A Genome-Wide Association Meta-Analysis of Circulating Sex Hormone-Binding Globulin Reveals Multiple Loci Implicated in Sex Steroid Hormone Regulation

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Published in:
PLoS Genetics

DOI:
10.1371/journal.pgen.1002805

2012

Link to publication

Citation for published version (APA):

Total number of authors:
99

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A Genome-Wide Association Meta-Analysis of Circulating Sex Hormone-Binding Globulin Reveals Multiple Loci Implicated in Sex Steroid Hormone Regulation


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Genetic Influences of Sex Hormone–Binding Globulin


Editor: Greg Gibson, Georgia Institute of Technology, United States of America

Received: May 19, 2012; Accepted: July 19, 2012

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Funding: This meta-analysis is a collaborative effort involving data from many individual studies and many sources of funding. The details of funding sources for each study are detailed in Text S1 as well as below. Framingham Heart Study (FHS): The phenotype-genotype association analyses were funded through grants from the NHLBI R01AG032598 (UM Murabito, KL Lunetta), R01HL094755 (AD Coviello, RS Vasan, S Bandinelli), and R01AG31206 (RS Vasan, S Bandinelli), R01 AR/AG 41398 (DP Kiel). This research was conducted in part using data and resources from the Framingham Heart Study of the National Heart, Lung, and Blood Institute of the National Institutes of Health and Boston University School of Medicine. The analyses reflect intellectual input and resource development from the Framingham Heart Study investigators participating in the SNP Health Association Resource (SHARE) project. This work was partially supported by the National Heart, Lung, and Blood Institute’s Framingham Heart Study (Contract No. N01-HC-25195) and its contract with Affymetrix for genotyping services (Contract No. N02-HL-6-4278). A portion of this research was supported by the Swedish Research Council (K2010-54X-09894-19, 2006-3832, and K2010-52X-20299-05-3), the Swedish Foundation for Strategic Research, the ALF/LUA research grant in Gothenburg, the Lundberg Foundation, the Torsten and Ragnar Söderberg’s Foundation, Petrus and Augusta Hedlunds Foundation, the Vastra Gotaland Foundation, the Göteborg Medical Society, the Novo Nordisk Foundation, and the European Commission grant HEALTH-F2-2008-201865-GEFOS. We would like to acknowledge Maria Netherall at the genomics core facility at University of Gothenburg for statistical analyses. We would also like to thank Dr. Tobias A. Knoch, Luc V. de Zeeuw, Anis Abuseiris, and Rob de Graaf as well as their institutions the Erasmus Computing Grid, Rotterdam, The Netherlands, and especially the national German MedGRID and ServicesMedGRID part of the German D-Grid, both funded by the German Bundesministerium fuer Forschung und Technologie under grants #01AK 803 A+H and #01IG 70015 G for access to their grid resources. We would also like to thank Karol Estrada, Department of internal Medicine, Erasmus MC, Rotterdam, The Netherlands, for advice regarding

Abstract

Sex hormone-binding globulin (SHBG) is a glycoprotein responsible for the transport and biologic availability of sex steroid hormones, primarily testosterone and estradiol. SHBG has been associated with chronic diseases including type 2 diabetes (T2D) and with hormone-sensitive cancers such as breast and prostate cancer. We performed a genome-wide association study (GWAS) meta-analysis of 21,791 individuals from 10 epidemiologic studies and validated these findings in 7,046 individuals in an additional six studies. We identified twelve genomic regions (SNPs) associated with circulating SHBG concentrations. Loci near the identified SNPs included SHBG (rs12150660, 17p13.1, p = 1.8 x 10⁻¹⁶), PRMT6 (rs17496332, 1p13.3, p = 1.4 x 10⁻¹⁵), KCNR1 (rs700933, 2p23.3, p = 2.2 x 10⁻⁷), ZBTB10 (rs440837, 8q21.13, p = 3.4 x 10⁻⁰⁷), JMD1C (rs7910927, 10q21.3, p = 6.1 x 10⁻⁸), SLC18A1 (rs4149056, 12p12.1, p = 1.9 x 10⁻⁰⁹), NR2F2 (rs8032350, 15q26.2, p = 8.3 x 10⁻¹²), ZNF652 (rs2411984, 17q21.32, p = 3.5 x 10⁻¹⁴), TDFG3 (rs1573036, Qq22.3, p = 4.1 x 10⁻¹⁴), LHCGR (rs10454142, 2p16.3, p = 1.3 x 10⁻⁰⁶), BAIAP2LI (rs3771915, 7q21.3, p = 2.7 x 10⁻⁰⁸), and UGT2B15 (rs293428, 4q13.2, p = 5.5 x 10⁻⁰⁸). These genes encompass multiple biologic pathways, including hepatic function, lipid metabolism, carbohydrate metabolism and T2D, androgen and estrogen receptor function, epigenetic effects, and the biology of sex steroid hormone-responsive cancers including breast and prostate cancer. We found evidence of sex-differentiated genetic influences on SHBG. In a sex-specific GWAS, the loci 4q13.2-UGT2B15 was significant in men only (men p = 2.5 x 10⁻⁰⁶, women p = 0.66, heterogeneity p = 0.003). Additionally, three loci showed strong sex-differentiated effects: 17p13.1-SHBG and Qq22.3-TDFG3 were stronger in men, whereas 8q21.12-ZBTB10 was stronger in women. Conditional analyses identified additional signals at the SHBG gene that together almost double the proportion of variance explained at the SNP. Using sex-differentiated or conditional analyses explained ~15.6% and ~8.4% of the genetic variation of SHBG concentrations in men and women, respectively. The evidence for sex-differentiated effects and allelic heterogeneity highlight the importance of considering these features when estimating complex trait variance.

the grid resources. Health, Aging, and Body Composition (Health ABC) Study: This Health ABC Study was supported by NIA contracts N01-AG26125, N01-AG26127, and N01-AG26137. The Wake Forest University andatted in part by the U.S. National Institutes of Mental Health Research and Development (NIMH) and the U.S. National Institute on Aging (Contract: N1-AG-1-1 and N1-AG-1-2111); the InCHIANTI study followed up 2 and 3 studies (2004–2010) were financed by the U.S. National Institute on Aging (Contract: N1-AG-5-5002), supported in part by the Intramural research program of the National Institute on Aging. National Institutes of Health, Baltimore, Maryland. JRB Perry is a Sir Henry Wellcome Postdoctoral Research Fellow (092447/2-10/2). Cooperative Health Research in the Region of Aarhus (KORA): The KORA data was initiated and financed by the Helmholtz Centre Munich, German Research Center for Environmental Health, which is funded by the German Federal Ministry of Education and Research (BMBF) and by the State of Bavaria. Part of this work was financed by the German National Genome Research Network (NGFN-2 and NGFNplus: 01GS08769-02 through the STAMPED program (1RL0M083286-01), NIH/NIMH (SR01MH6730602), ENGAGE project and grant agreement HEALTH-F4-2007-201413, the Medical Research Council of the United Kingdom (G050559, G086075, PrevMeta[rsyN(SALVE)], and the Wellcome Trust (grant G0906224). We acknowledge the support of U.S. National Heart, Lung, and Blood Institute grant HL087679 through the STAMPED program; grants MMH83266, GM033275-14, and U54 RR020278 from the U.S. National Institutes of Health; grant DMS-0239427 from the National Science Foundation; the Medical Research Council of the UK, EU-BLCS, QLG1-CT-2000-01643, NHLBI grant RO1HL087679-02 through the STAMPED program (1RL0M083286-01), NIH/NIMH (SR01MH6730602), ENGAGE project and grant agreement HEALTH-F4-2007-201413, the authors would like to thank the Center of Excellence inCommon Disease Genetics of the Academy of Finland and Nordic Center of Excellence in Disease Genetics, the Sydantasaatio (Finnish Finnish Institute for Cancer Research), the Broad Genomics Center, D. M. Korn, M. Z. Jorde, J. Y. Hopper, L. M. L. M. and J. C. N. for their support for this study. Funding for the Rotterdam study were supported by the Netherlands Organisation of Scientific Research NWO Investments (nr. 100.001.018). This study is funded by the Research Institute for Diseases in the Elderly (014-93-015; RIDE), the Netherlands Genomics Initiative (NGI)/Netherlands Organisation for Scientific Research (NWO) project nr. 050-060-810, and funding from the European Commission (HEALTH-F2-2008-201865, GEFOS - Protocol E02H35.4 TREAT) or the Medical Research Council of the UK, and Erasmus University Rotterdam, National Institute of Public Health and Research Development and ZonMw; the Research Institute for Diseases in the Elderly (RIDE); the Ministry of Education, Culture, and Science; the Ministry for Health, Welfare, and Sports; the European Commission (DG XII); and the Municipality of Rotterdam. We thank Pascal Arp, Mila Jhamai, Dr. Michael Moorhouse, Marijn Verkerk, and Sander Veroest for their help in creating the GWAS database. The authors are grateful to the study participants, the staff from the Rotterdam Study and the participating general practitioners and pharmacists. We would like to thank Dr. A. A. Koek, Karol Estrada, Luc V. de Zeeuw, Anis Abouzereb, and Rob de Graaf as well as their institutions: the Erasmus Computing Grid, Rotterdam, The Netherlands, and especially the national German MedIGRID and Services@MedIGRID part of the German D-Grid, both funded by the German Bundesminister fuer Forschung und Technology under grants #01 AK 803 A-H and #01 IG 07015 Gfb access to their grid resources. Study of Health in Pomerania (SHIP): SHIP is part of the Community Medicine Research Network of the University of Greifswald, Germany, which is funded by the Federal Ministry of Education and Research (Grant No. 01CBO4403, 01Z02013, and 01Z02043), and the Ministry of Cultural Affairs, as well as the Social Ministry of the Federal State of Mecklenburg-West Pomerania. Genome-wide data have been supported by the Federal Ministry of Education and Research (grant no. 03ZIK012) and a joint grant from Siemens Healthcare, Erlangen, Germany, and the Federal State of Mecklenburg-West Pomerania. The University of Greifswald is a member of the “Center of Knowledge Integration” program of the Siemens AG. This work is also part of the research project Greifswald Approach to Individualized Medicine (GAMI_MED). The GANI_MED consortium is funded by the Federal Ministry of Education and Research (grant-031ZIK012). We thank the support from the Genotyping Facilities at the Wellcome Trust Sanger Institute for sample preparation, quality control, and genotyping led by Leena Peltonen and Panos Deloukas; Le Centre National de Genotypage, France, for all direct genotyping; Duke University, North Carolina, USA, for all direct genotyping; David Goldstein, for genetic analysis and expert advice; the Finnish Institute of Molecular Medicine, Finnish Genome Center, University of Helsinki, for Aarno Palotie. Genotyping was also performed by CDR as part of an NIH-NIH project grant. The Cardiovascular Risk in Young Finns Study (YFS): The Young Finns Study has been financially supported by the Academy of Finland: grants 134309 (Eye), 126925, 128422, 129378 (Salve), 117787 (Gen), and 41071 (Skidi); the Social Insurance Institution of Finland; Kuopio, Tampere, and Turku University Hospital Medical Funds (grant 9904D for 90035 for T3); Juho Vainio Foundation; Paavo Nurmi Foundation; Finnish Foundation of Cardiovascular Research and Finnish Cultural Foundation; Tampere University Foundation; and Emil Aaltonen Foundation (T. Lehtimäki). The expert technical assistance in the statistical analyses by Inra Lissen and Ville Aalto are gratefully acknowledged. Women's Health Initiative (WHI): Genotyping was performed at the Broad Institute (Cambridge, MA) through the NHLHR-funded Genomics and Randomized Clinical Network (U01 HG005152 or GARNET). The WHI program is funded by the National Heart, Lung, and Blood Institute, National Institutes of Health, U.S. Department of Health and Human Services through contracts N01WH22110, 24152, 32105–6, 32108–9, 32111–13, 32115, 32118–32119, 32122, 42107– 26, 42129–32, and 44221. The authors thank the WHI investigators and staff for their dedication, and the study participants for making the program possible. A listing of WHI investigators can be found at http://www.whiscience.org/publications/WHI_investigators_shortlist.pdf. Coronary Artery Risk Development in Young Adults (CARDIA) Women's Study: The CARDIA study is funded by contracts N01-HC-59059, N01-HC-48070, N01-HC-48049, N01-HC-48049, N01-HC-45134, N01-HC-50174, N01-HC-45205, and N01-HC-45204 and by the CARDIA Women's study by R01-HL065611 from the National Heart, Lung, and Blood Institute to the CARDIA Investigators. Genotyping of the CARDIA participants was supported by grants U01-HG-004729, U01-HG-00446, and U01-HG-00424 from the National Human Genome Research Institute. Statistical analyses were supported by grants U01-HG-004729 and R01-HL-084099 to M. Fornage. M. Wellons is supported by the Career Development Award 5-K23- HC-45205, and N01-HC-45204 and by the CARDIA Women's study by R01-HL065611 from the National Heart, Lung, and Blood Institute to the CARDIA Investigators. We acknowledge the support from the Dutch Ministry of Health, Welfare, and Sports; the European Commission (FP7-2007–2013), ENGAGE project and grant agreement HEALTH-F4-2007-201413. The authors thank the National Heart, Lung, and Blood Institute grant HL087679 through the STAMPED program; grants MMH83266, GM033275-14, and U54 RR020278 from the U.S. National Institutes of Health; grant DMS-0239427 from the National Science Foundation; the Medical Research Council of the UK, EU-BLCS, QLG1-CT-2000-01643, NHLBI grant RO1HL087679-02 through the STAMPED program (1RL0M083286-01), NIH/NIMH (SR01MH6730602), ENGAGE project and grant agreement HEALTH-F4-2007-201413, the Medical Research Council of the UK, EU-BLCS, QLG1-CT-2000-01643, NHLBI grant RO1HL087679-02 through the STAMPED program; grants MMH83266, GM033275-14, and U54 RR020278 from the U.S. National Institutes of Health; grant DMS-0239427 from the National Science Foundation; the Medical Research Council of the UK, EU-BLCS, QLG1-CT-2000-01643, NHLBI grant RO1HL087679-02 through the STAMPED program; grants MMH83266, GM033275-14, and U54 RR020278 from the U.S. National Institutes of Health; grant DMS-0239427 from the National Science Foundation; the Medical Research Council of the UK, EU-BLCS, QLG1-CT-2000-01643, NHLBI grant RO1HL087679-02 through the STAMPED program.
Author Summary

Sex hormone-binding globulin (SHBG) is the key protein responsible for binding and transporting the sex steroid hormones, testosterone and estradiol, in the circulatory system. SHBG regulates their bioavailability and therefore their effects in the body. SHBG has been linked to chronic diseases including type 2 diabetes and to hormone-sensitive cancers such as breast and prostate cancer. SHBG concentrations are approximately 50%heritable in family studies, suggesting SHBG concentrations are under significant genetic control; yet, little is known about the specific genes that influence SHBG. We conducted a large study of the association of SHBG concentrations with markers in the human genome in ~22,000 white men and women to determine which loci influence SHBG concentrations. Genes near the identified genomic markers in addition to the SHBG protein coding gene included PRMT6, GCKR, ZBTB10, JMJD1C, SLCO1B1, NR2F2, ZNF652, TGF3, LHCGR, BAIAF2L1, and UGT2B15. These genes represent a wide range of biologic pathways that may relate to SHBG function and sex steroid hormone biology, including liver function, lipid metabolism, carbohydrate metabolism and type 2 diabetes, and the development and progression of sex steroid hormone-responsive cancers.

Introduction

Sex hormone-binding globulin (SHBG) is a protein secreted mainly by the liver that binds to the sex steroids, testosterone, dihydrotestosterone, and estradiol, transports them in the circulation, and influences their action in target tissues by regulating their bioavailability. SHBG thereby influences the expression of sex hormone sensitive phenotypes including sexual characteristics and reproductive function in men and women [1]. In addition to regulating sex steroid hormone effects, SHBG may exert independent effects through its own receptor [2]. Variation in SHBG concentration has also been associated with various chronic diseases including cancers [3], polycystic ovary syndrome (PCOS) [4,5] and type 2 diabetes (T2D) [6,7]. Although SHBG is estimated to have a heritable component (~50%) [8], little is known about the genetic regulation of SHBG. Polymorphisms at the SHBG gene locus have been associated with SHBG concentrations [9,10], but much remains unknown about specific genetic variants that may determine circulating SHBG concentrations. Identifying genetic factors that influence SHBG may provide insights into the biology of sex steroid hormone regulation, metabolism and tissue effects that underlie their relationship with chronic diseases such as T2D as well as hormone-sensitive cancers such as breast and prostate cancer.

Results

We identified nine loci associated with SHBG concentrations at the genome-wide significance threshold of $p = 5 \times 10^{-8}$ (Table 1 and Figure 1) in a genome-wide association study (GWAS) meta-analysis of circulating SHBG concentrations in 21,791 men and women from 10 studies (Table S1). All nine lead SNPs at these loci had effects in the same direction (seven with $p<0.05$) in the validation dataset of 7,046 men and women from six additional studies (Table S2). The strongest association was within the SHBG locus (rs12150660, $p = 2 \times 10^{-106}$). Together, these nine lead SNPs explained 7.2% of the genetic variance (assuming 50% heritability) in SHBG concentrations.

We next performed a series of additional analyses to explain more of the phenotypic variance (Figure 2). First, we hypothesized that genetic effects may be different in men and women, as SHBG concentrations are >50% higher in females than males, and may be differentially regulated between sexes. In a sex stratified analysis, three of the nine loci showed evidence of sex-differentiated effects at $p<0.02$ when we would not expect any signals to have reached this level of significance by chance. The associations at the 1p13.1-SHBG and Xq22.3 loci were stronger in males whereas the association at the 8q21.3 locus was stronger in females. To investigate the apparent differential sex effect for the X chromosome further we ran a recessive regression model for the X chromosome SNP rs1573036 in women in the Framingham Heart Study and found no association with SHBG suggesting the sex-differentiated effect is not the result of a recessive inheritance pattern. Sex stratified GWAS identified one novel signal in men, which showed no association in women (4q13.2: men $p = 2.5 \times 10^{-10}$, women $p = 0.66$, heterogeneity $p = 0.003$).

A series of conditional analyses were performed to identify statistically independent signals. At the SHBG locus, three apparently independent additional signals separate from the main index SNP were observed, based on low ($p<0.05$) pairwise correlations in HapMap (rs6258 $p = 2.7 \times 10^{-46}$, rs1625895 $p = 1.2 \times 10^{-14}$ and rs3833894 $p = 2.5 \times 10^{-11}$). A series of iterative conditional analyses (Table 2) involving SNPs at the SHBG locus generated a final regression model including six statistically independent SHBG SNPs. Four of these SNPs (#1–4 Table 2) retained GWS when conditioned against the other five, and were nominally associated (SNP#5 $p = 0.0001$, SNP#6 $p = 0.01$). Re-running the GWAS meta-analysis adjusting for these six SNPs revealed evidence for three additional statistically independent (pairwise HapMap $r^2<0.01$) signals at the SHBG locus (SNP#7 $p = 1.5 \times 10^{-7}$, SNP#8 $p = 1.6 \times 10^{-5}$, SNP#9 $p = 9.9 \times 10^{-6}$) (Figure 3). There were also two additional trans signals located at 2p16.3 and 7q21.3 (Table 1). Although the 2p16.3 signal dropped below GWS when combined with follow-up samples ($p = 1 \times 10^{-7}$), the index SNP at 2p16.3 is ~300 kb away from a strong candidate gene, the luteinizing hormone receptor gene (LHCGR).

The majority of pair-wise correlations for the nine SHBG locus SNPs highlighted by our conditional analyses showed very low HapMap $r^2$ values. However, the pairwise D’ values are often high (Table S3) indicating that no or few recombination events have occurred between some SNPs, and that combinations of SNPs may be tagging un-typed variants on a common haplotype. To investigate this possibility, we performed more extensive analyses in a single study (NFBC1966, n = 4467). We used a denser set of SNPs imputed from the June 2011 version of the 1000 Genomes data and performed model selection analyses. Model selection identifies a set of SNPs that best explain phenotypic variation, while simultaneously penalizing each SNP included in this set, and therefore correlated SNPs tend to be excluded from the final model. These analyses consistently included at least seven SNPs in the model, although it is hard to estimate the false-negative rate of using the reduced sample size. While we are underpowered to accurately pinpoint the exact number of independent signals, these analyses support the results of the conditional analysis and suggest that multiple variants at the SHBG locus are independently associated with SHBG concentrations.

Data from an independent study, the InCHIANTI study, was used to calculate the proportion of genetic variance in SHBG concentrations explained when accounting for sex specific effects, the multiple signals of association at the SHBG locus, and the additional trans signals identified post conditional analysis. In men and women we explained ~15.6% and ~8.4% of the heritable
Table 1. SNPs representing loci associated with circulating SHBG concentrations.

<table>
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<th>Index SNP</th>
<th>Analysis</th>
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<th>Chr</th>
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<td>4q13.2</td>
<td>UGT2B15</td>
<td>4</td>
<td>69623671</td>
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<td>-0.019</td>
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</table>

All SNPs are on the + strand and positions are based on build 36. EAF = 'effect allele frequency'. Beta units are per-allele effect estimates in natural log transformed nmol/L. Sex column gives the sex with the largest per-allele beta estimate. Missing values for conditional SNPs as sex-specific conditional analysis was not performed.
component respectively. The SHBG locus accounted for ~10% and ~6.6% of the genetic variation in men and women respectively with the lead SNP in isolation accounting for ~7.8% and ~3.3% of the variation in men and women, respectively.

We identified genes near the associated SNPs and explored their biologic relevance to SHBG. The genes associated with identified SNPs included the SHBG locus (rs12150660, 17p13.1, \( p = 1.8 \times 10^{-106} \)), PRMT6 (rs17496332, 1p13.3, \( p = 1.4 \times 10^{-11} \)), GCKR (rs780093, 2p23.3, \( p = 2.2 \times 10^{-16} \)), ZBTB10 (rs40837, 8q21.13, \( p = 3.4 \times 10^{-49} \)), JMJD1C (rs7910927, 10q21.3, \( p = 6.1 \times 10^{-35} \)), SLCO1B1 (rs4149056, 12p12.1, \( p = 1.9 \times 10^{-46} \)), NR2F2 (rs8023580, 15q26.2, \( p = 8.3 \times 10^{-12} \)), ZNF652 (rs2411984, 17q21.32, \( p = 3.5 \times 10^{-49} \)), TDGF3 (rs1573036, Xq22.3, \( p = 4.1 \times 10^{-28} \)), LHCGR (rs10454142, 2p16.3, \( p = 1.3 \times 10^{-47} \)), BAIAP2L1 (rs779195, 7q21.3, \( p = 2.7 \times 10^{-46} \)), and UGT2B15 (rs293428, 4q13.2, \( p = 5.5 \times 10^{-86} \)) (Figure 1).

We used the online tool STRING (www.string-db.org) to perform pathway analyses to explore possible interactions between the SHBG gene and the proteins encoded by the 11 most plausible genes near the 11 SNPs listed above. There was an interaction noted between GCKR and JMJD1C which were associated with the lipoprotein fractions VLDL and HDL, respectively [11]. In an expanded analysis, we assessed protein interactions among SHBG and 67 genes within 500 kb of our 11 identified SNPs and uncovered additional protein interaction pathways. An interaction between two proteins encoded by GTF2A1L and STON1 was found; these proteins are co-expressed in testicular germ cells in the mouse [12]. An interaction between LHCGR and BR3 encoded proteins that are associated with the G-protein coupled receptor complex in the human luteinizing hormone receptor was also identified [13]. Finally, an interaction between LHCGR and LAPP (amylin) proteins which are components of a ligand/G-protein receptor/G-protein alpha subunit complex was found (database: www.reactome.com).

Targeted analysis of two strong candidate genes, hepatocyte nuclear factor-4\(\alpha\) (HNF4\(\alpha\)) and peroxisome-proliferating receptor \(\gamma\) (PPAR\(\gamma\)) did not identify any SNPs at HNF4\(\alpha\) but did identify one SNP, rs2920502, at PPAR\(\gamma\) that reached statistical significance \(p = 9.9 \times 10^{-5}\) and a second SNP at PPAR\(\gamma\), rs13081389, that reached nominal significance \(p = 0.01\).

Discussion

In total, we identified 12 genomic regions associated with circulating SHBG concentrations, including extensive allelic heterogeneity at the SHBG locus itself. Conditional meta-analyses carried out at the SHBG locus, identified nine genome-wide significant SNPs with low correlation \(r^2 < 0.01\) between them. Two of these signals (rs6258 [10] and rs6259) are missense variants and two are low frequency variants (MAF ~ 2%). Furthermore, rs12150660 is highly correlated \(r^2 > 0.95\) [10] with a pentanu-
The proportion of genetic variance in SHBG serum concentrations explained when accounting for sex specific effects, the multiple signals of association at the SHBG locus, and the

Table 2. Statistically independent signals at the SHBG gene locus.

<table>
<thead>
<tr>
<th>SNP #</th>
<th>Model</th>
<th>Conditioned On SNP #</th>
<th>SNP</th>
<th>Position</th>
<th>Effect Allele</th>
<th>Other Allele</th>
<th>EAF</th>
<th>Beta</th>
<th>SE</th>
<th>p-value</th>
<th>Discovery p-value</th>
<th>Discovery Beta</th>
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<td>1</td>
<td>Full model</td>
<td>2–6</td>
<td>rs12150660</td>
<td>7462640</td>
<td>t</td>
<td>g</td>
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<td>0.082</td>
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<td>1.19E-79</td>
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<td>7475403</td>
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<td>g</td>
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All SNPs are on the +strand and positions are based on build 36. EAF = ‘effect allele frequency’. Beta units are per-allele effect estimates in natural log transformed nmol/L. ‘Full model’ SNPs were all included in a single regression model, where the effect estimates for each SNP are adjusted for the effect of the others in the model.

‘Conditional’ SNPs are SNPs with low pair-wise LD (HapMap r²<0.01) that were identified after conditioning on the full model SNPs.

doi:10.1371/journal.pgen.1002805.t002
Additional trans signals identified post conditional analysis was ~15.6% in men and ~8.4% in women. The SHBG locus accounted for ~10% and ~6.6% of the genetic variance in men and women, respectively, with the lead SNP explaining most of the genetic variation at ~7.8% for men and ~3.3% for women. Thus additional signals at the SHBG locus identified through conditional analyses approximately doubled the variance of the trait explained. While we provide evidence for multiple variants associated with SHBG concentrations, further studies are needed to pinpoint the causal loci and functional variants. For the 11 regions outside the SHBG locus, most have biologically plausible related genes within 300 kb.

**Biology of Plausible Genes near Identified SNPs**

Several genes near the identified SNPs regulate sex steroid production and function. The *NR2F2* locus (15q26.2) encodes a nuclear receptor important in testicular Leydig cell function, the primary source of gonadal testosterone production [15], and has been linked to male infertility [16]. *NR2F2* has also been associated with estrogen receptor alpha (ERα) signaling and may influence hormone responsivity in breast cancer [17]. *PRMT6* (1p13.3) also encodes a nuclear receptor regulatory protein that mediates estrogen signaling as a co-activator of the estrogen receptor [18]. *LHCGR* (2p16.3) encodes the luteinizing hormone receptor which was associated with polycystic ovary syndrome (PCOS) in a recent GWAS [19,20]. PCOS is both a reproductive and metabolic disorder characterized by higher testosterone serum concentrations as well as an increased prevalence of obesity, insulin resistance, and T2D in women. Inappropriate secretion of luteinizing hormone leads to increased ovarian production of testosterone. Coincident lower SHBG concentrations contribute to increased bioavailable testosterone concentrations and the expression of both reproductive and metabolic phenotypes in PCOS [21,22,23].

The *SLCO1B1* locus encodes a liver-specific transporter of thyroid hormone as well as estrogens which impact liver production of SHBG [24]. *JMJD1C* (10q21.3), also known as *TRIP 8* (thyroid hormone receptor interactor protein 8 [25]), may impact SHBG concentrations via thyroid hormone effects on liver protein production. Thyroid hormone may alter SHBG production through effects on *INF4* which is known to regulate *SHBG* transcription [26,27].

Many of the genes identified are involved in carbohydrate and lipid metabolism and liver function. The *GCKR* locus (2p23.3) encodes a protein that regulates glucokinase activity and has been associated with T2D in several ethnic populations [28,29,30,31]. *GCKR* has been associated with metabolic and inflammatory traits including triglyceride concentrations and other lipid fractions [30,32], fasting plasma glucose [33,34], insulin concentrations, uric acid, c-reactive protein (CRP), and non-alcoholic fatty liver disease which are all characteristic of the metabolic syndrome and T2D [28,35,36,37,39,40,41,42]. The *SLCO1B1* locus (12p12.1) codes for a protein, hepatocyte protein anion-transporting polypeptide 1B1, involved in liver metabolism of both endogenous and exogenous compounds [43]. Consistent with *SLCO1B1*’s role in liver metabolism, the same SNP (rs4149056) has been associated with circulating bilirubin concentrations in previous GWAS [44].

*BAXIP2L1* (7q21.3) encodes a protein important in cytoskeleton organization [45] that has been associated with the inflammatory marker CRP in patients with arthritis [46]. *BAXIP2L1* is also known as *IRTKS* (insulin receptor tyrosine kinase substrate) which...
is involved in insulin receptor signaling [47] and may relate to insulin resistant states including obesity and T2D [40,49,50,51,52,53,54]. We conducted a targeted analysis of \( PPAR \), a gene that influences SHBG gene expression in the liver [1,55] and is associated with T2D [56,57]. Our analysis identified one significant SNP (rs2920502, \( p = 9.9 \times 10^{-5} \)) and a second nominally significant SNP (rs13081389, \( p = 0.01 \)) at \( PPAR \). Some of the identified genes involved in hepatic metabolism of lipids and carbohydrates may affect SHBG concentrations indirectly through effects on the SHBG transcription regulator \( HNF4 \alpha \) although \( HNF4 \alpha \) itself was not identified in this meta-analyses [27,58,59,60].

The \( UGT2B15 \) locus (4q13.2) was significantly associated with SHBG concentrations in men but not women in this meta-analysis. \( UGT2B15 \) belongs to a family of genes (the \( UGT2B \) gene family) that code for enzymes involved in the metabolism of sex hormones through glucuronidation which allows for excretion of sex steroids through the kidney and the gut via bile excretion [61,62]. Primary clearance mechanisms for sex steroids [63]. \( UGT2B15 \) is involved in the conjugation and inactivation of testosterone [64]. An association between rs293428 in the \( UGT2B15 \) locus and circulating SHBG concentrations in men is supported by a previous study demonstrating that a non-synonymous SNP (rs1992023; D85Y) is associated with serum SHBG concentrations in younger adult men [65]. \( UGT2B15 \) is thought to play a significant role in local tissue inactivation of androgens in androgen dependent prostate cancer [66,67]. The mechanism behind the influence of genetic variants in \( UGT2B15 \) on SHBG concentrations is unknown, but one may speculate that \( UGT2B15 \) affects the local androgenic environment in selected tissues, which in turn results in regulation of SHBG concentrations.

In addition to \( UGT2B15 \), three other genes near the identified SNPs are associated with carcinogenesis, particularly in the prostate and breast. \( ZBTB10 \) (8q21.13), has been linked to breast cancer [68]. In breast cancer cell lines \( ZBTB10 \) is suppressed by ROS-microRNA27a thereby enhancing ER\( \alpha \) expression and mediating estrogen effects [17]. The \( ZNF652 \) (17q21.32) locus codes for a DNA binding protein thought to act as a tumor suppressor gene in breast cancer [69,70,71] that is also co-expressed with the androgen receptor in prostate cancer [72]. \( TDGF3 \), teratocarcinoma derived growth factor 3, is located on chromosome 3p23-p21 and has been associated with testicular germ cell tumors [73].

**Strengths and Limitations**

This GWAS meta-analysis incorporated data from approximately 22,000 men and women from 16 epidemiologic cohorts. The overall size of the study yields power but the meta-analysis of data from different epidemiologic studies requires the inclusion of different laboratory methods. The different studies used a variety of assay methodologies to measure serum SHBG concentrations although the vast majority were immunoassays (Tables S1 and S2, Text S1) with similar methodologies. Variation introduced by the use of different SHBG assays would result in loss of statistical power and likely bias toward the null. Additionally, the majority of women were post-menopausal as ascertained by self-report in all studies (Table S1). SHBG concentrations, like testosterone, decline only slightly across the menopause [74] so adjustment for menopause status is not necessary. SHBG may also increase with ovulation and be slightly higher in the luteal versus the follicular phase of the menstrual cycle in premenopausal women, but most studies did not collect data on menstrual phase at the time of SHBG measurement so adjustment for menstrual phase was not possible [75]. Finally, individuals were not excluded based on health status, therefore some individuals with chronic conditions that may affect hepatic production of or clearance of proteins including SHBG such as liver disease, renal disease, or severe malnutrition, may have been included in this analysis.

**Conclusion**

SHBG synthesis in the liver is known to be affected directly or indirectly by estrogens, androgens and thyroid hormones and has been observed to be inversely associated with the higher insulin concentrations characteristic of insulin resistant states such as T2D [1,6]. In summary, the results of this GWAS reflect these influences. Three regions map to proteins related to hepatic function [12p12.1-SLC01B1 [76], 2p23.3-GRK [77] and 10q21.3-3MJDIC [77]]. In addition, 2p23.3-GRK and 7q21.3-BMAL2P1 [alias insulin receptor tyrosine kinase substrate (IRTKS)] are involved in susceptibility to T2D [48] and insulin signaling [47], respectively. Two signals also mapped to loci involved in thyroid hormone regulation (10q21.3-3MJDIC and 12p12.1-SLC01B1). One signal mapped to the receptor for luteinizing hormone 2p16.3-LHGR [29], the hormone that stimulates testosterone production. Five regions mapped to genes previously implicated in androgen and estrogen signaling [1p13.3-PRMT6 [18], 8q21.13-ZBTB10 [17], 12p12.1-SLC01B1 [76], 15q26.2-NR2F2 [70], 4q13.2-UGT2B15 [63]].

We have combined a conventional GWAS approach with detailed additional analyses, including sex stratification, conditional analysis and imputation from 1000 Genomes. Our results demonstrate that these approaches can lead to an appreciable gain in heritable variance explained. It does however highlight the complexity of elucidating individual variant causality through statistical approaches. In addition to the extensive allelic heterogeneity at the SHBG locus, our data identify loci with a role in sex steroid hormone metabolism, which may help elucidate the role of sex steroid hormones in disease, particularly T2D and hormone-sensitive cancers.

**Methods**

We performed a genome wide association study (GWAS) meta-analysis of 21,791 individuals (Table S1: 9,390 women, 12,401 men) from ten observational studies. Data from an additional six studies totaling 7,046 individuals (Table S2: 4,509 women; 2,537 men) were used for validation. The proportion of variance explained was estimated in an independent study (InCHIANTI, \( n = 1,129 \)). The individual study protocols were approved by their respective institution’s ethics committee/institutional review board and all participants provided informed consent prior to participation. Individuals known to be taking hormonal contraceptives or hormone replacement therapy at time of SHBG measurement were excluded from analysis. Age, sex and body mass index (BMI) were included as covariates. After applying standard quality control measures, imputed genotypes were available for approximately 2.5 M SNPs. See Figure 2 for an overview of the analytic plan and the Text S1 for further information for individual studies included in this meta-analysis.

**GWAS Conditional Meta-Analysis Steps**

**Conditional analysis #1**. The initial starting point for the conditional analysis was the four SHBG locus SNPs that all showed low Hapmap LD (\( r^2<0.05 \)) with each other: rs12150660 [lead SNP Table 1], rs6258 \( p = 2.7 \times 10^{-16} \), rs1625895 \( p = 1.2 \times 10^{-14} \) and rs3853894 \( p = 2.5 \times 10^{-11} \). Each cohort fitted a single
regression model, fitting SHBG concentrations against these four genome-wide significant SHBG locus SNPs (rs12150660, rs6258, rs1625895 and rs3853894), in addition to age, sex and BMI. After meta-analyzing the results from all cohorts, three of the SNPs retained genome-wide significance when regressed against each other, with the fourth SNP narrowly missing that threshold (rs3853894, p = 4.1\times 10^{-6}).

**Conditional GWAS #1 (Table 1, conditional analysis).**  We next performed a conditional GWAS meta-analysis, where each study included, as additional covariates to the original analysis plan, the ten genome-wide significant autosomal SNPs (the eight ‘Main’ signals from Table 1 and the two unique SHBG locus signals described above in addition to the lead SNP rs12150660: rs6258 and rs1625895). Three additional signals (independence based on HapMap r^2<0.05) at the SHBG locus reached genome-wide significance (rs1641537 p = 7.8\times 10^{-12}, rs6259 p = 1.5\times 10^{-12} and rs10432092 p = 3\times 10^{-5}), giving a total of six independent signals in this region. In addition, two novel signals reached genome-wide significance in the conditional analysis, at 7q21.3 (rs3779195 p = 1.1\times 10^{-8}) and 2p16.3 (rs10454142 p = 3\times 10^{-5}). After replication, only rs3779195 at the BAIAP2LI locus retained genome-wide significance.

**Conditional analysis #2 (Table 2, full model).**  Given the six signals observed at the SHBG locus (three through conditional analysis #1: rs12150660, rs6258, rs1625895, three through LD estimates from conditional GWAS #1: rs1641537, rs6259, rs10432092), we sought to confirm which of these six were truly independent by a second round of conditional analysis. All discovery and replication cohorts fitted a single regression model of the six SNPs (SNPs # 1–6, Table 2) against SHBG concentrations, using the same parameters and covariates as conditional analysis #1. Four of the six SNPs (#1–4: rs12150660, rs6258, rs1641537, and rs1625895) retained genome-wide significance when conditioned against each other, with two showing nominal evidence of association (SNP #5 rs6259, p = 0.0001; SNP #6 rs10432092, p = 0.01).

**Conditional GWAS #2 (Table 2, conditional model).**  Finally, we performed a second conditional GWAS analysis, adjusting for the six SHBG locus SNPs which had evidence of association from conditional analysis #2. All the discovery cohorts were used in this analysis, in addition to three replication cohorts (total sample size 24,354). This analysis revealed evidence for a further three independent signals at the SHBG locus (based on HapMap r^2<0.01), SNP #7 rs9901675 p = 1.5\times 10^{-7}, SNP #8 rs8077824 p = 4.6\times 10^{-5}, and SNP #9 rs9393218 p = 9.9\times 10^{-6}.

**Sensitivity Analysis—Allelic Heterogeneity at the SHBG Locus**

We performed a sensitivity analysis using samples from the 1966 Northern Finland Birth Cohort (NFBC1966) study to further investigate allelic heterogeneity at the SHBG locus (Text S1). The conditional meta-analysis showed evidence for up to nine signals at the SHBG locus, but it is possible that these signals could be explaining a much smaller number of causal variants in the region. Since 1000 Genomes imputation allows us to assess the genetic variation associated with a phenotype across a much denser set of markers, it increases our power to detect allelic heterogeneity within a region. Therefore, 1000 Genomes imputation was carried out on all the samples in the NFBC1966 study and forward selection was used to identify the set of SNPs that best explain the variation in the SHBG phenotype. 1000 Genomes imputation was carried out using IMPUTE2. The mean genotype probabilities for each SNP were calculated and used in the model selection step. Only SNPs 250 kb upstream and 250 kb downstream from the SHBG locus (7283453–7786700 bp) were used in the analysis. All SNPs with MAF <0.1% or an imputation quality score less than 0.4 were excluded from the analysis. In total, 1978 SHBG region SNPs measured or imputed in 4467 samples from the NFBC1966 study were used in the sensitivity analysis. Forward selection was implemented in R (version 2.13.0) using the stepAIC package to estimate the Akaike Information Criterion (AIC), an inclusion parameter. Given the high degree of correlation between the SNPs in this region, we increased the penalty k on the number of terms included in the model to 12 (where it is usually two), to minimize possible over fitting. The final model included seven SNPs, adjusted for sex and BMI.

**Pathway Analysis**

We examined potential interactions among the proteins encoded by the SHBG locus and the proteins encoded by the 11 genes (ZBT10, TDFG1, ZNF632, PRMT6, JMJDC1, GCKR, BAIAP2LI, LIHCGR, SLC01B1, UGT2B15, NR2F2) closest to the 11 identified SNPs using pathway analysis with Search Tool for the Retrieval of Interacting Genes/Proteins (STRING) Pathways Analysis (www.string-db.org). The interactions explored by STRING include direct (physical) and indirect (functional) associations. We then expanded the analysis to examine protein interactions among the SHBG gene and the proteins encoded by 67 genes within 500 kb of the 11 identified SNPs.

**Targeted Candidate Gene Analysis**

We conducted targeted analysis of two strong candidate genes, hepatocyte nuclear factor-4a (HNF4a) and peroxisome-proliferating receptor γ (PPARγ). Statistical significance thresholds were set correcting for the number of SNPs tested in each gene region (±100 kb).

**Supporting Information**

**Table S1** Characteristics of 21,791 individuals from 10 discovery cohorts included in the meta-analysis.

**Table S2** Characteristics of 8,175 individuals from the six cohorts included in the validation analysis (WHI, CARDIA, Prospect-EPIC, MrOs, NHS, YFS) and the independent cohort used to estimate the proportion of genetic variance explained by the identified SNPs (IntChianti).

**Table S3** Hapmap (release 22) linkage disequilibrium estimates for the nine SHBG gene locus single nucleotide polymorphisms.

**Text S1** Supplementary Methods with Specific Cohort Information.

**Acknowledgments**

We are grateful to all study participants and staff in our participating studies. Full study acknowledgements are available in Text S1.

**Author Contributions**

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