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## Meta-analysis identifies 13 new loci associated with waist-hip ratio and reveals sexual dimorphism in the genetic basis of fat distribution

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## Abstract

Waist-hip ratio (WHR) is a measure of body fat distribution and a predictor of metabolic consequences independent of overall adiposity. WHR is heritable, but few genetic variants influencing this trait have been identified. We conducted a meta-analysis of 32 genome-wide association studies for WHR adjusted for body mass index (comprising up to 77,167 participants), following up 16 loci in an additional 29 studies (comprising up to 113,636 subjects). We identified 13 new loci in or near *RSPO3*, *VEGFA*, *TBX15-WARS2*, *NFE2L3*, *GRB14*, *DNM3-PIGC*, *ITPR2-SSPN*, *LY86*, *HOXC13*, *ADAMTS9*, *ZNRF3-KREMEN1*, *NISCH-STAB1* and *CPEB4* ( $P = 1.9 \times 10^{-9}$  to  $P = 1.8 \times 10^{-40}$ ) and the known signal at *LYPLALI*. Seven of these loci exhibited marked sexual dimorphism, all with a stronger effect on WHR in women than men ( $P$  for sex difference =  $1.9 \times 10^{-3}$  to  $P = 1.2 \times 10^{-13}$ ). These findings provide evidence for multiple loci that modulate body fat distribution independent of overall adiposity and reveal strong gene-by-sex interactions.

Central obesity and body fat distribution, as measured by waist circumference and WHR, are associated with individual risk of type 2 diabetes (T2D)<sup>1,2</sup> and coronary heart disease<sup>3</sup> and with mortality from all causes<sup>4</sup>. These effects are independent of overall adiposity as measured by body mass index (BMI). WHR is of particular interest as a measure of body fat distribution because it integrates the adverse metabolic risk associated with increasing waist circumference with the more protective role of gluteal fat deposition with respect to diabetes, hypertension and dyslipidemia<sup>5,6</sup>.

There is abundant evidence that body fat distribution is influenced by genetic loci distinct from those regulating BMI and overall adiposity. First, even after accounting for BMI, individual variation in WHR is heritable<sup>7,8</sup>, with heritability estimates ranging from 22%–61%<sup>7-10</sup>. Second, the striking abnormalities of regional fat deposition associated with lipodystrophic syndromes demonstrate that genetic variation can have dramatic effects on the development and maintenance of specific fat depots<sup>11,12</sup>. Third, in a previous genome-wide association analysis, we identified a locus near *LYPLALI* strongly associated with WHR independent of any effects on BMI<sup>13</sup>, providing proof of principle for the genetic control of body fat distribution distinct from that of overall adiposity.

Within the Genetic Investigation of Anthropometric Traits (GIANT) consortium, we performed a large-scale meta-analysis of genome-wide association studies (GWAS) informative for WHR using adjustment for BMI to focus discovery toward genetic loci associated with body fat distribution rather than overall adiposity<sup>14-16</sup>.

## RESULTS

### Genome-wide significant association of WHR with 14 SNPs

We conducted a two-stage study among individuals of European descent (Supplementary Table 1 and Online Methods). In the discovery stage, up to 2,850,269 imputed and genotyped SNPs were examined in 32 GWAS comprising up to 77,167 participants informative for anthropometric measures of body fat distribution. We performed a fixed-effects meta-analysis of WHR, employing study-specific linear regression adjusted for BMI and age, stratified by gender, and using an additive genetic model. After genomic control adjustment per each individual study and in the meta-analysis, these analyses revealed a substantial excess of low  $P$  values (Fig. 1a,b).

We selected SNPs representing the top 16 independent (defined as being located >1 Mb apart) regions of association (discovery  $P < 1.4 \times 10^{-6}$ ; Table 1) and evaluated them in 29 additional, independent studies (comprising up to 113,636 individuals) using a mixture of *in silico* data and *de novo* genotyping. In these follow-up studies, 14 of the 16 SNPs analyzed showed strong directionally consistent evidence for replication ( $P < 1.0 \times 10^{-3}$ ) and ten SNPs reached genome-wide significance ( $P < 5.0 \times 10^{-8}$ ). Joint analysis of the discovery and follow-up results revealed genome-wide significant associations for 14 signals (with  $P$  values between  $1.9 \times 10^{-9}$  and  $1.8 \times 10^{-40}$ ; Table 1). Between-study heterogeneity was low ( $I^2 < 30\%$ ) for all but two signals (*GRB14* and *LYPLAL1*; Supplementary Note), and all 14 associations remained genome-wide significant in a random-effects meta-analysis (Supplementary Table 2).

One of these SNPs, rs4846567, is in linkage disequilibrium (LD) ( $r^2 = 0.64$ ,  $D' = 0.84$ ; HapMap European CEU population) with the previously reported WHR-associated variant near *LYPLAL1* (rs2605100)<sup>13</sup>. The remaining 13 loci were in or near genes not previously associated with WHR or other measures of adiposity: *RSPO3*, *VEGFA*, *TBX15-WARS2*, *NFE2L3*, *GRB14*, *DNM3-PIGC*, *ITPR2-SSPN*, *LY86*, *HOXC13*, *ADAMTS9*, *ZNRF3-KREMEN1*, *NISCH-STAB1* and *CPEB4* (Fig. 2). These 14 loci explain 1.03% of the variance in WHR (after adjustment for BMI, age and sex), with each locus contributing from 0.02% (*ZNRF3-KREMEN1*) to 0.14% (*RSPO3*) of the variance based on effect estimates in the follow-up stage.

### Sexual dimorphism at several of the WHR loci

Given the known sexual dimorphism of WHR and the evidence from variance decomposition studies that this reflects sex-specific genetic effects<sup>17</sup>, we performed sex-specific meta-analyses for the 14 WHR-associated SNPs. These analyses included up to 108,979 women (42,735 in the discovery stage and 66,244 in the follow up) and 82,483 men (34,601 in the discovery and 47,882 in the follow up). In a joint analysis of discovery and follow-up data, 12 of the 14 SNPs reached genome-wide significance in women, but only three SNPs reached genome-wide significance in men (Table 2). At all but one locus (*TBX15-WARS2*), effect-size estimates were numerically greater in women. At seven of the loci (those near *RSPO3*, *VEGFA*, *GRB14*, *LYPLAL1*, *HOXC13*, *ITPR2-SSPN* and *ADAMTS9*), there were marked differences in sex-specific  $\beta$  coefficients (with  $P$  values ranging from  $1.9 \times 10^{-3}$  to  $1.2 \times 10^{-13}$ ). All loci displayed consistent patterns of sex-specific differences in both the discovery and follow-up studies (Table 2). These 14 loci explain 1.34% of the variance in WHR (after adjustment for BMI and age) in women but only 0.46% of the variance in WHR in men.

### Association with other anthropometric measures

By focusing on WHR after adjustment for BMI, our goal was to detect effects on body fat distribution independent of those influencing overall adiposity. As expected, we found very little evidence that known BMI-associated variants were detected in our WHR analysis. Of the ten loci shown to be associated with BMI in previous GWAS<sup>14,15,18</sup>, only two showed nominally significant ( $P < 0.05$ ) associations for BMI-adjusted WHR in the discovery analysis (*FTO*, rs8050136,  $P = 0.03$ ,  $n = 77,074$ ; *TMEM18*, rs6548238,  $P = 3.0 \times 10^{-3}$ ,  $n = 77,016$ ).

We also tested the 14 WHR-associated SNPs for their effect on BMI using data from up to 242,530 participants available from the GIANT consortium (including most of the studies available for WHR association). Of the 14 WHR loci, four (near *TBX15-WARS2*, *CPEB4*, *LYPLAL1* and *GRB14*) also showed evidence of association with BMI ( $4.1 \times 10^{-3} \leq P \leq 3.2 \times 10^{-6}$ ), with the WHR-increasing allele associated with decreased BMI (Supplementary

Table 3). After adding an interaction term of SNP with BMI into the model, we observed that BMI modified the WHR association at the *LY86* locus ( $P$  for interaction =  $9.5 \times 10^{-5}$ ), with a larger WHR effect among obese individuals compared to non-obese individuals (Supplementary Note).

To determine whether the WHR-associated signals exert their effects primarily through an effect on waist or hip circumference, we performed meta-analyses for these specific phenotypes in the discovery and follow-up studies (Supplementary Tables 1 and 3). Overall, we observed stronger associations for hip circumference than for waist circumference. Effect-size estimates were numerically greater for hip circumference than for waist circumference at 11 of the 14 loci, and there were nominal associations ( $P < 0.05$ ) with hip circumference for 12 of the WHR-associated loci but there were only four associations with waist circumference. In both sexes, the WHR-associated loci displaying nominal association with hip circumference always featured the WHR-increasing allele associated with reduced hip circumference. In contrast, we observed sexual dimorphism in the pattern of waist circumference associations. In women, the WHR-increasing allele at all 14 loci was associated with increased waist circumference, whereas this was only true for six of these loci in men (Fig. 3). At *GRB14*, for example, the WHR-increasing allele was associated with increased waist circumference in women ( $P = 3.6 \times 10^{-4}$ ) but with decreased waist circumference in men ( $P = 6.8 \times 10^{-3}$ ). These differences in the relationships between waist circumference, hip circumference and WHR underlie some of the sexual dimorphism in the patterns of WHR association.

### Enrichment of association with metabolic traits

We evaluated the 14 WHR-associated loci for their relationships with related metabolic traits using GWAS data provided by trait-specific consortia<sup>19-21</sup> as well as our *de novo* genotyped follow-up studies. As expected given the sample overlap between this GWAS data and our WHR GWAS data as well as information on known trait correlations (Supplementary Table 4), we observed directionally consistent enrichment of associations ( $P < 0.05$ ) between the 14 WHR-associated alleles and increased triglycerides, low-density lipoprotein (LDL) cholesterol, fasting insulin and homeostasis model assessment (HOMA)-derived measures of insulin resistance (binomial  $P$  from  $3.2 \times 10^{-4}$  to  $1.8 \times 10^{-8}$ ; Table 3 and Supplementary Table 5). For example, the WHR-increasing allele at *GRB14* showed strong associations with increased triglycerides ( $P = 7.4 \times 10^{-9}$ ), fasting insulin levels ( $P = 5.0 \times 10^{-6}$ ) and insulin resistance ( $P = 1.9 \times 10^{-6}$ ). Eleven of the 14 WHR-associated loci showed directionally consistent associations with T2D, with three of these loci (at *ADAMTS9*, *NISCH-STAB1* and *ITPR2-SSPN*) reaching nominal significance ( $P < 0.05$ ) (Table 3 and Supplementary Table 5). Because the association signals for correlated traits in this analysis were vulnerable to overestimation given the overlap in the GWAS samples examined, we repeated these analyses and restricted the samples included to those from our *de novo* genotyped follow-up studies. Although this also resulted in a lower sample size, similar patterns of enrichment were still observed (Supplementary Table 5).

### Pathway analysis and potential biological roles

To identify potential functional connections and pathway relationships between genes mapping at the WHR-associated loci, we focused on the 95 genes located in a 2-Mb interval centered around each of the 48 independent SNPs that attained  $P < 1.0 \times 10^{-5}$  in the WHR discovery studies.

First, we performed a survey of the published literature using GRAIL<sup>22</sup> to search for connectivity between the genes and specific keywords that describe these functional connections (Online Methods). Although there was no evidence after correcting for multiple

testing that the connectivity between these genes was greater than chance, we identified eight genes with nominal significance ( $P < 0.05$ ) for potential functional connectivity (*PLXND*, *HOXC10*, *TBX15*, *RSPO3*, *HOXC4*, *HOXC6*, *KREMEN1* and *HOXC11*). The keywords associated with these connections included ‘vegf’, ‘homeobox’, ‘patterning’, ‘mesenchyme’, ‘embryonic’, ‘development’ and ‘angiogenesis’.

Additionally, we performed pathway analyses using the PANTHER database<sup>23</sup> based on the same set of 95 genes (Online Methods and Supplementary Note). This analysis generated some evidence for over-representation of ‘developmental processes’ ( $P = 5.8 \times 10^{-8}$ ) and ‘mRNA transcription regulation’ ( $P = 2.7 \times 10^{-6}$ ) but neither of these factors retained nominal significance after adjustment for bias (for example, due to non-random SNP coverage in relation to genes) and the number of biological processes tested (Supplementary Note and Supplementary Table 6).

Finally, we examined the described functional roles of some of the most compelling candidates based on either proximity to the signal or the other analyses described in this paper. These analyses uncovered possible genetic roles in adipocyte development (*TBX15*), pattern formation during embryonic development (*HOXC13*), angiogenesis (*VEGFA*, *RSPO3* and *STAB1*), Wnt and  $\beta$ -catenin signaling (*RSPO3* and *KREMEN1*), insulin signaling (*ADAMTS9*, *GRB14* and *NISCH*), lipase activity (*LYPLALI*), lipid biosynthesis (*PIGC*) and intracellular calcium signaling (*ITPR2*) (Supplementary Note).

### Evaluation of copy number variants and non-synonymous changes

Both common and rare copy number variants (CNVs) have been reported to be associated with overall adiposity<sup>14,15,24,25</sup>, but the impact of CNVs on fat distribution has not been evaluated previously. To examine the potential contribution of common CNVs to variation in WHR, we looked for evidence of association in our genome-wide association discovery meta-analysis using a set of 6,018 CNV-tagging SNPs which collectively capture >40% of common CNVs that are greater than 1 kb in length<sup>26,27</sup> (Online Methods and Supplementary Note).

One CNV-tagging SNP (rs1294421 in *LY86*) was observed among our 14 WHR-associated loci. This SNP is in strong LD ( $r^2 = 0.98$ ) with a 2,832-bp duplication variant (CNVR2760.1)<sup>27</sup> located 12 kb from an expressed sequence tag (BC039678) and 87 kb from *LY86* such that the duplication allele is associated with reduced WHR. The duplicated region consists entirely of noncoding sequence but includes part of a predicted enhancer sequence (E.5552.1)<sup>28</sup>.

To identify other putatively causal variants in our associated regions, we searched for non-synonymous coding SNPs in strong LD (defined as  $r^2 > 0.7$ ) with the most strongly associated SNPs at each locus using data from the HapMap (Build 21) and 1000 Genomes Project (April and August 2009 releases). In this search, one lead SNP (rs6784615, at the *NISCH-STAB1* locus) was correlated with non-synonymous changes in two nearby genes, *DNAH1* (p.Val441Leu, p.Arg1285Trp and p.Arg3809Cys) and *GLYCTK* (p.Leu170Val). Fine-mapping and functional studies will be required to determine whether the *DNAH1* or *GLYCTK* SNPs or the *LY86* CNV are causal for the WHR associations at these loci.

### Effect of WHR associations on expression in relevant tissues

Expression quantitative trait locus (eQTL) data can implicate regional transcripts that mediate trait associations, and we therefore examined the 14 WHR-associated loci using eQTL data from human subcutaneous adipose tissue (SAT)<sup>29</sup> (two separate sample sets,  $n = 610$  and  $n = 603$ ), omental fat<sup>30</sup> ( $n = 740$ ), liver<sup>30</sup> ( $n = 518$ ), blood<sup>29</sup> ( $n = 745$ ) and lymphocytes<sup>31</sup> ( $n = 830$ ) (Online Methods and Supplementary Note).

At six of the loci, the WHR-associated SNP was either the strongest SNP associated with significant ( $P < 1.0 \times 10^{-5}$ ) expression of a local (within 1 Mb) gene transcript or explained the majority of the association between the most significant eQTL SNP and the gene transcript in conditional analyses (adjusted  $P > 0.05$ ; Table 4). For example, the WHR-associated SNP rs1011731 (near *DNM3-PIGC*) was strongly associated with expression of *PIGC* in lymphocytes ( $P = 5.9 \times 10^{-10}$ ); furthermore, rs1011731 is in high LD ( $r^2 = 1.00$ ,  $D' = 1.00$  from the HapMap CEU population) with the SNP with the strongest effect on *PIGC* expression (rs991790), and this *cis* eQTL association was abolished by conditioning on rs1011731. These analyses therefore indicate that these two signals are coincident and that *PIGC* is a strong candidate for mediating the WHR association at rs1011731. We found similar evidence for coincidence of the WHR signal with expression for rs984222 (*TBX15* in omental fat), rs1055144 (expressed sequence tag AA553656 in SAT), rs10195252 (*GRB14* in SAT), rs4823006 (*ZNRF3* in SAT and omental fat) and rs6784615 (*STAB1* in blood) (Table 4). Taken together, the overlap between trait association and gene expression at these loci suggests that the WHR associations may be driven through altered expression of *PIGC*, *TBX15*, AA553656, *GRB1*, *ZNRF3* and *STAB1*.

### RNA expression of gluteal and abdominal fat tissue

To determine whether genes within the WHR-associated loci showed evidence of differential transcription in distinct fat depots, we compared expression levels in gluteal or abdominal SAT in 49 individuals. We focused on the 15 genes with the strongest credentials for causal involvement (on the basis of proximity to the lead SNP and/or other biological or functional data; Table 1) for which expression data were available. Five of these genes (*RSPO3*, *TBX15*, *ITPR2*, *WARS2* and *STAB1*) were differentially expressed between the two tissues (using an *F* test, corrected for false discovery rate across the 15 expressed genes,  $P < 0.05$ ; Supplementary Table 7). This supports the hypothesis that, at some loci at least, the association with WHR reflects depot-specific differences in expression patterns.

## DISCUSSION

Overall, our findings demonstrate that the genetic regulation of body fat distribution involves loci and processes that are largely distinct from those that influence BMI and risk of obesity. This finding is consistent with the evidence that WHR displays substantial heritability even after adjustment for BMI. The loci that emerged from this study display no overlap with those shown to be associated with BMI either in previous reports<sup>14-16</sup> or in the expanded meta-analysis recently completed by the GIANT consortium<sup>32</sup>.

Another point of distinction between our findings and those for BMI relates to the evidence for sexual dimorphism that we observed at several of the WHR-associated loci. Sex differences in the regulation of body fat distribution have long been acknowledged without a clear understanding of the underlying molecular mechanisms. These differences become apparent during puberty and are generally attributed to the influence of sex hormones<sup>33</sup>. Consistent with our findings, variance decomposition studies have shown that the genetic contribution to the overall variance in WHR, waist and hip circumference is greater in women<sup>17</sup>. Although there is some evidence for loci with differential sex effects influencing lipids<sup>34</sup>, uric acid levels<sup>35</sup> and risk of schizophrenia<sup>36</sup>, we are unaware of prior reports indicating such strong enrichment of female-specific associations for any other phenotype, including BMI<sup>32</sup>.

The primary objective of genetic discovery efforts is to characterize the specific mechanisms involved in regulating the trait of interest. Despite the considerable challenges associated with moving from common variant association signals to defining causal alleles and pathways, we have identified strong candidates at several of the loci. For example, the *cis*



eQTL data implicate *GRB14* as a compelling candidate for the WHR association on chromosome 2, and we were able to show that the same *GRB14* variants are also associated with triglyceride and insulin levels, consistent with previous association of this locus with high-density lipoprotein (HDL) cholesterol<sup>37</sup>. These inferences about the role of *GRB14* are supported by evidence that *Grb14*-deficient mice exhibit improved glucose homeostasis despite lower circulating insulin levels, as well as enhanced insulin signaling in liver and skeletal muscle<sup>38</sup>. The signal near *ADAMTS9* overlaps a previously-reported T2D locus<sup>39</sup>, and the lead SNP for WHR in our study is identical to the SNP displaying the strongest T2D association in a previous expanded T2D meta-analysis<sup>40</sup>. Given evidence that *ADAMTS9* T2D risk alleles are associated with insulin resistance in peripheral tissues<sup>41</sup>, these findings are consistent with a primary effect of *ADAMTS9* variants on body fat distribution. At the chromosome 6 locus, *VEGFA* is the most apparent biological candidate given the presumed role of *VEGFA* as a mediator of adipogenesis<sup>42</sup> and evidence that serum levels of *VEGFA* are correlated with obesity<sup>43,44</sup>. Finally, at the *TBX15-WARS2* locus, *TBX15* emerges as the strongest candidate based on the *cis* eQTL data in omental fat, marked depot-specific differences in adipose tissue expression in mice and humans and associations between *TBX15* expression in visceral fat and WHR<sup>45,46</sup>.

Our efforts to use pathway- and literature-mining approaches to look for functional enrichment of the genes mapping to associated regions met with only limited success but did provide some support for over-representation of developmental processes. Developmental genes have been implicated in fat accumulation and distribution<sup>45,46</sup>, and recent evidence supports a link between developmental genes, including *HOXC13* (ref. <sup>47</sup>) and *TBX15* (refs. <sup>45,48</sup>), and body fat distribution. Developmental genes may in part determine the adipocyte-specific expression patterns that have been observed in different fat depots<sup>45</sup>. Taken together, our findings point to a set of genes influencing body fat distribution that have their principal effects in adipose tissue. This is in contrast to the predominantly central (hypothalamic) processes that are involved in the regulation of BMI and overall adiposity<sup>49</sup>.

By providing new insights into the regulation of body fat distribution, the present study raises a number of issues for future investigation. From the genetic perspective, re-sequencing, dense-array genotyping and fine-mapping approaches will be required to characterize causal variants at the loci we have identified and to support further discoveries that may account for the substantial proportion of genetic variance unexplained by our findings. From the clinical perspective, it will be important to explore the relationship of these variants to more refined measures of body fat distribution derived from detailed imaging studies, to use the variants identified to characterize the causal relationships between body fat distribution and related metabolic and cardiovascular traits and to explore population differences in patterns of body fat distribution. Efforts to tackle overall obesity through therapeutic or lifestyle-based modulation of overall energy balance have proved extremely challenging to implement, and the manipulation of processes associated with more beneficial patterns of fat distribution offers an alternative perspective for future drug discovery.

## METHODS

Methods and any associated references are available in the online version of the paper at <http://www.nature.com/naturegenetics/>.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

## Footnotes

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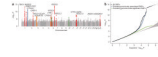
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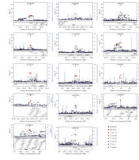
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**Figure 1.**

Genome-wide association analyses for WHR in discovery studies. (a) Manhattan plot shows results of the WHR association meta-analysis in discovery studies (with  $P$  values on the  $y$  axis and the SNP genomic position on the  $x$  axis). Colored genomic loci indicate significant association ( $P < 5 \times 10^{-8}$ ) detected previously (blue)<sup>13</sup>, in our GWAS stage (red) and after the meta-analysis combining GWAS data with that from the follow-up studies (orange). Two loci tested in the follow-up stage did not achieve genome-wide significance (green). (b) Quantile-quantile plot of SNPs for the discovery meta-analysis of WHR (black) and after removing SNPs within 1 Mb of either the recently reported *LYPLAL1* signal (blue) or the 14 significant associations (green). The gray area represents the 95% CI around the test statistic under the null distribution.



**Figure 2.**

Regional plots of 14 loci with genome-wide significant association. Shown is the SNP association with WHR in the meta-analysis of discovery studies for 14 loci (with  $-\log_{10} P$  values on the  $y$  axis and the SNP genomic position on the  $x$  axis). In each panel, an index SNP is denoted with a purple diamond and plotted using the  $P$  attained across discovery and follow-up data (Table 1). Estimated recombination rates are plotted in blue. SNPs are colored to reflect LD with the index SNP (pairwise  $r^2$  values from HapMap CEU). Gene and microRNA annotations are from the UCSC genome browser.



Table 1

Fourteen SNPs associated with WHr at genome-wide significant levels

SNP	Chr.	Position (b36)	Nearby genes	EA <sup>a</sup>	EAF <sup>b</sup>	Discovery			Follow-up			Combined		
						P	$\beta$	n	P	$\beta$	n	P	$\beta$	n
<b>SNPs evaluated in follow up achieving genome-wide significance</b>														
rs9491696	6	127,494,332	<i>RSPO3</i>	G	0.520	$2.10 \times 10^{-14}$	0.037	77,164	$3.27 \times 10^{-28}$	0.045	113,582	$1.84 \times 10^{-40}$	0.042	
rs6905288	6	43,866,851	<i>VEGFA</i>	A	0.562	$4.72 \times 10^{-10}$	0.033	77,129	$1.18 \times 10^{-16}$	0.039	95,430	$5.88 \times 10^{-25}$	0.036	
rs984222	1	119,305,366	<i>TBX15-WARS2</i>	G	0.365	$3.81 \times 10^{-14}$	0.037	77,167	$1.56 \times 10^{-12}$	0.031	109,623	$8.69 \times 10^{-25}$	0.034	
rs1055144	7	25,837,634	<i>NFE2L3</i>	T	0.210	$1.49 \times 10^{-8}$	0.034	77,145	$3.26 \times 10^{-18}$	0.043	113,636	$9.97 \times 10^{-25}$	0.040	
rs10195252	2	165,221,337	<i>GRB14</i>	T	0.599	$3.23 \times 10^{-10}$	0.031	77,119	$3.18 \times 10^{-16}$	0.036	102,449	$2.09 \times 10^{-24}$	0.033	
rs4846567	1	217,817,340	<i>LYPLAL1</i>	G	0.283	$2.37 \times 10^{-12}$	0.037	77,167	$3.15 \times 10^{-10}$	0.032	91,820	$6.89 \times 10^{-21}$	0.034	
rs1011731	1	170,613,171	<i>DNM3-PIGC</i>	G	0.572	$1.72 \times 10^{-10}$	0.031	77,094	$7.47 \times 10^{-9}$	0.026	92,018	$9.51 \times 10^{-18}$	0.028	
rs718314	12	26,344,550	<i>ITPR2-SSPN</i>	G	0.741	$2.41 \times 10^{-8}$	0.031	77,167	$1.49 \times 10^{-10}$	0.030	107,503	$1.14 \times 10^{-17}$	0.030	
rs1294421	6	6,688,148	<i>LY86</i>	G	0.387	$6.31 \times 10^{-9}$	0.029	77,154	$2.69 \times 10^{-10}$	0.028	102,189	$1.75 \times 10^{-17}$	0.028	
rs1443512	12	52,628,951	<i>HOXC13</i>	A	0.239	$3.33 \times 10^{-8}$	0.031	77,165	$2.92 \times 10^{-10}$	0.030	112,353	$6.38 \times 10^{-17}$	0.031	
rs6795735	3	64,680,405	<i>ADAMTS9</i>	C	0.406	$2.47 \times 10^{-7}$	0.025	77,162	$6.75 \times 10^{-8}$	0.026	84,480	$9.79 \times 10^{-14}$	0.025	
rs4823006	22	27,781,671	<i>ZNRF3-KREMEN1</i>	A	0.569	$4.47 \times 10^{-8}$	0.027	77,086	$2.41 \times 10^{-5}$	0.019	93,911	$1.10 \times 10^{-11}$	0.023	
rs6784615	3	52,481,466	<i>NISCH-STAB1</i>	T	0.941	$3.18 \times 10^{-7}$	0.052	76,859	$1.56 \times 10^{-4}$	0.036	109,028	$3.84 \times 10^{-10}$	0.043	
rs6861681	5	173,295,064	<i>CPEB4</i>	A	0.340	$1.40 \times 10^{-6}$	0.026	77,164	$2.13 \times 10^{-4}$	0.019	85,722	$1.91 \times 10^{-9}$	0.022	
<b>Further SNPs evaluated in follow up but not achieving genome-wide significance in the combined analysis</b>														
rs2076529	6	32,471,933	<i>BTNL2</i>	C	0.570	$2.22 \times 10^{-8}$	0.041	34,532	0.012	0.011	92,778	$3.71 \times 10^{-7}$	0.020	
rs7081678	10	32,030,629	<i>ZEB1</i>	A	0.085	$5.76 \times 10^{-7}$	0.045	76,270	0.094	0.013	100,527	$5.57 \times 10^{-6}$	0.027	

*P* values and  $\beta$  coefficients (per change of WHr-increasing allele) for the association with WHr on the inverse normal transformed ranked scale in the meta-analyses of discovery studies (up to 77,167 subjects), follow-up studies (up to 113,636 subjects) and both combined (up to 190,781 subjects). Fourteen of the sixteen SNPs examined in the follow-up samples showed genome-wide significant results ( $P < 5 \times 10^{-8}$ ) in the combined analysis. *P* values in the discovery stage were genomic control corrected per study and in the meta-analysis. Details on between-study heterogeneity are given in Supplementary Table 1c.

<sup>a</sup>EA, effect allele (WHR-increasing allele on the forward strand).

<sup>b</sup>EAF, effect allele frequency. Chr., chromosome.



Table 2

Evidence of sex-differences in the WHr association at seven of the 14 associated loci

SNP	Nearby genes	Men						Women						Sex difference	
		Discovery		Follow up		Combined		Discovery		Follow up		Combined		β	P
		P	β	P	β	P	β	P	β	P	P	β			
rs9491696	<i>RSPO3</i>	$1.68 \times 10^{-4}$	0.026	$6.97 \times 10^{-9}$	0.036	$1.05 \times 10^{-11}$	0.031	$1.62 \times 10^{-12}$	0.047	$8.84 \times 10^{-22}$	0.053	$1.93 \times 10^{-32}$	0.050	$1.94 \times 10^{-3}$	
rs6905288	<i>VEGFA</i>	0.066	0.013	$2.09 \times 10^{-4}$	0.025	$7.38 \times 10^{-5}$	0.020	$7.72 \times 10^{-13}$	0.052	$3.14 \times 10^{-15}$	0.051	$2.27 \times 10^{-26}$	0.052	$5.20 \times 10^{-6}$	
rs984222	<i>TBX15-WARS2</i>	$3.32 \times 10^{-9}$	0.041	$2.43 \times 10^{-5}$	0.029	$9.41 \times 10^{-13}$	0.035	$1.21 \times 10^{-7}$	0.036	$1.33 \times 10^{-8}$	0.033	$1.02 \times 10^{-14}$	0.034	0.951	
rs1055144	<i>NFE2L3</i>	$6.00 \times 10^{-4}$	0.029	$5.67 \times 10^{-8}$	0.040	$2.52 \times 10^{-10}$	0.035	$2.34 \times 10^{-6}$	0.040	$7.13 \times 10^{-12}$	0.046	$1.41 \times 10^{-16}$	0.044	0.270	
rs10195252	<i>GRB14</i>	0.201	0.009	0.114	0.011	0.043	0.010	$6.33 \times 10^{-15}$	0.053	$4.95 \times 10^{-21}$	0.054	$3.84 \times 10^{-34}$	0.054	$1.41 \times 10^{-11}$	
rs4846567	<i>LYPLALI</i>	0.191	0.010	0.982	0.000	0.358	0.005	$4.84 \times 10^{-18}$	0.064	$8.12 \times 10^{-17}$	0.055	$4.95 \times 10^{-33}$	0.059	$1.18 \times 10^{-13}$	
rs1011731	<i>DNM3-PIGC</i>	$4.88 \times 10^{-7}$	0.034	$1.95 \times 10^{-3}$	0.022	$7.81 \times 10^{-9}$	0.028	$2.13 \times 10^{-5}$	0.028	$7.03 \times 10^{-7}$	0.030	$6.90 \times 10^{-11}$	0.029	0.855	
rs718314	<i>ITPR2-SSPN</i>	0.177	0.010	$2.02 \times 10^{-3}$	0.022	$1.41 \times 10^{-3}$	0.017	$8.29 \times 10^{-10}$	0.047	$4.21 \times 10^{-9}$	0.038	$2.41 \times 10^{-17}$	0.042	$4.67 \times 10^{-4}$	
rs1294421	<i>LY86</i>	$4.18 \times 10^{-3}$	0.020	$7.00 \times 10^{-6}$	0.030	$1.63 \times 10^{-7}$	0.025	$3.44 \times 10^{-8}$	0.038	$7.32 \times 10^{-6}$	0.026	$2.40 \times 10^{-12}$	0.031	0.357	
rs1443512	<i>HOXC13</i>	0.184	0.011	$9.74 \times 10^{-4}$	0.024	$9.45 \times 10^{-4}$	0.018	$1.43 \times 10^{-9}$	0.048	$3.09 \times 10^{-8}$	0.035	$6.38 \times 10^{-16}$	0.040	$2.23 \times 10^{-3}$	
rs6795735	<i>AD-AMTS9</i>	0.011	0.017	0.614	0.004	0.027	0.011	$7.85 \times 10^{-7}$	0.033	$2.95 \times 10^{-11}$	0.042	$1.92 \times 10^{-16}$	0.038	$8.50 \times 10^{-5}$	
rs4823006	<i>ZNRF3-KRE</i>	$6.87 \times 10^{-3}$	0.019	0.094	0.012	$1.94 \times 10^{-3}$	0.015	$6.86 \times 10^{-8}$	0.037	$3.81 \times 10^{-5}$	0.024	$3.24 \times 10^{-11}$	0.030	0.032	
rs6784615	<i>NISCH-MEN1</i>	$1.51 \times 10^{-3}$	0.045	0.033	0.032	$1.68 \times 10^{-4}$	0.039	$6.23 \times 10^{-5}$	0.057	$1.72 \times 10^{-3}$	0.039	$6.01 \times 10^{-7}$	0.047	0.574	
rs6861681	<i>CPEB4</i>	$1.88 \times 10^{-3}$	0.023	0.045	0.015	$3.03 \times 10^{-4}$	0.019	$2.14 \times 10^{-4}$	0.027	$1.58 \times 10^{-3}$	0.021	$1.55 \times 10^{-6}$	0.024	0.555	

*P* values and  $\beta$  coefficients (per change of WHr-increasing allele in the sex-combined analysis as in Table 1) for the WHr association are given for the discovery (up to 34,601 men and 42,735 women), the follow-up (up to 47,882 men and 65,780 women) and the combined meta-analysis (up to 81,301 men and 107,429 women). Also given are the *P* values for testing for difference between sex-specific  $\beta$  coefficients in the combined meta-analysis; SNPs with *P* for sex difference  $< 3.6 \times 10^{-3}$  (0.05/14) were considered to show a significant sex difference.

**Table 3**

WHR signals show enrichment of association with other traits related to metabolic disorders

Trait	Sample size <sup>d</sup>	SNPs in concordant direction <sup>b</sup>		SNPs in concordant direction with $P < 0.05^c$	
		n	P	n	P
Triglycerides	43,826	14	$6.10 \times 10^{-5}$	7	$1.79 \times 10^{-8}$
HDL-C	45,561	13	$9.16 \times 10^{-4}$	4	$3.20 \times 10^{-4}$
LDL-C	43,889	10	0.090	1	0.298
Fasting glucose	63,849	10	0.090	1	0.298
Fasting insulin	54,883	13	$9.16 \times 10^{-4}$	5	$1.62 \times 10^{-5}$
HOMA-IR	53,625	13	$9.16 \times 10^{-4}$	6	$6.17 \times 10^{-7}$
2 h glucose	27,011	7	0.605	0	1.000
Type 2 diabetes	10,128 <sup>d</sup>	11	0.029	3	$4.62 \times 10^{-3}$

The 14 WHR SNPs were tested for association with other traits by meta-analysis of GWAS data from previous reports 19–21,39 together with our non-overlapping *de novo* genotyped follow-up studies. HDL-C, high density lipoprotein cholesterol; LDL-C, low density lipoprotein cholesterol; HOMA-IR, index of insulin resistance; 2 h glucose, glucose levels 2 h after an oral glucose challenge.

<sup>a</sup>Maximum number of subjects available for any of the 14 SNPs.

<sup>b</sup>Number of the 14 SNPs for which the WHR-increasing allele is associated with the trait in the concordant direction (that is, increased levels, except for HDL-C) and corresponding binomial  $P$  value to test whether this number is greater than that expected by chance and not accounting for the correlation between the traits.

<sup>c</sup>Number of SNPs in concordant direction that show  $P < 0.05$  for the association with the trait and the corresponding binomial  $P$  value as in <sup>b</sup>.

<sup>d</sup>4,549 cases, 5579 controls.

Table 4

Expression quantitative trait locus analysis for 11 of the 14 WHR signals

WHR SNP	Tissue	Gene	Effect <sup>a</sup>	WHR SNP association with transcript ( <i>P</i> )		Transcript peak SNP <sup>b</sup>	LD ( <i>r</i> <sup>2</sup> ) <sup>c</sup>	Peak SNP association with transcript ( <i>P</i> )	
				Unadj.	Adj. for peak SNP			Unadj.	Adj. for WHR SNP
rs9491696	SAT-D	<i>RSP03</i>	+	1.10 × 10 <sup>-7</sup>	0.03	rs1936795	0.26	2.20 × 10 <sup>-13</sup>	7.40 × 10 <sup>-8</sup>
<b>rs984222</b>	<b>Omental</b>	<b><i>TBX15</i></b>	+	<b>7.90 × 10<sup>-10</sup></b>	<b>1.00</b>	<b>rs984222</b>	<b>1.00</b>	<b>7.90 × 10<sup>-10</sup></b>	<b>1.00</b>
	Omental	<i>WARS2</i>	+	5.11 × 10 <sup>-36</sup>	0.03	rs10802075	0.27	1.31 × 10 <sup>-163</sup>	1.33 × 10 <sup>-88</sup>
	Subcutaneous fat	<i>WARS2</i>	+	1.67 × 10 <sup>-25</sup>	0.01	rs10802075	0.22	3.88 × 10 <sup>-110</sup>	1.01 × 10 <sup>-63</sup>
	Lymphocytes	<i>WARS2</i>	-	4.30 × 10 <sup>-18</sup>	5.47 × 10 <sup>-5</sup>	rs2645305	0.27	5.57 × 10 <sup>-40</sup>	6.88 × 10 <sup>-26</sup>
	Liver	<i>WARS2</i>	+	2.57 × 10 <sup>-17</sup>	0.07	rs1057990	0.26	6.69 × 10 <sup>-59</sup>	1.97 × 10 <sup>-32</sup>
	SAT-D	<i>WARS2</i>	+	1.10 × 10 <sup>-18</sup>	0.51	rs1057990	0.26	5.80 × 10 <sup>-130</sup>	5.80 × 10 <sup>-100</sup>
	Blood	<i>WARS2</i>	+	6.10 × 10 <sup>-17</sup>	0.11	rs1057990	0.26	6.30 × 10 <sup>-75</sup>	1.10 × 10 <sup>-54</sup>
<b>rs1055144</b>	<b>SAT-D</b>	<b><i>AA553656</i><sup>d</sup></b>	-	<b>1.20 × 10<sup>-11</sup></b>	<b>0.96</b>	<b>rs7798002</b>	<b>0.95</b>	<b>7.20 × 10<sup>-12</sup></b>	<b>0.32</b>
	SAT-M	<i>AA553656</i> <sup>d</sup>	-	2.46 × 10 <sup>-7</sup>	0.65	rs1451385	0.77	5.93 × 10 <sup>-8</sup>	0.38
<b>rs10195252</b>	<b>SAT-D</b>	<b><i>GRB14</i></b>	+	<b>4.40 × 10<sup>-11</sup></b>	<b>1.00</b>	<b>rs10195252</b>	<b>1.00</b>	<b>4.40 × 10<sup>-11</sup></b>	<b>1.00</b>
	SAT-M	<i>GRB14</i>	+	5.51 × 10 <sup>-6</sup>	1.00	rs10184004	1.00	5.51 × 10 <sup>-6</sup>	1.00
	<b>Omental</b>	<b><i>GRB14</i></b>	+	<b>1.02 × 10<sup>-13</sup></b>	<b>1.00</b>	<b>rs10195252</b>	<b>1.00</b>	<b>1.02 × 10<sup>-13</sup></b>	<b>1.00</b>
	SAT-M	<i>SLC38A11</i>	-	3.93 × 10 <sup>-6</sup>	0.66	rs10184126	0.18	7.76 × 10 <sup>-44</sup>	8.57 × 10 <sup>-34</sup>
	SAT-D	<i>SLC38A11</i>	-	3.70 × 10 <sup>-9</sup>	0.35	rs10184126	0.18	2.40 × 10 <sup>-94</sup>	7.40 × 10 <sup>-82</sup>
rs1011731	Blood	<i>C1orf105</i>	+	3.80 × 10 <sup>-16</sup>	0.20	rs2157451	0.28	1.30 × 10 <sup>-33</sup>	8.20 × 10 <sup>-18</sup>
	<b>Lymphocytes</b>	<b><i>PIGC</i></b>	-	<b>5.87 × 10<sup>-10</sup></b>	<b>1.00</b>	<b>rs991790</b>	<b>1.00</b>	<b>5.65 × 10<sup>-10</sup></b>	<b>1.00</b>
rs718314	Lymphocytes	<i>ITPR2</i>	+	1.79 × 10 <sup>-9</sup>	0.98	rs7976877	0.45	2.21 × 10 <sup>-18</sup>	1.91 × 10 <sup>-6</sup>
	Blood	<i>ITPR2</i>	-	2.40 × 10 <sup>-9</sup>	0.20	rs2570	0.41	2.40 × 10 <sup>-37</sup>	1.80 × 10 <sup>-28</sup>
rs1294421	SAT-M	<i>BC039678</i>	-	2.43 × 10 <sup>-7</sup>	0.38	rs1294404	0.64	1.89 × 10 <sup>-16</sup>	3.42 × 10 <sup>-4</sup>
	Omental	<i>BC039678</i>	-	1.09 × 10 <sup>-6</sup>	0.33	rs912056	0.71	8.28 × 10 <sup>-17</sup>	4.26 × 10 <sup>-5</sup>
rs6795735	SAT-D	<i>ADAMTS9</i>	-	1.50 × 10 <sup>-6</sup>	0.04	rs7372321	0.11	1.10 × 10 <sup>-9</sup>	2.30 × 10 <sup>-5</sup>
	Omental	<i>AK022320</i>	-	7.99 × 10 <sup>-15</sup>	0.64	rs4521216	0.02	5.15 × 10 <sup>-42</sup>	1.49 × 10 <sup>-19</sup>
	SAT-D	<i>AK022320</i>	-	2.24 × 10 <sup>-10</sup>	0.98	rs4521216	0.02	9.62 × 10 <sup>-37</sup>	7.58 × 10 <sup>-19</sup>

WHR SNP	Tissue	Gene	Effect <sup>d</sup>	WHR SNP association with transcript ( <i>P</i> )		Transcript peak SNP <sup>b</sup>	LD ( <i>r</i> <sup>2</sup> ) <sup>c</sup>	Peak SNP association with transcript ( <i>P</i> )	
				Unadj.	Adj. for peak SNP			Unadj.	Adj. for WHR SNP
rs4823006	SAT-D	<i>ZNRF3</i>	-	2.40 × 10 <sup>-8</sup>	0.63	rs3178915	0.81	6.70 × 10 <sup>-11</sup>	8.90 × 10 <sup>-4</sup>
	SAT-M	<i>ZNRF3</i>	-	1.08 × 10 <sup>-18</sup>	0.93	rs6005975	0.79	1.59 × 10 <sup>-19</sup>	0.50
	Omental	<i>ZNRF3</i>	-	9.13 × 10 <sup>-18</sup>	0.98	rs6005975	0.79	6.07 × 10 <sup>-21</sup>	0.27
rs6784615	Blood	<i>STAB1</i>	+	2.80 × 10 <sup>-9</sup>	0.32	rs9846089	0.83	9.40 × 10 <sup>-10</sup>	0.08
	Lymphocytes	<i>CPEB4</i>	+	3.79 × 10 <sup>-22</sup>	0.89	rs7705502	0.87	4.95 × 10 <sup>-29</sup>	2.00 × 10 <sup>-3</sup>
	Blood	<i>HMP19</i>	+	1.60 × 10 <sup>-16</sup>	0.97	rs10516107	0.83	1.10 × 10 <sup>-21</sup>	4.30 × 10 <sup>-6</sup>

Association between the 14 WHR SNPs and expression of transcripts located within 1 Mb of the WHR SNP in two sets of abdominal subcutaneous adipose tissue (SAT-D from deCODE and SAT-M from Massachusetts General Hospital), omental fat, liver, lymphocytes and blood (Supplementary Note). Results are given if the unadjusted WHR SNP association showed  $P < 1.00 \times 10^{-5}$ . Findings are highlighted in bold font where the WHR SNP was the transcript peak SNP or where the WHR signal and the *cis* eQTL signal were considered coincident (that is, the transcript peak SNP was highly correlated with the WHR SNP,  $r^2 > 0.7$  and the transcript peak association disappeared by adjusting on the WHR SNP,  $P > 0.05$ ); see also Online Methods. Unadj., unadjusted; Adj., adjusted.

<sup>d</sup>Effect direction for the WHR-increasing allele.

<sup>b</sup>SNP with the strongest association with the transcript in the region (transcript peak SNP).

<sup>c</sup>Correlation (HapMap CEU, build 36) between the WHR SNP and the transcript peak SNP.

<sup>d</sup>The transcript labeled AA553656 was detected as Contig27623\_RC and corresponds to chromosome 7 locations 25,854,143–25,854,203 (HapMap build 36).