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# Identification of multiple risk loci and regulatory mechanisms influencing susceptibility to multiple myeloma

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Genome-wide association studies (GWAS) have transformed our understanding of susceptibility to multiple myeloma (MM), but much of the heritability remains unexplained. We report a new GWAS, a meta-analysis with previous GWAS and a replication series, totalling 9974 MM cases and 247,556 controls of European ancestry. Collectively, these data provide evidence for six new MM risk loci, bringing the total number to 23. Integration of information from gene expression, epigenetic profiling and in situ Hi-C data for the 23 risk loci implicate disruption of developmental transcriptional regulators as a basis of MM susceptibility, compatible with altered B-cell differentiation as a key mechanism. Dysregulation of autophagy/apoptosis and cell cycle signalling feature as recurrently perturbed pathways. Our findings provide further insight into the biological basis of MM.

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**M**ultiple myeloma (MM) is a malignancy of plasma cells primarily located within the bone marrow. Although no lifestyle or environmental exposures have been consistently linked to an increased risk of MM, the two- to four-fold increased risk observed in relatives of MM patients provides support for inherited genetic predisposition<sup>1</sup>. Our understanding of MM susceptibility has recently been informed by genome-wide association studies (GWAS), which have so far identified 17 independent risk loci for MM<sup>2–5</sup>, with an additional locus being subtype-specific for t(11;14) translocation MM<sup>6</sup>. Much of the heritable risk of MM, however, remains unexplained and statistical modelling indicates that further common risk variants remain to be discovered<sup>7</sup>.

To gain a more comprehensive insight into MM aetiology, we performed a new GWAS followed by a meta-analysis with existing GWAS and replication genotyping (totalling 9974 cases and 247,556 controls). Here, we report the identification of six new MM susceptibility loci as well as refined risk estimates for the previously reported loci. In addition, we have investigated the possible gene regulatory mechanisms underlying the associations seen at all 23 GWAS risk loci by analysing in situ promoter Capture Hi-C (Chi-C) in MM cells to characterise chromatin interactions between predisposition single-nucleotide polymorphism (SNPs) and target genes, integrating these data with chromatin immunoprecipitation-sequencing (ChIP-seq) data generated in house and a range of publicly available genomics data. Finally, we have quantified the contribution of both new and previously discovered loci to the heritable risk of MM and implemented a likelihood-based approach to estimate sample sizes required to explain 80% of the heritability.

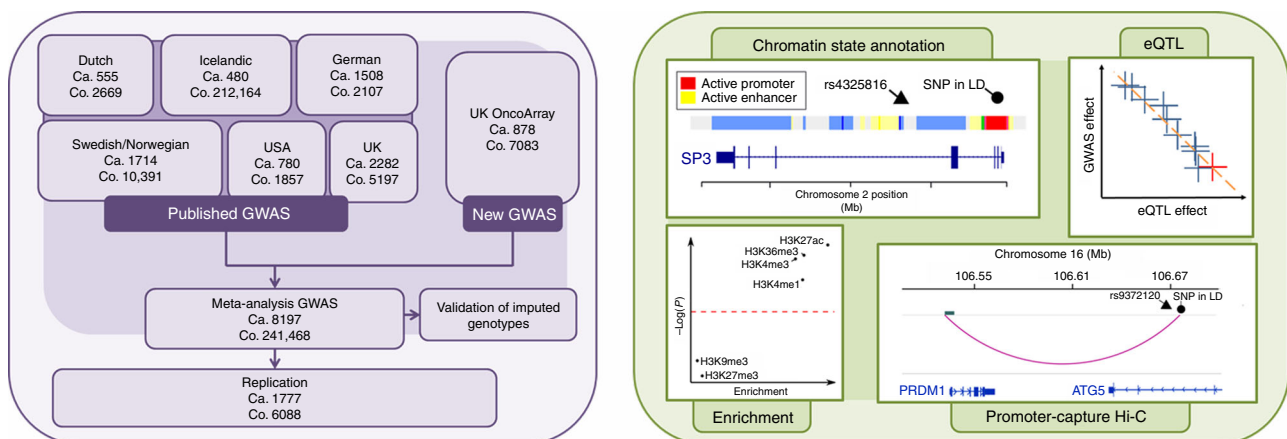
## Results

**Association analysis.** We conducted a new GWAS using the OncoArray platform<sup>8</sup> (878 MM cases and 7083 controls from the UK), followed by a meta-analysis with six published MM GWAS data sets (totalling 7319 cases and 234,385 controls) (Fig. 1, Supplementary Tables 1–3)<sup>2–5</sup>. To increase genomic resolution, we imputed data to >10 million SNPs. Quantile–quantile (Q–Q) plots for SNPs with minor allele frequency (MAF) >1% after imputation did not show evidence of substantive over-dispersion for the OncoArray GWAS ( $\lambda = 1.03$ ,  $\lambda_{1000} = 1.02$ , Supplementary Fig. 1). We derived joint odds ratios (ORs) under a fixed-effects model for each SNP with MAF >1%. Finally, we sought validation of nine SNPs associated at  $P < 1 \times 10^{-6}$  in the meta-analysis, which did not map to known MM risk loci and displayed a

consistent OR across all GWAS data sets, by genotyping an additional 1777 cases and 6088 controls from three independent series (Germany, Denmark and Sweden). After meta-analysis of the new and pre-existing GWAS data sets and replication series, we identified genome-wide significant associations (i.e.  $P < 5 \times 10^{-8}$ )<sup>9</sup> for six new loci at 2q31.1, 5q23.2, 7q22.3, 7q31.33, 16p11.2 and 19p13.11 (Table 1, Supplementary Table 4 and 5, Fig. 2). Additionally, borderline associations were identified at two loci with  $P$  values of  $5.93 \times 10^{-8}$  (6p25.3) and  $9.90 \times 10^{-8}$  (7q21.11), which have corresponding Bayesian false-discovery probabilities (BFDP)<sup>10</sup> of 4% and 6%, respectively (Supplementary Table 4 and 5). We found no evidence for significant interactions between any of the 23 risk loci. Finally, we found no evidence to support the existence of the putative risk locus at 2p12.3 (rs1214346), previously proposed by Erickson et al.<sup>11</sup> (GWAS meta-analysis  $P$  value = 0.32).

**Risk SNPs and myeloma phenotype.** We did not find any association between sex or age at diagnosis and the 23 MM risk SNPs using case-only analysis (Supplementary Table 6 and 7). Aside from previously reported relationships between the risk loci at 11q13.3 and 5q15 with t(11;14) MM<sup>6</sup> and hyperdiploid MM<sup>12</sup>, respectively, we found no evidence for subtype-specific associations (Supplementary Table 8–11) or an impact on MM-specific survival (Supplementary Table 12). A failure to demonstrate additional relationships may, however, be reflective of limited study power. Collectively, these data suggest that the risk variants are likely to have generic effects on MM development.

**Contribution of risk SNPs to heritability.** Using linkage disequilibrium adjusted kinships (LDAK)<sup>13</sup>, the heritability of MM ascribable to all common variation was 15.6% ( $\pm 4.7$ ); collectively the previously identified and new risk loci account for 15.7% of the GWAS heritability (13.6% and 2.1%, respectively). To assess the collective impact of all identified risk SNPs, we constructed polygenic risk scores (PRS) considering the combined effect of all risk SNPs modelled under a log-normal relative risk distribution<sup>14</sup>. Using this approach, an individual in the top 1% of genetic risk has a threefold increased risk of MM when compared to an individual with median genetic risk (Supplementary Fig. 2). We observed an enrichment of risk variants among familial MM compared with both sporadic MM cases and population-based controls comparable to that expected in the absence of a strong monogenic predisposition (respective  $P$  values 0.027 and



**Fig. 1** GWAS study design. Details of the new and existing GWAS samples, including recruitment centres or trials and quality control, are provided in Supplementary Tables 1 and 2. Trials or centres from which replication samples were recruited are detailed in Supplementary Table 3. Ca.: cases, Co.: controls, eQTL: expression quantitative trait loci, SNP: single-nucleotide polymorphism, LD: linkage disequilibrium

**Table 1 Summary of genotyping results for all 23 risk SNPs**

SNP	Locus	Pos. (b37)	Risk Allele	RAF	OncoArray		Previous data		Replication		Combined meta		$I^2$
					OR	$P_{trend}$	OR	$P_{trend}$	OR	$P_{trend}$	OR	$P_{meta}$	
rs7577599	2p23.3	25613146	T	0.81	1.22	$2.63 \times 10^{-3}$	1.24	$1.24 \times 10^{-16}$	-	-	1.23	$1.29 \times 10^{-18}$	0
<b>rs4325816</b>	2q31.1	<b>174808899</b>	<b>T</b>	<b>0.77</b>	<b>1.16</b>	<b><math>1.23 \times 10^{-2}</math></b>	<b>1.11</b>	<b><math>1.30 \times 10^{-5}</math></b>	<b>1.16</b>	<b><math>3.00 \times 10^{-3}</math></b>	<b>1.12</b>	<b><math>7.37 \times 10^{-9}</math></b>	<b>9</b>
rs6599192	3p22.1	41992408	G	0.16	1.24	$1.35 \times 10^{-3}$	1.26	$8.75 \times 10^{-18}$	-	-	1.26	$4.96 \times 10^{-20}$	0
rs10936600	3q26.2	169514585	A	0.75	1.18	$5.12 \times 10^{-3}$	1.20	$5.94 \times 10^{-15}$	-	-	1.20	$1.20 \times 10^{-16}$	0
rs1423269	5q15	95255724	A	0.75	1.09	0.125	1.17	$1.57 \times 10^{-11}$	-	-	1.16	$8.30 \times 10^{-12}$	23
<b>rs6595443</b>	5q23.2	<b>122743325</b>	<b>T</b>	<b>0.43</b>	<b>1.14</b>	<b><math>9.87 \times 10^{-3}</math></b>	<b>1.10</b>	<b><math>4.69 \times 10^{-6}</math></b>	<b>1.10</b>	<b>0.022</b>	<b>1.11</b>	<b><math>1.20 \times 10^{-8}</math></b>	<b>0</b>
rs34229995	6p22.3	15244018	G	0.02	1.05	0.781	1.40	$1.76 \times 10^{-8}$	-	-	1.36	$5.60 \times 10^{-8}$	0
rs3132535	6p21.3	31116526	A	0.29	1.26	$2.67 \times 10^{-5}$	1.20	$2.97 \times 10^{-17}$	-	-	1.21	$6.00 \times 10^{-21}$	0
rs9372120	6q21	106667535	G	0.21	1.18	$7.74 \times 10^{-3}$	1.20	$8.72 \times 10^{-14}$	-	-	1.19	$2.40 \times 10^{-15}$	0
rs4487645	7p15.3	21938240	C	0.65	1.23	$1.06 \times 10^{-4}$	1.24	$5.30 \times 10^{-25}$	-	-	1.24	$2.80 \times 10^{-28}$	0
<b>rs17507636</b>	7q22.3	<b>106291118</b>	<b>C</b>	<b>0.74</b>	<b>1.12</b>	<b><math>5.71 \times 10^{-2}</math></b>	<b>1.12</b>	<b><math>5.54 \times 10^{-7}</math></b>	<b>1.10</b>	<b>0.036</b>	<b>1.12</b>	<b><math>9.20 \times 10^{-9}</math></b>	<b>50</b>
<b>rs58618031</b>	7q31.33	<b>124583896</b>	<b>T</b>	<b>0.72</b>	<b>1.17</b>	<b><math>7.61 \times 10^{-3}</math></b>	<b>1.11</b>	<b><math>4.70 \times 10^{-6}</math></b>	<b>1.10</b>	<b>0.061</b>	<b>1.12</b>	<b><math>2.73 \times 10^{-8}</math></b>	<b>0</b>
rs7781265	7q36.1	150950940	A	0.12	1.33	$3.23 \times 10^{-4}$	1.20	$1.82 \times 10^{-7}$	-	-	1.22	$4.82 \times 10^{-10}$	49
rs1948915	8q24.21	128222421	C	0.32	1.19	$1.68 \times 10^{-3}$	1.14	$3.14 \times 10^{-10}$	-	-	1.15	$2.53 \times 10^{-12}$	26
rs2811710	9p21.3	21991923	C	0.63	1.13	$1.76 \times 10^{-2}$	1.14	$6.50 \times 10^{-10}$	-	-	1.14	$3.64 \times 10^{-11}$	0
rs2790457	10p12.1	28856819	G	0.73	1.09	0.124	1.12	$8.44 \times 10^{-7}$	-	-	1.11	$2.66 \times 10^{-6}$	0
<b>rs13338946</b>	16p11.2	<b>30700858</b>	<b>C</b>	<b>0.26</b>	<b>1.17</b>	<b><math>7.90 \times 10^{-3}</math></b>	<b>1.12</b>	<b><math>2.22 \times 10^{-7}</math></b>	<b>1.26</b>	<b><math>2.5 \times 10^{-7}</math></b>	<b>1.15</b>	<b><math>1.02 \times 10^{-13}</math></b>	<b>26</b>
rs7193541	16q23.1	74664743	T	0.58	1.14	$9.01 \times 10^{-3}$	1.12	$1.14 \times 10^{-8}$	-	-	1.12	$3.68 \times 10^{-10}$	34
rs34562254	17p11.2	16842991	A	0.10	1.32	$7.63 \times 10^{-4}$	1.30	$3.63 \times 10^{-17}$	-	-	1.30	$1.18 \times 10^{-19}$	29
<b>rs11086029</b>	19p13.11	<b>16438661</b>	<b>T</b>	<b>0.24</b>	<b>1.26</b>	<b><math>1.02 \times 10^{-4}</math></b>	<b>1.12</b>	<b><math>1.69 \times 10^{-6}</math></b>	<b>1.15</b>	<b><math>5.00 \times 10^{-3}</math></b>	<b>1.14</b>	<b><math>6.79 \times 10^{-11}</math></b>	<b>42</b>
rs6066835	20q13.13	47355009	C	0.08	1.13	0.162	1.24	$1.16 \times 10^{-9}$	-	-	1.23	$6.58 \times 10^{-10}$	38
rs138747	22q13.1	35700488	A	0.66	-	-	1.21	$2.58 \times 10^{-8}$	-	-	1.21	$2.58 \times 10^{-8}$	0
rs139402	22q13.1	39546145	C	0.44	1.11	$4.146 \times 10^{-2}$	1.23	$4.98 \times 10^{-26}$	-	-	1.22	$3.84 \times 10^{-26}$	56

Newly identified risk loci are emboldened.<sup>1</sup> Where >10 TF were implicated at a locus, only those that overlap with TF which demonstrated enrichment in GM12878 are shown here. A full list of TFs localising to loci are detailed in Supplementary Table 17

$1.60 \times 10^{-5}$ ; Supplementary Fig. 3). Undoubtedly, the identification of further risk loci through the analysis of larger GWAS are likely to improve the performance of any PRS model. To estimate the sample size required to explain a greater proportion of the GWAS heritability, we implemented a likelihood-based approach using association statistics in combination with LD information to model the effect-size distribution<sup>15,16</sup>. The effect-size distributions for susceptibility SNPs were best modelled using the three-component model (mixture of two normal distributions) (Supplementary Fig. 4). Under this model, to identify SNPs explaining 80% of the GWAS heritability is likely to require sample sizes in excess of 50,000 (Supplementary Fig. 5).

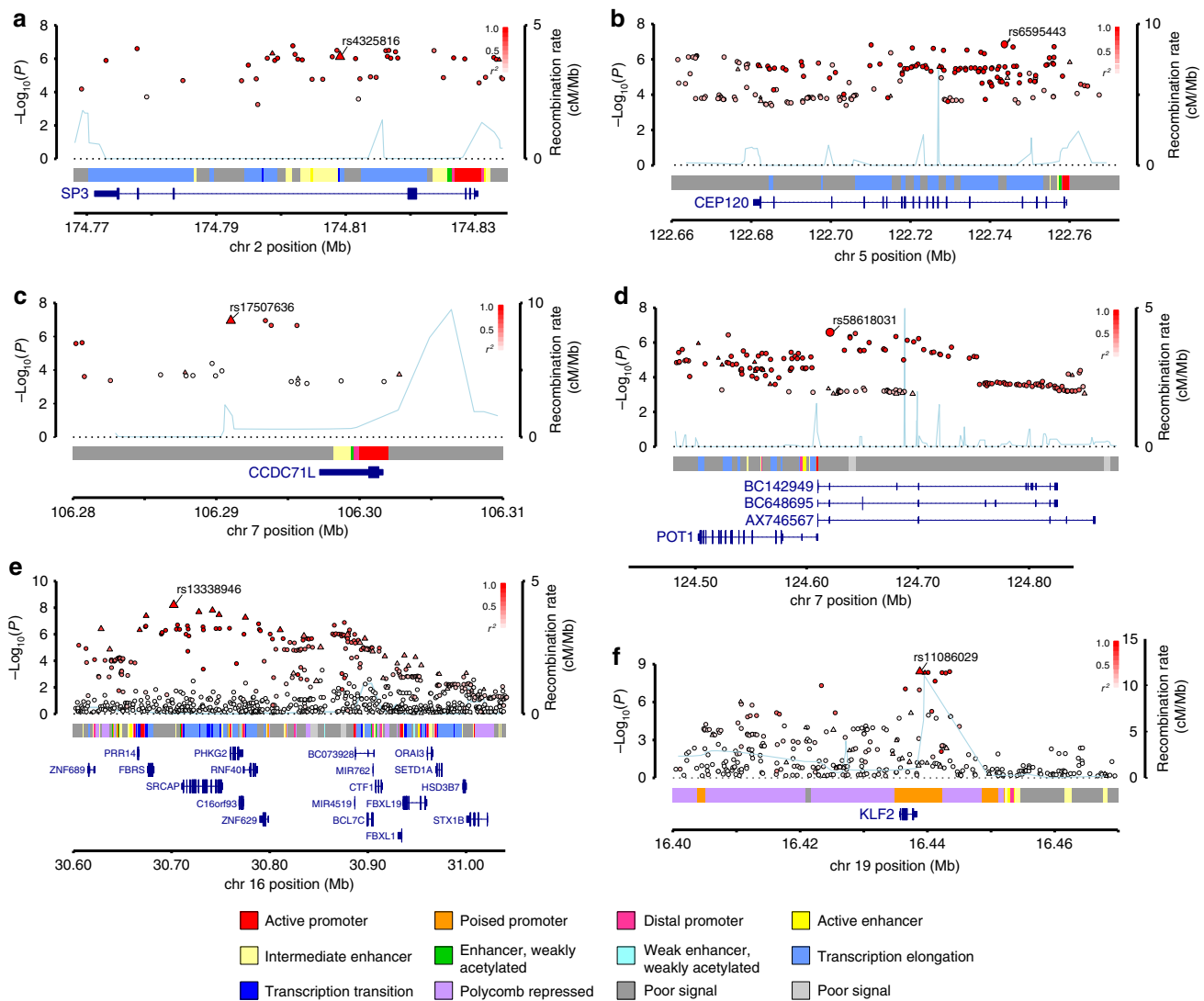
**Functional annotation and biological inference of risk loci.** To the extent that they have been studied, many GWAS risk SNPs localise to non-coding regions and influence gene regulation<sup>17</sup>. To investigate the functional role of previously reported and new MM risk SNPs, we performed a global analysis of SNP associations using ChIP-seq data generated on the MM cell line KMS11 and publicly accessible naive B-cell Blueprint Epigenome Project data<sup>18</sup>. We found enrichment of MM SNPs in regions of active chromatin, as indicated by the presence of H3K27ac, H3K4Me3 and H3K4Me1 marks (Supplementary Fig. 6). We also observed an enrichment of relevant B-cell transcription factor-(TF) binding sites using ENCODE GM12878 lymphoblastoid cell line data (Supplementary Fig. 7). Collectively these data support the tenet that the MM predisposition loci influence risk through effects on *cis*-regulatory networks involved in transcriptional initiation and enhancement.

Since genomic spatial proximity and chromatin looping interactions are key to the regulation of gene expression, we interrogated physical interactions at respective genomic regions in KMS11 and naive B-cells using CHi-C data<sup>19</sup>. We also sought to gain insight into the possible biological mechanisms for associations by performing an expression quantitative trait locus

(eQTL) analysis using mRNA expression data on CD138-purified MM plasma cells; specifically, we used Summary data-based Mendelian Randomisation (SMR) analysis<sup>20</sup> to test for pleiotropy between GWAS signal and *cis*-eQTL for genes within 1 Mb of the sentinel SNP to identify a causal relationship. We additionally annotated risk loci with variants mapping to binding motifs of B-cell-specific TFs. Finally, we catalogued direct promoter variants and non-synonymous coding mutations for genes within risk loci (Supplementary Data 1 and Fig. 1).

Although preliminary, and requiring functional validation, our analysis delineates four potential candidate disease mechanisms across the 23 MM risk loci (Supplementary Data 1). Firstly, four of the risk loci contain candidate genes linked to regulation of cell cycle and genomic instability, as evidenced by Hi-C looping interactions in KMS11 cells to *MTAP* (at 9p21.3) and eQTL effects for *CEP120* (at 5q23.2). *CEP120* is required for microtubule assembly and elongation with overexpression of *CEP120* leading to uncontrolled centriole elongation<sup>21</sup>. rs58618031 (7q31.33) maps 5' of *POT1*, the protection of telomeres 1 gene. *POT1* is part of the shelterin complex that functions to protect telomeres and maintain chromosomal stability<sup>22,23</sup>. While mutated *POT1* is not a feature of MM, it is commonly observed in B-cell chronic lymphocytic leukaemia<sup>24–26</sup>. The looping interaction from the rs58618031 annotated enhancer element implicates *ASB15*. Members of the ASB family feature as protein components of the ubiquitin–proteasome system, intriguingly a therapeutic target in MM<sup>27–29</sup>.

Second, candidate genes encoding proteins involved in chromatin remodelling were implicated at three of the MM risk loci, supported by promoter variants at 2q31.1, 7q36.1 and 22q13.1. The new locus at 2q31.1 implicates *SP3*, encoding a TF, which through promoter interaction, has a well-established role in B-cell development influencing the expression of germinal centre genes, including activation-induced cytidine deaminase *AID*<sup>30,31</sup>.



**Fig. 2** Regional plots of the six new risk loci. Regional plots of loci **a** 2q31.1, **b** 5q23.2, **c** 7q22.3, **d** 7q31.33, **e** 16p11.2 and **f** 19p13.11. Plots show results of the meta-analysis for both genotyped (triangles) and imputed (circles) single-nucleotide polymorphisms (SNPs) and recombination rates.  $-\log_{10}(P)$  (y axes) of the SNPs are shown according to their chromosomal positions (x axes). The sentinel SNP in each combined analysis is shown as a large circle or triangle and is labelled by its rsID. The colour intensity of each symbol reflects the extent of LD with the top SNP, white ( $r^2 = 0$ ) through to dark red ( $r^2 = 1.0$ ). Genetic recombination rates, estimated using 1000 Genomes Project samples, are shown with a light blue line. Physical positions are based on NCBI build 37 of the human genome. Also shown are the relative positions of genes and transcripts mapping to the region of association. Genes have been redrawn to show their relative positions; therefore maps are not to physical scale. The middle track represents the chromatin-state segmentation track (ChromHMM) for KMS11

Third, the central role *IRF4-MYC*-mediated apoptosis/autophagy in MM oncogenesis is supported by variation at five loci, including eQTL effects WAC (at 10p12.1) and Hi-C looping interactions (at 8q24.21 and 16q23.1). The 7p15.3 association ascribable to rs4487645 has been documented to influence expression of *c-MYC*-interacting *CDCA7L* through differential *IRF4* binding<sup>32</sup>. Similarly, the long-range interaction between *CCAT1* (colon cancer-associated transcript 1) and *MYC* provides an attractive biological basis for the 8q24.21 association, given the notable role of *MYC* in MM<sup>33,34</sup>. It is noteworthy that the promising risk locus at 6p25.3 contains *IRF4*. At the new locus 19p13.11, the missense variant (NP\_057354.1:p.Leu104Pro) and the correlated promoter SNP rs11086029 implicates *KLF2* in MM biology. Demethylation by KDM3A histone demethylase sustains *KLF2* expression and influences *IRF4*-dependent MM cell survival<sup>35</sup>. The new 16p11.2 risk locus contains a number of genes including Proline-Rich Protein 14 (*PRR14*), which is

implicated in PI3-kinase/Akt/mTOR signalling, a therapeutic target in myelomatous plasma cells<sup>36</sup>.

Fourth, loci related to B cell and plasma cell differentiation and function are supported by variation at three loci, including eQTL effects (*ELL2* at 5q15) and Hi-C looping interactions (at 6q21). As previously inferred from GM12878 cell line data, the region at 6q21 (rs9372120, *ATG5*) participates in intra-chromosome looping with the B-cell transcriptional repressor *PRDM1* (alias *BLIMP1*)<sup>4</sup>. Additionally, SNP rs34562254 at 17p11.2 is responsible for the amino acid substitution (NP\_036584.1:p.Pro251Leu) in *TNFRSF13B*, a key regulator of normal B-cell homeostasis, which has an established role in MM biology<sup>37–42</sup>.

## Discussion

Our meta-analysis of a new GWAS series in conjunction with previously published MM data sets has identified six novel risk



loci. Together, the new and previously reported loci explain an estimated 16% of the SNP heritability for MM in European populations. Ancestral differences in the risk of developing MM are well recognised, with a greater prevalence of MM in African Americans as compared with those with European ancestry<sup>43</sup>. It is plausible that the effects of MM risk SNPs may differ between Europeans and non-Europeans and hence contribute to differences in prevalence rates. Thus far, there has only been limited evaluation of this possibility with no evidence for significant differences<sup>44</sup>.

Integration of Hi-C data with ChIP-seq chromatin profiling from MM and lymphoblastoid cell lines and naive B cells and eQTL analysis, using patient expression data, has allowed us to gain preliminary insight into the biological basis of MM susceptibility. This analysis suggests a model of MM susceptibility based on transcriptional dysregulation consistent with altered B-cell differentiation, where dysregulation of autophagy/apoptosis and cell cycle signalling feature as recurrently modulated pathways. Specifically, our findings implicate mTOR-related genes *ULK4*, *ATG5* and *WAC*, and by virtue of the role of *IRF4-MYC* related autophagy, *CDCA7L*, *DNMT3A*, *CBX7* and *KLF2* in MM development (Supplementary Data 1). Further investigations are necessary to decipher the functional basis of risk SNPs, nevertheless we highlight mTOR signalling and the ubiquitin-proteasome pathway, targets of approved drugs in MM. As a corollary of this, genes elucidated via the functional annotation of GWAS that discovered MM risk loci may represent promising therapeutic targets for myeloma drug discovery. Finally, our estimation of sample sizes required to identify a larger proportion of the heritable risk of MM attributable to common variation underscore the need for further international collaborative analyses.

## Methods

**Ethics.** Collection of patient samples and associated clinico-pathological information was undertaken with written informed consent and relevant ethical review board approval at respective study centres in accordance with the tenets of the Declaration of Helsinki. Specifically for the Myeloma-IX trial by the Medical Research Council (MRC) Leukaemia Data Monitoring and Ethics committee (MREC 02/8/95, ISRCTN68454111), the Myeloma-XI trial by the Oxfordshire Research Ethics Committee (MREC 17/09/09, ISRCTN49407852), HOVON65/GMMG-HD4 (ISRCTN 644552890; METC 13/01/2015), HOVON87/NMSG18 (EudraCTnr 2007-004007-34, METC 20/11/2008), HOVON95/EMN02 (EudraCTnr 2009-017903-28, METC 04/11/10), University of Heidelberg Ethical Commission (229/2003, S-337/2009, AFmu-119/2010), University of Arkansas for Medical Sciences Institutional Review Board (IRB 202077), Lund University Ethical Review Board (2013/54), the Norwegian REK 2014/97, the Danish Ethical Review Board (no. H-16032570) and Icelandic Data Protection Authority (2,001,010,157 and National Bioethics Committee 01/015).

The diagnosis of MM (ICD-10 C90.0) in all cases was established in accordance with World Health Organization guidelines. All samples from patients for genotyping were obtained before treatment or at presentation.

**Primary GWAS.** We analysed constitutional DNA (EDTA-venous blood derived) from 931 cases ascertained through the UK Myeloma XI trial; detailed in Supplementary Table 1. Cases were genotyped using the Illumina OncoArray (Illumina Inc. San Diego, CA 92122, USA). Controls were also genotyped using the OncoArray and comprised: (1) 2976 cancer-free men recruited by the PRACTICAL Consortium—the UK Genetic Prostate Cancer Study (UKGPCS) (age <65 years), a study conducted through the Royal Marsden NHS Foundation Trust and SEARCH (Study of Epidemiology & Risk Factors in Cancer), recruited via GP practices in East Anglia (2003–2009) and (2) 4446 cancer-free women across the UK, recruited via the Breast Cancer Association Consortium (BCAC).

Standard quality-control measures were applied to the GWAS<sup>45</sup>. Specifically, individuals with low SNP call rate (<95%) as well as individuals evaluated to be of non-European ancestry (using the HapMap version 2 CEU, JPT/CHB and YRI populations as a reference) were excluded (Supplementary Fig. 8). For apparent first-degree relative pairs, we excluded the control from a case-control pair; otherwise, we excluded the individual with the lower call rate. SNPs with a call rate <95% were excluded as were those with a MAF <0.01 or displaying significant deviation from Hardy-Weinberg equilibrium ( $P < 10^{-5}$ ). GWAS data were imputed to >10 million SNPs using IMPUTE2 v2.3<sup>46</sup> software in conjunction with

a merged reference panel consisting of data from 1000 Genomes Project<sup>47</sup> (phase 1 integrated release 3 March 2012) and UK10K<sup>48</sup>. Genotypes were aligned to the positive strand in both imputation and genotyping. We imposed predefined thresholds for imputation quality to retain potential risk variants with MAF >0.01 for validation. Poorly imputed SNPs with an information measure <0.80 were excluded. Tests of association between imputed SNPs and MM was performed under an additive model in SNPTESTv2.5<sup>49</sup>. The adequacy of the case-control matching and possibility of differential genotyping of cases and controls was evaluated using a Q-Q plot of test statistics (Supplementary Fig. 1). The inflation  $\lambda$  was based on the 90% least-significant SNPs<sup>50</sup> and assessment of  $\lambda_{1000}$ . Details of SNP QC are provided in in Supplementary Table 2.

**Published GWAS.** The data from six previously reported GWAS<sup>2–5</sup> are summarised in Supplementary Table 1. All these studies were based on individuals with European ancestry and comprised: UK-GWAS (2282 cases, 5197 controls), Swedish-GWAS (1714 cases, 10,391 controls), German-GWAS (1508 cases, 2107 controls), Netherlands-GWAS (555 cases, 2669 controls), US-GWAS (780 cases, 1857 controls) and Iceland (480 cases, 212,164 controls).

**Replication studies and technical validation.** To validate promising associations, we analysed three case-control series from Germany, Sweden and Denmark, summarised in Supplementary Table 3. The German replication series comprised 911 cases collected by the German Myeloma Study Group (Deutsche Studiengruppe Multiples Myeloma (DSMM)), GMMG, University Clinic, Heidelberg and University Clinic, Ulm. Controls comprised 1477 healthy German blood donors recruited between 2004 and 2007 by the Institute of Transfusion Medicine and Immunology, University of Mannheim, Germany. The Swedish replication series comprised 534 MM cases from the Swedish National Myeloma Biobank and the Danish replication series comprised 332 MM cases from the University Hospital of Copenhagen. As controls, we analysed 2382 Swedish blood donors and 2229 individuals from Denmark and Skåne County, Sweden (the southernmost part of Sweden adjacent to Denmark). Replication genotyping of German and Scandinavian samples was performed using competitive allele-specific PCR KASPar chemistry (LGC, Hertfordshire, UK). Call rates for SNP genotypes were >95% in each of the replication series. To ensure the quality of genotyping in all assays, at least two negative controls and duplicate samples (showing a concordance of >99%) were genotyped at each centre. The fidelity of imputation was assessed by directly sequencing a set of 147 randomly selected samples from the UK OncoArray case series. Imputation was found to be robust; concordance was >90% (Supplementary Table 13). Genotyping and sequencing primers are detailed in Supplementary Table 14 and 15, respectively.

**Meta-analysis.** Meta-analyses were performed using the fixed-effects inverse-variance method using META v1.6<sup>51</sup>. Cochran's Q-statistic to test for heterogeneity and the  $I^2$  statistic to quantify the proportion of the total variation due to heterogeneity was calculated. Using the meta-analysis summary statistics and LD correlations from a reference panel of the 1000 Genomes Project combined with UK10K, we implemented Genome-wide Complex Trait Analysis<sup>52</sup> to perform conditional association analysis. Association statistics were calculated for all SNPs conditioning on the top SNP in each loci showing genome-wide significance. This was carried out step-wise.

For borderline associations, the BFDP<sup>10</sup> was calculated based on a plausible OR of 1.2 and a prior probability of association of 0.0001. For both promising associations, the BFDP was <10%.

**Fluorescence in situ hybridisation.** Fluorescence in situ hybridisation (FISH) and ploidy classification of UK and German samples were performed as previously described<sup>53,54</sup>. Logistic regression in case-only analyses was used to assess the relationship between SNP genotype and IgH translocations or tumour ploidy.

**eQTL analysis.** eQTL analyses were performed using CD138-purified plasma cells from 183 UK MyIX trial patients and 658 German GMMG patients<sup>32</sup>. Briefly, German and UK data were pre-processed separately, followed by analysis using a Bayesian approach to probabilistic estimation of expression residuals to infer broad variance components, accounting for hidden determinants influencing global expression. The association between genotype of SNPs and expression of genes within 1 Mb either side of each MM risk locus was evaluated based on the significance of linear regression coefficients. We pooled data from each study under a fixed-effects model.

The relationship between SNP genotype and gene expression was carried out using SMR analysis as per Zhu et al.<sup>2</sup> Briefly, if  $b_{xy}$  is the effect size of  $x$  (gene expression) on  $y$  (slope of  $y$  regressed on the genetic value of  $x$ ),  $b_{zx}$  is the effect of  $z$  on  $x$  and  $b_{zy}$  be the effect of  $z$  on  $y$ ,  $b_{xy}$  ( $b_{zy}/b_{zx}$ ) is the effect of  $x$  on  $y$ . To distinguish pleiotropy from linkage where the top associated *cis*-eQTL is in LD with two causal variants, one affecting gene expression and the other affecting a trait, we tested for heterogeneity in dependent instruments (HEIDI), using multiple SNPs in each *cis*-eQTL region. Under the hypothesis of pleiotropy,  $b_{xy}$  values for SNPs in LD with the causal variant should be identical. For each probe that passed significance

threshold for the SMR test, we tested the heterogeneity in the  $b_{xy}$  values estimated for multiple SNPs in the *cis*-eQTL region using HEIDI.

GWAS summary statistics files were generated from the meta-analysis. We set a threshold for the SMR test of  $P_{SMR} < 1 \times 10^{-3}$  corresponding to a Bonferroni correction for 45 tests, i.e. 45 probes which demonstrated an association in the SMR test. For all genes passing this threshold, we generated plots of the eQTL and GWAS associations at the locus, as well as plots of GWAS and eQTL effect sizes (i.e. input for the HEIDI heterogeneity test). HEIDI test  $P$  values  $< 0.05$  were considered as reflective of heterogeneity. This threshold is, however, conservative for gene discovery because it retains fewer genes than when correcting for multiple testing. SMR plots for significant eQTLs are shown in Supplementary Fig. 9 and 10, and a summary of results are shown in Supplementary Table 16.

**Promoter CHI-C.** To map risk SNPs to interactions involving promoter contacts and identify genes involved in MM susceptibility, we analysed publicly accessible promoter CHI-C data on the naive B cells downloaded from Blueprint Epigenome Project. Additionally, we also analysed promoter CHI-C data that we have previously generated for the MM cell line KMS11<sup>12</sup>. Interactions were called using the CHICAGO pipeline to obtain a unique list of reproducible contacts<sup>55</sup> and those with a  $-\log(\text{weighted } P) \geq 5$  were considered significant.

**Chromatin state annotation.** Variant sets (i.e. sentinel risk SNP and correlated SNPs,  $r^2 > 0.8$ ) were annotated for putative functional effect based upon histone mark ChIP-seq data for H3K27ac, H3K4Me1, H3K27Me3, H3K9Me3, H3K36Me3 and H3K27Me3 from KMS11 cell lines, generated in-house and naive B cells from Blueprint Epigenome Project<sup>56</sup>. We used ChromHMM to infer chromatin states by integrating information on these histone modifications, training the model on three MM cell lines; KMS11, MM1S and JLN3. Genome-wide signal tracks were binarized (including input controls for ChIP-seq data), and a set of learned models were generated using ChromHMM software<sup>57</sup>. A 12-state model was suitable for interpretation, and biological meaning was assigned to the states based on chromatin marks that use putative rules as previously described (Supplementary Fig. 11).

**TF and histone mark enrichment analysis.** To examine enrichment in specific TF binding across risk loci, we adapted the method of Cowper-Salari et al.<sup>58</sup>. Briefly, for each risk locus, a region of strong LD (defined as  $r^2 > 0.8$  and  $D' > 0.8$ ) was determined, and these SNPs were considered the associated variant set (AVS). Publicly available data on TF ChIP-seq uniform peak data were obtained from ENCODE for the GM12878 cell line, including data for 82 TF and 11 histone marks<sup>59</sup>. In addition, ChIP-seq peak data for six histone marks from KMS11 cell line were generated in-house, and naive B-cell ChIP-seq data were downloaded from Blueprint Epigenome Project<sup>56</sup>. For each mark, the overlap of the SNPs in the AVS and the binding sites was assessed to generate a mapping tally. A null distribution was produced by randomly selecting SNPs with the same characteristics as the risk-associated SNPs, and the null mapping tally calculated. This process was repeated 10,000 times, and  $P$  values were calculated as the proportion of permutations, where null mapping tally was greater or equal to the AVS mapping tally. An enrichment score was calculated by normalising the tallies to the median of the null distribution. Thus, the enrichment score is the number of standard deviations of the AVS mapping tally from the median of the null distribution tallies. Enrichment plots are shown in Supplementary Fig. 6 and 7.

**Functional annotation.** For the integrated functional annotation of risk loci, variant sets (i.e. all SNPs in LD  $r^2 > 0.8$  with the sentinel SNP) were annotated with: (i) presence of a Hi-C contact linking to a gene promoter, (ii) presence of an association from SMR analysis, (iii) presence of a regulatory ChromHMM state, (iv) evidence of transcription factor binding and (v) presence of a nonsynonymous coding change. Candidate causal genes were then assigned to MM risk loci using the target genes implicated in annotation tracks (i), (ii), (iii) and (iv). If the data supported multiple gene candidates, the gene with the highest number of individual functional data points was considered as the candidate. Where multiple genes have the same number of data points, all genes are listed. Direct non-synonymous coding variants were allocated additional weighting. Competing mechanisms for the same gene (e.g. both coding and promoter variants) were permitted.

**Heritability analysis.** We used LDAK to estimate the polygenic variance (i.e. heritability) ascribable to all genotyped and imputed GWAS SNPs from summary statistic data. SNP-specific expected heritability, adjusted for LD, MAF and genotype certainty was calculated from the UK10K and 1000 Genomes data. Samples were excluded with a call rate  $< 0.99$  or if individuals were closely related or of divergent ancestry from CEU. Individual SNPs were excluded if they showed deviation from HWE with  $P < 1 \times 10^{-5}$ , an individual SNP genotype yield  $< 95\%$ , MAF  $< 1\%$ , SNP imputation score  $< 0.99$  and the absence of the SNP in the GWAS summary statistic data. This resulted in a total 1,254,459 SNPs which were used to estimate the heritability of MM.

To estimate the sample size required for a given proportion of the GWAS heritability, we implemented a likelihood-based approach to model the effect-size distribution<sup>15</sup> using association statistics, from the MM meta-analysis, and LD information, obtained from individuals of European ancestry in the 1000 Genomes

Project Phase 3. LD values were based on an  $r^2$  threshold of 0.1 and a window size of 1 MB. The goodness of fit of the observed distribution of  $P$  values against the expected from a two-component model (single normal distribution) and a three-component model (mixture of two normal distributions) were assessed<sup>15</sup>, and a better fit was observed for the latter model (Supplementary Figure 4). The percentage of GWAS heritability explained for a projected sample size was determined using this model and is based on power calculations for the discovery of genome-wide significant SNPs. The genetic variance explained was calculated as the proportion of total GWAS heritability explained by SNPs reaching genome-wide significance at a given sample size. The 95% confidence intervals were determined using 10,000 simulations.

PRS for familial MM ( $n = 38$ ) from 25 families were compared with sporadic MM ( $n = 1530$ ) and population-based controls ( $n = 10,171$ ); first as a simple sum of risk alleles and secondly as sum of risk alleles weighted by their log-transformed ORs. Family member scores were averaged. A one-sided Student's  $t$ -test was used to assess difference between groups. The genetic data have been previously described<sup>5,60</sup> with the familial MM cases having been identified by linkages of Swedish registry information.

**Data availability.** SNP genotyping data that support the findings of this study have been deposited in Gene Expression Omnibus with accession codes GSE21349, GSE19784, GSE24080, GSE2658 and GSE15695; in the European Genome-phenome Archive (EGA) with accession code EGAS0000000001; in the European Bioinformatics Institute (Part of the European Molecular Biology Laboratory) (EMBL-EBI) with accession code E-MTAB-362 and E-TABM-1138; and in the database of Genotypes and Phenotypes (dbGaP) with accession code phs000207.v1.p1. Expression data that support the findings of this study have been deposited in GEO with accession codes GSE21349, GSE2658, GSE31161 and EMBL-EBI with accession code E-MTAB-2299. The remaining data are contained within the paper and Supplementary Files or are available from the author upon request. KMS11 Hi-C data used in this manuscript are deposited in EGA under accession number EGAS00001002614. The accession number for the KMS11 ChIP-seq data reported in this paper is EGA: S00001002414. Naive B-cell HiC data used in this work is publicly available from Blueprint Epigenome Project [<https://osf.io/u8tzp/>]. ChIP-seq data for H3K27ac, H3K4Me1, H3K27Me3, H3K9Me3, H3K36Me3 and H3K27Me3 from naive B cells are publicly available and were obtained from Blueprint Epigenome Project [<http://www.blueprint-epigenome.eu/>].

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## Author contributions

M.W. and R.S.H. designed the study. M.W. and R.S.H. drafted the manuscript with contributions from K.H., B.N., A.S. and N.W. M.W. performed principal statistical and bioinformatics analyses. A.S., N.L., P.L., D.C.J., J.S.M. and G.O. performed additional bioinformatics analyses. S.K. performed in situ Chi-C. P.B. coordinated UK laboratory analyses. M.W. and A.H. performed sequencing of UK samples. D.C.J. managed and prepared Myeloma IX and Myeloma XI Case Study DNA samples. M.K., G.J.M., F.E.D., W.A.G. and G.H.J. performed ascertainment and collection of Case Study samples. B.A. W. performed UK expression analyses. F.M.R. performed UK fluorescence in situ hybridisation analyses. The PRACTICAL consortium, D.E., P.P., A.D., J.P., F.C., R.E., Z. K.-J., K.M. and N.P. provided control samples for the UK OncoArray GWAS. H.G., U.B., J.H., J.N. and N.W. coordinated and managed Heidelberg samples. C.L. and H.E. coordinated and managed Ulm/Wurzburg samples. A.F. coordinated German genotyping. C.C. and O.R.B. performed German genotyping. P.H. and M.M.N. performed GWAS of German cases and controls. H.T., B.C. and M.I.d.S.F. carried out statistical analysis. K.H. coordinated the German part of the project. M.M.N. generated genotype data from the Heinz-Nixdorf recall study. K.-H.J. contributed towards the Heinz-Nixdorf control data set. N.N. from Bonn and K.-H.J. provided samples for the German GWAS. M.H. and B.N. coordinated the Swedish/Norwegian part of the project. M.A. and B.-M. H. performed data analysis. E.J., A.-K.W., U.-H.M., H.N., A.V., N.F.A., A.W., I.T. and U. G. performed sample acquisition, sample preparation, clinical data acquisition and additional data analyses of Sweden/Norway samples. In Iceland, G.T. and D.F.G. performed statistical analysis. S.Y.K. provided clinical information. T.R. performed additional analyses. U.T. and K.S. performed project oversight. M.v.D., P.S., A.B. and R.K. coordinated and prepared HOVON65/GMMG-HD4, HOVON87/NMSG18 and HOVON95/EMN02 studies for participating in this study, and coordinated genotyping and pre-processing. At the Myeloma Institute, University of Arkansas for Medical Sciences, N.W. coordinated the US part of the project and performed statistical and eQTL analyses. O.W.S. and N.W. managed Case Study samples. G.J.M. and F.E.D. performed ascertainment and collection of Case Study samples.

## Additional information

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