



Rothman, N; Garcia-Closas, M; Chatterjee, N; Malats, N; Wu, X; Figueroa, JD; Real, FX; Van Den Berg, D; Matullo, G; Baris, D; Thun, M; Kiemeny, LA; Vineis, P; De Vivo, I; Albanes, D; Purdue, MP; Rafnar, T; Hildebrandt, MA; Kiltie, AE; Cussenot, O; Golka, K; Kumar, R; Taylor, JA; Mayordomo, JI; Jacobs, KB; Kogevinas, M; Hutchinson, A; Wang, Z; Fu, YP; Prokunina-Olsson, L; Burdett, L; Yeager, M; Wheeler, W; Tardn, A; Serra, C; Carrato, A; Garca-Closas, R; Lloreta, J; Johnson, A; Schwenn, M; Karagas, MR; Schned, A; Andriole, G. Andriole G Jr, Grubb R 3rd, Black A, Jacobs, EJ; Diver, WR; Gapstur, SM; Weinstein, SJ; Virtamo, J; Cortessis, VK; Gago-Dominguez, M; Pike, MC; Stern, MC; Yuan, JM; Hunter, DJ; McGrath, M; Dinney, CP; Czerniak, B; Chen, M; Yang, H; Vermeulen, SH; Aben, KK; Witjes, JA; Makkinje, RR; Sulem, P; Besenbacher, S; Stefansson, K; Riboli, E; Brennan, P; Panico, S; Navarro, C; Allen, NE; Bueno-de-Mesquita, HB; Trichopoulos, D; Caporaso, N; Landi, MT; Canzian, F; Ljungberg, B; Tjonneland, A; Clavel-Chapelon, F; Bishop, DT; Teo, MT; Knowles, MA; Guarrera, S; Polidoro, S; Ricceri, F; Sacerdote, C; Allione, A; Cancel-Tassin, G; Selinski, S; Hengstler, JG; Dietrich, H; Fletcher, T; Rudnai, P; Gurzau, E; Koppova, K; Bolick, SC; Godfrey, A; Xu, Z; Sanz-Velez, JI; D Garca-Prats, M; Sanchez, M; Valdivia, G; Porru, S; Benhamou, S; Hoover, RN; Fraumeni, JF Jr; Silverman, DT; Chanock, SJ (2010) A multi-stage genome-wide association study of bladder cancer identifies multiple susceptibility loci. *Nature genetics*, 42 (11). pp. 978-84. ISSN 1061-4036 DOI: <https://doi.org/10.1038/ng.687>

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A multi-stage genome-wide association study of bladder cancer identifies multiple susceptibility loci

Nathaniel Rothman^{1,67}, Montserrat Garcia-Closas^{1,67}, Nilanjan Chatterjee^{1,67}, Nuria Malats^{2,67}, Xifeng Wu^{3,67}, Jonine Figueroa^{1,68}, Francisco X Real^{2,4,68}, David Van Den Berg^{5,68}, Giuseppe Matullo^{6,7,68}, Dalsu Baris^{1,68}, Michael Thun^{8,68}, Lambertus A Kiemeneij^{9,10,11,68}, Paolo Vineis^{12,7,68}, Immaculata De Vivo^{13,68}, Demetrius Albanes^{1,68}, Mark P Purdue^{1,68}, Thorunn Rafnar^{14,68}, Michelle A T Hildebrandt^{3,68}, Anne E Kiltie^{15,68}, Olivier Cussenot^{16,17,68}, Klaus Golka^{18,68}, Rajiv Kumar^{19,68}, Jack A Taylor^{20,68}, Jose I Mayordomo^{21,68}, Kevin B Jacobs^{22,68}, Manolis Kogevinas^{23,24,25,26,68}, Amy Hutchinson²², Zhaoming Wang²², Yi-Ping Fu¹, Ludmila Prokunina-Olsson¹, Laurie Burdette²², Meredith Yeager²², William Wheeler²⁷, Adonina Tardón^{25,28}, Consol Serra²⁹, Alfredo Carrato³⁰, Reina García-Closas³¹, Josep Lloreta³², Alison Johnson³³, Molly Schwenn³⁴, Margaret R Karagas³⁵, Alan Schned³⁵, Gerald Andriole Jr³⁶, Robert Grubb III³⁶, Amanda Black¹, Eric J Jacobs⁸, W Ryan Diver⁸, Susan M Gapstur⁸, Stephanie J Weinstein¹, Jarmo Virtamo³⁷, Victoria K Cortessis⁴, Manuela Gago-Dominguez⁴, Malcolm C Pike^{4,38}, Mariana C Stern⁴, Jian-Min Yuan³⁹, David Hunter⁴⁰, Monica McGrath⁴⁰, Colin P Dinney⁴¹, Bogdan Czerniak⁴², Meng Chen³, Hushan Yang³, Sita H Vermeulen^{43,9}, Katja K Aben^{9,10}, J Alfred Witjes¹¹, Remco R Makkinje⁴³, Patrick Sulem¹⁴, Soren Besenbacher¹⁴, Kari Stefansson^{14,44}, Elio Riboli¹², Paul Brennan⁴⁵, Salvatore Panico⁴⁶, Carmen Navarro^{47,25}, Naomi E Allen⁴⁸, H Bas Bueno-de-Mesquita⁴⁹, Dimitrios Trichopoulos^{50,51}, Neil Caporaso¹, Maria Teresa Landi¹, Federico Canzian⁵², Borje Ljungberg⁵³, Anne Tjønneland⁵⁴, Francoise Clavel-Chapelon⁵⁵, David T Bishop⁵⁶, Mark T W Teo⁵⁶, Margaret A Knowles⁵⁶, Simonetta Guarrera⁷, Silvia Polidoro⁷, Fulvio Ricceri^{6,7}, Carlotta Sacerdote^{7,57}, Alessandra Allione⁷, Geraldine Cancel-Tassin¹⁷, Silvia Selinski¹⁸, Jan G Hengstler¹⁸, Holger Dietrich⁵⁸, Tony Fletcher⁵⁹, Peter Rudnai⁶⁰, Eugen Gurzau⁶¹, Kvetoslava Koppova⁶², Sophia C E Bolick²⁰, Ashley Godfrey²⁰, Zongli Xu²⁰, José I Sanz-Velez⁶³, María D García-Prats⁶³, Manuel Sanchez²¹, Gabriel

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Correspondence should be addressed to: Montserrat Garcia-Closas, Section of Epidemiology, Institute of Cancer Research, 15 Cotswold Rd, Belmont, Sutton, Surrey SM2 5NG, United Kingdom, Montse.GarciaClosas@icr.ac.uk.

⁶⁷These authors contributed equally to this work.

⁶⁸These authors are the designated representatives of the participating studies.

⁶⁹These authors jointly directed this work.

Author Contributions

N.R., M.G-C., N.C., J.F., D.T.S., and S.J.C. organized and designed the study.

S.J.C., K.B.J., A.H., Z.W., Y-P.F., J.P-O., L.B., X.W., M.A.T.H., M.C., D.V.D.B., S.G., S.P., R.R.M., I.D.V., T.R., D.T.B., G.C-T., J.G.H., R.K., S.C.E.B., and A.G. conducted and supervised genotyping of samples.

M.G-C., N.C., N.R., K.B.J., M.Y., N.M., D.T.S., and S.J.C. contributed to the design and execution of statistical analysis.

M.G-C., N.R., N.C., N.M., J.F., F.X.R., J.F.F., D.T.S., and S.J.C. wrote the first draft of the manuscript.

N.R., M.G-C., N.M., X.W., J.F., D.V.D.B., F.X.R., G.M., D.B., M.T., L.A.K., P.V., I.D.V., D.A., M.P.P., T.R., M.A.T.H., A.E.K., O.C., K.G., R.K., J.A.T., J.I.M., M.K., A.T., C.S., A.C., R.G-C., J.L., A.J., M.S., M.R.K., A.S., G.A., R.G., A.B., E.J.J., W.R.D., S.M.G., S.J.W., J.V., V.K.C., M.G-D., M.C.P., M.C.S., J.Y., D.H., M.M., C.P.D., B.C., M.C., H.Y., S.H.V., K.K.A., J.A.W., R.R.M., P.S., S.B., K.S., E.R., P.B., S.P., C.N., N.E.A., H.B.B., D.T., N.C., M.T.L., F.C., B.L., A.T., F.C-C., D.T.B., M.T.W.T., M.A.K., S.G., S.P., F.R., C.S., A.A., G.C-T., S.S., J.G.H., H.D., T.F., P.R., E.G., K.K., S.C.E.B., A.G., Z.X., J.I.S-V., M.D.G-P., M.S., G.V., S.P., S.B., R.N.H., J.F.F., D.T.S., and S.J.C. conducted the epidemiologic studies and contributed samples to the bladder cancer GWAS and/or replication.

All authors contributed to the writing of the manuscript.

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Valdivia²¹, Stefano Porru⁶⁴, Simone Benhamou^{65,66}, Robert N Hoover¹, Joseph F Fraumeni Jr.¹, Debra T Silverman^{1,69}, and Stephen J Chanock^{1,69}

¹Division of Cancer Epidemiology and Genetics, National Cancer Institute, Bethesda, Maryland, USA ²Spanish National Cancer Research Centre, Madrid, Spain ³Department of Epidemiology, The University of Texas M.D. Anderson Cancer Center, Houston, Texas, USA ⁴Departament de Ciències Experimentals i de la Salut, Universitat Pompeu Fabra, Barcelona, Spain ⁵Department of Preventive Medicine, Keck School of Medicine, University of Southern California, Los Angeles, California, USA ⁶Department of Genetics, Biology and Biochemistry, University of Torino, Torino, Italy ⁷Human Genetics Foundation (HuGeF), Torino, Italy ⁸Epidemiology Research Program, American Cancer Society, Atlanta, Georgia, USA ⁹Department of Epidemiology, Biostatistics and Health Technology Assessment, Radboud University Nijmegen Medical Centre, Nijmegen, The Netherlands ¹⁰Comprehensive Cancer Center East, Nijmegen, The Netherlands ¹¹Department of Urology, Radboud University Nijmegen Medical Centre, Nijmegen, The Netherlands ¹²Imperial College London, London, UK ¹³Channing Laboratory, Department of Medicine, Brigham and Women's Hospital, Boston, Massachusetts, USA ¹⁴deCODE Genetics, Reykjavik, Iceland ¹⁵Gray Institute for Radiation Oncology and Biology, University of Oxford, Oxford, UK ¹⁶Department of Urology, Assistance Publique-Hôpitaux de Paris, Tenon Hospital, Paris, France ¹⁷Centre de Recherche sur les Pathologies Prostatiques, Paris, France ¹⁸Leibniz Research Centre for Working Environment and Human Factors, Dortmund, Germany ¹⁹Division of Molecular Genetic Epidemiology, German Cancer Research Center, Heidelberg, Germany ²⁰National Institute of Environmental Health Sciences, Research Triangle Park, North Carolina, USA ²¹University of Zaragoza, Zaragoza, Spain ²²Core Genotype Facility, SAIC-Frederick, Inc., NCI-Frederick, Frederick, Maryland, USA ²³Centre for Research in Environmental Epidemiology (CREAL), Barcelona, Spain ²⁴Municipal Institute of Medical Research, Barcelona, Spain ²⁵CIBER Epidemiología y Salud Pública (CIBERESP), Barcelona, Spain ²⁶National School of Public Health, Athens, Greece ²⁷Information Management Services, Inc., Rockville, Maryland, USA ²⁸Instituto Universitario de Oncología, Universidad de Oviedo, Oviedo, Spain ²⁹Universitat Pompeu Fabra, Barcelona, Spain ³⁰Ramón y Cajal University Hospital, Madrid, Spain ³¹Unidad de Investigación, Hospital Universitario de Canarias, La Laguna, Spain ³²Hospital del Mar-IMIM, Univesitat Pompeu Fabra, Barcelona, Spain ³³Vermont Cancer Registry, Burlington, Vermont, USA ³⁴Maine Cancer Registry, Augusta, Maine, USA ³⁵Dartmouth Medical School, Hanover, New Hampshire, USA ³⁶Department of Urology, Washington University School of Medicine, St. Louis, Missouri, USA ³⁷National Institute for Health and Welfare, Helsinki, Finland ³⁸Department of Epidemiology and Biostatistics, Memorial Sloan-Kettering Cancer Center, New York, USA ³⁹Masonic Cancer Center, University of Minnesota, Minneapolis, Minnesota, USA ⁴⁰Department of Epidemiology, Program in Molecular and Genetic Epidemiology, Harvard School of Public Health, Boston, Massachusetts, USA ⁴¹Department of Urology, University of Texas M.D. Anderson Cancer Center, Houston, Texas, USA ⁴²Department of Pathology, University of Texas M.D. Anderson Cancer Center, Houston, Texas, USA ⁴³Department of Human Genetics, Radboud University Nijmegen Medical Centre, Nijmegen, The Netherlands ⁴⁴Faculty of Medicine, University of Iceland, Reykjavik, Iceland ⁴⁵International Agency for Research on Cancer, Lyon, France ⁴⁶Department of Clinical and Experimental Medicine, Federico II University, Naples, Italy ⁴⁷Department of Epidemiology, Murcia Regional Health Council, Murcia, Spain ⁴⁸Cancer Epidemiology Unit, Nuffield Department of Clinical Medicine University of Oxford, Oxford, UK ⁴⁹National Institute for Public Health and the Environment (RIVM), Bilthoven, The Netherlands ⁵⁰Department of Epidemiology, Harvard School of Public Health, Boston, Massachusetts, USA ⁵¹Bureau of Epidemiologic Research, Academy of Athens, Greece ⁵²Genomic Epidemiology Group, German Cancer Research Center, Heidelberg, Germany ⁵³Department of Surgical and Perioperative Sciences, University of Umea, Umea, Sweden ⁵⁴Danish Cancer Society, Institute of Cancer Epidemiology, Copenhagen, Denmark ⁵⁵Centre for Research in Epidemiology and Population Health, Institute Gustave-Roussy, Paris South University, Villejuif, France ⁵⁶Leeds

Institute of Molecular Medicine, St James's University Hospital, University of Leeds, Leeds, UK
⁵⁷Unit of Cancer Epidemiology, University of Torino, Torino, Italy ⁵⁸Department of Urology, Paul Gerhardt Foundation, Wittenberg, Germany ⁵⁹Institute of Hygiene and Tropical Medicine, London, UK ⁶⁰National Institute of Environmental Health, Budapest, Hungary ⁶¹Babes Bolyai University, Environmental Health Center, Cluj-Napoca, Romania ⁶²State Health Institute, Banská Bystrica, Slovakia ⁶³San Jorge University Hospital, Huesca, Spain ⁶⁴Department of Experimental and Applied Medicine, Section of Occupational Medicine and Industrial Hygiene, University of Brescia, Italy ⁶⁵Institut national de la santé et de la recherche médicale, U946, Fondation Jean Dausset-CEPH, Paris, France ⁶⁶Centre national de la recherche scientifique, UMR8200, Institut Gustave-Roussy, Villejuif, France

Abstract

We conducted a multi-stage, genome-wide association study (GWAS) of bladder cancer with a primary scan of 589,299 single nucleotide polymorphisms (SNPs) in 3,532 cases and 5,120 controls of European descent (5 studies) followed by a replication strategy, which included 8,381 cases and 48,275 controls (16 studies). In a combined analysis, we identified three new regions associated with bladder cancer on chromosomes 22q13.1, 19q12 and 2q37.1; rs1014971, ($P=8\times 10^{-12}$) maps to a non-genic region of chromosome 22q13.1; rs8102137 ($P=2\times 10^{-11}$) on 19q12 maps to *CCNE1*; and rs11892031 ($P=1\times 10^{-7}$) maps to the *UGT1A* cluster on 2q37.1. We confirmed four previous GWAS associations on chromosomes 3q28, 4p16.3, 8q24.21 and 8q24.3, validated previous candidate associations for the *GSTM1* deletion ($P=4\times 10^{-11}$) and a tag SNP for *NAT2* acetylation status ($P=4\times 10^{-11}$), as well as demonstrated smoking interactions with both regions. Our findings on common variants associated with bladder cancer risk should provide new insights into mechanisms of carcinogenesis.

Bladder cancer is the fourth most common incident cancer in men¹ and its frequent recurrence requires regular screening and interventions. Cigarette smoking and occupational exposure to aromatic amines have been strongly linked to bladder cancer risk.¹ A family history of bladder cancer is associated with an approximately two-fold increase in risk; however, multiple-cancer families are rare and no high-penetrance genes have been identified to date²⁻⁴. Large meta-analyses of candidate gene studies have provided support for associations between *NAT2* slow acetylation phenotype⁵ (defined by *NAT2* haplotypes) and a common gene deletion of *GSTM1* with bladder cancer risk⁷⁻⁸. Further, gene-environment interactions have been shown for smoking and *NAT2* acetylation, with an increased risk in slow acetylators, apparent only among cigarette smokers⁷⁻⁸.

Previous genome-wide association studies (GWAS) in bladder cancer have identified common variants in four genomic regions on chromosomes 3q289 (*TP63*), 4p16.3 (*TMEM129*, *TACC3-FGFR3*)¹⁰, 8q24.219, and 8q24.311 (*PSCA*) that are associated with risk. Interestingly, the variants on 8q24.21 map to a region centromeric to *MYC* that has been identified in GWAS of breast, colorectal and prostate cancers, as well as chronic lymphocytic leukemia¹²⁻¹⁸. Also, in follow-up analyses, an association with bladder cancer risk has been suggested for variants near the *TERT-CLPTMIL* locus on chromosome 5p15.33, which has also been associated by GWAS with risk for basal cell carcinoma, cutaneous melanoma, lung, brain and pancreatic cancers¹⁹⁻²³. However, the previously reported association with bladder cancer did not achieve genome-wide significance.

We conducted a multi-stage GWAS involving 3,532 cases and 5,120 controls of self-described European descent in stage I, and followed up the most notable signals in two stages of replication (stages IIa/b and III) totaling 8,381 cases and 48,275 controls (Figure 1

and Online Methods). Individuals with scan data in stage I were participants in two case-control studies carried out in Spain and the USA (Maine and Vermont component of the New England Bladder Cancer Study) and three prospective cohort studies in the USA and Finland (see Supplementary Table 1 online for details). Replication analyses in stage II were carried out using existing scan data from two earlier studies. First, we evaluated the 100 most significant SNPs (excluding previously reported loci and SNPs with pairwise $r^2 > 0.8$) in 969 cases and 957 controls from the Texas Bladder Cancer study in the USA (stage IIa)¹¹. Five of these SNPs were further evaluated in a second scan of 1,274 cases and 1,832 controls in The Netherlands (stage IIb)⁹. Three of the five SNPs were included or tagged at a pair-wise $r^2 > 0.8$ in the Dutch scan, and risk associations were confirmed for all three. In stage III, the three SNPs plus a tagging SNP for the *NAT2* acetylation status were evaluated in 6,141 cases and 45,486 controls from 11 case-control and 3 prospective cohort studies in the USA and Europe (see Figure 1 and Supplementary Table 1).

After quality control analysis of genotypes, we combined the data sets in stage I resulting in 589,299 SNPs available for analysis (based on the common SNPs called from both the Illumina Human1M and Human 610-Quad) in 3,532 cases and 5,120 controls (**Online Methods**). A logistic regression model was fit for genotype trend effects (1 d.f.) adjusted for study center, age, sex, smoking status (current, former or never) and DNA source (blood/buccal). The quantile-quantile (Q-Q) plot showed little evidence for inflation of the test statistics as compared to the expected distribution (corrected λ_{1000} subjects=1.021), which minimizes the likelihood of substantial hidden population substructure or differential genotype calling between cases and controls²⁴ (**Online Methods and Supplementary Figure 1**). A Manhattan plot displays the results of the combined GWAS in stage I (Supplementary Figure 2).

Data from the first stage confirm the associations reported with tag SNPs in the four previously identified genomic regions on chromosomes 3q28 (rs710521)⁹, 8q24.21 (rs9642880)⁹, 8q24.3 (rs2294008)¹¹ and 4p16.3 (rs798766)¹⁰ as well as a suggested region in 5p15.33 (rs401681; a neighboring SNP, rs2736098, was also reported but data were not available in our study)¹⁹ (Table 1 and Supplementary Figure 3). Consistent with prior reports⁹⁻¹⁰, rs9642880 on 8q24.21 and rs798766 on 4p16.3 were most strongly associated with tumors of low grade/low risk of progression (Supplementary Table 2). A stronger association with low grade/low risk disease was also suggested for rs401681 on 5p15.33 (Supplementary Table 2). In addition, we used a copy number variation TaqMan assay⁷ to assess the presence of *GSTM1* on 1p13.3 to genotype stage I samples, and confirmed an association with increased bladder cancer risk (Table 1).

In a combined analysis based on case/control counts by genotype and study, we estimated odds ratios (ORs) using logistic regression analyses adjusted for study center. Meta-analyses of estimated ORs adjusted for age, sex, smoking status and DNA source produced comparable point estimates (Supplementary Table 3). Our combined analysis of stages I, II and III identified three novel genomic regions on chromosomes 22q13.1, 19q12 and 2q37.1 that were associated with bladder cancer risk below the threshold for genome-wide significance ($P < 5 \times 10^{-7}$)²⁵ (Table 2 and Supplementary Figure 4 for study and stage specific estimates, Figure 2). We also confirmed a signal below genome-wide significance for rs1495741, which tags the *NAT2* acetylator status²⁶ previously reported as a bladder cancer susceptibility locus on 8p227.8. The new SNP is located approximately 10kb of the 3' end of the gene.

The locus on chromosome 22q13.1, rs1014971 ($P_{trend} = 8.4 \times 10^{-12}$; OR per C allele = 0.88, 95% CI 0.85-0.91), was primarily associated with high-risk tumors (Supplementary Table 2). The locus is located in a non-genic region, approximately 25 kb centromeric of the

catalytic polypeptide-like 3A (*APOBEC3A*) and 64 kb telomeric of the chromobox homolog 6 (*CBX6*). *APOBEC3A* is an apolipoprotein B mRNA editing enzyme that belongs to the cytidine deaminase gene family, which can play a role in the initiation of tumorigenesis by deamination of cytosine (C) to uracil (U)²⁷. *CBX6* is a component of the chromatin – associated polycomb complex involved in transcriptional repression.

In the combined analysis, we observed an association with rs8102137 on chromosome 19q12 ($P_{trend}=1.7\times 10^{-11}$; OR per C allele =1.13, 95%CI 1.09-1.17), which maps to the cyclin E1 gene (*CCNE1*). *CCNE1* is a key member of the cyclin/cyclin-dependent kinase (Cdk)/retinoblastoma protein (pRB) pathway which determines the rates of cell cycle transition from G1 to S phase, and is commonly altered in bladder cancer and other tumors²⁸. Cyclin E1 expression in bladder cancer has been associated with high grade or muscle invasive tumors and poor clinical outcome²⁹. Consistently, rs8102137 was most strongly associated with risk of high grade/high risk tumors (Supplementary Table 2).

A third locus is marked by rs11892031 ($P=1.0\times 10^{-7}$; OR per C allele =0.84, 95%CI 0.79-0.89) on chromosome 2q37.1 and resides in an intronic region of the UDP-glucuronosyltransferase (*UGT*) 1A gene locus, which encodes the *UGT1A* family of proteins. Glucuronidation by UGTs facilitates solubility and removal of substrates such as endo- and xenobiotics (including carcinogens in tobacco smoke) via bile or urine³⁰. Genetic variation in *UGT1A* has been associated with predisposition to severe gastrointestinal toxicity of the anticancer drug irinotecan³¹. The *UGT1A* locus is represented by at least nine highly homologous transcripts, collectively known as UGTs, generated by alternative splicing. Tissue-specific loss or decreased expression of UGTs has been associated with several gastrointestinal cancers and bladder cancer³²⁻³⁴, as well as experimentally induced bladder cancer in animal models³⁵.

Previously, a promising signal in the *CLPTMIL-TERT* locus on chromosome 5p15.33 was reported in a region in which common variants have been associated with multiple cancers in recent GWAS¹⁹⁻²³. In addition, rare mutations in *TERT* have been linked to dyskeratosis congenita (a bone marrow failure syndrome), idiopathic pulmonary fibrosis, acute myelogenous leukemia and chronic lymphocytic leukemia³⁶⁻³⁹. In the first stage of this GWAS, we observed a moderately significant effect for rs401681 ($P=2.9\times 10^{-3}$), which was at genome-wide significance when combined with the Rafnar et al. data ($P=5.0\times 10^{-7}$; OR per C allele 1.11, 95% CI 1.07-1.16) (Table 1, Supplementary Figure 3).

The risk associated with *GSTM1* and *NAT2* varied in strength across categories of cigarette smoking, whereas genotype risk associations by smoking categories were of similar magnitude for the eight susceptibility loci identified by GWAS (Supplementary Table 4). In a combined analysis, the risk association with *GSTM1* deletion was strongest in never smokers (OR=1.75, 95%CI=1.44-2.13), and progressively weaker in former (OR=1.55, 95%CI=1.35-1.78) and current smokers (OR=1.25, 95%CI=1.07-1.46; $P_{interaction}=0.008$ for current vs. never smokers; Table 3). The stronger association of the *GSTM1* deletion among non-smokers is a novel observation that was not evident in previous case-only meta-analyses⁷. rs1495741 located on the 3' end of *NAT2* is a marker of the *NAT2* phenotype associated with bladder cancer risk²⁶. The rs1495741 GG genotype marking the slow acetylation phenotype, compared to the combined AG/AA genotypes corresponding to the intermediate/rapid acetylation phenotypes, showed a highly significant ($P=5.5\times 10^{-7}$) association with increased bladder cancer risk that was limited to cigarette smokers (OR=1.24, 95% CI=1.16-1.32 $P=4.3\times 10^{-11}$; $P_{interaction}=6.3\times 10^{-5}$) (Supplementary Figure 5 and Supplementary Table 3). This interaction is consistent with the role of *NAT2* in the detoxification of bladder carcinogens such as aromatic amines from tobacco smoke.

Our three-stage study had adequate power to detect variants of moderate effect sizes over a range of common allele frequencies. For the newly discovered SNP markers, the power to detect the observed associations at a level of genome-wide significance was at 54%, 30%, 30% and 6% for rs104971, rs1495741, rs8102137 and rs11892031, respectively. In light of the limited power to discover SNPs with modest effect sizes, additional loci with similar effect sizes will likely be identified with larger scale GWAS. Based on a recent estimator⁴⁰ that incorporates novel and previously reported loci together, we estimate that approximately two dozen additional bladder cancer susceptibility SNP markers of similar magnitude and frequencies might be discovered. Future studies should be powered with adequate sample size to detect additional variants.

With the exception of the *GSTM1* deletion, relative risk estimates for novel loci are based on associations using tag SNPs, which most likely underestimate the association with biologically important alleles. Accordingly, further studies are needed to define the functional variants and the clinical utility of risk models that combine genetic markers with epidemiologic risk factors for bladder cancer (i.e. smoking, occupational and environmental exposures, family history). Our combined analysis of 12,254 individuals with bladder cancer and 53,395 controls has uncovered three new genomic regions associated with bladder cancer risk. Fine-mapping studies of these three regions are needed to identify candidate variants for functional studies that should shed light into biological mechanisms for the associations reported through GWAS. This knowledge could establish the foundation for developing improved preventive, diagnostic and/or therapeutic approaches.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Appendix

Online Methods

Study Participants

Participants were drawn from 21 studies (Supplementary Table 1). For stage I, cases were defined as histologically confirmed primary carcinoma of the urinary bladder including carcinoma *in situ* (ICD-0-2 topography codes C67.0-C67.9 or ICD9 codes 188.1-188.9). Each participating study obtained informed consent from study participants and approval from its Institutional Review Board (IRB) for this study. For stage I only, participating studies obtained institutional certification permitting data sharing in accordance with the NIH Policy for Sharing of Data Obtained in NIH Supported or Conducted Genome-Wide Association Studies (GWAS).

Genotyping and Quality Control

For stage I, genome-wide genotyping was conducted using three chips, SBCS (HumanHap 1 Million), NEBC-ME/VT (Human Hap 610-Quad), ATBC, CPS-II and PLCO (cases)

(Human Hap 610) and controls from CGEMS/GEI for PLCO (Human Hap 550-r equivalent). DNA samples were selected for genotyping based on pre-genotyping quality control measures performed for GWAS at the Core Genotyping Facility of the NCI 4,089 blood samples and 2,813 buccal samples were analyzed. Repeat genotyping was performed on 38 blood samples (19 cases and 19 controls) and 10 buccal samples (2 cases and 8 controls) on Illumina 1M chips after suitable metrics identified performance issues. Cancer free controls (N=2003) were previously scanned in CGEMS18 and a lung cancer GWAS21.

Genotype clusters were estimated with samples by study with preliminary completion rates greater than 98% per individual study (namely SBCS, NEBC-ME/VT, PLCO, ATBC and CPS-II). Genotypes for the analytical build were based on study specific clustering. SNP assays with locus call rates lower than 90% were excluded.

SNPs with extreme departures from Hardy-Weinberg proportions ($P < 1 \times 10^{-7}$) were excluded from the association analysis due to the increased likelihood of spurious associations due to problematic assays or genotyping calling.⁴² Additional participants were excluded based on: 1) completion rates lower than 94-96% (n=203 samples); 2) heterozygosity of less than 22% or >35% (n=12); 3) inter-study unexpected duplicates (n=5); 4) phenotype exclusions (due to ineligibility or incomplete information) (n=94).

Assessment of population structure of study participants was performed with STRUCTURE⁴³ by seeding the analysis with founder genotypes from three HapMap populations (Phase I and II build 26).⁴⁴ A set of 12,898 SNPs with extremely low pair-wise correlation ($r^2 < 0.004$) was selected for this analysis.⁴⁵⁻⁴⁷ A total of 55 participants (43 cases and 12 controls) were estimated to have less than 85% HapMap CEU admixture (Supplementary Figure 6). Principal component analysis (PCA) of scanned subjects (excluding inferred sib and half-sib pairs) was performed with GLU (a similar procedure to EIGENSTRAT)^{45,46} and did not reveal notable eigenvectors. Consequently, a study-specific indicator was used for the stage I analysis⁴⁶.

We estimated the inflation of the test statistic, λ , adjusted to a sample size of 1000 cases/1000 controls as per the method of de Bakker et. al: $\lambda(\text{corrected}) = 1 + (\lambda - 1) \times [n_{\text{case}}^{-1} + n_{\text{cont}}^{-1}] / [2 \times 10^{-3}]$.⁴⁸ The corrected estimated λ_{1000} is 1.021 while the uncorrected λ is 1.086 (Supplementary Figure 1).

Twenty participant pairs were identified as potential relatives based on genotyping sharing in excess of theoretical expectations. A set of 4,546 SNPs were selected (with completion rates >95%, MAF>0.3 and $r^2 < 0.01$ in the three HapMap populations) and used to run PREST⁴⁹ to formally test for cryptic relatedness. 19 unexpected full-sib and 1 parent-child pairs were identified and excluded from PCA (but included in the association analysis). 243 expected duplicates (including 6 triplicates in ATBC) were evaluated and yielded a concordance rate of 99.99%.

The final participant count for stage I analysis was 3,532 cases and 5,120 controls (Supplementary Table 1). The number of SNPs available for association analysis in all studies but SBCS was 589,299. In the SBCS, genotyped with the Infinium HumanHap 1 M chip, after quality control metrics were applied, 1,002,634 SNPs were available and 571,643 overlapped exactly with the 610Quad/550k data.

TaqMan custom genotyping assays (ABI, Foster City, CA) were designed and optimized for 4 SNPs, including the tag SNP for *NAT2*. In an analysis of 1,107 samples from three studies, the comparison of the Illumina calls with the TaqMan assays showed an average concordance rate of 99.4% (range 99.2-99.8%); no shifts from wild type to homozygotes

were observed. The Illumina Infinium cluster plots for the four novel associations, rs1014971, rs8102137, rs11892031 and rs1495741 are shown in Supplementary Figure 7.

Association Analysis

Association analyses for stage I were conducted using logistic regression, adjusted for age (in five-year categories), sex, smoking (current, former or never), DNA source (buccal/ blood) and study. Each SNP genotype was coded as a count of minor alleles, with the exception of X-linked SNPs among men that were coded as 2 if the participant carried the minor allele and 0 if he carried the major allele.⁵⁰ A score test with one degree of freedom was performed on all genetic parameters in each model to determine statistical significance. We assessed heterogeneity in genetic effects across studies using the I^2 statistic. For the inclusion of stage II and III data, we used genotype counts by case-control status and study, and conducted a fixed effects meta-analysis. We also conducted a meta-analysis based on estimates of allelic odds ratio adjusted by age, sex, smoking status, DNA source and study; the estimates did not materially differ from the fixed-effects meta-analysis (Supplementary Table 3).

Polytomous logistic regression was used to obtain estimates of effect for different tumor subtypes. Case-only analyses with tumor type as an outcome were used to test for differences in effect size across subtypes. Models for tumor grade constrained the effect size to increase linearly across levels. Genotype-smoking interactions were assessed using logistic regression for grouped data adjusted by study and including interaction terms. Forest plots by smoking, including summary estimates from fixed effects meta-analyses, are also shown for rs1495741.

Data analysis and management was performed with GLU (Genotyping Library and Utilities version 1.0), a suite of tools available as an open-source application for management, storage and analysis of GWAS data, and STATA.

Estimate of Recombination Hotspots

SequenceLDhot⁵¹ that uses an approximate marginal likelihood method⁵² was used to compute likelihood ratio (LR) statistics for a set of putative hotspots across the region of interest. We sequentially analyzed subsets of 100 controls of European descent (by pooling 5 controls from each study). We used Phasev2.1 to infer the haplotypes as well as background recombination rates. The analysis was repeated with five non-overlapping sets of 100 pooled controls.

Data Access

The CGEMS data portal provides access to individual level data for investigators from certified scientific institutions after approval of their submitted Data Access Request.

URLs:

CGEMS portal: <http://cgems.cancer.gov/>

CGF: <http://cgf.nci.nih.gov/>

GLU: <http://code.google.com/p/glu-genetics/>

EIGENSTRAT: <http://genepath.med.harvard.edu/~reich/EIGENSTRAT.htm>

SNP500Cancer: <http://snp500cancer.nci.nih.gov/>

STRUCTURE: <http://pritch.bsd.uchicago.edu/structure.html>

Tagzilla: <http://tagzilla.nci.nih.gov/>

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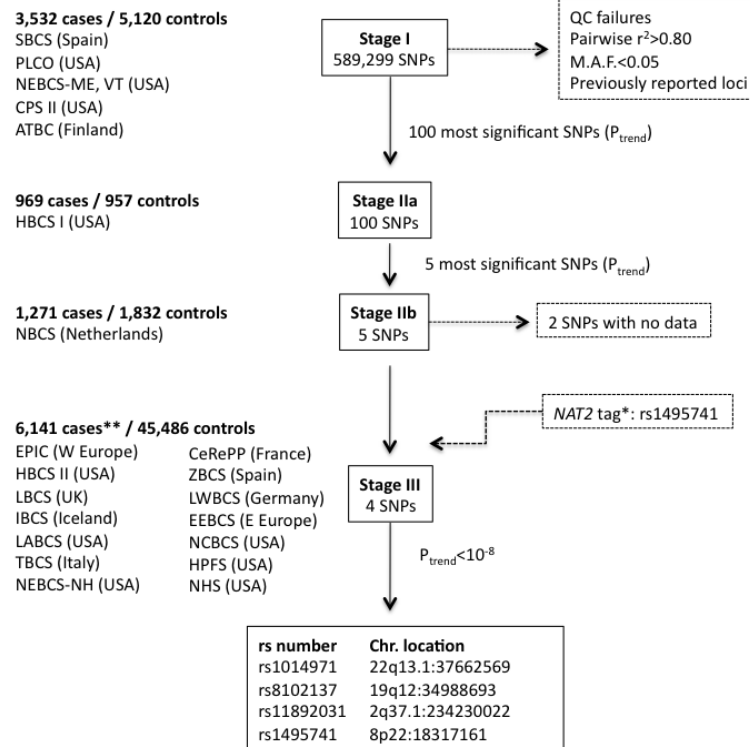


Figure 1. Study design of multi-stage GWAS of bladder cancer

See **Online Methods** and Supplementary Table 1 for details of study designs and sample sizes. *The tag SNP, rs1495741 located 3' of *NAT241* was genotyped in subjects in stage II and III studies as well as on the Illumina bead chips used in stage I. **Includes 338 additional cases from NBSC that were added to the final combined analyses.

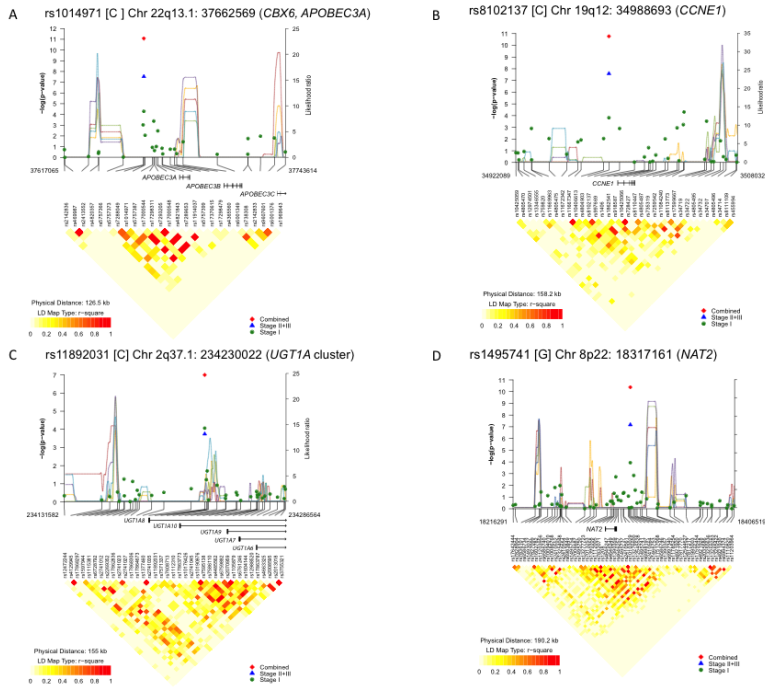


Figure 2. Association results, recombination and linkage disequilibrium plots for four regions on chromosomes 22q13.1, 19q12, 2q37.1 and 8p22
 Results of stage I (green circles), combined stages II and III (blue diamonds) and combined data from the three stages (red diamonds) with P-values for log-additive association results with recombination rates (cm/Mb) based on HapMap phase II data. Pairwise r^2 values based on control populations are displayed at the bottom for all SNPs included in the GWAS analysis. **Panel A** depicts chromosome 22q13.1 region (37,617,065 to 37,743,614). **Panel B** depicts the region of chromosome 19q12 (34,922,089 to 35,080,325). **Panel C** depicts the region of 2q37.1 (234,131,582 to 234,286,564). **Panel D** depicts the region of 8p22 (18,216,291 to 18,406,519). Genomic coordinates are based on NCBI Human Genome Build 36.3.

Table 1
Previously reported genetic variants associated with bladder cancer risk

Results of meta-analyses of allelic OR estimates for the markers reported to achieve genome-wide significance²⁵. Studies in Kiemeny et al. 20089 include: NBCS, LBCS, IBCS, TBCS, Sweden, Belgium, EEBCS, BBBCS, ZBCS. Studies in Wu et al. 200911 include: NBCS, TXBCS1/2, New Hampshire, LBCS, IBCS, TBCS, Sweden, EEBCS, Belgium, BBBCS, ZBCS, MSKCC. Studies in Rafnar et al 200919 include NBCS, IBCS, LBCS, Sweden, TBCS, EEBCS, Belgium, ZBCS, BBBCS.

Marker ^a , risk allele ^b , chr ^c , location ^c and gene ^d	Groups of studies	N ^e	Cases	Controls	Freq ^f	Allelic OR (95%CI) ^g	P value ^h
rs9642880 [T]	Previously reported ⁱ 9	9	3,855	37,985	0.45	1.22 1.15 1.29	7.8E-12
Chr 8q24.21:	Stage I	5	3,525	5,108	0.45	1.21 1.13 1.29	4.6E-08
128787250	Combined	14	7,380	43,093	1.21 1.16 1.27	2.0E-18	
<i>MYC</i>							
rs710521 [A]	Previously reported ⁱ 9	9	3,855	37,985	0.73	1.19 1.12 1.27	1.1E-07
Chr 3q28:	Stage I	5	3,519	5,110	0.72	1.15 1.07 1.25	3.3E-04
191128627	Combined	14	7,374	43,095	1.18 1.12 1.24	1.8E-10	
<i>TP63</i>							
rs2294008 [T]	Previously reported ⁱ 11	13	6,667	39,590	0.46	1.15 1.10 1.20	2.0E-10
Chr 8q24.3:	Stage I	5	3,529	5,115	0.45	1.08 1.01 1.16	2.2E-02
143758933	Combined	18	10,196	44,705	1.13 1.09 1.17	4.4E-11	
<i>PSCA</i>							
rs401681 [C]	Previously reported ⁱ 19	9	4,147	34,988	0.54	1.12 1.06 1.18	5.1E-05
Chr 5p15.33:	Stage I	5	3,526	5,117	0.55	1.11 1.04 1.19	2.9E-03
1375087	Combined	14	7,673	40,105	1.11 1.07 1.16	5.0E-07	
<i>TERT-CLPTMIL</i>							
rs798766 [T]	Previously reported ⁱ 10	11	4,580	45,269	0.19	1.24 1.17 1.32	9.5E-12
Chr 4p16.3:	Stage I	5	3,531	5,118	0.19	1.14 1.05 1.23	2.6E-03
1704037	Combined	16	8,111	50,387	1.20 1.14 1.26	3.9E-13	
<i>TMEM129</i> <i>TACC3-FGFR3</i>							
Deletion Assay	Previously reported ⁱ 7-9	28	5,072	6,466	0.51	1.46 1.35 1.58	1.9E-21

Marker ^a , risk allele ^b , chr ^c , location ^c and gene ^d	Groups of studies	N ^e	Cases	Controls	Freq ^f	Allelic OR (95%CI) ^g	P value ^h		
Chr 1p13.3	Stage I ^j	4	2,480	3,222	0.49	1.49	1.33	1.68	3.7E-11
<i>GSTM1</i>	Combined	32	7,552	9,688	0.53	1.47	1.38	1.57	5.0E-31

^aNCBI dbSNP identifier.

^bRisk allele shown in [].

^cChromosome and NCBI Human Genome Build 36.3 location.

^dGene neighborhood closest to the most notable SNP.

^eN: number of studies

^fRisk allele frequency in control populations.

^gEstimate assuming multiplicative odds model. OR, odds ratio; CI, 95% confidence interval.

^h₁ d.f. trend test.

ⁱSummary estimates differ slightly from previously published because we used a fixed effects meta-analysis.

^jData from SBCS was excluded from stage I because it had been included in the previous meta-analyses published in Garcia-Closas et al. 20057. Data from NEBCS are reported separately (Unpublished data to appear in "GSTM1 null and NAT2 Slow Acetylation Genotypes, Smoking Intensity, and Bladder Cancer Risk: Results from the New England Bladder Cancer Case-Control Study and Meta-Analyses" by Moore LE, Barris D, Figueroa J, Garcia-Closas M, Karagas M, Schwenn M, Johnson A, Lubin J, Hein DW, Dagnall C, Colt J, Kita M, Jones M, Schmed A, Cherela S, Chanock S, Cantor K, Silverman D, Rothman N)

Table 2

Novel SNPs identified in a multi-stage GWAS of bladder cancer

Results of the meta-analysis of genotype counts included in combined stages I, II and III. The initial scan results were adjusted by age, gender, smoking status, study site and DNA source.

Marker ^a , minor allele ^b , chr ^c , location ^c and gene ^d	Studies included	N ^e	Cases	Controls	Freq ^f	Allelic OR (95%CI) ^g	P value ^h
rs1014971 [C]	Stage I	5	3,529	5,092	0.38	0.87	0.93
	Stage II and III	15	8,277	47,517	0.37	0.89	0.93
	Combined	20	11,806	52,609	0.38	0.88	0.91
<i>CBX6, APOBEC3A</i>							
rs8102137 [C]	Stage I	5	3,530	5,114	0.33	1.13	1.06
	Stage II and III	15	8,261	47,708	0.32	1.13	1.08
	Combined	20	11,791	52,822	0.33	1.13	1.09
<i>CCNE1</i>							
rs11892031 [C]	Stage I	5	3,524	5,108	0.08	0.79	0.70
	Stage II and III	15	8,284	47,727	0.08	0.86	0.80
	Combined	20	11,808	52,835	0.08	0.84	0.79
<i>UGT1A</i> cluster ^j							
rs1495741 [G]	Stage I	5	3525	5116	0.24	0.86	0.80
	Stage II and III	15	8,279	47,744	0.22	0.87	0.83
	Combined	20	11,804	52,860	0.24	0.87	0.83
18317161							
<i>NAT2</i>							

^aNCBI dbSNP identifier.

^bMinor allele shown in [].

^cChromosome and NCBI Human Genome Build 36.3 location.

^dGene neighborhood closest to the most notable SNP.

^eN: number of studies

^fMinor allele frequency in control populations.

^gEstimate assuming multiplicative odds model. OR, odds ratio; CI, 95% confidence interval.

h_1 d.f. trend test.

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Table 3
Interaction of NAT2 tagSNP (rs1495741) and GSTM1 deletion with cigarette smoking in bladder cancer risk

Results from logistic regression analyses of aggregated data adjusted by study. rs1495741 genotypes were classified according to an established approach based on the phenotype of NAT2 acetylators; AA/AG were collapsed to tag rapid/intermediate acetylation, and GG tags slow acetylation.²⁶

	Cases			Controls			P			
	AA/AG	GG	AA/AG	GG	AA/AG	GG	OR	95% CI	P	Interaction
NAT2 (rs1495741)										
All subjects	3,784	6,915	5,233	8,182	1.15	1.09	1.22	2.9E-07		
By smoking status										
Never smoker	760	1,202	1,679	2,758	0.95	0.85	1.06	3.3E-01	Ref.	
Ever smoker	3,024	5,713	3,554	5,424	1.24	1.16	1.32	4.3E-11	6.3E-05	
Former smoker	1,859	3,455	2,300	3,559	1.20	1.11	1.29	6.8E-06	6.7E-04	
Current smoker	1,165	2,258	1,254	1,865	1.27	1.14	1.40	7.2E-06	1.6E-04	
GSTM1 Del										
Present		Null	Present	Null						
All subjects	1,319	1,995	1,717	1,726	1.47	1.33	1.62	4.4E-14		
By smoking status										
Never smoker	210	346	519	510	1.71	1.38	2.12	6.9E-07	Ref.	
Ever smoker	1,109	1,649	1,198	1,216	1.47	1.30	1.67	2.1E-09	1.1E-01	
Former smoker	564	961	622	653	1.62	1.39	1.89	4.7E-10	6.9E-01	
Current smoker	545	688	576	563	1.19	1.00	1.40	4.5E-02	8.1E-03	