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SOURCE (OR PART OF THE FOLLOWING SOURCE):
Type Dissertation
Title Hypertrophic cardiomyopathy: towards an optimal strategy
Author I. Christiaans
Faculty Faculty of Medicine
Year 2010
Pages 189
ISBN 978-90-9025445-6

FULL BIBLIOGRAPHIC DETAILS:
http://dare.uva.nl/record/340634

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List of publications

1. Kolder ICRM*, Michels M*, **Christiaans I**, ten Cate FJ, Majoor-Krakauer D, Danser AHJ, Tanck MWT, Wilde AAM, Dooijes D, Bezzina CR. Septum thickness in Dutch MYBPC3 mutation carriers is influenced by the M235T-AGT polymorphism, increasing age, and male sex. Submitted.
2. **Christiaans I**, Birnie E, van Langen IM, Bonsel GJ, Wilde AAM. How to screen asymptomatic hypertrophic cardiomyopathy mutation carriers: a cost-effectiveness analysis. Submitted.
3. **Christiaans I**, Birnie E, Mannens MMAM, Michels M, Majoor-Krakauer D, Dooijes D, van Tintelen JP, van den Berg MP, Volders P, Arens Y, van den Wijngaard A, Atsma DE, Helderman-van den Enden ATJM, Houweling AC, van der Smagt JJ, Hauer RN, Marcelis CLM, Timmermans J, Bonsel GJ, van Langen IM, Wilde AAM. Risk factors for sudden cardiac death and follow-up in a large nationwide cohort of predictively tested hypertrophic cardiomyopathy mutation carriers. Submitted.
4. **Christiaans I**, Carrier L. Chapter 5 Hypertrophic Cardiomyopathy. In: Baars HF, van der Smagt JJ, Doevendans PAFM (Editors) Clinical Cardiogenetics. From basic science to clinical medicine. Springer 2010. In press.
5. **Christiaans I**, van Engelen K, van Langen IM, Birnie E, Bonsel GJ, Elliott PM, Wilde AAM. Risk stratification for sudden cardiac death in hypertrophic cardiomyopathy: systematic review of clinical risk markers. Europace 2010;12:313-21.
6. **Christiaans I**, Birnie E, van Langen IM, van Spaendonck-Zwarts KY, van Tintelen JP, van den Berg MP, Atsma DE, Helderman-van den Enden ATJM, Pinto YM, Hermans-van Ast JF, Bonsel GJ, Wilde AAM. The yield of risk stratification for sudden cardiac death in hypertrophic cardiomyopathy Myosin-Binding Protein C gene mutation carriers: focus on predictive screening. Eur Heart J 2010;31:842-8.
7. **Christiaans I**, Kok TM, van Langen IM, Birnie E, Bonsel GJ, Wilde AAM, Smets EMA. Obtaining insurance after DNA diagnostics: a survey among hypertrophic cardiomyopathy mutation carriers. Eur J Hum Genet 2010;18:251-3.
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10. **Christiaans I**, Lekanne Dit Deprez RH, van Langen IM, Wilde AA. Ventricular fibrillation in MYH7-related hypertrophic cardiomyopathy before onset of ventricular hypertrophy. Heart Rhythm 2009;6:1366-9.
11. **Christiaans I**, van Langen IM, Birnie E, Bonsel GJ, Wilde AAM, Smets EMA. Genetic counseling and cardiac care in predictively tested hypertrophic cardiomyopathie mutation carriers: The patients' perspective. Am J Med Genet Part A 2009;149A:1444–51.
12. Alders M*, Koopmann TT*, **Christiaans I**, Postema PG, Beekman L, Tanck MW, Zeppenfeld K, Loh P, Kock KT, Demolombe S, Mannens MM, Bezzina CR, Wilde AA. Haplotype-sharing analysis implicates chromosome 7q36 harboring DPP6 in familial idiopathic ventricular fibrillation. Am J Hum Genet 2009; 84:468-76.
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and psychological distress in hypertrophic cardiomyopathy mutation carriers: a cross-sectional cohort study. Am J Med Genet A 2009;149:602-12.
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