

Published in final edited form as:

Nat Genet. 2013 June ; 45(6): . doi:10.1038/ng.2635.

Identification of nine new susceptibility loci for testicular cancer, including variants near *DAZL* and *PRDM14*

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Abstract

Testicular germ cell tumor (TGCT) is the most common cancer in young men and is notable for its high familial risks^{1,2}. To date, six loci associated with TGCT have been reported³⁻⁷. From GWAS

Author Contributions C.T and N.R designed the study. C.T, N.R., R.A.H. and U.K.T.C.C.. coordinated the studies that provided case samples. D.F.E., J.P., R.E, Z.K-J. and K.M. coordinated studies that provided control samples. C.T, H.M., S.S, A.E., and S.DVD coordinated sample management, selection and transfer. M.R.S., D.F.E., D.T.B. J.N. and E.Ra. designed and executed the original case genome-wide genotyping. C.T. designed the statistical analyses. C.T. and E.Ru., conducted statistical analyses with assistance from K.W.L and F.Z.. C.T. drafted the manuscript with assistance from N.R., E.Ru., J.S, J.N, D.T.B. and E.P. All authors contributed to the final paper.

Competing financial interests The authors declare no competing financial interests

URLs. Illuminus algorithm, <https://www.sanger.ac.uk/resources/software/illuminus/>

PLINK, <http://pngu.mgh.harvard.edu/~purcell/plink/>;

CaTS Power Calculator, <http://www.sph.umich.edu/csg/abecasis/CaTS/>;

Haploview, <http://www.broadinstitute.org/haploview/haploview/>;

SNAP, <http://www.broadinstitute.org/mpg/snap/>;

Wellcome Trust Case Control Consortium 2, <http://www.wtccc.org.uk/cc2/>

COGS <http://www.cogseu.org/>

GenCall http://www.illumina.com/documents/products/technotes/technote_gencall_data_analysis_softwa_re.pdf

NHGRI Catalog of Published Genome-Wide Association Studies, <http://www.genome.gov/gwastudies/>

Haploreg, <http://www.broadinstitute.org/mammals/haploreg/haploreg.php>

Genevar, <http://www.sanger.ac.uk/resources/software/genevar/>

analysis of 307,291 SNPs in 986 cases and 4,946 controls, we selected for follow-up 694 SNPs, which we genotyped in a further 1,064 TGCT cases and 10,082 controls from the UK. We identified SNPs at nine new loci showing association with TGCT ($P < 5 \times 10^{-8}$), at 1q22, 1q24.1, 3p24.3, 4q24, 5q31.1, 8q13.3, 16q12.1, 17q22 and 21q22.3, which together account for an additional 4–6% of the familial risk of TGCT. The loci include genes plausibly related to TGCT development. *PRDM14*, at 8q13.3, is essential for early germ cell specification⁸ whilst *DAZL*, at 3p24.3, is required for regulation of germ cell development⁹. Furthermore, *PITX1*, at 5q31.1 regulates TERT expression, and is the third TGCT locus implicated in telomerase regulation¹⁰.

Testicular germ cell tumor (TGCT) is the most common malignancy in men aged 15–45 years¹. Studies in families have estimated the risk to brothers of TGCT cases to be increased 8- to 10- fold and to sons of cases 4- to 6-fold, and these risks are substantially higher than the equivalent familial relatives risk of ~2 typical for common cancers such as breast, colorectal and prostate². Recent genome-wide association studies (GWAS) of TGCT have identified eight associated SNPs at six loci, which together account for >11% of the genetic risk of TGCT³⁻⁷. These loci lie on chromosome 12q21, encompassing *KITLG*, 5q31 (*SPRY4*), 6p21 (*BAK1*), 5p15 (*TERT* and *CLPTMIL*), 12p13 (*ATF7IP*) and 9p24 (*DMRT1*)¹¹.

To identify further susceptibility loci for TGCT, we identified the most strongly associated 1,050 SNPs from our previous GWAS, which comprised 307,291 SNPs genotyped in 986 cases of TGCT and 4,946 controls from the UK^{3,4}. These SNPs were included on a custom Illumina iSelect genotyping array (the iCOGS array). The iCOGS array includes 211,155 cancer-related SNPs selected for further evaluation in different cancers and has already yielded 80 new loci for breast, ovarian and prostate cancer^{12,13}. Using the iCOGS array, we genotyped 1,094 cases of TGCT from the UK, which had not been included in the original GWAS. We compared these genotypes to those of three cancer-free UK control series, also genotyped on the iCOGS array. Following quality control exclusions, we obtained data for 694 of the 1,050 SNPs in 1,064 cases and 10,082 controls (the ‘iCOGS replication series’).

We tested association between each SNP and TGCT risk using a 1df trend test, adjusted for six principle components. Inflation in the test statistic, quantified using 36,879 uncorrelated non-TGCT-related SNPs, revealed evidence of only modest inflation of the test statistics (Supplementary Figure 1, $\lambda = 1.14$, $\lambda_{1000} = 1.07$; following adjustment for six principle components, $\lambda = 1.05$, $\lambda_{1000} = 1.02$). There was clear evidence of association of the six previously reported loci in the iCOGS replication series ($P_{\text{trend}} = 3.88 \times 10^{-9}$ to 9.31×10^{-28}), with odds ratios consistent with those previously published (Supplementary Table 1).

In a combined fixed-effects meta-analysis of the GWAS and the adjusted iCOGS replication for the remaining 640 SNPs outside of these six loci, we identified association of $P < 5 \times 10^{-8}$ for a further 18 SNPs at nine loci: 1q22, 1q24.1, 3p24.3, 4q24, 5q31.1, 8q13.3, 16q12.1, 17q22 and 21q22.3. Following multiple logistic regression to test for independence of effects, there remained one SNP at each of the nine loci showing association with TGCT (Table 1). The associations at 1q24.1 and 17q22 have been replicated concurrently in other independent TGCT series^{14,15}.

For each SNP, we examined for evidence of departure from a log-additive (multiplicative) model: only rs2072499 at 1q22 showed evidence for departure from the log-additive model, indicating a recessive effect (heterozygote OR (OR_{het}): 1.00 (0.87-1.16), homozygote OR (OR_{hom}): 1.53 (1.27-1.84), $P_{2\text{df}} = 5.18 \times 10^{-6}$, Supplementary Table 2). This observation was supported by genotypic (2df) analysis of the GWAS data for rs2072499 (OR_{het} 1.09 (0.94-1.26) OR_{hom} 1.64 (1.24-2.01), $P_{2\text{df}} 1.25 \times 10^{-6}$). We found no evidence for statistical interaction between any pair of these new nine SNPs or with the previous eight SNPs; for

every combination of SNPs, the combined risk was consistent with the product of the individual risks.

We investigated whether the newly identified loci were associated with different risks in subgroups of TGCT cases characterized by specific phenotypic characteristics (Supplementary Tables 3,4,5). Consistent with our findings at the previous six loci, none of the newly identified loci showed a significant difference in effect when comparing cases of the two TGCT histological subtypes, seminoma and nonseminoma. Likewise, there was no clear evidence of differential association by age of diagnosis, presence of testicular maldescent, family history of TGCT or unilateral versus bilateral disease, although power to detect differences in effects was limited on account of the case distribution.

Encyclopedia of DNA Elements (ENCODE) data indicated that rs2072499, rs3790672, 3805663, rs7010162, rs9905704 and rs2839186 may have effects on transcription on account of evidence of location within a DNase I hypersensitivity site, local modification of histones indicating promoter/enhancer activity, influence on binding of transcription factors and/or effect on a regulatory motif. Functional annotations from the ENCODE data for all correlated variants within each of the nine LD block ($r^2 > 0.8$), are shown in Supplementary Table 6. We then investigated whether the genotype at these nine SNPs was associated with differential expression of genes lying within 500kb of the SNP: there were no compelling patterns evident from expression data available from fibroblast, skin, adipose and lymphoblastoid cells (Supplementary Table 7). Further examination in testicular germ cells both for evidence of transcriptional regulation and of gene expression would be of interest.

Evidence supports TGCT developing from primordial germ cell or gonocytes, with tumor initiation described *in utero* through a pre-invasive stage termed intra germ cell neoplasia unclassified (IGCNU) or carcinoma *in situ* (CIS)¹⁶. It is therefore noteworthy that a number of the genes identified at loci identified in this study relate to pathways of early differentiation and development of the testicular germ cell¹¹ (Table 1).

rs10510452 lies in an LD block of 82kb at 3p24.3 which contains only one gene, *DAZL* (deleted in azoospermia-like). *DAZL* encodes an RNA-binding protein which has been shown from expression studies in human primordial germ cells to have a central role in early differentiation of primordial germ cells¹⁷. *Dazl* knockout mice are infertile, with differentiation of the germ cells halted at the A spermatogonia level⁹ and expression of *DAZL* is testis-specific^{18,19}. Sequence analysis indicates that the DAZ cluster on the Y-chromosome arose during primate evolution via transposition of the *DAZL* gene²⁰. Of note, *DAZ* has been implicated as the critical gene within the Y-chromosome AZF (azoospermia factor) region, so-named as deletions of this region cause non-obstructive spermatogenetic failure (azoospermia) in humans²¹.

rs7010162 lies in an LD block of 61kb at 8q13.3 in which there is one gene, *PRDM14*, which is required for both the reacquisition of potential pluripotency and the genomewide epigenetic reprogramming necessary for primordial germ cell specification^{8,22,23}. *PRDM14* encodes a transcriptional regulator which controls expression of key pluripotency genes such as *POU5F1* (*OCT4*), *NANOG* and *SOX2*, which are highly expressed in IGCNU and TGCT^{24,25}. Whilst normal *PRDM14* expression is restricted to germ cell and stem cell lineages, amplification and/or over-expression of *PRDM14* have been reported in a number of different cancer types, including TGCT²⁶⁻²⁹.

rs3805663 is located in a 51kb LD block at 5q31.1 which contains two genes: *CATSPER3* and *PITX1*. *CATSPER3* (Cation-Channel, Sperm-associated 3) is one of a family of genes which together form a functional hetero-tetrameric cation channel, exclusively expressed in the testis and essential for hyperactivated motility of the differentiated germ cell

(spermatozoa); accordingly *Catsper3*^{-/-} male mice are completely infertile³⁰⁻³². An equally plausible candidate gene at this locus is *PITX1*, (paired-like homeodomain transcription factor 1), which binds to specific *PITX1*-binding sites in the promoter region of *TERT*, thus regulating expression of telomerase¹⁰. Thus, along with the previously identified TGCT-associated loci at 5p15 (*TERT*) and 12p13 (*ATF7IP*, which regulates expression of *TERT*), the signal at 5q31.1 may also be mediated via effects on telomerase regulation. Whilst the 5p15 (*TERT*) region is notable for its associations with multiple cancers³³⁻³⁹, to date there have been no associations reported with other cancers for the loci at 5q31.1 (*PITX1*) or 12p13 (*ATF7IP*).

rs9905704 lies at 17q22 in an LD block of 813kB in which there are several genes including *RAD51C* and *TEX14* (testis expressed 14). *TEX14* is a protein kinase highly expressed in the human male germ cell^{40,41}. Male *Tex14*^{-/-} mice are infertile whilst the female knockouts are normally fertile⁴². Cellular studies have demonstrated that *TEX14* regulates kinetochore-microtubule assembly and spindle assembly checkpoint in the germ cells of the testis and requires recruitment of key proteins such as *CENPE*, a kinesin-like motor protein which associates with the centromere⁴¹⁻⁴³. It is thus noteworthy that *CENPE* (Centromere-associated protein E) is located within the 214kb TGCT-associated LD block at 4q24. Furthermore, *PMF1*, one of the two genes located in the TGCT-associated region at 1q22, is part of the *MIS12* complex which is required for normal chromosome alignment, segregation and kinetochore formation during mitosis⁴⁴. Therefore, regulation of microtubule assembly may be the mechanism linking the signals at 17q22, 4q24 and 1q22.

In summary, we have identified nine new loci for TGCT, which brings the total number of TGCT-associated loci identified to date to fifteen. These fifteen loci have provided considerable new insights into TGC tumorigenesis, implicating genes involved in germ cell specification and/or differentiation (*DAZL*, *PRDM14*) including the *KIT-KITL* signalling pathway (*KITL*, *SPRY4*, *BAKI*) and genes involved in sex-determination (*DMRT1*), microtubule assembly (*TEX14*, *CENPE*, *PMF1*) and telomerase regulation (*TERT*, *ATF7IP*, *PITX1*). The nine new susceptibility alleles account for ~4% of the excess familial risk to brothers and ~6% to sons of men with TGCT: this brings the cumulative totals to ~15% and ~22% respectively. The power to detect these loci in this combined analysis was modest (29-84%): accordingly there are likely to be several additional loci for TGCT of equivalent or lesser effects, which may be identifiable via still larger follow-up studies of these data and meta-analyses with other GWAS data. Moreover, imputation and fine-scale association mapping of the loci may uncover more of the missing heritability, either through identifying more strongly associated variants or by identifying additional signals, including rare variants, at these loci. Integration of these genetic variants together with non-genetic TGCT risk factors into risk models may enable clinically useful risk profiling for TGCT in unaffected men; which in turn may provide a rational basis for application of strategies for screening and targeted prevention of TGCT.

Methods

Samples

See Supplementary Note

GWAS analysis

Cases were genotyped on the Illumina HumanCNV370-Duo bead array and controls were genotyped on the Illumina Infinium 1.2M array at the Wellcome Trust Sanger Institute (see³ and⁴ for full methods). The genotypes were re-called since the analysis reported in⁴, such that SNPs on both sex chromosomes were included in the current analysis. We used data on

314,861 SNPs that were successfully genotyped on both arrays. We excluded individuals: with low call rate (<95%), with abnormal autosomal heterozygosity or with >10% non-European ancestry (based on multi-dimensional scaling). We filtered out all SNPs with (i) minor allele frequency <1%, (ii) a call rate of <95% in cases or controls or (iii) minor allele frequency of 1–5% and a call rate of <99% or (iv) deviation from Hardy-Weinberg equilibrium (10^{-12} in controls and 10^{-5} in cases). 307,291 SNPs passed the QC filters and we put forward for replication the 1050 SNPs with the lowest P-value ($P < 1.3 \times 10^{-3}$), based on the 1-d.f. Cochran-Armitage trend test.

iCOGS Genotyping

Genotyping was conducted using a custom Illumina Infinium array (iCOGS array) comprising 211,155 SNPs selected across multiple consortia within the COGS (Collaborative Oncological Gene-environment Study, see ¹² for details of SNP inclusion). Of the 1,050 SNPs submitted from our analysis, 740 attained an Illumina design score of 0.8, and were included on the array. Genotyping was performed at Genome Quebec, one of the seven genotyping centres providing genotyping for the COGS collaboration. Genotypes were called using Illumina's proprietary GenCall algorithm. Pan-COGS initiatives were employed across experiments to optimize calling algorithms for the iCOGS array: firstly, initial calling employed a cluster file generated using samples from Hapmap2; secondly, a cluster file based on 3,018 individuals selected across genotyping centers, across participating consortia and across ethnicities, combined with 380 samples of European, Asian or African ancestry genotyped as part of the Hapmap and 1000 genomes project was applied to call the genotypes for the subsequent samples. Thirdly, the performance of GenCall was compared to two other calling algorithms: Illuminus54 and GenoSNP55: all three algorithms were >99% concordant in their calling for 91% of the SNPs on the array. Manual inspection of a sample of the discrepant SNPs indicated that the GenCall calls were almost invariably superior (because Illuminus and GenoSNP frequently attempted to call SNPs that clustered poorly) ¹².

Quality Control

Using the full SNP set of 211,155 SNPs on the iCOGS array, we applied quality control exclusions as follows to subjects: i) subjects with overall call rate <95% or low or high heterozygosity ($P < 10^{-6}$) (5 cases), ii) using identity-by-state estimates based on 37,046 uncorrelated SNPs, we identified “cryptic” duplicates and related samples and the sample with the lower call rate was excluded (7 cases), iii) we identified ethnic outliers by multi-dimensional scaling by combining the iCOGS data with the three Hapmap2 populations using 37,046 uncorrelated markers and removed individuals with >10% non-Western European ancestry (18 cases). We included 1,064 cases and 10,082 controls in the final analysis.

We applied quality control exclusions as follows to SNPs: i) discrepant calls in more than 2% of duplicate samples across COGS consortia, ii) call rate <95%, MAF<1%, call rate <99% if MAF=1-5%, iii) deviation from Hardy-Weinberg ($P < 10^{-5}$ in controls, $P < 10^{-12}$ in cases). Following quality control exclusions applied individually to case and control data, we included genotypes from 694 SNPs in subsequent analyses. We checked genotype intensity cluster plots manually for SNPs in each new region for which a genome-wide significant association was achieved.

Statistical Analysis

For the 694 SNPs in the TGCT iCOGS replication experiment, we estimated the per-allele odds ratio (OR), 1df, using logistic regression, adjusting for principal components as covariates. We generated eight principle components and examined reduction in the

inflation factor using iterative combinations of principle components. Inclusion of principle components 1-6 achieved optimal reduction in inflation with no further reduction achieved by incorporating additional principle components into the regression model. We found no significant differences in the point estimates of effect size on performing sensitivity analysis between the control groups used in the TGCT iCOGS replication experiment.

The inflation factor (λ) was derived by dividing the median of the lowest 90% of the 1-d.f. statistics by the 45% percentile of a 1-d.f. χ^2 distribution (0.357), utilizing 36,879 uncorrelated SNPs selected to not include TGCT-associated SNPs. The inflation was converted to an equivalent inflation for a study with 1000 cases and 1000 controls ($\lambda_{1,000}$):

$$\lambda_{1,000} = 1 + 500(1/N_{\text{cases}} + 1/N_{\text{controls}})^* (\lambda - 1)$$

We excluded from further analysis the 54 SNPs lying within the six previously identified TGCT-associated loci, defining these LD blocks via the Oxford recombination rates. We included the 640 remaining SNPs in the replication analysis.

Because the inflation factor in the GWAS data had been modest ($\lambda = 1.078$, $\lambda_{1,000} = 1.045$), we did not adjust the analysis of the GWAS data. We obtained overall significance levels by combining the estimates from the GWAS (unadjusted) and the iCOGS replication (adjusted for principle components 1-6) using a fixed effects meta-analysis, to derive a 1df test, calculating an I^2 statistic to evaluate heterogeneity between the GWAS and replication. We used a threshold of 5×10^{-8} to denote genome-wide significance. Of note we had previously examined the loci at 1q24.1 and 4q24 in our first GWAS replication experiment³: whilst there had been evidence of replication of these signals via Taqman genotyping of 565 cases and 1,758 controls, these associations had failed to achieve genome-wide significance.

We also computed genotype-specific ORs (2 df) for the iCOGS replication, adjusted for principle components 1-6 (Supplementary Table 2). We assessed each SNP for dose response ($P_{\text{heterogeneity}}$) in the replication series by comparing 1-d.f. and 2-d.f. logistic regression models, both adjusted for principle components 1-6, using a likelihood ratio test (evaluated using a significance threshold of $P_{\text{heterogeneity}} < 0.006$ to account for 9 tests). We examined for statistical interaction between TGCT predisposition SNPs (nine new SNPs and the previously published eight SNPs) by evaluating the effect of adding an interaction term to the regression model, adjusted for stage, using a likelihood ratio test (using a significance threshold of $P < 0.0003$ to account for 144 tests).

We assessed for modification of the odds ratios by covariate phenotype and risk factors using a case-only analysis incorporating both GWAS and iCOGS replication data (unadjusted). We evaluated the effect of age on SNP genotype via polytomous regression: we divided age at diagnosis into 5 ordinal categories and fitted maximum-likelihood multinomial logistic models, executed using the `mlogit` command in Stata12 (using a significance threshold of $P < 0.006$ to account for 9 tests, Supplementary Tables 3, 4, 5).

We identified variants from within each LD block reported within the 1000 genomes project ($r^2 > 0.8$ and $< 200\text{kb}$ from sentinel SNP) and used HaploReg and ENCODE data to apply functional annotations relevant to the regulation of transcription: (i) whether the variant lies in a region in which modification of histone proteins is suggestive of enhancer and other regulatory activity (*H3K4Me1* and *H3K27A* histone modification) or promoter activity (*H3K4Me3* histone modification), (ii) whether the variant lies in a region where the chromatin is hypersensitive to cutting by the DNase enzyme (suggestive of regulatory region), (iii) whether the variant lies in a region of binding of transcription factor proteins

(as assayed by chromatin immunoprecipitation with antibodies specific to the transcription factor followed by sequencing of the precipitated DNA (ChIP-seq)), (iv) whether the variant affects a specific regulatory motif, as evaluated from position weighted matrices assembled from TRANSFAC, JASPAR and protein-binding microarray experiments (Supplementary Table 6)^{45,46}.

We investigated associations of the nine TGCT-associated SNPs with gene expression using GENEVAR, which includes gene expression profiling and genotypic data from four datasets: (i) lymphoblastoid cell lines (LCL) from 726 HapMap individuals adipose (HapMap3), (ii) LCL and skin collected from 856 healthy female twins of the MuTHER resource (Muther pilot Twin 1), (iii) adipose, LCL and skin derived from a subset of 160 MuTHER healthy female twins (Muther pilotTwin2) and (iv) fibroblast, LCL and T-cells derived from umbilical cords of 75 Geneva GenCord individuals (Gencord). Spearman Rank Correlation between normalized gene expression levels and the count of one of the alleles of the SNP (0, 1 or 2) was analysed with significance assessed by permutation (10,000 permutations). Expression of genes within 500kB of the sentinel SNP was examined (using a significance threshold of $P < 0.00001$ to account for 440 tests, Supplementary Table 7).

LD matrices between SNPs reported in HapMap were based on Data Release 27/phaseII + III (Feb 2009) on NCBI B36 assembly, dbSNP b126, viewed using Haploview software (v4.2) and plotted using SNAP. LD blocks were evaluated using the HapMap recombination rates (cM/Mb) and defined using the Oxford recombination hotspots⁴⁷. GWAS associations for other diseases were evaluated using the NHGRI (National Human Genome Research Institute) Catalog of Published Genome-Wide Association Studies. All genomic references are based on NCBI Build 36. Analyses were performed using R (v2.6), Stata12 (State College) and PLINK (v1.07) software.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

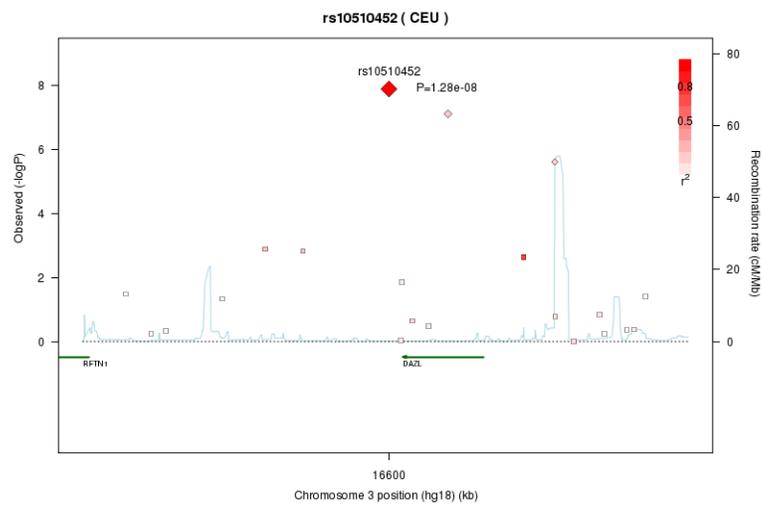
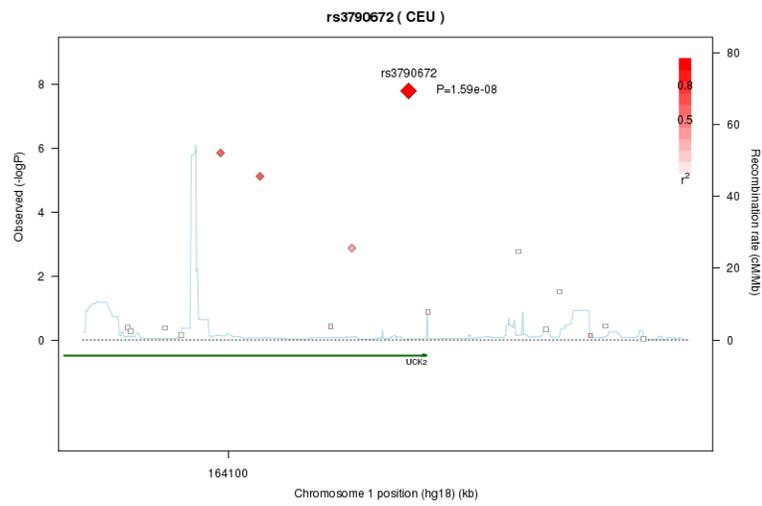
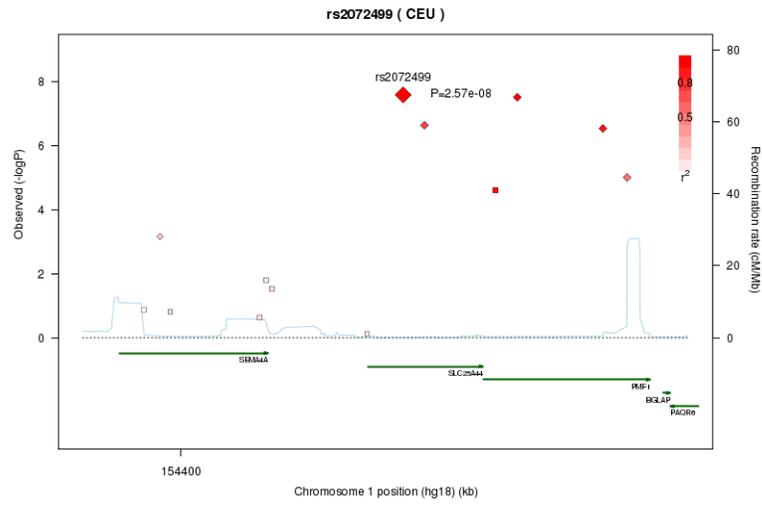
We thank subjects with TGCT and the clinicians involved in their care for participation in this study. We also thank M. Warren Perry, D. Dudakia, J. Pugh, R. Linger, J. Marke, D. Hughes and D. Pernet for recruitment of subjects and database entry for the TGCT collections and the UKGPCS study teams for recruitment of the UKGPCS controls. We acknowledge National Health Service funding to the National Institute for Health Research Biomedical Research Centre. The COGS research initiative, leading to these results has received support from the European Community's Seventh Framework Programme under grant agreement n° 223175 (HEALTH-F2-2009-223175), Cancer Research UK (Grants C1287/A10710 and C5047/A7357) and Prostate Action. This study makes use of data generated by the Wellcome Trust Case Control Consortium (WTCCC) 2. A full list of the investigators who contributed to the generation of the data is available from the WTCCC website. D.F.E. is a Principal Research Fellow of Cancer Research UK. The initial GWAS study was supported by the Institute of Cancer Research, Cancer Research UK and the Wellcome Trust. This study was supported by the Institute of Cancer Research and Movember.

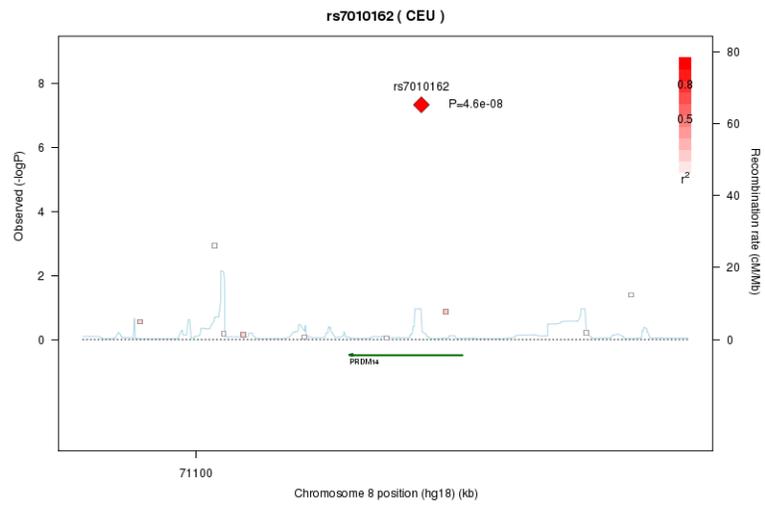
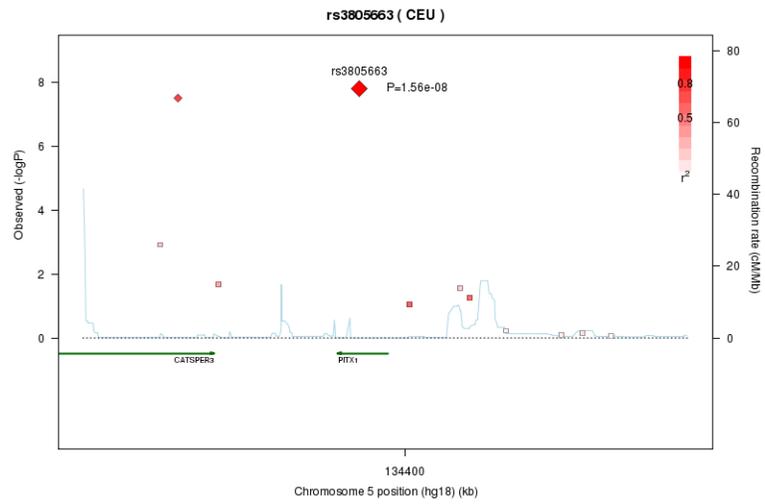
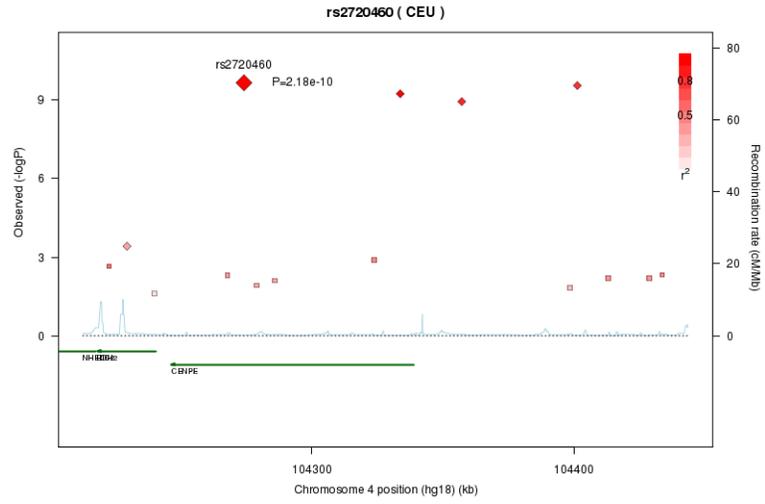
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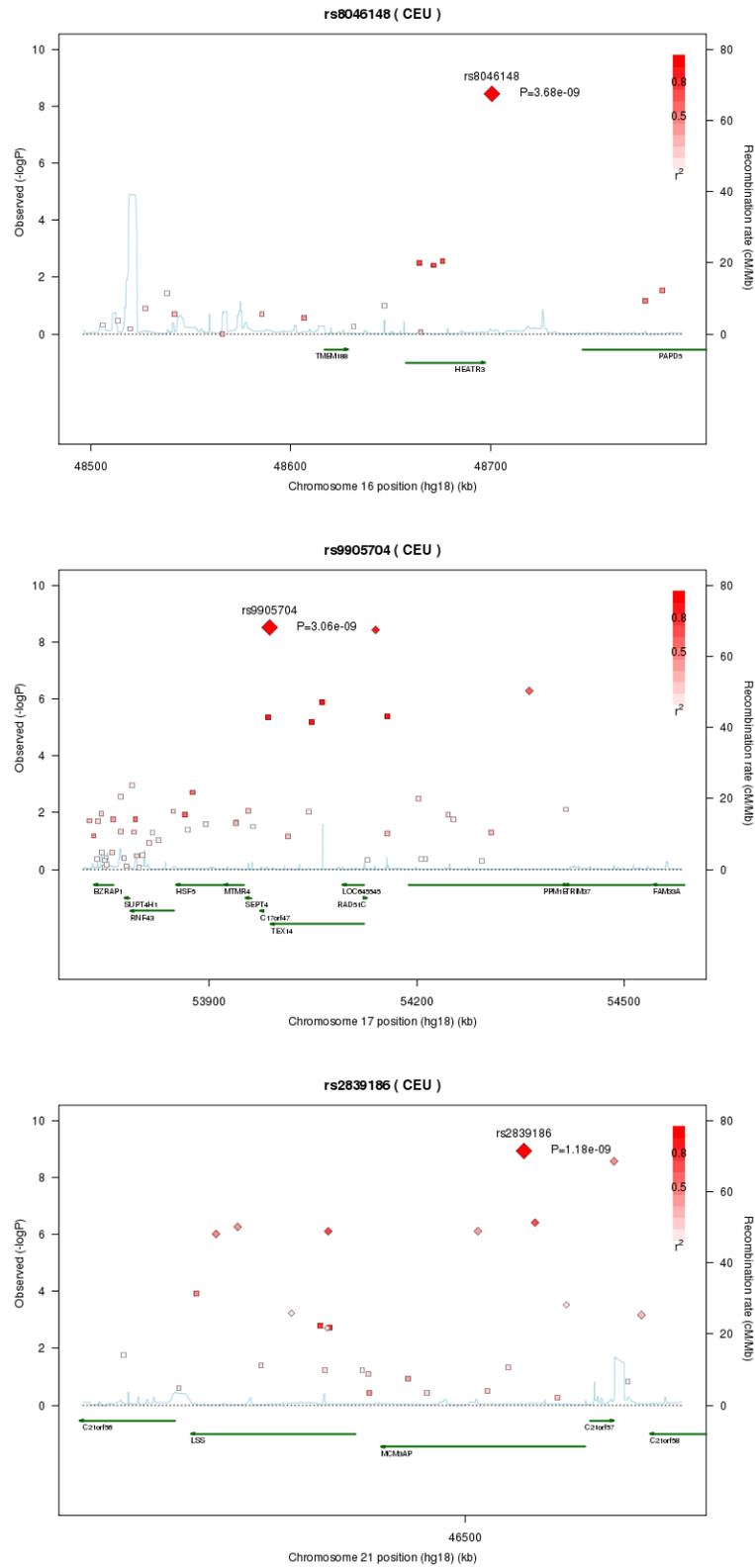


Figure 1. Regional plots of the nine new TGCT loci (a-i)

Plots show the genomic regions of association with TGCT on chromosome (a), 1q22 (b) 1q24.1, (c) 3p24.3, (d) 4q24, (e) 5q31.1, (f) 8q13.3, (g) 16q12.1, (h) 17q22 and (i) 21q22.3. Shown by **diamonds** are the $-\log_{10}$ association P values of SNPs in 986 cases and 4,946 controls from the GWAS, combined with 1,064 cases of TGCT and 10,082 controls from the iCOGs replication. Shown by squares are SNPs genotyped in the GWAS alone. The intensity of red shading indicates the strength of LD with the index SNP (labeled). Also shown are the SNP build 36 coordinates in kilobases (kb), recombination rates in centimorgans (cM) per megabase (Mb) (in blue) and the genes in the region (in green).

Table 1

Summary results for nine SNPs showing genome-wide association ($P < 5 \times 10^{-8}$) in the combined analysis.

SNP ¹	Location ²	Alleles ³	RAF ⁴	GWAS		iCOGS replication		Combined
				OR ⁵ (95% CI)	P _{trend} ⁶	adjusted OR ⁷ (95% CI)	P _{trend} ⁶	P _{meta} ⁸
rs2072499	154,436,234 1q22	G/A	0.35	1.24 (1.12-1.37)	1.84×10 ⁻⁵	1.19 (1.08-1.3)	2.98×10 ⁻⁴	2.57×10 ⁻⁸
rs3790672	164,140,016 1q24.1	C/T	0.28	1.26 (1.14-1.4)	1.10×10 ⁻⁵	1.20 (1.09-1.33)	2.92×10 ⁻⁴	1.59×10 ⁻⁸
rs10510452	16,600,052 3p24.3	A/G	0.69	1.24 (1.12-1.39)	7.06×10 ⁻⁵	1.24 (1.12-1.37)	4.69×10 ⁻⁵	1.28×10 ⁻⁸
rs2720460	104,274,135 4q24	A/G	0.61	1.28 (1.16-1.42)	2.25×10 ⁻⁶	1.24 (1.12-1.36)	1.97×10 ⁻⁵	2.18×10 ⁻¹⁰
rs3805663	134,394,099 5q31.1	T/C	0.63	1.20 (1.08-1.33)	4.86×10 ⁻⁴	1.25 (1.13-1.38)	7.14×10 ⁻⁶	1.56×10 ⁻⁸
rs7010162	71,139,059 8q13.3	G/A	0.61	1.21 (1.09-1.34)	2.03×10 ⁻⁴	1.22 (1.11-1.34)	6.05×10 ⁻⁵	4.60×10 ⁻⁸
rs8046148	48,700,445 16q12.1	G/A	0.79	1.29 (1.13-1.46)	1.03×10 ⁻⁴	1.32 (1.17-1.48)	8.75×10 ⁻⁶	3.68×10 ⁻⁹
rs9905704	53,987,542 17q22	T/G	0.67	1.29 (1.16-1.43)	3.74×10 ⁻⁶	1.21 (1.1-1.34)	1.50×10 ⁻⁴	3.06×10 ⁻⁹
rs2839186	46,514,496 21q22.3	T/C	0.46	1.20 (1.09-1.32)	3.01×10 ⁻⁴	1.26 (1.15-1.38)	7.43×10 ⁻⁷	1.18×10 ⁻⁹

¹ dbSNP rs number

² Build 36 position, chromosomal region

³ Risk/non-risk associated alleles.

⁴ RAF: Frequency of the risk allele

⁵ OR: per allele odds ratio

⁶ P_{trend}: p-value for trend, via logistic regression

⁷ Adjusted OR: per allele odds ratio, adjusted for six principle components

⁸ P_{meta}: P-value for fixed effects meta-analysis