CORA Cork Open Research Archive
Cartlann Taighde Oscailte Chorcaí

Title	Study protocol: a randomised controlled trial on the clinical effects of levothyroxine treatment for subclinical hypothyroidism in people aged 80 years and over
Authors	Du Puy, R. S.;Postmus, I.;Stott, D. J.;Blum, M. R.;Poortvliet, R. K. E.;Den Elzen, W. P. J.;Peeters, R. P.;van Munster, B. C.;Wolffenbuttel, B. H. R.;Westendorp, R. G. J.;Kearney, Patricia M.;Ford, I.;Kean, S.;Messow, C. M.;Watt, T.;Jukema, J. W.;Dekkers, O. M.;Smit, J. W. A.;Rodondi, N.;Gussekloo, J.;Mooijaart, S. P.
Publication date	2018-09-19
Original Citation	Du Puy, R.S., Postmus, I., Stott, D.J., Blum, M.R., Poortvliet, R.K.E., Den Elzen, W.P.J., Peeters, R.P., van Munster, B.C., Wolffenbuttel, B.H.R., Westendorp, R.G.J. and Kearney, P.M., 2018. Study protocol: a randomised controlled trial on the clinical effects of levothyroxine treatment for subclinical hypothyroidism in people aged 80 years and over. BMC endocrine disorders, 18(1), 67 (14pp). DOI:10.1186/s12902-018-0285-8
Type of publication	Article (peer-reviewed)
Link to publisher's version	https://bmcendocrdisord.biomedcentral.com/articles/10.1186/s12902-018-0285-8 - 10.1186/s12902-018-0285-8
Rights	© The Author(s) 2018. This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated. - https://creativecommons.org/licenses/by/4.0/
Download date	2026-11-07 23:24:28
Item downloaded from	https://hdl.handle.net/10468/8669



University College Cork, Ireland
Coláiste na hOllscoile Corcaigh

The IEMO 80+ Thyroid Trial

- I've read the information letter. I was able to ask additional questions. My questions have been answered satisfactorily. I have had enough time to consider my participation.
- I understand that my participation is completely voluntary and that I am free to withdraw at any time, without giving any reason.
- I agree that my GP and/or treating specialist are informed about my study
- I agree that my GP and/or treating specialist are informed about the results of the blood tests
- I agree to using my data for the aims described in the information letter
- I agree with participating in the selection phase of this study.

Last name and initials: ……...

Address: ……

Postal code: ……

City: ……

Sex : **… male … female**

Date of birth: ……...

Date: …. …. …. **Phone:** ……

Signature: ……..