in 1993 it became obligatory, in Brazil, to screen blood donors for HBV and HCV [3]. Furthermore, HBV and HCV are also the most frequent causes of chronic hepatic diseases in the world, and their transmission occurs, mainly, through direct contact with blood, through the utilization of intravenous drugs, blood transfusions and/or

hemocomponents, and through sexual relations. However, sexual relations seem not to be the most frequent mode of HCV transmission [4].

Infection by these two viruses may induce chronic hepatitis, which may progress to cirrhosis, and eventually to hepatocellular carcinoma [5-8]. It is estimated that around 350 million people in the world are chronic carriers of HBV, which represents approximately 7% of the total population [9], whereas infection with HCV is found in approximately 3% of the world population, which represents 160 million people [10-12]. This high prevalence of HBV and/or HCV certainly results in high medical costs, due to a great number of cases of fulminating hepatitis, hepatic cirrhosis and carcinoma, and also provokes the death of a significant part of the population by these pathologies.

Studies show that a co-infection by HBV and HCV is as frequent in Asia [13,14] as it is in western countries [15,16], varying from 10% to 15% in patients who are chronically infected by HBV. Patients

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infected simultaneously by HBV and HCV are more frequently infected with the more severe hepatic diseases and are at a significantly greater risk of developing fulminant hepatitis, hepatic cirrhosis and carcinoma [17,18], particularly in the underdeveloped and/or developing countries, where the prevalence of simultaneous infections by these viruses appears to be higher. Also, the high prevalence of HBV and HCV may result in an increase in the risk of transmission of these viruses through the transfusion of hemocomponents, since it is not possible to totally guarantee the absence of these infections among blood donors through the serologic tests utilized routinely in the screening of blood donors [19].

We evaluated the prevalence of HbsAg, anti-HBc and anti-HCV markers among blood donors of the different regions of the state of Santa Catarina, Brazil, in 1999, 2000 and 2001.

Materials and Methods

The data were obtained from the data processing centers of the blood banks of the Centro Hemoterápico Blumenau (Blumenau Hemotherapeutic Center - Blumenau-SC), Hospital Dona Helena (Dona Helena Hospital - Joinville-SC), Hemocentro Jaraguaense LTDA (Jaraguaense Hemocenter Ltd - Jaraguá do Sul-SC), Banco de Sangue do Hospital Universitário (University Hospital Blood Bank - Florianópolis - SC), Centro de Hematologia e Hemoterapia de Santa Catarina - Hemosc - (Hematology and Hemotherapy Center of Santa Catarina - Florianópolis-SC) and Agência de Vigilância Sanitária ANVISA - (National Health Agency) [20].

The research on markers of infection by HBV and by HCV was carried out by immunoenzymatic tests according to Ministry of Health directives No. 1376 of 1993 and No. 121 of 1995.

Results

The prevalence of HbsAg and anti-HBc markers varied among the different regions of the state of Santa

Catarina; from 1999 and 2001 the mean prevalence in the state declined progressively. The highest prevalence of these two markers was found in the region of the town of Chapecó, in western Santa Catarina (Tables 1 and 2).

The mean prevalence of anti-HCV in the state of Santa Catarina did not change significantly from 1999 to 2001, though it varied among regions (Table 3).

Discussion

Blood donors are interviewed with screening questionnaires, which have evolved over the years [21,22]. In the United States, since 1992, following recommendations from the Food and Drug Administration, donors are directly questioned about behavioral habits [23]. This modification in the screening of the donors resulted in a decrease in the prevalence of HCV from 0.63% in 1992 to 0.40% in 1996 among new blood donors [24], whereas among the general population, at the same time, the prevalence of this virus was 1.8% [25]. In the case of HbsAg, a prevalence close to 0.2% was found, and there was no significant change from 1991 to 1996 [22]. In Brazil, where there is greater endemicity of HBV, as we know that the prevalence of HbsAg and anti-HBc among the blood donors in 2001 was similar to the prevalence among the general population of Germany [26], it appears that the anti-HBc research has assisted in the identification of HBV carriers who are seronegative for HbsAg [27].

The prevalence of infection by HCV among the blood donors in the state of Santa Catarina in 1999 was similar to that found among new blood donors in the United States in 1996 [24,28]. In Santa Catarina, the prevalence of HCV among blood donors decreased considerably between 1994/1995 and 1999 [19,28], but there was only a small reduction between 1999 and 2001. Among blood donors in the state of Santa Catarina there was a significant decrease in the prevalence of HbsAg and anti-HBc between 1999 and 2001 (Tables 1 and 2). However, the mean frequency of the markers of infection by HBV (HbsAg and anti-HBc) among blood donors in the state of Santa Catarina from 1999 to 2001 only decreased to levels similar to

Table 1. Prevalence of HbsAg among blood donors of the main regional blood banks of Santa Catarina

Year and Location	1999		2000		2001	
	N	%	N	%	N	%
Florianópolis	176/24.570	0.71	280/24.274	1.15	159/27.396	0.58
Lages	17/6.626	0.26	16/5.899	0.27	24/6.509	0.37
Joaçaba	64/8.036	0.80	30/5.633	0.53	27/5.668	0.48
Chapecó	375/11.713	3.20	149/9.132	1.63	122/7.909	1.54
Criciúma	36/7.193	0.50	20/6.858	0.29	23/7.493	0.31
Joinville	86/14.715	0.58	63/13.957	0.45	79/14.994	0.53
Tubarão	27/2.420	1.12	7/2.899	0.24	22/2.804	0.78
Blumenau	92/13.704	0.67	106/13.407	0.79	85/12.580	0.85
Jaraguá do Sul	20/2.201	0.91	17/2.317	0.73	20/2.888	0.69
Total	893/91.358	0.98	688/84.376	0.84	561/88.241	0.64

Table 2. Prevalence of anti-HBc among blood donors of the main regional blood banks of Santa Catarina

Year Location	1999		2000		2001	
	N	%	N	%	N	%
Florianópolis	1.303/24.570	5.30	1.097/24.274	4.52	968/27.396	3.53
Lages	157/6.626	2.37	106/5.899	1.80	138/6.509	2.12
Joaçaba	736/8.036	9.16	354/5.633	6.28	247/5.668	4.36
Chapecó	3.401/11.713	29.04	1.632/9.132	18.09	1.066/7.909	12.72
Criciúma	469/7.193	6.52	263/6.858	3.83	221/7.493	2.95
Joinville	1.055/14.715	7.17	951/13.957	6.81	635/14.994	4.23
Tubarão	134/2.420	5.54	79/2.899	2.73	110/2.804	3.92
Blumenau	565/13.704	4.12	1.251/13.407	9.33	1.164/12.580	9.25
Jaraguá do Sul	244/2.201	11.10	17/2.312.317	9.97	234/2.888	8.10
Total	8.064/91.358	8.83	5.984/84.376	7.09	4.723/88.241	5.35

Table 3. Prevalence of anti-HCV among blood donors of the main regional blood banks of Santa Catarina

Year Location	1999		2000		2001	
	N	%	N	%	N	%
Florianópolis	133/24.570	0.54	80/24.274	0.33	102/27.396	0.46
Lages	18/6.626	0.27	10/5.899	0.17	17/6.509	0.26
Joaçaba	17/8.036	0.21	10/5.633	0.18	23/5.668	0.41
Chapecó	39/11.713	0.33	17/9.132	0.19	17/7.909	0.21
Criciúma	42/7.193	0.58	27/6.858	0.39	30/7.493	0.40
Joinville	27/14.715	0.58	26/13.957	0.19	27/14.994	0.18
Tubarão	25/2.420	0.18	10/2.899	0.34	9/2.804	0.32
Blumenau	36/13.704	1.03	81/13.407	0.60	74/12.580	0.59
Jaraguá do Sul	9/2.201	0.26	1/2.317	0.04	5/2.888	0.17
Total	346/91.358	0.38	262/84.376	0.31	304/88.241	0.34

those found in Germany population during the same period, while the frequency of the marker of HCV infection (anti-HCV) was approximately 50% lower [29], though it was approximately five times greater than that found between 1991 and 1993 in blood donors in the United States [30].

The reduction in the prevalence of markers of HBV and HCV infection, which signified a significant reduction in the residual risk of transmission of these infections through blood transfusion, may be attributed to the introduction of the third generation ELISA tests, which increased the detection sensitivity for these viruses, reducing the window of detection, and also due to the effort employed by the blood banks in recruiting a greater number of volunteers who donate blood periodically. This is also a consequence of the efforts aimed at detecting behavior that increases the risk of infection by these viral agents and by HIV [21,31], which results in the exclusion and/or auto-exclusion of suspect donors. This reduction is also due to recent efforts to vaccinate children and fertile age women against hepatitis B.

Nevertheless, a residual risk of transmission of HBV and HCV by transfusion of contaminated blood persists, as a consequence of the infectious window period [21]. This risk could be reduced even more through the introduction of tests based on the polymerase chain reaction (PCR) or the increase in nucleic acids (NAT), since these tests, developed at the end of the last decade [32-35] have greater sensitivity and specificity, thereby decreasing the amplitude of the infectious window period and the residual risk of transmission via blood transfusion. However, although the cost-benefit relation may seem low for developed countries, developing countries are evaluating their utilization in areas where high prevalence of these infections has been found [21].

The average residual risks of transmission of HBV and HCV infection, 1:2077 and 1:13721, respectively, is still very high in Florianópolis, SC, Brasil, [21], when we compare these figures to those known from the United States in 1996, 1:63,000 and 1:103,000, respectively [36].

Given the high prevalence of HBV markers in the different regions of the state of Santa Catarina, the high risk of transmission of HBV infection through transfusions and the high degree of success attained by the mass vaccination programs against hepatitis B, implemented in Taiwan and Saudi Arabia, which resulted in drastic reductions in the prevalence of HbsAg and anti-HBc markers in the populations of these two countries, after 8 and 15 years, respectively [37,38], efforts should be made to implement mass hepatitis B vaccination campaigns for the population from 0-15 years of age and for fertile age women, in order to reduce the prevalence of HBV among the population and thereby more rapidly reduce the risk of transmission of this virus. This type of measure seems to be even more urgent in regions of high prevalence, such as Chapecó, located in western Santa Catarina, where the prevalence of HbsAg and anti-HBc markers among blood donors, found from 1999 to 2001, was approximately three times greater than the state average; sexual transmission and transmission through blood transfusions may be interacting in this region, increasing the total risk of transmission of HBV and thereby progressively increasing the prevalence of the markers of this infection.

References

1. Schreiber G.B., Busch M.P., Kleinman S.H., Korelitz J.J. The risk of transfusion-transmitted viral infections. The Retrovirus Epidemiology Donor Study. *N Engl J Med* **1996**;334(26):1685-90.
2. Kleinman S., Busch M.P., Korelitz J.J., Schreiber G.B. The incidence/window period model and its use to assess the risk of transfusion-transmitted human immunodeficiency virus and hepatitis C virus infection. *Transfus Med Rev* **1997**;11(3):155-72.
3. Gonçalves Junior F.L. Prevenção das hepatites pós-transfusionais. In: Covas, D.T. & Zago, M.A. eds. *Atualização em Hemoterapia*, vol. 5, Ribeirão Preto: Gráfica Canavaci **1998**.
4. Wright T.L., Hollander H., Pu X., Held M.J. Hepatitis C in HIV-infected patients with and without AIDS: prevalence and relationship to patient survival. *Hepatology* **1994**;20(5):1152-5.

5. Benvegna L., Fattovich G., Noventa F., et al. Concurrent hepatitis B and C virus infection and risk of hepatocellular carcinoma in cirrhosis. A prospective study. *Cancer* **1994**;74(9):2442-8.
6. Kaklamani E., Trichopoulos D., Tzonou A., et al. Hepatitis B and C viruses and their interaction in the origin of hepatocellular carcinoma. *JAMA* **1991**;265(15):1974-6.
7. El-Refaie A., Savage K., Bhattacharya S., et al. HCV-associated hepatocellular carcinoma without cirrhosis. *J Hepatol* **1996**;24(3):277-85.
8. Simonetti R.G., Camma C., Fiorello F., et al. Hepatitis C virus infection as a risk factor for hepatocellular carcinoma in patients with cirrhosis. A case-control study. *Ann Intern Med* **1992**;116(2):97-102.
9. Kao J.H., Chen P.J., Lai M.Y., Chen D.S. Occult hepatitis B virus infection and clinical outcomes of patients with chronic hepatitis C. *J Clin Microbiol* **2002**;40(11):4068-71.
10. Sánchez N.M., González H.B., Gómez R.H.S., et al. Prevalência de hepatitis B y C em donadores de sangue em um hospital de tercer veç de Ciudad de México. *Salud Publica de México*. **1999**;41(6):475-8.
11. Frider B. Epidemiologia de laof hepatitis C. *Acta Gastroenterol Latinoam* **2000**;30(2):142-4.
12. Zou S., Tepper M., Giulivi A. [Current status of hepatitis C in Canada.] *Can J Public Health* **2000**;91 Suppl 1:S10-5, S10-6.
13. Sato S., Fujiyama S., Tanaka M., et al. Coinfection of hepatitis C virus in patients with chronic hepatitis B infection. *J Hepatol* **1994**;21(2):159-66.
14. Ohkawa K., Hayashi N., Yuki N., et al. Hepatitis C virus antibody and hepatitis C virus replication in chronic hepatitis B patients. *J Hepatol* **1994**;21(4):509-14.
15. Fattovich G., Tagger A., Brollo L., et al. Hepatitis C virus infection in chronic hepatitis B virus carriers. *J Infect Dis* **1991**;Feb;163(2):400-2.
16. Crespo J., Lozano J.L., de la Cruz F., et al. Prevalence and significance of hepatitis C viremia in chronic active hepatitis B. *Am J Gastroenterol* **1994**;89(8):1147-51.
17. Chu C.M., Sheen I.S., Liaw Y.F. The role of hepatitis C virus in fulminant viral hepatitis in an area with endemic hepatitis A and B. *Gastroenterology* **1994**;Jul;107(1):189-95.
18. Zarski J.P., Bohn B., Bastie A., et al. Characteristics of patients with dual infection by hepatitis B and C viruses. *J Hepatol*. **1998**;28(1):27-33.
19. Kupek E.J. Residual transfusion risk for hepatitis B and C in southern Brazil, 1991-99. *J Viral Hepat* **2001**;8(1):78-82.
20. www.anvisa.org.br. consulted June **2002**
21. Zuck T.F. Transfusion-transmitted AIDS reassessed. *N Engl J Med* **1988**;318(8):511-2.
22. Glynn S.A., Kleinman, S.H., Schreiber, G.B., et al. Trends in incidence and prevalence of major transfusion-transmissible viral infections in US blood donors, 1991 to 1996. Retrovirus Epidemiology Donor Study (REDS) *JAMA* **2000**;284(2):229-35.
23. Kleinman S. Blood donor screening: principles and policies. In: Petz, L.D., Swisher, S.N., Kleinman, S., Spence, R.K., Strauss, R.G., eds. *Clinical Practice of Transfusion Medicine*. 3rd ed. New York, NY: Churchill Livingstone INC; **1996**;245-70.
24. Alter H.J., Conry-Cantilena C., Melpolder J., et al. Hepatitis C in asymptomatic blood donors. *Hepatology*. **1997**;26(3 Suppl 1):29S-33S.
25. Alter M.J., Kruszon-Moran D., Nainan O.V., et al. The prevalence of hepatitis C virus infection in the United States, 1988 through 1994. *N Engl J Med* **1999**;341(8):556-62.
26. Jilg W., Hottentrager B., Weinberger K., et al. Prevalence of markers of hepatitis B in the adult German population. *J Med Virol* **2001**;63(2):96-102.
27. Martelli C.M.T., Turchi M.D., Souto F.J.D. et al. Anti-HBc testing for blood donations in areas with intermediate hepatitis B endemicity. *Rev Panam Salud Publica* **1999**;6(1):69-73.
28. Treitinger A., Spada C., Ferreira L.A., et al. Hepatitis B and hepatitis C prevalence among blood donors and HIV-1 infected patients in Florianopolis—Brazil. *Braz J Infect Dis* **2000**;4(4):192-6.
29. Palitzsch K.D., Hottentrager B., Schlottmann K., et al. Prevalence of antibodies against hepatitis C virus in the adult German population. *Eur J Gastroenterol Hepatol*. **1999**;11(11):1215-20.
30. Korelitz J.J., Busch M.P., Kleinman S.H., et al. Relationship between antibody to hepatitis B core antigen and retroviral infections in blood from volunteer donors. *Transfusion* **1996**;36(3):232-7.
31. Kupek E.J. HIV seroprevalence among blood donors in southern Brazil in the decade of 1990. *Braz J Infect Dis* **2000**;4(5):217-25.
32. Schottstedt V., Tuma W., Bunger G., Lefevre H. PCR for HBV, HCV and HIV-1 experiences and first results from a routine screening programme in a large blood transfusion service. *Biologicals* **1998**;26(2):101-4.
33. Yerly S., Pedrocchi M., Perrin L. The use of polymerase chain reaction in plasma pools for the concomitant detection of hepatitis C virus and HIV type 1 RNA. *Transfusion* **1998**;38(10):908-14.
34. Roth W.K., Weber M., Seifried E. Feasibility and efficacy of routine PCR screening of blood donations for hepatitis C virus, hepatitis B virus, and HIV-1 in a blood-bank setting. *Lancet* **1999**;353(9150):359-63.
35. Bush M.P., Stramer S.L., Kleinman S.H. Evolving applications of nucleic acid amplifications assays for prevention of virus transmission by blood components and derivatives. In: Garratty, G., ed. *Applications of Molecular Biology in Blood Transfusion*. Bethesda, Md: American Association of Blood Banks **1997**:123-176.

36. Goodnough L.T., Brecher M.E., Kanter M.H., AuBuchon J.P. Transfusion medicine. First of two parts—blood transfusion. *N Engl J Med* **1999**;340(6):438-47.
37. Ni Y.H., Chang M.H., Huang L.M., et al. Hepatitis B virus infection in children and adolescents in a hyperendemic area: 15 years after mass hepatitis B vaccination. *Ann Intern Med* **2001**;135(9):796-800.
38. Al-Faleh F.Z., Al-Jeffri M., Ramia S., et al. Seroepidemiology of hepatitis B virus infection in Saudi children 8 years after a mass hepatitis B vaccination programme. *J Infect* **1999**;38(3):167-70.