

University of Groningen

Psychosocial and medical determinants of long-term patient outcomes

Prihodová, Lucia

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version

Publisher's PDF, also known as Version of record

Publication date:

2014

[Link to publication in University of Groningen/UMCG research database](#)

Citation for published version (APA):

Prihodová, L. (2014). *Psychosocial and medical determinants of long-term patient outcomes: A specific focus on patients after kidney transplantation and with haemophilia*. [Thesis fully internal (DIV), University of Groningen]. [S.n.].

Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: <https://www.rug.nl/library/open-access/self-archiving-pure/taverne-amendment>.

Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): <http://www.rug.nl/research/portal>. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.

Chapter 10

A SURVEY OF THE OUTCOME
OF PROPHYLAXIS, ON-DEMAND OR
COMBINED TREATMENT IN 20-35 YEAR OLD
MEN WITH SEVERE HAEMOPHILIA IN FOUR
EUROPEAN COUNTRIES

Declan Noone, Brian O'Mahony, Lucia Prihodova

Haemophilia 2011; 17, E842-E843. DOI: 10.1111/j.1365-2516.2011.02582.x

For children with severe haemophilia, prophylaxis is recognized as the optimum standard of care.^{4,2} It is also one of the principles of haemophilia care espoused by the European Association for Haemophilia and Allied Disorders (EAHAD).³ However, the continuation of prophylactic therapy into adulthood is being more closely scrutinized on the grounds of benefit and cost. This study was carried out to examine the long-term effects of prophylaxis and the continuing benefit of the therapy in adulthood. National Haemophilia patient organisations in Ireland, UK, France and Sweden were asked to participate by randomly selecting 20 severe haemophilia patients between 20 and 35 years from their database. Of the total of 80 questionnaires, 58 (72.5%) responses were received either by mail or by phone interview. Ireland provided 19, UK provided nine, France provided 10 and Sweden provided 20 responses. The four countries were chosen based on their access to treatment for patients with severe haemophilia. Swedish patients with severe haemophilia have been treated with prophylaxis since the late 1970s. Ireland, France and the UK introduced prophylaxis in the mid-1990s. The study examined the differences in medical outcomes and quality-of life in patients who had full access to primary prophylaxis, entirely on-demand or combinations of therapies. Information on age, country and employment status and responses to an EQ5D questionnaire were collected. Medical data was also collected on type and severity of haemophilia, treatment regime (prophylaxis vs. on-demand), bleeds per year, target joints, major bleeds (e.g. ilio-psoas or intracranial) and days missed from work/college per year. The primary analysis evaluated the differences between the four countries, and the secondary analysis examined the differences as a proportion of life spent on prophylaxis in comparison to on-demand therapy. The mean age was 27.5 ± 4.7 years. In the primary analysis of the individual countries, patients in Sweden have spent a significantly higher percentage of their life on prophylaxis ($p \leq 0.05$), showed the lowest number of bleeds/year, lowest presence of target joints ($p \leq 0.001$) and major bleeds ($p \leq 0.005$), lowest number of days missed from work/college, higher scores in Mobility (EQ5D) ($p \leq 0.05$) and highest utility value. The secondary analysis (Table 10.1) confirmed these findings. Patients always treated with prophylaxis reported significantly lower number of bleeds/year than patients treated entirely or primarily on-demand ($p \leq 0.05$), significantly lower presence of target joints ($p \leq 0.001$) and higher score in mobility ($p \leq 0.005$). Of the respondents who had received prophylaxis all of their lives, five reported major bleeds at some point, and five reported target joints without specifying that most bleeds occur in these joints. Three of each of these reported both target joints and major bleeds. No information on lifestyle or compliance was collected. Patients treated on-demand reported significantly higher number of days missed from work than all other groups ($p \leq 0.05$) and significantly lower score in EQ5D dimension Self-care ($p \leq 0.05$). The reported average number of bleeds/year for Sweden and France of 3.2 and 20.1, respectively, are broadly consistent with previous studies.^{4,5} The results in the primary analysis for France, UK and Ireland and in the secondary analysis for

the on-demand group are also similar to other published work.^{6,7} Hence, the results in this study support the view that prophylaxis started at a young age and continued into adulthood is an extremely effective treatment for patients with severe haemophilia. Four patients who were treated using prophylaxis switched to on-demand therapy and subsequently reverted to prophylaxis. Nine patients (22%), who had been treated primarily with on-demand therapy, have now changed to prophylaxis treatment. Respondents in both of these groups reported that this change took place due to increased bleeding episodes and/or joint problems that were developing when treated with on-demand therapy. When asked about days missed from work, there was a significant difference between countries. Swedish respondents reported a mean 0.5 days missed/year from work or college for reasons related to haemophilia. In Ireland and the UK, the days missed were 5 and 6.6 days respectively. With the French respondents, the mean time missed from work or college for reasons related to haemophilia was 15 days. This number was dramatically increased by two patients in the group not being able to attend work or college for 6 months following orthopaedic operations. For adults who have been treated with on-demand therapy, a requirement for orthopaedic surgery or joint replacement is not uncommon. This is further supported when the treatment regime was examined. Patients on prophylaxis missed a mean of 0.7 days per year and patients on-demand missed 19.2 days. The results from the EQ5D demonstrate the clear benefits of long-term prophylaxis over on-demand therapy. A number of studies on cost effectiveness 8–10 have reported the difference in utility values between prophylaxis and on-demand of 0.03 and 0.09. This study has shown that the benefit of prophylaxis continued into adulthood increases the utility value by a more significant 0.16–0.20. Overall, on-demand treatment results in a lower utility value in relation to quality-of-life for people with severe haemophilia. Prophylaxis started at an early age and continued into adulthood results in less bleeding, less damage to joints and less time missed at work. Prophylaxis increases mobility and the ability to do everyday activities and improves the health-related quality of life of people with severe haemophilia. It would be beneficial to extend this survey in the future to gather data on a larger number of people with Haemophilia from a larger number of countries and the authors plan to do this. It would be interesting to extend this survey to countries where distinctly different prophylaxis regimes are used and to countries that use low levels of FVIII per capita to assess what may well be, in effect, a baseline utility figure.

Table 10.1 Results by treatment regime

	Reported Mean Number of Bleeds per Year (n)	Presence of target joints (%)	Occurrence of Major Bleeds (%)	Mean Days missed per year (n)	Mean EQ-5D Utility Value
Group 1 (100% on Prophylaxis)	3.2	26.3	26.3	0.9	0.88
Group 2 (50-99% on Prophylaxis)	11.5	81.0	59.1	3.6	0.77
Group 3 (1-50% on Prophylaxis)	20.1	93.8	56.2	3.6	0.79
Group 4 (0% on Prophylaxis)	26.5	88.9	48.5	19.2	0.72

REFERENCES

1. Medical and Scientific Advisory Council (MASAC). MASAC Recommendation Concerning. Prophylaxis (Regular Administration of Clotting Factor Concentrate to Prevent Bleeding). The following recommendation was approved by the Medical and Scientific Advisory Council (MASAC) on April 22, 2006, and adopted by the NHF Board of Directors on June 3, 2006.
2. Keeling D, Tait C, Makris M. Guideline on the selection and use of therapeutic products to treat haemophilia and other hereditary bleeding disorders - A United Kingdom Haemophilia Center Doctors' Organisation (UKHCDO) guideline - Approved by the British Committee for Standards in Haematology. *Haemophilia* 2008;14(4):671-84.
3. Ludlam CA, Mannucci PM, Powderly WG. Addressing current challenges in haemophilia care: consensus recommendations of a European Interdisciplinary Working Group. *Haemophilia* 2005;11(5):433-7.
4. Van den Berg HM, Fischer K, Van der Bom JG. Comparing outcomes of different treatment regimens for severe haemophilia. *Haemophilia* 2003;9:27-31.
5. Petrini P, Chambost H, Nemes L. Towards the goal of prophylaxis: experience and treatment strategies from Sweden, France and Hungary. *Haemophilia* 2004;10:94-6.
6. Manco-Johnson M. Comparing prophylaxis with episodic treatment in haemophilia A: implications for clinical practice. *Haemophilia* 2007;13:4-9.
7. Carcao M, Chambost H, Ljung R. Devising a best practice approach to prophylaxis in boys with severe haemophilia: evaluation of current treatment strategies. *Haemophilia* 2010;16:4-9.
8. Lippert B, Berger K, Berntorp E, et al. Cost-effectiveness of haemophilia treatment: a cross-national assessment. *Blood Coagulation & Fibrinolysis* 2005;16(7):477-85.
9. Risebrough N, Oh P, Blanchette V, et al. Cost-utility analysis of Canadian tailored prophylaxis, primary prophylaxis and on-demand therapy in young children with severe haemophilia A. *Haemophilia* 2008 July;14(4):743-52.
10. Royal S, Schramm W, Berntorp E, et al. Quality-of-life differences between prophylactic and on-demand factor replacement therapy in European haemophilia patients. *Haemophilia* 2002;8(1):44-50.

