

Irish Compensation Scheme for Hepatitis C and / or HIV: Non Compensation Elements

Brian O Mahony

2022

Non Compensation Provisions


AN HILLE AMBONNE C'ESTER L'ADJAL LE DEPTERAN C
EASU 2006
REPARER C'COMPENSATION TRIBUNAL
(AMBUDENE) BELL 2006
Her e nithash of Pail Tinnan
As passed by Pail Tinnan
ARRANGEMENT OF SECTIONS

Section
1. Amendment of section 1 of Hospital C Compensation Tribunal Act 1997 (interpretation).
2. Amendment of section 4 of Hospital C Compensation Tribunal Act 1997 (interpretation).
3. Amendment of section 7 of Hospital C Compensation Tribunal Act 1997 (interpretation).
4. Insertion of new section 1A (appeals against relevant decisions of relevant authorities) in the Hospital C Compensation Tribunal Act 1997.
5. Amendment of section 12 of Hospital C Compensation Tribunal Act 1997 (interpretation).
6. Amendment of section 1 of Hospital C Compensation Tribunal Act 1997 (interpretation).
7. That all, subject to the provisions of this Act, shall be deemed to be the provisions of this Act.

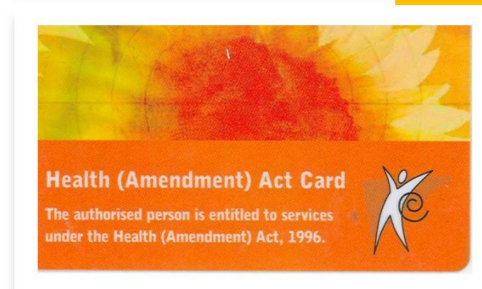
[No. 26 of 2006]

- 1996: Health Amendment act- provided list of free additional health care cover for PWH
- 2006-2007: Government subsidised Insurance
 - : Life
 - :Mortgage
 - : Travel
- 2007 : Tax concessions negotiated with revenue Dept for tribunal claimants – if > 50% of total income annually was from tribunal award- no tax due on income

Health Amendment Card 1996

Free and prioritised access to healthcare for all conditions for life

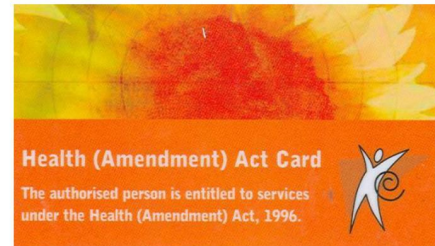
- General practitioner
- Prescription medications
- Priority access to appointments with new Specialists if required
- Counselling, Physiotherapy ,alternative medicine
- Glasses, hearing appliances
- Home workers
- Home nursing



Health Amendment Card 1996

Free and prioritised access to healthcare for all conditions for life

- Chiropody/Podiatry
- Complementary therapies- acupuncture, hydrotherapy, reflexology.... Referred by GP
- Priority access to Hospital bed and to Specialists
- Counselling- for PWH and immediate family members
- Dental services
- Travel vaccinations
- Aids and appliances
- Specific Liaison officers
to manage provisions and deal with queries





AN BILLE UM BINSÉ CÚDÚIH I ADÁIL LE HEPATITEAS C
(HEASÚ) 2008
HEPATITIS C COMPENSATION TRIBUNAL
(AMENDMENT) BILL 2008

Mae a rhwyddiwyd Ddiol Eirannu
As passed by Dáil Éireann

ARRANGEMENT OF SECTIONS

- Section
1. Amendment of section 1 of Hepatitis C Compensation Tribunal Act 1997 (interpretation).
 2. Amendment of section 4 of Hepatitis C Compensation Tribunal Act 1997 (claims before Tribunal).
 3. Amendment of section 7 of Hepatitis C Compensation Tribunal Act 1997 (regulations to give effect to Act).
 4. Insertion of new section 7A (appeals against relevant decisions of scheme administrator), 7B (special account for relevant insurance scheme), 7C (power of scheme administrator to specify forms) and 7D (confidentiality of matters relating to relevant claimants) into Hepatitis C Compensation Tribunal Act 1997.
 5. Amendment of section 8B of Hepatitis C Compensation Tribunal Act 1997 (special account).
 6. Amendment of section 2 of Health (Amendment) Act 1995 (provision of health services without charge to certain persons who have contracted hepatitis C).
 7. Short title, collective citation, construction and commencement.

[No. 35a of 2008]



STATUTORY INSTRUMENTS

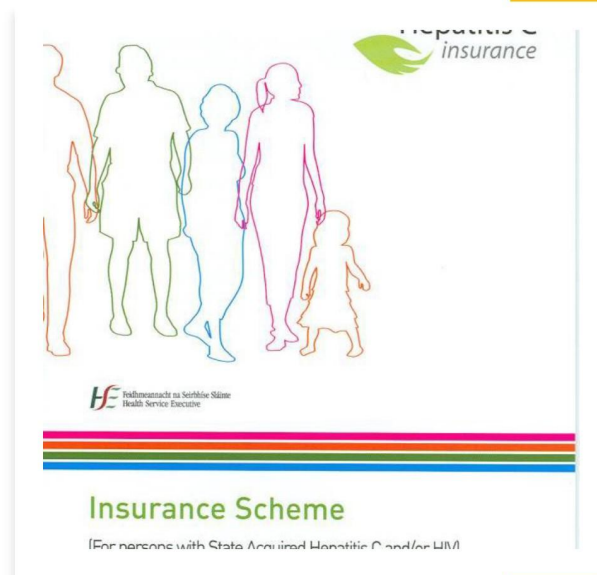
S.I. No. 364 of 2008

HEPATITIS C COMPENSATION TRIBUNAL (INSURANCE SCHEME
FOR RELEVANT CLAIMANTS) (AMENDMENT) REGULATIONS 2008

(Prt. A83489)

Other Elements: Insurance

- Life insurance: up to € 525,000
- Mortgage: up to € 475,000
- Travel: Annual
- PWH pays normal premium for healthy (non- smoker) person of their age- Government pays loading
- No non- insurable- all conditions covered
- Cover to age 65 (75 if taken out in first year of scheme)



Tax Concessions

- **Section 191 of Taxes Consolidation Act (TCA) 1997**- payments from the Hepatitis C Tribunal are exempt from income tax and CGT

- **Section 189 of Taxes Consolidation Act 1997**

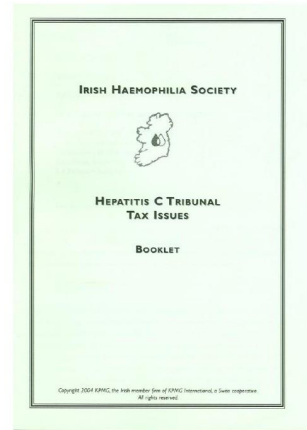
Deals with Income and gains from the investment of compensation received, which has been exempted under Section 191 TCA 1997

“ an individual who is permanently and totally incapacitated by reason of a mental or physical infirmity from maintaining himself or herself”

- Revenue agreed that this applied to all infected with HIV or Hepatitis C via blood or blood products

Other Elements: Tax Concessions

- Annual calculation
- Investment income and capital gains from compensation income exempt from tax if > 50% of income from Tribunal award
- Society provide tax advice via a professional and investment advice via meetings
- Can reclaim automatically deducted taxes- deposit interest/dividend taxes



Tax Concessions and Confidentiality

- HAA card holders can apply to have their tax affairs dealt with by a Confidential tax inspector
- He deals annually with the 50/50 applications and issues tax refunds
- Members fall into 1 of 3 categories:
 - : always exempt- Most or all of their income from award
 - : never or rarely exempt: >50% income always from non- award
 - : annual calculation- can be close to 50% each year depending on income and investment decisions
- Confidential service- much appreciated by members
- Society and our financial/Tax advisor have been able to make representations to the inspector and tax unit

Universal Social Charge (USC)

- Effectively an additional tax introduced in 2011 and maintained since.
- Levied at 2% of taxable income for first €10,036
- Levied at 4% of taxable income for next €5,980
- Levied at 7% of taxable income for remainder of income
- For HAA card holders with a taxable income < €60,000, USC capped at 4% of taxable income

Impact of Non Compensation Provisions

- Life insurance very valued
- Mortgage insurance rarely used
- Travel insurance universally used
- HAA card invaluable
- Role of liaison officers very much appreciated
- Tax concessions have also enabled many to retire early

National Hepatitis C Database

for infection acquired through blood and blood products



2012 Report



ORIGINAL ARTICLE

Progression of hepatitis C in the haemophilic population in Ireland, after 30 years of infection in the pre-DAA treatment era

NIAMH MURPHY,¹ BRIAN O'MAHONY,¹ PAULA FLANAGAN,² DECLAN NOONE,¹ BARRY WHITE,¹ COLM BERGIN,¹ SUZANNE NORRIS¹ and LILLA THORNTON¹ ON BEHALF OF THE NATIONAL HEPATITIS C DATABASE SCIENTIFIC AND TECHNICAL COMMITTEE¹

¹HSC Health Protection Surveillance Centre, Irish Haemophilia Society and ²St James's Hospital, Dublin, Ireland

Introduction: Prior to the introduction of viral inactivation of factor concentrates and screening of blood, 222 people with haemophilia became infected with hepatitis C (HCV) in Ireland. After 30 years we assess liver disease progression and mortality in this population after 30 years of infection. **Methods:** Demographic and clinical data were extracted from medical records to the hepatitis units and one infectious disease unit retrospectively in 2005, and on four subsequent occasions. Results: The participation rate was 75% (163/215). Eighty-three percent of patients, who had been tested for DNA in a WHOIS, developed chronic HCV infection. Thirty-four percent were co-infected with HIV. All-cause mortality, after approximately 30 years of infection with chronic HCV, was 44% in HIV-positive patients and 29% in HIV-negative patients. Liver-related mortality was 12.5% and did not vary significantly by HIV status. Thirty-seven percent of patients had developed advanced liver disease, including 20% with cirrhosis and 9% with hepatocellular carcinoma. In the pre-inactivation era direct acting antiviral use, 57% ($n = 60/106$) of patients were treated for HCV, 65% of whom achieved a sustained virological response. Incidentally treated patients had low adverse liver outcomes. **Conclusion:** After 30 years of infection, 40% of the patients who had evidence of chronic HCV had developed advanced liver disease, such as cirrhosis and HCC, or had died from liver-related causes. This proportion is high relative to similar immunosuppressed chronic hepatitis C patients and anti-HCV treatment uptake and response.

Keywords: cirrhosis, haemophilia, HCV, hepatitis C, hepatocellular carcinoma, HIV

Introduction

Blood clotting factor concentrates were used in Ireland from 1967 and represented a significant improvement in treatment for bleeding disorders over fresh plasma and whole blood. However, the lack of availability of a commercial blood test for the human immunodeficiency virus (HIV) before 1983, and the hepatitis C virus (HCV) before 1975, and the absence of viral inactivation, combined with the use of commercially fractionated factor concentrates from large pooled donations, meant that Irish haemophiliacs were exposed to, and infected with, HIV and HCV [1, 2].

Correspondence: Niamh Murphy, HSC Health Protection Surveillance Centre, 22-27 Maudsley Road, Dublin, Ireland. Tel: +353 1 436 9386; fax: +353 1 436 9386; e-mail: niamh.murphy@hpsc.ie

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least 222 people with haemophilia and other inherited bleeding disorders in Ireland became infected with HCV [3]. 104 became infected with HIV and it has been estimated [4] that most had severe haemophilia (5% normal clotting factor activity in blood). As there were approximately 250 people with severe haemophilia in Ireland at the time (personal communication: Brian O'Mahony, Irish Haemophilia Society), almost all were infected with HCV; more than 40% were infected with HIV and over a quarter were co-infected with HIV and HCV.

Overall, approximately 1700 people were infected with HCV through the administration of blood or blood products in Ireland [5]. Other cohorts included women infected through anti-D immunoglobulin and people infected through blood transfusions. All were referred to specialist hospital hepatology units and all are entitled to hospital and primary care services, free of charge, under the Health (Assessment) Act, 1996 [6(A)]. A national

CLINICAL PRACTICE

Hepatitis C and bleeding disorders in Europe

Laure Savits, Barbara Vaccaro, Declan Noone, Paul Cargnelli, Geoffrey Durrheim, Brian O'Mahony

In the 1980s and 1990s, thousands of people with bleeding disorders (PWBDs) across the world were infected with HIV and hepatitis C virus (HCV) through contaminated treatment products. The extent of the infection, as well as the needs of those still living with HCV, were never properly assessed. The purpose of our survey was to identify how many PWBDs were infected with HCV in Europe, as well as their health status and needs. HCV infection was defined as any person with a bleeding disorder who was exposed to the virus and seroconverted to become anti-HCV antibody positive.

The survey also looked at testing and treatment availability. Between December 2010 and March 2012, the survey was distributed to 46 national patient organisations in the European Haemophilia Consortium (EHC), who were encouraged to respond with the support of a local hepatologist. The data gathered had us estimate that some 12,000 people with bleeding



Image: HCV diagnosis with medication and syringe

disorders were infected with HCV in the 30 countries that responded. Although some countries have detailed records of patients with HCV, most – including some with national haemophilia registries – were unable to provide exact numbers of initial infections, HIV co-infection, survival and still rates. Responding countries reported varying degrees of monitoring for disease progression, as well as extremely divergent access to new direct-acting antivirals, with only eight countries prioritising PWBDs for treatment. With liver disease and hepatocellular carcinoma being among the main causes of death in an ageing bleeding disorder population, this survey identifies a clear gap in care. It is a troubling paradox that today in many European countries PWBDs, such as haemophiliacs, may live long and productive lives, due to much-improved access to factor replacement therapy, yet die prematurely of a curable disease such as hepatitis C. It has been demonstrated that HCV eradication in PWBDs can be achieved through national commitment, especially when the patient population is limited and HCV eradication could be achieved in the short term. The eradication of HCV in PWBDs in Europe is achievable and time has come.

Keywords: Bleeding disorders, Haemophilia, Hepatitis C, Monitoring, Treatment

Authors' disclosures of potential conflicts of interest and author contributions are found at the end of this article.
Address correspondence to: Niamh Murphy, HSC Health Protection Surveillance Centre, 22-27 Maudsley Road, Dublin, Ireland. Tel: +353 1 436 9386; fax: +353 1 436 9386; e-mail: niamh.murphy@hpsc.ie

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