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# Hepatitis C Virus Seroconversion Among Hemodialysis Patients and the Role of Hepatitis C Virus Positive Patient's Isolation in Benha, Egypt

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**Abstract:** The prevalence of hepatitis C virus (HCV) infection in hemodialysis units (HD) is higher than among normal population. Seroconversion was included in many previous studies which constitute a great problem against infection control policies. The aim of this study was to evaluate seroconversion rate and the effect of isolation of hepatitis C positive patients as infection control method. This is controlled prospective study that included 90 patients for 2 years. Isolation policy of hepatitis C positive patients was implemented in the second year of the study. The prevalence of HCV was 48.9% among hemodialysis patients. Seroconversion rate decreased from 15.2% in the first year to 5.1% in the second year after application of isolation. The duration of hemodialysis in months, positive history of blood transfusion and the amount of transfused blood were considered significant factors affecting seroconversion. so we concluded that Isolation of hepatitis C positive patients as an infection control policy is mandatory to control HCV seroconversion in Egypt.

**Keywords:** HCV, Isolation, Seroconversion and HD

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## 1. Introduction

End stage renal disease (ESRD) has become a public Health problem worldwide as the total number of patients was increasing due to the increased prevalence of hypertension and diabetes mellitus [1]. Also ESRD patients are requiring different modality of renal replacement therapy (RRT), which put more burdens on health budget especially in developing countries [2].

Egypt has the largest burden of HCV infection worldwide which is the most common chronic blood-borne infection [3, 4]. The at risk groups include patients that receive multiple blood transfusions as hemophiliacs, individuals who are intravenous and inhalant drug users and hemodialysis patients [5].

The prevalence of HCV infection among HD patients is generally much higher than general population due to underlying impaired cellular immunity which increases their susceptibility to infection. Also HD requires prolonged vascular access and exposure to contaminated equipment. In

addition HD patients required blood transfusion, frequent hospitalization and surgery, which increase opportunities forgetting nosocomial infection exposure [6]. HD staff was found also to be an important factor in transmission of HCV infections among HD patients [7].

The prevalence of HCV infection among HD patients in developed countries ranges from 3.6 to 20% and is higher in the developing countries [8] and in Egypt, it ranges from 49% to 64% [9].

HCV is responsible for over one million patient's deaths from cirrhosis and liver cancer every year [4]. It has been associated with high morbidity and mortality rates and the management of these infections among HD patients with specific antiviral agents is associated with high rates of adverse effect [10].

Isolation of HCV positive patients in HD units is recommended by many authors [11, 12].

The rationale of this study was to declare that isolation of hepatitis C positive patients as an infection control policy is mandatory to control HCV seroconversion in Egypt.

## 2. Patients and Methods

This controlled prospective study was conducted at HD unit, Benha university hospital from October 2014 to October 2016. We included all the patients who were on HD and at the end of the study and exclude all patients who transferred from our unit, died before the end of the study or received any HD sessions outside our unit. Also we did not include new HD patient who started HD after the study had been started. Final number of the study was 90 patients.

The aim of this study was to evaluate seroconversion rate and the effect of isolation of hepatitis C positive patients as infection control method.

All new patients who will start HD sessions must do virology testing for Human immunodeficiency virus (HIV), hepatitis B virus (HBV) and HCV, and it must be repeated every 3 months. But we don't accept HBV or HIV positive patients.

For all patients, complete history was taken including age, gender, duration of HD, any surgical procedure, family history of HCV infection, vascular access and history of blood transfusion.

For HCV diagnosis we used third generation enzyme-linked immunosorbant Assay (ELISA) for detection of HCV antibodies (AB). Follow up of patients was carried out without isolation in the first year; then we applied isolation policy of HCV positive patients in October 2015, and follow up the patients continued for another year.

Annual seroconversion (any patient who was HCV AB negative and became HCV AB positive ) rate was calculated as number of HCV positive seroconverted patients in one year subdivided by number of HCV negative patient and multiply by 100.

## 3. Statistical Analysis

The collected data were summarized in terms of mean  $\pm$ SD for quantitative data and frequency and percentage for categorical data. The test of proportion (Z-test), chi square test and fisher exact test were used to compare categorical variables while student t test was used to compare quantitative variables. A P-value <0.05 was considered statistically significant. All statistical analysis was carried out

using the computerized Statistical Package for Social Science (SPSS; Version 20.0 for Windows, SPSS Inc., Chicago, IL).

## 4. Results

This study included 90 patients on regular HD in Benha University Hospital, Egypt (56.7% were male and 43.3% were female). They had a mean age of 54.41 years (standard deviation 10.81). The prevalence of HCV infection was 48.9% (44 patients). Among the negative group for HCV infection (46 patients) who started HD, 9 patients seroconverted at the end of the study. Participants had been on dialysis for a mean duration of 44.39 months (standard deviation 18.1). About one third of them had positive history of blood transfusion (more than 50% of them get  $\geq 3$  units (Table 1).

Table 1. Characteristics of patients.

	Value
<b>Age in years</b>	
Mean $\pm$ SD (range)	54.41 $\pm$ 10.81 (21-76)
Sex n (%)	
Male	51(56.7)
Female	39(43.3)
HCV n (%)	
Positive	44(48.9)
Negative	46(51.1)
Vascular access	
AVF	63(70.0)
Catheter	27(30.0)
Duration of HD in months	
Mean $\pm$ SD (range)	44.39 $\pm$ 18.1 (12-120)
Blood transfusion n (%)	
Yes	30(33.3)
No	60(66.7)
Amount of blood transfusion (32)	
<3 units	14(46.7)
$\geq 3$ units	16(53.3)
Conversion from negative HCVAB to positive	
Yes	9/46
No	37/46

Table 2 showed that there were no significant difference between cohorts with HCV infection versus cohorts without it regarding age, sex and vascular access (p>0.05).

Table 2. Comparison between HCV positive and negative patients.

	Positive HCV (44)	Negative HCV (46)	test	P*
Age in years	53.07 $\pm$ 10.95	55.70 $\pm$ 10.62		
Mean $\pm$ SD (range)	(21-76)	(30-71)	#1.17	0.25
Sex n (%)				
Male	25(56.8)	26(56.5)	\$0.01	0.98
Female	19(43.2)	20(43.5)		
Vascular access				
AVF	31(70.5)	32(69.6)	\$0.01	0.93
Catheter	13(29.5)	14(30.4)		

#= student t test \$=chi square test

The cohort was divided into two groups: Group I consisted of 9 patients (66.7% male and 33.3% female) who were negative for HCV when they started HD and seroconverted

during HD to HCV positive and group II consisted of 37 patients (54.1% male and 45.9% female) who were negative for HCV when they started HD and remained negative.

Comparison between the studied groups with regard to the risk factors for HCV seroconversion showed that the duration of HD in months ( $67.11 \pm 12.26$  in group I and  $40.73 \pm 17.29$  in group II) and a positive history for blood transfusion and amount of transfused blood were the significant factors

between the two studied groups.

A positive family history of HCV infections and surgical operation history were non-significant among the studied groups (Table 3).

**Table 3.** Seroconversion to HCV seropositive patients among HCV seronegative patients.

	Group I Positive seroconversion (9)	Group II Negative seroconversion (37)	Test	P*
Age in years	$51.89 \pm 10.86$	$56.62 \pm 10.51$	#1.20	0.24
Mean $\pm$ SD (range)	(30-67)	(30-71)		
Sex n (%)				
Male	6(66.7)	20(54.1)	$\wedge$ 0.10	0.76
Female	3(33.3)	17(45.9)		
Duration of HD in months	$67.11 \pm 12.26$	$40.73 \pm 17.29$	#4.31	0.001**
Mean $\pm$ SD (range)	(49-90)	(12-90)		
Blood transfusion n (%)				
Yes	6(66.7)	29(78.4)	$\wedge$ 4.97	0.03*
No	3(33.3)	8(21.6)		
Amount of blood transfusion				
<3 units	1(16.7)	7(87.5)	$\wedge$ 4.43	0.04*
$\geq$ 3 units	5(83.3)	1(12.5)		
Surgery n (%)				
Yes	8(88.9)	22(59.5)	$\wedge$ 1.62	0.13
No	1(11.1)	15(40.5)		
Family history				
Yes	7(77.8)	23(62.2)	\$0.78	0.38
No	2(22.2)	14(37.8)		
After one year (before isolation)	7/46(15.2)	39(84.8)	£6.57	0.001**
After 2 years (after isolation)	2/39(5.1)	37(94.9)	£12.70	0.001**

#= student t test  $\wedge$ = fisher exact test \$= chi square test £=z test \*=sig \*\*=highly sig

After one year (before isolation) 15.2% of the negative group for HCV infection had converted while only 5.1% converted in the second year of the study (after isolation) which was statistically significant ( $p=0.02$ ) (Table 4).

**Table 4.** Seroconversion before and after isolation.

	Before isolation (46)	After isolation (39)	Z test	P*
Positive conversion	7(15.2)	2(5.1)	2.0	0.02*
Negative conversion	39(84.8)	37(94.9)	0.23	0.41

## 5. Discussion

Contaminated blood products and needles and instruments were considered as major sources for transmission of HCV [13]. Also, inadequate application and or breakdown of infection control policies in HD units (contamination of dialysis machines and, improper decontamination and sterilization, inadequately trained staff and unawareness of the value of hand washing) increase the transmission of HCV [14].

Several studies have reported nosocomial patient to patient transmission of HCV infection among HD patients [15, 16].

This study evaluated the incidence of seroconversion of HCV among HD patients in Benha university hospital HD unit, and the effect of isolation of HCV positive patients on HCV seroconversion.

Isolation included dedicated HD machines, personnel, area and other barrier precautions (aprons, gowns, or gloves), used by healthcare professionals take care of HCV positive

patients.

As regard general Characteristics of this study, it was found that the mean age of starting HD was 54.41 years. This was in agreement with some Egyptian studies [17, 18, and 19]. The increasing mean age of HD starting reflects the improvement of health care; but, we are still away from developed countries as the mean age in the United State was 61.1 years [20] and the median age in the United Kingdom was 65.9 years [21], but we are better than other developing countries as in Sudan (45.78 years) [22].

According to the gender, more than half of patients was male (56.7). This was in accordance with previous Egyptian studies [18, 19] which found that male constituted 53.7% and 61.0% respectively. Also in United States the male patients were more than female patients on HD as males represented about 55.0 % to 61.0 % according to different areas in United States [23].

In our study patients used only 2 types of vascular access, arterio-venous fistula (AVF) and temporary catheter. According to National Kidney Foundation, AVF is the best choice for HD, followed by arteio- venous graft and lastly catheter [24]. In this study there were 63 % of patients uses AVF and 27.0% uses catheters.

In the present study, the prevalence of HCV among HD patients was 48.9 % which was higher than Al Gharbiyah governorate, Egypt (35.0%) [19], Kafer El-Shakh governorate, Egypt (39.7%) [25] and much higher than in developed countries (3.6 to 20.0%) [8] and Morocco (32.0%) [26]. High prevalence of HCV among HD patients in Egypt is coinciding with that Egypt has the largest epidemic for

hepatitis C virus Worldwide [27].

As a comparison between HCV positive and HCV negative patients, we found that the mean age of HCV positive patients (53.07 years) was lower than HCV negative patients (55.7 years). The same result was reported in Libya [28]. But in Saudi Arabia they found that HCV is more common among the elderly as the immunity decreased with aging [29].

In our study the number of males was more than females in both HCV positive patients (56.8 %) and HCV negative patients (56.6 %). This was in agreement with Ayman, 2007 who added that Most of the studies did not find any effect of the sex on the incidence or prevalence of HCV infection [30].

The Dialysis Outcome and Practice Pattern Study (DOPPS) showed that high HCV seroconversion was associated with a longer duration on dialysis, and seroconversion was associated with an increase in the HCV prevalence, but not with the isolation of HCV-infected patients [31].

In our study there were 46/90 HCV negative patients at the start and at the end of the study after two years 9/46 patients had been seroconverted to be HCV positive with seroconversion rate 15.2 % in first year and 5.1% in second year.

As regard comparison between seroconverted HCV patients (group I) and none seroconverted HCV patients (group II), we found that mean age of group I was insignificantly lower than mean age of group II. This wasn't in agreement with another study done in Egypt as they found the mean age in seroconverted HCV patient was higher than non-seroconverted patient [32]. This can be explained as in our study that the mean age of HCV positive patients was lower than HCV negative patients.

In this study males were more than females in seroconverted HCV patients. This was in accordance with another study in Egypt in which males were more in seroconverted and females were more in non-seroconverted [18].

According to duration of HD in months, group I had significantly longer duration than group 2 ( $p < 0.001$ ). And this was in agreement with Soliman et al., who added that the duration on regular HD was found to be a significant predictor for HCV seroconversion in HD patients [32]. Also in another study done in Menoufia governorate, Egypt, they found that long duration of HD led to more exposure in HD units, with increased risk of HCV nosocomial infection [18].

The history of blood transfusion and total number of units (more than 3 units) were significant risk factors for HCV seroconversion in this study. Blood transfusion was still a significant relative risk for HCV seroconversion in HD patients [32] even after the introduction of nucleic acid amplification testing for the screening of blood donors which has markedly reduced the risk of HCV transmission through blood product transfusion [33]. But another study reported that blood transfusion had no significance as a risk factor in HCV seroconversion [18].

In this study the surgical history was more insignificantly in group 1 than group 2. This also was reported in another study as it found that surgical history had no significance as a risk factor in HCV seroconversion [18].

As regard family history, positive family history was more common in the seroconverted HCV patients; but this was not significant. This wasn't in agreement with another study that found that positive family history of HCV infection was significant risk factor for HCV seroconversion in HD patients [18].

HCV seroconversion rate (15.2 %) in first year (no isolation) was compared with a study done at Al Gharbiyah governorate, Egypt where HCV seroconversion rate was 11% [19] but the prevalence of HCV-Ab was 35% in all HD patients in comparison with our study which was 48.9 %.

Improving Global Outcome (KDIGO guidelines) didn't recommend the isolation policy for HCV-infected patients and did not even recommend the use of dedicated machines for them [34]. But in 2 studies in Saudi Arabia and Spain, they concluded that complete isolation of HCV-negative and HCV-positive patients and machines with strict adherence to infection control policies and procedures, can even eliminate nosocomial transmission and obtain reduction in prevalence and seroconversion of HCV [35, 36].

Finally, HCV seroconversion in first year (before isolation) was 15.2 % (7 patients from 46 HCV negative patients) and decreased to 5.1 % (2 patients from 39 HCV negative patients), and this decrease was statistically significant ( $p < 0.02$ ). This was agreed by Egyptian study compare between centers uses isolation of HCV positive patients and others don't, and found that seroconversion seen only in units not use isolation [17]. Other studied in Egypt in three different governorates found that incidence of HCV seroconversion is significantly lower in the group of patients within units implementing isolation programs of the HCV infected patients [32]. Also studies done in Saudi Arabia and Spain supported our finding and added that, strict isolation is mandatory in HD units [35, 36]. But in another study done in Peru, they did not found differences in terms of the number of participants developing HCV infection when comparing the use of dedicated HD machines for HCV infected patients with the use of non-dedicated machines [11].

## 6. Conclusion

In areas with high prevalence of HCV like Egypt, the prevalence of HCV in HD patients is high. So application of infection control policies and procedures which including isolation of HCV positive patients helps in decreasing HCV seroconversion.

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## References

- [1] Perico N, Cattaneo D, Bikbov B and Remuzzi G: Hepatitis C infection and chronic renal diseases. *Clin J Am Soc Nephrol.* 2009;4 (1): 207-20.
- [2] Bello AK, Nwankwo E and El Nahas AM: Prevention of chronic kidney disease: a global challenge. *Kidney Int* 2005; 98: 11-17.

- [3] Paez Jimenez A, Mohamed MK, SharafEldin N, AbouSeif H, El Aidi S, Sultan Y, Elsaid N, Rekacewicz C, El-Hoseiny M, El-Daly M, Abdel-Hamid M and Fontanet A: Injection drug use is a risk factor for HCV infection in urban Egypt. *PLoS One*. 2009; 4: e7193.
- [4] Hahn JA: Sex, drugs and hepatitis C Virus. *J Infect Dis*. 2007 Jun 1; 195 (11): 1556-9.
- [5] Galperim B, Mattos AA, Stein AT, Schneider NC, Buriol A, Fonseca A, Lunge V and Ikuta N: Hepatitis C in hemodialysis: the contribution of injection drug use. *The Brazilian Journal of Infectious Diseases*. 2010; 14 (4): 422-6.
- [6] Khan S, Attaullah S, Ali I, Ayaz S, Naseemullah, Khan SN, Siraj S and Khan J: Rising burden of Hepatitis C Virus in hemodialysis patients. *J Virol*. 2011; 8: 438.
- [7] Fabrizi F, Messa P. and Martin P.: Transmission of hepatitis C virus infection in hemodialysis: current concepts. *Int J Artif Organs*. 2008; 31 (12): 1004-16.
- [8] Afifi A.: The Egyptian Renal Registry. The 9th annual report for the year 2008 Published on 29th Annual congress of nephrology of Egyptian Society of Nephrology and Transplantation ESNT Hurghada Egypt 2009.
- [9] Egyptian Renal Registry: 2008 report, ESNT congress, Hurghada Egypt, February, 2009.
- [10] Jadoul M, Poinet JL, Geddes C, Locatelli F, Medin C, Krajewska M, Barril G, Scheuermann E, Sonkodi S and Goubau P; HCV Collaborative Group: The changing epidemiology of hepatitis C virus (HCV) infection in hemodialysis, European multicentre study. *Nephrol Dial Transplant*. 2004; 19 (4): 904-9.
- [11] Bravo Zuñiga JI, LozaMunárriz C and López-Alcalde J: Isolation as a strategy for controlling the transmission of hepatitis C virus (HCV) infection in haemodialysis units. *Cochrane Database of Systematic Reviews* 2016, 8. Art. No.: CD006420.
- [12] Delarocque-Astagneau E, Baffoy N, Thiers V, Simon N, de Valk H, Laperche S, Couroucé AM, Astagneau P, Buisson C, Desenclos JC.: Outbreak of hepatitis C virus infection in a hemodialysis unit: potential transmission by the hemodialysis machine? *Infect Control HospEpidemiol* 2002; 23 (6): 328-34.
- [13] Al-Jiffri AM, Fadeg RB, Ghabrah TM and Ibrahim A.: Hepatitis C virus infection among patients on hemodialysis in Jeddah: A single center experience. *Saudi J Kidney Dis Transplant*. 2003; 14 (1): 84-9.
- [14] Goldberg D and Anderson E.: Hepatitis C: Who is at risk and how do we identify them? *J Viral Hepatitis*. 2004; 11 (1): 12-8.
- [15] Iwasaki Y, Esumi M, Hosokawa N, Yanai M, and Kawano K: Occasional infection of hepatitis C virus occurring in hemodialysis units identified by serial monitoring of the virus infection, *Journal of Hospital Infection*, 2000; 45 (1): 54-61.
- [16] Schneeberger PM, Keur I, Van Loon AM, Mortier D, De Coul KO, Van Haperen AV, Sanna R, Van Der Heijden TG, Van Den Hoven H, Van Hamersvelt HW, Quint W and Van Doorn LJ: The prevalence and incidence of hepatitis C virus infections among dialysis patients in the Netherlands: a nationwide prospective study, *Journal of Infectious Diseases*, 2000; 182 (5): 1291-9.
- [17] Nasser ME, Younes KM, Sany DH, Youssef SS, Mahmoud M and El-Sayed BS: HCV Seroconversion in two Egyptian Hemodialysis Units: Role of Detection Method and Patients Isolation, *Macedonian Journal of Medical Sciences*, 2014; 2 (1): 124-127.
- [18] Zahran AM: Prevalence of seroconversion of hepatitis C virus among hemodialysis patients in Menoufia Governorate, Egypt. *Arab J Nephrol Transplant* 2014, 7 (2): 133-5.
- [19] Khodir SA, Alghateb M, Okasha KM and Shalaby SS: Prevalence of HCV Infections Among Hemodialysis Patients in Al Gharbiyah Governorate, Egypt, *Arab J Nephrol Transplant*. 2012; 5 (3): 145-7.
- [20] Collins AJ, Foley RN, Chavers B, Gilbertson D, Herzog C, Johansen K, Kasiske B, Kutner N, Liu J, St Peter W, Guo H, Gustafson S, Heubner B, Lamb K, Li S, Li S, Peng Y, Qiu Y, Roberts T, Skeans M, Snyder J, Solid C, Thompson B, Wang C, Weinhandl E, Zuan D, Arko C, Chen SC, Daniels F, Ebben J, Frazier E, Hanzlik C, Johnson R, Sheets D, Wang X, Forrest B, Constantini E, Everson S, Eggers P, Agodoa L. United States Renal Data System, USRDS 2011. Annual Data Report: Atlas of chronic kidney disease & end-stage renal disease in the United States. *AJKD* 2012; 59 (1): A7.
- [21] Stel VS, van de Luijngaarden MW, Wanner C and Jager KJ; on Behalf of the European Renal Registry Investigators. The 2008 ERA-EDTA registry annual report-a précis. *NDT Plus* 2011; 4 (1): 1-13.
- [22] Elsharif ME and Elsharif EG: Causes of End-Stage Renal Disease in Sudan: A Single-Center Experience. *Saudi J Kidney Dis Transpl* 2011; 22 (2): 373-6.
- [23] United States Renal Data System public health surveillance of chronic kidney disease and end-stage renal disease *Kidney Int Suppl*. 2015; 5 (1): 2-7.
- [24] National Kidney Foundation, 2015: [www.kidney.org](http://www.kidney.org)
- [25] Ahmed HA, Yassinea YS, Tawafea AR and Ebazaway MM. Epidemiological study of patients on regular haemodialysis at the Kafer El-Shakh Governorate, Egypt. *Menoufia Med J* 2015; 28 (2): 267-71.
- [26] Abdelaali B, Omar M, Taoufik D, Samir A, Saad M and Benyahia M: Hepatitis C Viral Prevalence and Seroconversion in Moroccan Hemodialysis Units: Eight Year Follow Up. *J Med Diagn Meth* 2013; 2 (5): 2-5.
- [27] Frank C, Mohamed MK, Strickland GT, Lavanchy D, Arthur RR, Magder LS, El Khoby T, Abdel-Wahab Y, Aly Ohn ES, Anwar W and Sallam I: The role of parenteral antischistosomal therapy in the spread of hepatitis C virus in Egypt. *Lancet*. 2000; 355 (9207): 887-91.
- [28] Alashek WA, McIntyre CW and Taal MW: Hepatitis B and C infection in haemodialysis in Libya: prevalence, incidence and risk factors. *BMC Infectious Diseases* 2012, 12:265.
- [29] Saxena AK and Panhotra BR: The vulnerability of middle aged and elderly patients to hepatitis C virus infection in a high-prevalence hospital-based hemodialysis setting. *J Am Geriatr Soc*. 2004; 52 (2): 242-6.
- [30] Ayman KARKAR: Hepatitis C in dialysis units: The Saudi Experience. *Hemodialysis International* 2007; 11 (3): 354-67.
- [31] Fissell RB, Bragg-Gresham JL, Woods JD, Jadoul M, Gillespie B, Hedderwick SA, Rayner HC, Greenwood RN, Akiba T and Young EW: Patterns of hepatitis C prevalence and seroconversion in hemodialysis units from three continents: the DOPPS. *Kidney Int* 2004; 65 (6): 2335-42.

- [32] Soliman AR, Abdelaziz MM and El lawindi MI: Evaluation of an Isolation Program of Hepatitis C Virus Infected Hemodialysis Patients in Some Hemodialysis Centers in Egypt. *ISRN Nephrology* 2013; Article ID 395467: 1-5.
- [33] O'Brien SF, Yi QL, Fan W, Scalia V, Kleinman SH and Vamvakas EC: Current incidence and estimated residual risk of transfusion-transmitted infections in donations made to Canadian Blood Services," *Transfusion* 2007; 47 (2): 316–25.
- [34] Kidney Disease: Improving Global Outcomes (KDIGO). KDIGO clinical practice guidelines for the prevention, diagnosis, evaluation, and treatment of hepatitis C in chronic kidney disease. *Kidney Int* 2008; 73 (109): 1–99.
- [35] Saxena AK, Panhotra BR, Sundaram DS, Naguib M, Venkateshappa CK, Uzzaman W and MulhimKA: Impact of dedicated space, dialysis equipment, and nursing staff on the transmission of hepatitis C virus in a hemodialysis unit of the Middle East. *Am J Infect Control*. 2003; 31 (1): 26–33.
- [36] Gallego E, López A, Pérez J, Llamas F, Lorenzo I, López E, Illescas ML, Andrés E, Olivas E, Gómez-Roldan C.: Effect of isolation measures on the incidence and prevalence of hepatitis C virus infection in hemodialysis. *Nephron ClinPract*. 2006; 104 (1): 1-6.