

PUBLICATION APPENDIX

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Full publication list based on archive at google scholar.

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- 1. Lyon, M., Andrews, S., Elsworth, B., Gaunt, T., Hemani, G. & Marcora, E. The variant call format provides efficient and robust storage of gwas summary statistics. *Genome Biology* (2021).
- 2. Baird, D., Liu, J., Zheng, J., Sieberts, S., Perumal, T., Elsworth, B. & others. Identifying drug targets for neurological and psychiatric disease via genetics and the brain transcriptome. *PLoS genetics* (2021).
- 3. Hemani, G., Thomas, A., Walker, J., Trickey, A., Nixon, E., Ellis, D. & others. Modelling pooling strategies for sars-cov-2 testing in a university setting. *medRxiv* (2021).
- 4. Wootton, R., Richmond, R., Stuijfzand, B., Lawn, R., Sallis, H. & others. Evidence for causal effects of lifetime smoking on risk for depression and schizophrenia: A mendelian randomisation study. *Psychological medicine* (2020).
- 5. Zhao, Q., Wang, J., Hemani, G., Bowden, J. & Small, D. Statistical inference in two-sample summary-data mendelian randomization using robust adjusted profile score. *Annals of Statistics* (2020).
- 6. Griffith, G., Morris, T., Tudball, M., Herbert, A., Mancano, G., Pike, L. & others. Collider bias undermines our understanding of covid-19 disease risk and severity. *Nature communications* (2020).
- 7. Brumpton, B., Sanderson, E., Heilbron, K., Hartwig, F., Harrison, S., Vie, G. & others. Avoiding dynastic, assortative mating, and population stratification biases in mendelian randomization through within-family analyses. *Nature communications* (2020).
- 8. Lawson, D., Davies, N., Haworth, S., Ashraf, B., Howe, L., Crawford, A. & others. Is population structure in the genetic biobank era irrelevant, a challenge, or an opportunity? *Human Genetics* (2020).
- 9. Zheng, J., Haberland, V., Baird, D., Walker, V., Haycock, P., Hurle, M. & others. Phenome-wide mendelian randomization mapping the influence of the plasma proteome on complex diseases. *Nature Genetics* (2020).
- 10. Morris, T., Davies, N., Hemani, G. & Smith, G. Population phenomena inflate genetic associations of complex social traits. *Science Advances* (2020).
- 11. Anderson, E., Howe, L., Wade, K., Ben-Shlomo, Y., Hill, W., Deary, I. & others. Education, intelligence and alzheimer's disease: Evidence from a multivariable two-sample mendelian randomization study. *International journal of epidemiology* (2020).
- 12. Richardson, T., Hemani, G., Gaunt, T., Relton, C. & Smith, G. A transcriptome-wide mendelian randomization study to uncover tissue-dependent regulatory mechanisms across the human phenome. *Nature communications* (2020).
- 13. Cho, Y., Haycock, P., Sanderson, E., Gaunt, T., Zheng, J., Morris, A. & others. Exploiting horizontal pleiotropy to search for causal pathways within a mendelian randomization framework. *Nature communications* (2020).
- 14. Zheng, J., Brion, M., Kemp, J., Warrington, N., Borges, M., Hemani, G. & others. The effect of plasma lipids and lipid⊠Lowering interventions on bone mineral density: A mendelian randomization study. *Journal of Bone and Mineral Research* (2020).

- 15. Neumeyer, S., Hemani, G. & Zeggini, E. Strengthening causal inference for complex disease using molecular quantitative trait loci. *Trends in molecular medicine* (2020).
- 16. Banos, D., McCartney, D., Patxot, M., Anchieri, L., Battram, T., Christiansen, C. & others. Bayesian reassessment of the epigenetic architecture of complex traits. *Nature communications* (2020).
- 17. Elsworth, B., Lyon, M., Alexander, T., Liu, Y. & Matthews, P. et al. The mrc ieu opengwas data infrastructure. bioRxiv (2020).
- 18. Christensen, H., Turner, K., Trickey, A., Booton, R., Hemani, G., Nixon, E. & others. COVID-19 transmission in a university setting: A rapid review of modelling studies. *medRxiv* (2020).
- 19. Reed, Z., Suderman, M., Relton, C., Davis, O. & Hemani, G. The association of dna methylation with body mass index: Distinguishing between predictors and biomarkers. *Clinical epigenetics* (2020).
- 20. Dooley, H., Lee, K., Freidin, M., Hemani, G., Roberts, A., Lochlainn, M. & others. ACE inhibitors, arbs and other anti-hypertensive drugs and novel covid-19: An association study from the covid symptom tracker app in 2,215,386 individuals. (2020).
- 21. Brooks-Pollock, E., Christensen, H., Trickey, A., Hemani, G., Nixon, E. & others. High covid-19 transmission potential associated with re-opening universities can be mitigated with layered interventions. *medRxiv* (2020).
- 22. Howe, L., Hemani, G., Lesseur, C., Gaborieau, V., Ludwig, K., Mangold, E. & others. Evaluating shared genetic influences on nonsyndromic cleft lip/palate and oropharyngeal neoplasms. *Genetic Epidemiology* (2020).
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- 24. Min, J., Hemani, G., Hannon, E., Dekkers, K., Castillo-Fernandez, J., Luijk, R. & others. Genomic and phenomic insights from an atlas of genetic effects on dna methylation. *medRxiv* (2020).
- 25. McCartney, D., Min, J., Richmond, R., Lu, A., Sobczyk, M., Davies, G. & others. Genome-wide association studies identify 137 loci for dna methylation biomarkers of ageing. *BioRxiv* (2020).
- 26. Russell, A., Ford, T., Gunnell, D., Heron, J. & Joinson, C. *et al.* Investigating evidence for a causal association between inflammation and self-harm: A multivariable mendelian randomisation study. *Brain, behavior, and immunity* (2020).
- 27. Lawson, D., Davies, N., Haworth, S., Ashraf, B., Howe, L., Crawford, A. & others. Is population structure in the genetic biobank era irrelevant, a challenge, or an opportunity?(Vol 138, pg 2321, 2019). *HUMAN GENETICS* (2020).
- 28. Nixon, E., Trickey, A., Christensen, H., Finn, A., Thomas, A., Relton, C. & others. Contacts and behaviours of university students during the covid-19 pandemic at the start of the 2020/21 academic year. *medRxiv* (2020).
- 29. Battram, T., Gaunt, T., Speed, D., Timpson, N. & Hemani, G. Exploring the variance in complex traits captured by dna methylation assays. *bioRxiv* (2020).
- 30. Liu, Y., Elsworth, B., Erola, P., Haberland, V. & Hemani, G. *et al.* EpiGraphDB: A database and data mining platform for health data science. *BioRxiv* (2020).
- 31. Sanderson, E., Richardson, T., Hemani, G. & Smith, G. The use of negative control outcomes in mendelian randomisation to detect potential population stratification or selection bias. *BioRxiv* (2020).
- 32. Bonilla, C., Bertoni, B., Min, J., Hemani, G. & others. Investigating dna methylation as a potential mediator between pigmentation genes, pigmentary traits and skin cancer. *Pigment cell & melanoma research* (2020).

- 33. Howe, L., Battram, T., Morris, T., Hartwig, F., Hemani, G., Davies, N. & others. Assortative mating and within-spouse pair comparisons. *BioRxiv* (2020).
- 34. Howard, D., Adams, M., Clarke, T., Hafferty, J., Gibson, J., Shirali, M. & others. Genome-wide meta-analysis of depression identifies 102 independent variants and highlights the importance of the prefrontal brain regions. *Nature neuroscience* (2019).
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- 38. Bandres⊠Ciga, S., Noyce, A., Hemani, G., Nicolas, A., Calvo, A., Mora, G. & others. Shared polygenic risk and causal inferences in amyotrophic lateral sclerosis. *Annals of neurology* (2019).
- 39. Carter, A., Gill, D., Davies, N., Taylor, A., Tillmann, T., Vaucher, J. & others. Understanding the consequences of education inequality on cardiovascular disease: Mendelian randomisation study. *bmj* (2019).
- 40. Morris, A., Le, T., Wu, H., Akbarov, A., Most, P., Hemani, G. & others. Trans-ethnic kidney function association study reveals putative causal genes and effects on kidney-specific disease aetiologies. *Nature communications* (2019).
- 41. Kraja, A., Liu, C., Fetterman, J., Graff, M. & Have, C. *et al.* Associations of mitochondrial and nuclear mitochondrial variants and genes with seven metabolic traits. *The American Journal of Human Genetics* (2019).
- 42. Taylor, D., Jackson, A., Narisu, N., Hemani, G., Erdos, M., Chines, P. & others. Integrative analysis of gene expression, dna methylation, physiological traits, and genetic variation in human skeletal muscle. *Proceedings of the National Academy of Sciences* (2019).
- 43. Noyce, A., Bandres⊠Ciga, S., Kim, J., Heilbron, K. & Kia, D. *et al.* The parkinson's disease mendelian randomization research portal. *Movement Disorders* (2019).
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- 45. Walker, V., Davies, N., Hemani, G., Zheng, J., Haycock, P., Gaunt, T. & others. Using the mr-base platform to investigate risk factors and drug targets for thousands of phenotypes. *Wellcome open research* (2019).
- 46. Moen, G., Hemani, G., Warrington, N. & Evans, D. Calculating power to detect maternal and offspring genetic effects in genetic association studies. *Behavior genetics* (2019).
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- 48. Howe, L., Lawson, D., Davies, N., Pourcain, B., Lewis, S., Smith, G. & others. Genetic evidence for assortative mating on alcohol consumption in the uk biobank. *Nature communications* (2019).
- 49. Howe, L., Richardson, T., Arathimos, R., Alvizi, L., Passos-Bueno, M. & others. Evidence for dna methylation mediating genetic liability to non-syndromic cleft lip/palate. *Epigenomics* (2019).

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- 67. Hemani, G., Bowden, J. & Smith, G. Evaluating the potential role of pleiotropy in mendelian randomization studies. *Human molecular genetics* (2018).
- 68. Xue, A., Wu, Y., Zhu, Z., Zhang, F. & Kemper, K. *et al.* Genome-wide association analyses identify 143 risk variants and putative regulatory mechanisms for type 2 diabetes. *Nature communications* (2018).
- 69. Võsa, U., Claringbould, A., Westra, H., Bonder, M., Deelen, P., Zeng, B. & others. Unraveling the polygenic architecture of complex traits using blood eQTL metaanalysis. *BioRxiv* (2018).
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