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Inherited coding variants at the CDKN2A locus influence susceptibility to acute lymphoblastic leukaemia in children

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There is increasing evidence from genome-wide association studies for a strong inherited genetic basis of susceptibility to acute lymphoblastic leukaemia (ALL) in children, yet the effects of protein-coding variants on ALL risk have not been systematically evaluated. Here we show a missense variant in CDKN2A associated with the development of ALL at genome-wide significance (rs3731249, $P = 9.4 \times 10^{-23}$, odds ratio = 2.23). Functional studies indicate that this hypomorphic variant results in reduced tumour suppressor function of p16INK4A, increases the susceptibility to leukaemic transformation of haematopoietic progenitor cells, and is preferentially retained in ALL tumour cells. Resequencing the CDKN2A–CDKN2B locus in 2,407 childhood ALL cases reveals 19 additional putative functional germline variants. These results provide direct functional evidence for the influence of inherited genetic variation on ALL risk, highlighting the important and complex roles of CDKN2A–CDKN2B tumour suppressors in leukaemogenesis.
The risk of developing acute lymphoblastic leukaemia (ALL) is highest between 2 and 5 years after birth\(^1,2\), with initiating sentinel somatic genomic lesions (for example, chromosomal translocations) detectable at the time of birth in many cases\(^3,4\). This early disease onset suggests a strong inherited genetic basis for ALL susceptibility, and recent genome-wide association studies (GWAS) have discovered at least six risk loci: ARID5B, IKZF1, CEBPE, PIP4K2A-BMI1, GATA3 and CDKN2A–CDKN2B\(^5–10\). These ALL risk genes are directly involved in haematopoietic stem cell function, lymphocyte differentiation and development, and cell cycle regulation\(^11–13\), several of which are also commonly targeted by somatic genomic lesions. In particular, the CDKN2A–CDKN2B locus is one of the most frequently deleted genomic regions in childhood ALL with focal copy number loss in both B- and T-cell ALL\(^14,16\). The vast majority of variants examined in previous ALL GWAS are intronic or intergenic. Although it is now evident that non-coding variants related to disease traits are significantly over-represented in regulatory DNA and often function as modulators of local or distal gene transcription\(^17,18\), questions also arise whether coding variants within ALL susceptibility genes might confer even greater effects on disease development. Moreover, a large number of low-frequency and rare-coding germline variants have been discovered by exome-sequencing efforts\(^19\), but their contributions to ALL pathogenesis have yet to be examined systematically.

In the present study, we perform an exome-focused GWAS to systematically examine the impact of germline-coding variants on the development of ALL in children of European descent, experimentally explore the functional consequences of the genome-wide significant variant in the CDKN2A gene, and comprehensively characterize coding variation at this locus by targeted resequencing.

**Results**

**Exome-focused GWAS of ALL susceptibility.** In the discovery GWAS, we genotyped 1,773 children with B-ALL and 10,448 non-ALL controls of European descent\(^20,21\) for 247,505 variants using the Illumina Infinium HumanExome array. Three loci with genome-wide significant association signals were observed: ARID5B (10q21.2), IKZF1 (7q11.2) and CDKN2A (9p21.3) (Fig. 1). Non-coding variants rs10821936 in ARID5B and rs4132601 in IKZF1 showed the strongest association (\(P=9.9 \times 10^{-46}\) and \(4.3 \times 10^{-37}\)) the logistic regression test, respectively; Fig. 1 and Supplementary Table 1), confirming previous GWAS findings from our group and others\(^5,6\). No coding variants in ARID5B and IKZF1 were significantly associated with ALL susceptibility. The third genome-wide significant hit was a missense SNP at the CDKN2A locus (rs3731249, \(P=9.4 \times 10^{-23}\), the logistic regression test, Fig. 1, Table 1). The T allele at rs3731249 was over-represented in ALL compared with controls (6.8% versus 3.0%, Table 1), with every copy of the allele conferring 2.3-fold increase in disease risk (95% confidence interval 1.90–2.61). The C-to-T nucleotide substitution at rs3731249 (c.C442T) resulted in an alanine-to-threonine change in amino-acid sequence (p.A148T) for tumour suppressor p16\(^{INK4A}\). This variant also locates in the 3′ untranslated region (3′-UTR) of the p14\(^{ARF}\) transcript, an alternative open reading frame at this locus encoding a different tumour suppressor. Interestingly, previous GWAS had identified an intronic variant in CDKN2A (rs3731217) to be strongly associated with susceptibility to ALL in populations of European descent\(^9\). Genotype correlation between the coding variant rs3731249 and the intronic rs3731217 is exceedingly low (\(r^2<0.01\) in Europeans, Supplementary Fig. 1), and multivariate analyses including both SNPs indicated their independent contribution to ALL risk (Supplementary Table 2). In the replication cohort of 409 childhood ALL cases and 1,599 non-ALL controls of European descent in Denmark, the association signal at rs3731249 was validated (\(P=5.2 \times 10^{-4}\), odds ratio = 1.73 (1.27–2.36), the logistic regression test, Table 1) and this variant also remained significant after adjusting for rs3731217.

**Functional characterization of the rs3731249 variant.** To experimentally evaluate the effects of rs3731249 on ALL leukemogenesis, we directly compared the effect of wildtype versus variant allele p16\(^{INK4A}\) (p.148A versus p.148T) on BCR-ABL1-mediated leukaemic transformation in vitro. We chose mouse haematopoietic progenitor Ba/f3 cell line because it is inherently p16\(^{INK4A}\)-defective due to methylation at the Ink4a-Arf locus\(^22\), and ectopic expression of BCR-ABL1 in Ba/f3 cells efficiently induces exogenous cytokine (interleukin 3 (IL3))-independent proliferation. Over-expression of wild-type p16\(^{INK4A}\) (p.148A) significantly inhibited leukaemic transformation by BCR-ABL1 (Fig. 2a, Supplementary Fig. 2), consistent with its role as a critical tumour suppressor in ALL. In contrast, Ba/f3 cells overexpressing variant p16\(^{INK4A}\) (p.148T) were significantly more susceptible to BCR-ABL1 transformation measured by IL3-independent growth, suggesting that the p.148T variant is likely hypomorphic with reduced tumour suppressor function. In Ba/f3 cells transfected with both variant and wild-type p16\(^{INK4A}\), the relative ratio of the p.148T (variant) to p.148A (wildtype) C2/C0 (log10 axis) were plotted against respective chromosomal position of each SNP (x axis). Gene, symbols were indicated for 3 loci achieving genome-wide significance threshold (\(P<5 \times 10^{-8}\), dashed blue line): ARID5B (10q21.2), IKZF1 (7p12.2) and CDKN2A (9p21.3). Blue dots indicated SNPs within 2M bp of the top ALL susceptibility variants at the ARID5B (rs10821936) and IKZF1 (rs4132601) loci, the red dots indicated SNPs in the 2M-bp region around the novel ALL risk variant rs3731249 in CDKN2A.

**Figure 1 | GWAS results of ALL susceptibility in European Americans.** Association between genotype and ALL was evaluated for 35,802 SNPs in 1,773 ALL cases and 10,448 non-ALL controls. P-values (the logistic regression test, –log10 \(P, y\) axis) were plotted against respective chromosomal position of each SNP (x axis). Gene, symbols were indicated for 3 loci achieving genome-wide significance threshold (\(P<5 \times 10^{-8}\), dashed blue line): ARID5B (10q21.2), IKZF1 (7p12.2) and CDKN2A (9p21.3). Blue dots indicated SNPs within 2M bp of the top ALL susceptibility variants at the ARID5B (rs10821936) and IKZF1 (rs4132601) loci, the red dots indicated SNPs in the 2M-bp region around the novel ALL risk variant rs3731249 in CDKN2A.
CDKN2A gene (p16INK4A and p14ARF) in children with ALL, a higher burden of rare missense variants relative to controls in the NHLBI GO Exome Sequencing Project (ESP), there was a trend for comparing with 4,300 European American individuals from the damaging based on combined annotation dependent depletion23.

To further examine the potential susceptibility to ALL conferred by the rs3731249 variant, we performed targeted resequencing of the CDKN2A locus, the variant p16INK4A, or transfected with control vector was transduced with p16INK4A (p.148T, green) significantly potentiated leukaemic transformation by BCR–ABL1, compared with cells expressing wild-type p16INK4A (p16INK4A, p.148A, blue), consistent with the presence of cytokine IL3 was measured daily as an indicator of leukaemic blasts was determined by comparing the number of sequence reads for each allele.

**Table 1 | Genome-wide significant association and replication of novel coding B-ALL susceptibility variant at CDKN2A locus.**

<table>
<thead>
<tr>
<th>SNP ID</th>
<th>Chr</th>
<th>Position</th>
<th>Gene</th>
<th>Alleles</th>
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<tr>
<td>rs3731249</td>
<td>9</td>
<td>21970916</td>
<td>CDKN2A</td>
<td>C/T</td>
<td>TT</td>
<td>8 (0.45%)</td>
<td>Control (N = 1,773)</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>CT</td>
<td>224 (12.63%)</td>
<td>7 (0.07%)</td>
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<tr>
<td></td>
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<td></td>
<td></td>
<td>CC</td>
<td>1,541 (86.92%)</td>
<td>619 (5.93%)</td>
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</table>

**Discussion**

Encoding three tumour suppressor proteins (p16INK4A, p14ARF and p15INK4B), the CDKN2A–CDKN2B locus at 9p21 is promiscuously associated with tumorigenesis and commonly

![Figure 2 | Functional characterization of ALL risk variant at rs3731249.](image)

(a) Mouse haematopoietic progenitor cell Ba/f3 overexpressing wildtype, variant p16INK4A, or transfected with control vector was transduced with leukaemia oncogenic BCR–ABL1 fusion gene. Cell proliferation in the absence of cytokine IL3 was measured daily as an indicator of leukaemic transformation. Ectopic expression of p16INK4A (p.148 T) in 15 childhood ALL cases with heterozygous germline genotype at this SNP, six exhibited somatic deletion of one copy of CDKN2A, all of which retained the risk allele in tumour cells. Even in cases not affected by somatic copy number loss at this locus, the variant p16INK4A (c.442C) was preferentially transcribed relative to wildtype (c.442C), with allele-biased expression ranging from 61 to 100%, Fig. 2b). Altogether, these results pointed to the possibility that cells carrying the hypomorphic risk allele at rs3731249 might have been enriched during leukaemogenesis.

Targeted resequencing of CDKN2A and CDKN2B in childhood ALL. To comprehensively identify putative functional ALL susceptibility variants at this locus, we resequenced the coding region of the CDKN2A and CDKN2B genes in germline DNA from 2,407 childhood ALL cases (1,450 of which were also included in the discovery GWAS). In addition to rs3731249, we observed another 13 germline exonic variants in tumour suppressors p16INK4A and p14ARF encoded by the CDKN2A gene, 12 of which result in amino-acid sequence changes (Fig. 3, Supplementary Table 3). These missense variants were all singletons, except for the p.D125H variant in p16INK4A, and the p.A121T variant in p14ARF observed in two and five cases, respectively. Five variants were predicted to be damaging based on combined annotation dependent depletion23 (CADD score > 13, Supplementary Table 3), and we did not observe germline insertions or deletions in CDKN2A in our ALL cohort.

Comparing with 4,300 European American individuals from the NHLBI GO Exome Sequencing Project (ESP), there was a trend for a higher burden of rare missense variants in relative to controls the CDKN2A gene (p16INK4A and p14ARF) in children with ALL (0.71% versus 0.23%, P = 0.0045; Fisher’s exact test, Fig. 3). In addition, we identified six germline-coding variants in the adjacent CDKN2B gene in this cohort of children with ALL, although there was no significant over-representation compared with European controls in the ESP cohort (0.83% versus 0.79%, Fig. 3).

![Table 1 | Genome-wide significant association and replication of novel coding B-ALL susceptibility variant at CDKN2A locus.](image)

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targeted by somatic mutation, deletion and/or hypermethylation in various cancers. p16INK4A and p15INK4B are highly homologous inhibitors of cyclin-dependent kinase and function mainly as master regulators of cell cycle entry via the Rb–E2F signalling axis. Although also encoded by the CDKN2A gene, p14ARF utilizes a completely different reading frame with distinct tumour suppression functions by inhibiting MDM2 and activating p53. Suppressed during normal haematopoiesis, p16INK4A and p14ARF expression is activated on oncogenic stimuli (for example, constitutive expression of BCR–ABL1 fusion) to trigger cell cycle exit (senescence) or apoptosis as a means of eliminating oncogene-stressed cells. In fact, the CDKN2A–CDKN2B locus is either bi- or monoallelically deleted in 64% of BCR–ABL1-positive ALL cases and in 32–72% of T- or B-ALL cases without the BCR–ABL1 translocation, suggesting positive selection for cells with defective p16INK4A, p14ARF and p15INK4B (or some combinations thereof) during leukaemogenesis.

The previously reported ALL susceptibility variant rs3731217 is located in a non-coding region downstream of exon 1B (specific for p14ARF), but distal to exon 1z (specific for p16INK4A) of the CDKN2A gene. The germline genotype at this SNP was not associated with overall CDKN2A expression in lymphoblastoid cell lines but transcript-specific analyses may be needed to definitively determine the effects of this variant on p14ARF versus p16INK4A expression. In contrast, the genome-wide significant variant rs3731249 in our current GWAS localizes to exon 2 of CDKN2A. While this exon is shared by both p16INK4A and p14ARF, the C-to-T nucleotide transition causes a missense change for the p16INK4A open reading frame but is in the UTR of the p14ARF, therefore, likely to have a more direct effect on the former. This hypothesis is supported by the fact that haematopoietic progenitor cells (Ba/f3) expressing variant p16INK4A were substantially more susceptible to BCR–ABL1-mediated leukaemic transformation compared with cells with the wild-type protein (Fig. 2a), pointing to rs3731249 as a possible functional variant directly contributing to the association with ALL risk. The structural basis of the hypomorphic effects of the p.A148T variant is unclear, since this residue is not directly involved in binding to CDK4 or CDK6. However, there was evidence that the variant p16INK4A (p.148T) is preferentially retained in the nucleus compared with the wild-type p16INK4A (p.148A), compromising its ability to inhibit CDKs in the cytoplasm. The relative contribution of p16INK4A versus p14ARF to ALL pathogenesis is not unequivocal because somatic deletions at this locus almost always lead to the loss of both genes. Although the rs3731249 variant also results in sequence changes in the 3′-UTR of the p14ARF transcript, bioinformatic prediction did not identify any potential effects on mRNA stability or microRNA binding and no difference was observed in reporter gene transcription under the influence of 3′-UTR containing either the wildtype or variant allele at rs3731249 (Supplementary Fig. 4), suggesting minimal effects of this variant on p14ARF transcription. Finally, rs3731249 is also observed in non-European populations, for example, there was a trend for a higher frequency of the risk allele in African American children with ALL than that in individuals from this racial background in the NHLBI ESP cohort (0.58% in 260 ALL cases versus 0.38% in 2,203 controls), although a much larger sample size is needed to rigorously examine the statistical significance of such differences. It should be noted that we and others previously showed that the

Figure 3 | Targeted resequencing of CDKN2A–CDKN2B locus identified additional germline coding variants in children with ALL. CDKN2A and CDKN2B genes were sequenced using Illumina HiSeq platform following capture-based enrichment of this genomic region in 2,407 ALL cases of European descent. Variants in non-ALL controls were based on publicly available data from the individuals of European descent within the NHLBI Exome Sequencing Project (N = 4,300). Exonic variants are classified as silent or missense (grey or purple solid circles) and are mapped to three distinct open reading frames at this locus: p16INK4A, p14ARF and p15INK4B, for ALL cases (red vertical lines) and non-ALL controls (blue vertical lines), and functional domains are indicated by colour based on Pfam annotation. Each circle represents a unique individual carrying the indicated variant (heterozygous or homozygous), except for variants recurring in more than 10 individuals for which the number in the circle indicates the exact frequency of the observed variant.
non-coding ALL risk variants (rs17756311 and rs3731217) at this locus had much stronger effects in European Americans than in other race/ethnic groups22,23, suggesting potential racial differences in genetic susceptibility to ALL. We subsequently identified additional coding variants in p16INK4A, p14ARF and p15INK4B by resequencing, most of which were rare and were present at a frequency of $<0.046 \%$. Interestingly, rs199886003 is the only variant that is located in the coding region of both p16INK4A and p14ARF, resulting in an alanine-to-threonine change in p14ARF (p.A121T) with synonymous effect on p16INK4A. This is also the most frequent germline missense variant in p14ARF in our cohort and was over-represented in ALL compared with non-ALL controls (0.21% versus 0.046%, respectively, Fig. 3). The substitution of threonine in p14ARF adds a possible glycosylation and phosphorylation site and also introduces a phosphoprotein-binding FHA domain implicated in DNA damage response and cell cycling31. Future studies are warranted to determine the exact consequences of this variant on p14ARF functions. To systemically evaluate the contribution of low-frequency variants to ALL risk, we also performed genome-wide gene-level burden test but did not observe any genome-wide significant associations (Supplementary Table 4). Of the six known ALL risk loci, we noted two coding variants in CEBPE (rs141903485 and rs146580935, Supplementary Table 5) nominally associated with ALL susceptibility.

In conclusion, we comprehensively evaluated exonic genetic variations for association with ALL susceptibility and identified novel coding risk variants at the CDKN2A–CDKN2B locus that may directly affect tumour suppressor functions and potentiate leukaemic transformation. These results provided functional evidence for the influence of inherited genetic variants on ALL leukaemogenesis, further indicating that a continuum of genetic variations in both host and tumour genomes contribute to malignant transformation and cancer risk.

Genotype calls (coded as 0, 1, and 2 for AA, AB and BB genotypes) were determined using the Illumina GenomeStudio Software. For the ALL cases, samples for which genotypes were not called at $<99\%$ SNP on the array failed and were excluded from the analyses. Quality control procedures were performed for both samples and SNPs on the basis of call rate, minor allele frequency (MAF), and Hardy–Weinberg equilibrium (Supplementary Fig. 5). Detailed quality control for the non-ALL controls from the ARIC study was performed at the University of Texas Health Science Center following established protocols.28

We performed principal component analysis of cases and controls in the discovery GWAS to characterize population structure (Supplementary Fig. 6).

**Methods**

**Subjects and samples.** The discovery GWAS consisted of 1,773 childhood B-ALL cases and 10,448 non-ALL controls of European descent (>90% European genetic ancestry) as estimated using STRUCTURE22,23). ALL cases were from the Children’s Oncology Group (COG) AALL0232 study (N = 1,277)5, the COG P9906 protocol (N = 115)52 and St Jude Total Therapy XIIIIB and XV protocols (N = 381)5. Unrelated individuals of European descent from the Atherosclerosis Risk in Communities (ARIC) study20,21,26 were used as non-ALL controls because the prevalence of adult survivors of childhood ALL is less than 1 in 10,000 in the US. The replication series included 409 children with ALL from NOPHO ALL92, ALL2000 and ALL2008 protocols53 and 1,599 unrelated non-ALL controls from Danish Childhood Obesity Biobank study (clinicaltrials.gov: NCT00928473) in Holbæk and at random schools in Zealand, Denmark. ALL cases were selected only on the basis of sample availability, and we did not observe any statistically significant differences in demographic or clinical features of children included versus not included in this genetic study. We elected to focus on individuals of European descent to minimize population stratification.

Germline DNA for cases was extracted from peripheral blood or bone marrow samples obtained during clinical remission (<5% ALL blasts by morphology). This study was approved by the Institutional Review Board at St Jude Children’s Research Hospital and COG member institutions and the Ethics Committee at the Danish Data Protection Agency, Region Zealand and the University Hospital Rigshospitalet, Denmark. Informed consent was obtained from parents, guardians, or patients, as appropriate.

Genotyping and quality control. SNP genotyping was performed in germline DNA using the Illumina Infinium HumanExome Array v1.0 in the discovery GWAS, and using Illumina HumanCoreExome chip for the replication series.

Leukemic transformation assay in Ba/f3 cells. The full-length CDKN2A was purchased from GE Healthcare. The p.A148T variant (rs3731249) was introduced using site-directed mutagenesis (forward primer: 5'-TGGCCGCGATATGGAGGAGGTTGCCCCTCAAG-3', reverse primer: 5'-TCTGAGAGGACCTTCCGTGGC-3'. The PCR product was cloned into the pCI–CDKN2A region on pSp61. Quantitative PCR was used to determine the appropriate capture probe/tissue necessary to efficiently populate an Illumina HiSeq 2000 flowcell for paired-end 2 × 101 bp sequencing. Each sequence pool of 96 samples was demultiplexed, with coverages of >20 × depth across >90% of the targeted regions for nearly all samples. Sequence reads in FASTQ format were filtered and aligned using the Bowtie2 and the ‘-X’ option was set with the default setting. SNP calls were made using the GATK pipeline version 3.1 (ref. 38). We also performed gene-level analyses to evaluate the aggregated effects of low-frequency variants on ALL susceptibility, using the SKAT test27. Missense, stop codon-altering and splice-site variants with MAF $<5\%$ were included. In total, 12,678 genes with at least two variants were tested. R (version 3.0) statistical software was used for all analyses unless indicated otherwise.

CDKN2A–CDKN2B resequencing and rare variant analyses. Germline DNA from 2,407 children with ALL was used to create individual Illumina dual-indexed libraries. These libraries were pooled in sets of 96 and hybridized with a custom version of the Roche NimbleGen SeqCap EZ custom probes to capture the CDKN2A–CDKN2B region on Sp61. Quantitative PCR was used to determine the appropriate capture probe/tissue necessary to efficiently populate an Illumina HiSeq 2000 flowcell for paired-end 2 × 101 bp sequencing. Each sequence pool of 96 samples was demultiplexed, with coverages of >20 × depth across >90% of the targeted regions for nearly all samples. Sequence reads in FASTQ format were filtered and aligned using the Bowtie2 and the ‘-X’ option was set with the default setting. SNP calls were made using the GATK pipeline version 3.1 (ref. 38). We also performed gene-level analyses to evaluate the aggregated effects of low-frequency variants on ALL susceptibility, using the SKAT test27. Missense, stop codon-altering and splice-site variants with MAF $<5\%$ were included. In total, 12,678 genes with at least two variants were tested. R (version 3.0) statistical software was used for all analyses unless indicated otherwise.
For quantitative reverse transcription-PCR (qRT-PCR), total RNA was extracted using the RNeasy Micro kit (Qiagen) according to the manufacturer’s protocol. Total RNA (500 ng) was reverse transcribed into cDNA using oligoT primers and the SuperScript III reverse transcriptase kit (Invitrogen). Quantitative real-time PCR was performed by using ABI Prism 7900HT detection system (Applied Biosystems) with FastStart SYBR Green master mix (Roche). Relative expression was calculated as a ratio of BCIR-ABL1 to Hprt. Primer sequences of BCIR-ABL1 and Hprt were as follows: BCIR-ABL1 (forward: 5'-CTGGCCAACAGATGGGCA-3'; reverse: 5'-CATCAGACCGTTAGGCTCA-3'); Hprt (forward: 5'-GAGCAATGTTGCTTCTC-3'; reverse: 5'-TTCTTTCCTGGTATGGGAATT-3').

To co-express rs3731249 variant and wild-type p16INK4a Beta 3 cells were transduced with equal molar cl20c-p16INK4a-1.48A-IRES-GFP and cl20c-p16INK4a-1.48A-IRES-lenti virus and cells successfully transfected with both were selected by flow cytometry sorting for GFP/FPP double positivity. BCIR-ABL1-mediated transformation was performed as described above. Genomic DNA and RNA samples were collected at day 0, 2, 4 and 5 after IL-3 removal, p.145A and p.148T transcript in RNA was quantified using allele-specific Tagman genotyping assay and normalized to allele ratio in matched DNA samples at respective time points. Each experiment was performed three times and each sample was assayed in triplicate.

Luciferase reporter assays. The p14INK4a 3'-UTR vector (3'-UTR for Human NM_005189.5 was placed downstream of luciferase reporter gene on the pEXMT01 backbone) was purchased from GeneCopoeia and the T variant at rs3731249 was introduced by site-directed mutagenesis (forward primer: 5'-CCATGGGCCG ATATCCGCGGTGTCGAGGTGCTC-3'; reverse primer: 5'GGATGTCGTGAGGACCTTCAGCTACGTGGGATGG-3'). For reporter gene assay, 2.5x10^5 293 T cells cultured in 96-well plate were transiently transfected with 100 ng empty vector, variant, or wild-type p14INK4a 3'UTR constructs using Lipofectamine 2000 (Invitrogen). Firefly luciferase activities were measured 24 h later using the Dual Luciferase Assay (Promega). The results were normalized against Renilla luciferase. Each reporter construct transfection was replicated at least three times, and each assay was completed in triplicate.

References

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The authors also thank the staff and participants of the ARIC study for their important contributions.

**Author contributions**


**Additional information**

**Accession codes.** The RNA-seq data have been deposited in European Genome Phenome archive under the accession codes EGAS00001000654.

[Supplementary Information](http://www.nature.com/naturecommunications) accompanies this paper at [http://www.nature.com/naturecommunications](http://www.nature.com/naturecommunications).

**Competing financial interests:** The authors declare no competing financial interests.

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