

How to Cite:

Kareem, H. F., Abdullah, M. A. K., & Kadim, A. M. (2022). Hepatitis screen in children with malignancy at child's central teaching hospital. *International Journal of Health Sciences*, 6(S8), 2415–2422. <https://doi.org/10.53730/ijhs.v6nS8.12370>

Hepatitis screen in children with malignancy at child's central teaching hospital

Dr. Hind Fadhil Kareem

Babylon health directorate, Babylon, Iraq
Corresponding author email: qaisajam1981@gmail.com

Dr. Mohammed Abdul Kareem Abdullah

Babylon health directorate, Babylon, Iraq

Abbas Mehdi Kadim

Babylon health directorate, Babylon, Iraq

Abstract---Introduction: Transfusion-transmitted infections continue to be a threat to the safety of the blood supply, in particular the risk is high for parenterally transmitted viral hepatitis in pediatric malignancy. The aim was to estimate the prevalence of hepatitis in children with malignancy, identify some variables that could affect the prevalence of hepatitis in these patients, and to have an idea about the effect of vaccination in controlling hepatitis infection. Method: A Cross sectional study of (180) children between the age of [1-15] years with malignancy, in the period seven months, diagnosed and treated in CCTH were studied. Among those children with malignancy, testing for HBV and HCV were done through blood samples taken from the patients and sent to lab. of CCTH. Results: patients were referred from different parts of the country, from (180) patients with malignancy, 48(27%) were HBV positive &33(20%) with HCV positive. History of clinical jaundice was reported in 33 of cases (18%). The no. of blood transfusion had significant impact on prevalence of hepatitis virus infection. Conclusion: There is a high prevalence of HBV&HCV infection in patients with malignancy treated in Child's Central Teaching Hospital (CCTH).

Keywords---hepatitis screen, children, malignancy, child's central teaching hospital.

Introduction

Viral hepatitis is a major Health problem in both developing and developed countries. This disorder is caused by at least five pathogenic hepatotropic viruses

are designated hepatitis A, B, C, D, E and G, many other viruses can cause hepatitis as one component of multisystem disease which includes herpes simplex virus, cytomegalo virus, Epstein-Barr virus, varicella-zoster virus, HIV, rubella, adenoviruses, enteroviruses, parvovirus B19 and arboviruses. Hepatitis G virus (HGV) and transfusion transmissible virus (TTV) often infect the liver as a co-infection with another hepatotropic virus, and may produce acute or chronic viremia but rarely produce hepatocellular injury on their own ⁽¹⁾. Transfusion-transmitted infections (TTI) continue to be a threat to the safety of the blood supply. Viral infections cause the major part of mortality and morbidity in blood recipients ⁽²⁾. Patients treated for pediatric malignancy are at high risk for parenterally transmitted viral hepatitis ⁽³⁾. Blood product transfusion is the major risk factor. Moreover, when compared with immunocompetent patients, the immunodepression caused by chemotherapy increases the chronicity rate of viral hepatitis ⁽⁴⁾. During the last 2 decades, screening blood donors for the hepatitis B virus (HBV) has resulted in a remarkable reduction of post-transfusion B virus hepatitis. Thus, HCV has become the major cause of the parenterally transmitted hepatitis ⁽⁵⁾. Over the two last decades, much attention has been given to the prevention of transfusion-transmitted viral infections such as HIV-1 and -2, human T cell lymphotropic virus (HTLV) I and II, hepatitis C virus (HCV), hepatitis B virus (HBV) and West Nile Virus (WNV). Given the potential transmission of viruses during the 'immunological window period' [i.e. the period of early infectivity when an immunologic test is non-reactive] ⁽⁶⁾. However, it has some limitations in blood components with very low levels of viremia, which can even escape detection by NAT. Despite this limitation, the combination of both serological testing and NAT has considerably reduced the risk of viral transmission by blood transfusion ⁽⁷⁾. In order to maintain the integrity, purity and adequacy of the blood supply new donor screening assays, donor deferral and pathogen inactivation of blood components need to be balanced against the undue loss of potential donors because of overly stringent exclusion criteria ⁽⁸⁾. The aim of study is to find out the overall prevalence of HBV and HCV among patients with malignancy in CCTH and to find out the effect of frequent blood transfusion or blood product transfusion on the prevalence of hepatitis B and C.

Method

A cross sectional study was carried out in the oncology department of child's central teaching hospital in Baghdad city. A total of (180) children between the age of [1-15] years with cancer, over seven months between December 1st, 2009 to June 30th, 2010, diagnosed before 6 months and more and treated in CCTH were studied, (from a total of 180 patients, 3 patients were not tested for HBs Ag. And 15 patients were not tested for anti HCV antibody). Information regarding (name of patient, date of birth, sex, date of diagnosis, hepatitis screen at time of diagnosis, number of hepatitis vaccines given, number and place of blood transfusion, and history of jaundice) were obtained. These data were taken from hospital records, questionnaire of the parents & from the patient's own medical notebook (provided to parents during first hospitalization) which contain all information regarding inpatient & outpatient visits with the detailed management. The results of hepatitis screen at time of diagnosis of cancer were taken from the hospital records. Among those children with malignancy, evaluation for HBV and HCV were done through blood samples taken from them and sent to lab. of CCTH.

Hepatitis B Surface Antigen and Anti Hepatitis C Virus Antibody were investigated by commercially available ELISA diagnostic kit techniques. For Hepatitis B; Bioelisa HBsAg 5,6 (Biokit) for screening tests made in Spain. For Hepatitis C; Bioelisa HCV 4,0 (Biokit) for screening tests made in Spain. No confirmatory tests were available for both HBV and HCV.

Results

The age (years) was [1-4 (42 patients 23.3%), >4-9 (81,45%), >9-15(57,31.7%)] mean age was 7.6 years, females were more predominant than males [93(51.7%)], the patients were referred from different parts of the country; however the majority were from other governorates [114(63.3%)], No. of HB vaccines were [3 vaccines 78(43.3%), <3 vaccines 76(42.2%), not given 26(14.4%)].(Table 3)

Table 3: Demographic & clinical data of 180 patients

	No	%
Age (years) 1—4	42	23.3
>4—9	81	45.0
>9—15	57	31.7
Mean±SD (Range)	7.68±3.73	
Sex Male	87	48.3
Female	93	51.7
Residence Baghdad	66	36.7
Others	114	63.3
Number of Hep B vaccines		
3 vaccines	78	43.3
<3 vaccines	76	42.2
Not given	26	14.4

Of total 180 patients, only 177 were tested for HBsAg. while three patients were not tested. The impact of some variables on prevalence of HBV infection, these variables include age, sex, and residence. The most predominant age was between > 4-9 years [21(26.9%)]. The residence has significant effect on prevalence of HBV [p value (0.0001)].(Table2)

Table 2: Impact of Age, Sex & Residence of patients on the prevalence of HBV infection

	HBsAg positive		Negative		P value
	No	%	No	%	
Age (years) 1—4	12	28.6	30	71.4	0.968
>4—9	21	26.9	57	73.1	
>9--15	15	26.3	42	73.7	
Mean±SD (Range)	7.92		7.65		
Sex Male	27	32.1	57	67.9	0.153
Female	21	22.6	72	77.4	

Residence	Baghdad	6	9.1	60	90.9	0.0001*
	Others	42	37.8	69	62.2	

*Significant using Pearson Chi-squared test at 0.05 level of significance

Of total 180 patients, only 165 were tested for HCV. while 15 patients were not tested. The impact of some variables on prevalence of HCV infection, The variables had no impact on prevalence of HCV infection[age(p value 0.094), sex(p value 0.185), residence(p value 0.326)].(Table 3)

Table 3: Impact of Age, Sex & Residence of patients on the prevalence of HCV infection

	Anti HCV Ab positive		Negative		P value	
	No	%	No	%		
Age (years)	1-4	5	13.9	31	86.1	0.094
	>4--9	12	16.0	63	84.0	
	>9--15	16	29.6	38	70.4	
	Mean±SD	9.97±3.07		7.24±3.71		
Sex	Male	19	24.4	59	75.6	0.185
	Female	14	16.1	73	83.9	
Residence	Baghdad	9	15.8	48	84.2	0.326
	Others	24	22.2	84	77.8	

Of 33 patients with jaundice, 21 patients were positive for only HBV, 9 patients were positive for only HCV and 3 patients were positive for both HBV & HCV. So the prevalence of HBV & HCV is more among patients with jaundice than those without jaundice (Table 4, 5).

Table 4: Impact of Jaundice & Vaccination on the prevalence of HBV infection

	HBsAg positive		Negative		P value	
	No	%	No	%		
Jaundice						
	Positive	24	72.7	9	27.3	0.0001*
	Negative	24	16.7	120	83.3	
Number of HB vaccines						
	3 vaccines	13	16.7	65	83.3	0.019*
	<3 vaccines	25	34.2	48	65.8	
	Not given	10	38.5	16	61.5	

Table 5: Impact of Jaundice on the prevalence of HCV infection

	Anti HCV Ab positive		Negative		P value	
	No	%	No	%		
Jaundice						
	Positive	12	36.4	21	63.6	0.009*

Negative	21	15.9	111	84.1
----------	----	------	-----	------

Of total 180 patients, only 162 patients received blood (18 patients were not received blood).The prevalence of HBV were high among patients who received blood more than three times therefore the number of blood transfusion had significant effect, while the place had no significant effect on prevalence of HBV(Table 6, 7).

Table 6: Number &place of blood transfusions

Number of blood transfusions	No	%
No transfusion	18	10.0
1--3	90	50.0
>3	72	40.0
Place of blood transfusion		
CCTH	123	75.9
Others	39	24.1

Table 7: Impact of Number &Place of blood transfusions on the prevalence of HBV infection

	HBsAg positive		Negative		P value
	No	%	No	%	
Number of blood transfusions					
No	3	16.7	15	83.3	0.034*
1—3	18	20.7	69	79.3	
>3	27	37.5	45	62.5	
Place of blood transfusion					
CCTH	32	26.0	91	74.0	0.237
Others	13	36.1	23	63.9	
Anti HCV Ab					
Positive	9	30.0	21	70.0	0.961
Negative	39	29.5	93	70.5	

The results show the number & place of blood transfusion had no correlation with the prevalence of HCV infection [No. of blood transfusions (no p value), place of blood transfusion (p value 0.412), HBsAg (p value 0.961)].(Table9).

Table 8: Impact of Number &Place of blood transfusions on the prevalence of HCV infection

	Anti HCV Ab positive		Negative		P value
	No	%	No	%	
Number of blood transfusions					
No	-	-	12	100.0	-
1--3	18	22.2	63	77.8	

>3	15	20.8	57	79.2	
Place of blood transfusion					
CCTH	23	20.0	92	80.0	0.412
Others	10	26.3	28	73.7	
HBsAg					
Positive	9	18.8	39	81.3	0.961
Negative	21	18.4	93	81.6	

Discussion

The study has taken a sample of 180 patients with malignancy attending the oncological department of child's central teaching hospital so the prevalence of HBS Ag. among those patients was (27.1%), and the prevalence of anti-HCV was (20%). These results are comparably similar to the results to the study done in CWTH⁽⁹⁾ that found the prevalence of HBS Ag. was (27.3%) but higher than the prevalence of HCV that was (7.8%). Another study carried by Ali S.M.⁽¹⁰⁾ included 607 patients with leukemia& lymphoma at CCTH, hepatitis B surface antigen was positive in 20/607 (3.29%) and Anti-HCV was 6/607 (0.98%). Mollah AH⁽¹¹⁾ from Bangladesh showed that The HBV and HCV-markers were found significantly more often among multi-transfused thalassaemic children than among the controls in terms of HBsAg (13.8% vs 6.5%, $p < 0.04$) and anti-HCV (12.5% vs 0.9%, $p < 0.0001$) Another study done in Brazil showed that the overall prevalence of HCV, HIV, HBV and co-infection among MTP were 16.7%, 1.7%, 0.8% and 1.7% respectively⁽¹²⁾. Children with cancer require multiple transfusions during intensive therapy are at an increased risk of blood transmissible infections, such as HBV and HCV infections. The need for frequent blood counts, intravenous therapy and surgery in addition to the immunosuppressed status of these patients further increase the risk⁽¹³⁾. The variables are age, sex and residence which may have an impact on the prevalence on HBV &HC V infection. The effect of age may have related to two directions; first: whether the prevalence of hepatitis in the younger age group is lower due to high efficacy of hepatitis vaccine received through the (NIP), second: whether chronic hepatitis is affected by age group as mentioned previously, the older the age of acquisition, the lower the risk of chronic disease⁽¹⁴⁾. This study didn't show any significant association between prevalence of hepatitis and age, also similar results regarding the sex, this results approximately similar to the results of study carried in CWTH⁽⁹⁾, But different from the results of the other study carried by Ali S.M⁽¹⁰⁾. This study shows that patients with history of clinical jaundice were (18%), among those with jaundice were HBsAg positive in (72.7%) and HCV Ab positive in (36.4%), these results higher than the results carried in CWTH⁽⁹⁾. It showed that patients received three doses of HBV vaccine had better protection against HBV in comparison with those who received less than three doses or never taken any dose of HBV vaccine, on the other hand some children who received three doses of vaccine but do not get protection against infection with HBV, which might raise the question of the efficacy of the vaccines given, the possibility of attenuation of immune system during period of receiving chemotherapy and the reliability of the history taken from the parents . This results were different from the results of study occurred in CWTH⁽⁹⁾. This study showed the risk of blood transfusion for transmission of HBV increase with the

increased in number of blood transfusion, while risk of transfusion transmitted viral infection (HCV, HBV) in France for the 2001-2003 periods was estimated at 1 in 10 million for HCV and at 1 in 640 000 for HBV ⁽¹⁵⁾.

Conclusion

There is a high prevalence of HBV&HCV infection in patients with malignancy treated in Child's Central Teaching Hospital (CCTH). But there is lower incidence of HCV infections than HBV. Blood transfusions (more than 3 times) is significantly increasing the incidence of HBV infection and HBV vaccine is significantly reducing the incidence of HBV infection.

References

1. Ali S.M .The prevalence of hepatitis B and C serological markers among patients with leukemia and lymphoma at child's central teaching hospital. Athesis Submitted to the scientific council of pediatrics in partial fulfillment for the degree of fellowship of the Iraqi Board for medical specialization in pediatrics. 2007.
2. AL-khafagi M.A. Hepatitis screen in children with cancer at Children Welfare Teaching Hospital. A thesis Submitted to the Scientific Council of Pediatrics in Partial Fulfillment of the requirement for the degree of Fellowship of the Iraqi Board for Medical Specializations.2008.
3. Andreu G, Morel P, Forestier F. et al Hemovigilance network in France: organization and analysis of immediate transfusion incident reports from 1994 to 1998. *Transfusion*. 2002; 42:1356–1364.
4. Brickman H., Oldenburg J., Eis-Hubinger A.M., et al. Hepatitis A virus infection among the hemophilus population at the Bonn Hemophilia centre. 1994: 67 (suppl.), 3-8.
5. De Paula EV, Gonçalves NS, Xueref S, Addas-Carvalho M, Gilli SC, Angerami RN, Veríssimo MP, Gonçalves FL. Transfusion-transmitted infections among multi-transfused patients in Brazil. *J Clin Virol*. 2005 Dec;34 Suppl 2:S27-32.
6. Fanning L.,Kenny E., Sheehan M.: viral load and clinicopathological features of chronic hepatitis C(Ib) in a homogeneous patient population. *Gut*, 1994, 44:563-7.
7. Fioredda,-F; Plebani,-A; Hanau,-G; Haupt,-R; Giacchino,-M; Barisone,-E; Blbo,-L; Castagnola,-E.Re- Immunisation schedule in leukaemic children after intensive chemotherapy: a possible strategy. *Eur-J-Haematol*. 2005 Jan.; 74 (1): 3-20.
8. Kebudi R, Ayan I, Yilmaz G, Akıcı F, Görün O, Badur S. Seroprevalence of hepatitis B, hepatitis C, and human immunodeficiency virus infections in children with cancer at diagnosis and following therapy in Turkey. *Med Pediatr Oncol*. 2000 Feb;34(2):102-5.
9. Kebudi R., Aya I. and Yilmaz G.: Seroprevalence of hepatitis B, hepatitis C and HIV virus infection with cancer at diagnosis following therapy in Turkey. *Med. Pediatr. Oncol.*, 2000, 34(2): 102-5.
10. Locascuilli A., Alberti A. and de bock R.: Impact of liver disease and hepatitis infections on allogeneic bone marrow transplantation in Europe. *Bone Marrow Transplant*, 1994: 14: 833-7.

11. Mohammed, A. A., Abdulsattar, S. A., & Al-bayati, S. (2022). Viral load and B-catenin level correlation study in patients with chronic hepatitis B virus. *International Journal of Health Sciences*, 6(S2), 3278–3287. <https://doi.org/10.53730/ijhs.v6nS2.5815>
12. Mollah AH, Nahar N, Siddique MA, Anwar KS, Hassan T, Azam MG. Common transfusion-transmitted infectious agents among thalassaemic children in Bangladesh. *J Health Popul Nutr*. 2003; 21(1): 67-71.
13. Pillonel J, Laperche; Groupe” Agents Transmissibles par Transfusion”de la Société française de transfusion sanguine; Etablissement français dusang; Centre de transfusion sanguine des arméesç. Trends in residual risk of transfusion-transmitted viral infections (HIV, HCV, HBV) in France between 1992 and 2002 and impact of viral genome screening (Nucleic Acid Testing). *Transfus Clin Biol*. 2004 Apr;11(2):81-6.
14. Schuttler CG, Caspari G, Jursch CA. et al Hepatitis C virus transmission by a blood donation negative in nucleic acid amplification tests for viral RNA. *Lancet*. 2000; 355:41–42.
15. Suryatika, I. B. M., Anggarani, N. K. N., Poniman, S., & Sutapa, G. N. (2020). Potential risk of cancer in body organs as result of torak CT-scan exposure. *International Journal of Physical Sciences and Engineering*, 4(3), 1–6. <https://doi.org/10.29332/ijpse.v4n3.465>
16. tramer SL, Glynn SA, Kleinman SH. et al Detection of HIV-1 and HCV infections among antibody-negative blood donors by nucleic acid-amplification testing. *N Engl J Med*. 2004;351:760–768.
17. Widana, I.K., Sumetri, N.W., Sutapa, I.K., Suryasa, W. (2021). Anthropometric measures for better cardiovascular and musculoskeletal health. *Computer Applications in Engineering Education*, 29(3), 550–561. <https://doi.org/10.1002/cae.22202>
18. YaZigi N., Balistreri WF. Viral Hepatitis . Nelson Text Book Of Pediatrics . 18th edition. Saunders. Philadelphia; 2007 . 1680-1687.