RPL5 on 1p22.1 is recurrently deleted in multiple myeloma and its expression is linked to bortezomib response

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Chromosomal region 1p22 is deleted in ≥20% of multiple myeloma (MM) patients, suggesting the presence of an unidentified tumor suppressor. Using high-resolution genomic profiling, we delimit a 58 kb minimal deleted region (MDR) on 1p22.1 encompassing two genes: ectopic viral integration site 5 (EVI5) and ribosomal protein L5 (RPL5). Low mRNA expression of EVI5 and RPL5 was associated with worse survival in diagnostic cases. Patients with 1p22 deletion had lower mRNA expression of EVI5 and RPL5, however, 1p22 deletion status is a bad predictor of RPL5 expression in some cases, suggesting that other mechanisms downregulate RPL5 expression. Interestingly, RPL5 but not EVI5 mRNA levels were significantly lower in relapsed patients responding to bortezomib and; both in newly diagnosed and relapsed patients, bortezomib treatment could overcome their bad prognosis by raising their progression-free survival to equal that of patients with high RPL5 expression. In conclusion, our genetic data restrict the MDR on 1p22 to EVI5 and RPL5 and although the role of these genes in promoting MM progression remains to be determined, we identify RPL5 mRNA expression as a biomarker for initial response to bortezomib in relapsed patients and subsequent survival benefit after long-term treatment in newly diagnosed and relapsed patients.
INTRODUCTION

Multiple myeloma (MM) accounts for up to 10% of all hematologic malignancies. Patient outcome has improved significantly in the last decade, partially due to the introduction of novel agents, such as immunomodulatory drugs (IMiDs) as well as the proteasome inhibitors (bortezomib and second generation agents carfilzomib and ixazomib). Nevertheless, not all patients respond to these new drugs, and factors determining response are still poorly understood. \(^1\)\(^2\) In the light of high costs and higher occurrence of severe side-effects such as peripheral neuropathy associated with bortezomib treatment \(^1\)\(^3\), future efforts should be directed to developing biomarkers that can identify patients that will benefit from a particular drug or drug scheme.

MM, like other cancers, is caused by stepwise accumulation of genetic abnormalities. Chromosomal translocations involving the immunoglobulin heavy (IgH) locus causing overexpression of targeted oncogenes are primary events. The most prevalent of these translocations are t(11;14)(q13;q32) and t(4;14)(p16;q32), each present in 15% of cases. In the absence of IgH translocations, the disease is usually characterized by hyperdiploidy, specifically trisomy of odd chromosomes. Secondary hits then cause progression from asymptomatic monoclonal gammopathy of undetermined significance (MGUS), to smoldering myeloma, and finally to symptomatic myeloma with organ damage and bone lesions. Secondary events consist of mutations, chromosomal translocations and/or copy number changes. The mutational spectrum of MM has recently been characterized using genome wide next generation sequencing, which has revealed a heterogeneous mutational landscape with few recurrently affected genes. Only three genes have been found to be mutated in more than 10% of patients: \(KRAS\) (22% of cases), \(NRAS\) (20%) and \(FAM46C\) (12%).\(^4\)\(^-\)\(^7\) With regard to chromosomal abnormalities, secondary hits consist of t(8;14)(q24;q32) causing \(MYC\) activation, as well as copy number changes, with the most common ones being gains on 1q, 3p, 6p, 9p, 11q, 19p, 19q and 21q along with deletions of 1p, 4q, 16q and 22q. For some of these regions, candidate oncogenes and tumor suppressors contributing to disease progression have been identified.\(^8\)\(^-\)\(^10\)
Among the deleted regions, 1p is one of the most prevalent with up to 30% of myeloma patients carrying the deletion. Four distinct minimally deleted regions (MDRs) have been identified on 1p (1p12, 1p21.3p22.1, 1p31.1 and 1p32.3), of which 1p21.3p22.1 is the most common one (15-22% of patients).\textsuperscript{11-13} This region is associated with higher incidence of t(4;14)(p16;q32) and deletion of 17p and 13q14, and is a negative prognostic factor for progression-free survival (PFS) and overall survival (OS) in newly diagnosed patients.\textsuperscript{13} Low expression of several genes on 1p22 is part of a high risk MM gene signature.\textsuperscript{14} So far, however, no tumor suppressor has been identified on this cytoband. Previously, an MDR of 1p21.3p22.1 encompassing 35 genes was defined and MTF2 and TMED5 were proposed as candidate tumor suppressors in this region, as they show differential expression between deleted and non-deleted cases. However, no mutations were found in these genes, low MTF2 expression does not affect PFS or OS, and the association between low TMED5 expression and shorter survival is only borderline significant.\textsuperscript{11,12}

To uncover potentially clinically relevant tumor suppressors on 1p22, we performed an in-depth genetic analysis of 1p22 in MM. We delineate an MDR encompassing only 2 genes (EVI5 and RPL5) and show for each of these genes that low expression is associated with lower survival in newly diagnosed but not in relapse patients. RPL5 expression is significantly lower in relapse patients with initial response to bortezomib and both newly diagnosed and relapse patients with low RPL5 expression have better PFS when bortezomib is included in their treatment scheme. We thus identify RPL5 expression levels as a novel clinical biomarker for response to bortezomib.

**MATERIALS AND METHODS**

**Patient samples**

We studied 35 advanced diagnostic MM bone marrow samples available at UZ Leuven with at least 70% plasma cells (Supplementary Table 1). All cases were routinely characterized by FISH, as described previously.\textsuperscript{15} This study was approved by the ethical committee of the UZ Leuven. Copy number, mutation and gene expression data of the Multiple Myeloma Research Consortium (MMRC) (https://www.broadinstitute.org/mmgp/home) were also used.
survival analyses, we analyzed data from the phase III HOVON-65/ GMMG-HD4 and APEX 
trials.\textsuperscript{16-18} In the phase III HOVON-65/ GMMG-HD4 trial, newly diagnosed MM patients were 
treated with an experimental PAD protocol (bortezomib, doxorubicin, dexamethasone with 
bortezomib maintenance) or a conventional VAD protocol (vincristine, doxorubicin, 
dexamethasone with thalidomide maintenance). In the APEX trial, bortezomib versus high-
dose dexamethasone monotherapy was tested in relapsed patients. All study participants 
provided informed consent to use their data for research purposes and all studies were 
conducted according to the Declaration of Helsinki.

High resolution copy number arrays
UZ Leuven cases were analyzed on Cytoscan HD arrays (Affymetrix). Data were processed 
using Chromosome Analysis Suite (ChAS) software (Affymetrix) with hg19 as reference. In 
order to call a deletion, we required at least 20 consecutive markers with a weighted log2 ratio 
of -0.15 or less. Raw data is available at NCBI as GEO accession GSE73976.

Analysis of gene expression levels in patients
The 245 MMRC cases with copy number and expression data available on 
(https://www.broadinstitute.org/mmgp/home) were analyzed. Patients with a log2 copy 
number ratio >-0.1 for 1p22.1 were considered non-deleted; ratios <-0.6 were considered 
deleted. Patients with values between -0.1 and -0.6 were excluded.

Bortezomib response analysis
Gene expression array data from the APEX trial were used (GSE9782). Only the bortezomib 
arm of the trial was used in this analysis. Patients with complete remission, partial response 
or minimal response were assigned to the group of responders. Patients with no change or 
progressive disease were considered non-responders. Probe sets with differential signal 
between responders and non-responders were calculated using limma (Bioconductor).\textsuperscript{19} The 
entire list of probe sets was ranked according to log2 fold changes and used as input for 
Gene Set Enrichment Analysis (GSEA) against the MSigDB C2 KEGG and C1 positional 
gene sets.\textsuperscript{20,21} Only GSEA results with a FDR q-value <0.2 were considered.
Survival analysis

For the analysis on the phase III HOVON-65/ GMMG-HD4 trial, 327 patients with gene expression array profiling were included (GEO ID: GSE19784). For the APEX trial, 264 patients from which gene expression array data were available were analyzed (GEO ID: GSE9782). On each of the trial datasets, Cox regression (including both arms of the trial) testing association between the indicated gene levels and survival was performed if the proportionality criterion was met (tested using covariate time analysis). Subsequently, patients were divided into groups with expression levels above the median (high expression) and below the median (low expression) or according to the specified threshold. Kaplan-Meier analysis was then performed on patients with either low or high expression to test the prognostic value of the indicated gene expression levels on survival and to test whether these patients differ in the benefit they get from each of the treatments.

More details on methods are available in the supplement.

RESULTS

Delineation of a 58 kb MDR on 1p22.1 encompassing the \textit{EVI1} and \textit{RPL5} genes.

We analyzed 35 advanced MM samples on high resolution copy number arrays (Supplementary table 1). The 1p22 cytoband, or part of it, was deleted in 15 out of 35 cases (43%), confirming the high incidence of 1p22 deletions in this disease (Figure 1A; Supplementary table 2).\textsuperscript{11-13} Loss of 1p22 was detected in cases with hyperdiploid karyotype (7/15) or \textit{IGH}-mediated translocation/deletion (5/15). The most frequent aberration associated with the 1p22 loss was del(13q14/\textit{RB1}) (10/15) (Supplementary table 1). For 2 cases with deletions in bone marrow at diagnosis, buccal swab DNA was available and absence of the lesion in the buccal swab was confirmed by MLPA, supporting the somatic nature of these lesions (Supplementary figure 1).

To determine the MDR at 1p22, the average weighted log2 array values were calculated for each of the genes on 1p22 in our cohort (Figure 1B, Supplementary table 3), with the genes with the lowest value defining the MDR. Chromosome band 1p22.1 had lower values than
1p22.2 and 1p22.3. Of interest, patients MM05 and MM02, showed highly focal deletions of 304 kb and 366 kb with an overlap of only 58 kb. The deletion in MM05 affected the 5' parts of the EVI5 and MTF2 genes, as well as the entire RPL5 and FAM69A genes and was confirmed by FISH (Figure 1C-D). For patient MM02, it was clear that the entire GFI1 and EVI5 genes were deleted. The centromeric border in MM02 was hard to define on the array and by FISH, although it seemed within RPL5 (Figure 1C-D). To better delineate this centromeric border, the copy number status of the different exons of RPL5 was analyzed using MLPA (Figure 1E). This assay revealed that the deletion breakpoint was right within RPL5, with exons 1-4 deleted, whereas the 3’ of the gene was unaffected. We thus confirmed that RPL5 was part of the MDR. In conclusion, the MDR on 1p22.1 was restricted to a 58 kb region in our cohort, encompassing the 5’ parts of EVI5 and RPL5. EVI5 is a modulator of cell cycle progression, cytokinesis, and cellular membrane traffic. RPL5 encodes ribosomal protein L5, one of the 81 protein components of the ribosome.

**RPL5 and EVI5 are the genes on 1p22.1 with most predicted functionally impairing mutations**

We reasoned that a relevant tumor suppressor might also be targeted by mutations. Therefore, integration of the copy number data with mutation data might help in pinpointing relevant genes. Exome-wide mutation screening in large MM cohorts has been performed.\(^4\-7\) Whereas mutations in genes on 1p22 are rare in all of these studies, RPL5 was significantly mutated in one study with an incidence of 2/84 (2.4%).\(^4\) We analyzed the mutational load of all 1p22.1 genes in the MMRC exome data of 203 patients\(^6\) and calculated a mutation score for each gene on 1p22.1 (with a higher score referring to more mutations in a gene that are likely to impair protein function) (Figure 2A, Supplementary tables 4-5). Whereas EPHX4, a gene outside the MDR, had a high mutation score because of a relatively high number of gene size corrected mutations with low predicted functional impact, the EVI5 and RPL5 genes had the highest mutation scores (Figure 1B, Figure 2A).

**EVI5 and RPL5 mRNA levels are lower in 1p22.1 deleted cases**
Next, we tested if deletion of 1p22 reduces *RPL5* and/or *EVI5* expression levels. No high
quality RNA was available from our patients analyzed by copy number arrays. Therefore, we
analyzed the association between gene expression and copy number status in the MMRC
cohort. For both genes, expression levels were significantly lower in 1p22 deleted cases as
compared to non-deleted cases (p<0.0001) (Figure 2B). However, for *EVI5*, there was less
overlap in expression levels between 1p22 wild type and deleted cases than for *RPL5*, as
reflected by a slightly larger fold change in expression for *EVI5* than for *RPL5* when
comparing deleted versus non-deleted cases (0.59 versus 0.71).

**Low *EVI5* and *RPL5* expression correlates with shorter PFS and OS in newly diagnosed
but not in relapse cases**

1p22 deletion is associated with lower PFS and OS in newly diagnosed MM. Since 1p22
deletion status was not available for the trial data we had access to, we could not confirm this
previously described correlation. We tested however if *EVI5* and *RPL5* expression levels
can mirror the bad prognosis of 1p22 deletion. Cox regression analysis on the data from the
phase III HOVON-65/ GMMG-HD4 trial (referred to as HOVON-65 later on) on newly
diagnosed cases demonstrated that lower *RPL5* levels were associated with shorter PFS and
OS (Table 1). In agreement with this, Kaplan-Meier analysis showed a lower median PFS and
OS in patients with *RPL5* expression below median (’*RPL5* low’) than in patients with *RPL5*
expression above median (’*RPL5* high’) (Figure 3A). However, in the APEX trial on relapsed
patients, no association between *RPL5* levels and survival was found (Supplementary figure
2; Table 1). Similarly, low *EVI5* expression was associated with worse PFS and OS in the
HOVON-65 but not in the APEX trial (Figure 3B and Supplementary figure 2; Table 1).

Optimal cutoff for *RPL5* expression in relation to survival in the HOVON-65 trial was
determined at the 22.5% lowest *RPL5* expressers. Using this cutoff instead of median *RPL5*
expression gave a superior separation of patients in the HOVON-65 trial as well as in a
validation cohort for which OS data were available in the R2: Genomics Analysis and
Visualization Platform (Supplementary figure 3A-D). For *EVI5* expression median expression
was close to the best cutoff for PFS in the HOVON-65 trial while for OS, the optimal cutoff
was determined at the 31.5% lowest EVI5 expressers. Application of this cutoff in the validation cohort in R2 again confirmed the superior separation (Supplementary figure 3E-G).

**Bortezomib responders express lower levels of RPL5 and other ribosomal protein and translation genes**

Proteasome inhibitors are now included in most therapeutic schemes of MM patients. Cellular protein metabolism and homeostasis, which are probably affected by proteasome inhibitors, might also be altered by reduced expression levels of a ribosomal protein like RPL5. To investigate this potential association, data from the APEX clinical trial were analyzed. In this trial, relapse patients were treated with bortezomib or dexamethasone as single agent. In the bortezomib arm of the trial (n=169), RPL5 mRNA expression was significantly lower in the bortezomib responders than in non-responders (fold change responders versus non-responders: 0.68, p<0.0001) (Figure 4A). When analyzing all differentially expressed genes between responders and non-responders in this trial arm, 1211 probe sets were significant (adjusted p-value < 0.2). Interestingly, the 2 probe sets that reliably detect RPL5 ranked on the 8th and 387th position when listing the genes by significance (adjusted p-values 0.014 and 0.112) (Supplementary table 6). EVI5 was not present in this list of significant probe sets. The top 20 of differentially expressed probe sets seemed enriched for genes involved in translation (RPS7, RPL5, RPS21, EIF3M, RPS29 and EIF3H) (Table 2). In agreement with this and with previous results, GSEA revealed the ribosome as top downregulated KEGG pathway in bortezomib responders versus non-responders (Figure 4B, Supplementary table 7). Downregulation of the other ribosomal and translation associated genes besides RPL5 in our top 20 did not seem to be caused by 1p22 deletion associated downregulation of RPL5, because these genes did not differ in expression level between 1p22 deleted and non-deleted cases (Supplementary figure 4). However, expression of RPL5 did correlate with expression of each of these other genes (Supplementary figure 5), suggesting that other mechanisms besides 1p22 deletion are regulating expression of this entire ribosome – translation gene set in MM.

Based on Cox regression analysis, only RPL5 expression was significantly associated with both PFS and OS (Supplementary figure 6A). We also ran GSEA comparing the genes
differentially expressed between bortezomib responders and non-responders versus gene
sets corresponding to each chromosome cytoband. Besides 1p22, region 14q was the only
other one in the list showing recurrent deletions in MM (Supplementary Table 8). On 1p22,
four additional genes besides RPL5 were in the list of differentially expressed genes in
bortezomib responders, but none ranked as high as RPL5 (SH3GLB1: position 841 in ranked
list, LRRC8S: 157th, DR1: 342th and 643th and ZNF644: 756th) (Supplementary Table 6).
These results suggest that although some other genes on 1p22 correlate with bortezomib
response, expression of RPL5 is the best singular predictor. This is probably due to additional
effects regulating the expression of RPL5 in MM patients, possibly regulating a set of
ribosomal/translational genes as a whole. As such, in relation to bortezomib response, RPL5
expression acts as a biomarker independent of 1p22 deletion. ROC analysis was performed to
find an optimal cutoff to predict Bortezomib response (Supplementary figure 7).

RPL5 expression levels are associated with the survival benefit of bortezomib
The association between RPL5 expression levels and clinical bortezomib response raised the
question whether RPL5 levels also influence the survival benefit of bortezomib treatment. To
address this question, we divided the patients from the HOVON-65 trial in two groups
according to RPL5 expression levels below or above median, and compared survival in the
bortezomib versus non-bortezomib arm in this trial. PFS of patients with low RPL5 expression
was significantly raised when they were treated on the bortezomib arm to the point that their
PFS did not differ significantly anymore with PFS of RPL5 high patients (median PFS
bortezomib protocol 30 months versus 19 months for non-bortezomib, p=0.03; Figure 5A,
left). On the other hand PFS of RPL5 high patients was not influenced by bortezomib (median
PFS 34 months versus 33 months, p=0.94; Figure 5A, right). These findings were confirmed
on the PFS data from the APEX trial (Figure 5B). We performed the same analyses for EVI5,
but low levels of this gene did not significantly correlate with better PFS upon bortezomib
treatment (Supplementary figure 8). Cox regression analysis was also performed on the other
ribosome/translation components found in the GSEA analysis. Besides low RPL5 expression,
only low RPS7 expression correlated with improved PFS on bortezomib treatment in the
HOVON-65 and APEX trials (Supplementary figure 6B-C).
DISCUSSION

Cytoband 1p22 is deleted in ≥20% of MM patients, although no tumor suppressors have been identified. Previously, an MDR encompassing 35 genes was defined with MTF2 and TMED5 proposed as candidate tumor suppressors. However, no mutations were found in these genes, low MTF2 expression does not affect survival, and the association between low TMED5 expression and shorter survival is only borderline significant.11,12

We studied a cohort of 35 advanced MM patients and found that 40% of them carried a 1p22 deletion with two patients having a focal deletion in the region. As such, we delineated a 58 kb MDR on 1p22.1 in our cohort encompassing only two genes: EVI5 and RPL5. Interestingly, the genes in our MDR are also part of the most commonly deleted region on 1p22 in the public MMRC cohort (Supplementary figure 9). The lower resolution of the arrays used to analyze the MMRC cohort may however prohibit detection of smaller lesions affecting these genes. It is also worth noting that EVI5 and RPL5 are still part of the MRD in our cohort when removing the two cases with highly focal lesions (MM02 and MM05) from our analysis, supporting that our MRD is not purely determined by only these two cases. Although mutations in EVI5 and RPL5 are rare, they are the genes in the region with the highest frequency of mutations which are predicted to impair protein function. Additionally, as discussed below, we do find low expression of EVI5 and RPL5 to be correlated with lower PFS and OS.

Data on a potential role of EVI5 in cancer are scarce and support both tumor suppression and oncogenic functions.14,22-26 Data linking RPL5 to cancer are piling up and consistently support a tumor suppressor role. First, congenital inactivating mutations and deletions in RPL5 occur in Diamond Blackfan anemia, a rare bone marrow failure syndrome with elevated cancer risks.27,28 Also, inactivating mutations in RPL5 were recently described in T-ALL and glioblastoma29,30 and RPL5 is the only gene on 1p22 that was identified as recurrently mutated in the pan-cancer project.30-32

It is intriguing that RPL5 and EVI5 deletions are much more common than mutations. This may indicate that inactivation of both genes together is required to drive MM progression. On the other hand, the plot of RPL5 mRNA expression in 1p22 deleted versus non-deleted cases (Figure 2B) indicates that there are cases in which deletion status is a bad predictor of RPL5
expression level, suggesting that other mechanisms besides 1p22 deletion can downregulate
RPL5 expression in MM.

In the second part of this study, we looked into the clinical relevance of RPL5 and EVI5
eexpression levels in MM. First of all, we observed a worse PFS and OS in cases with low
RPL5 and EVI5 mRNA levels in the phase III HOVON-65/ GMMG-HD4 trial on newly
diagnosed patients, but not in the APEX trial on relapsed patients. These data are in line with
the known poor prognosis associated with 1p22 deletion at diagnosis. As far as we are
aware, no data are available on the prognostic value of 1p22 lesions in relapse.

Secondly, we found an association between low RPL5 mRNA levels and initial response to
bortezomib in relapse patients. Due to the lack of copy number data of the patients in the
HOVON-65 and APEX trials, we could not test association between 1p22 deletion and
bortezomib response. Our GSEA analysis for chromosomal regions enriched in the signature
of responders did identify cytoband 1p22, but this enrichment was not as convincing as the
enrichment we saw for downregulation of ribosome components. These results again indicate
that the expression of ribosomal proteins is downregulated by additional mechanisms other
than 1p22 deletion, which may make expression levels of ribosomal proteins and RPL5 in
particular, more reliable to stratify patients for bortezomib response than 1p22 deletion.

Although RPL5 expression may be useful to predict response to bortezomib in relapse
patients, validation in an independent dataset is required. In addition, in the relapse patients
tested, RPL5 levels were not perfectly associated with response, suggesting that adding as
yet unknown markers may achieve this.

Thirdly, survival analysis showed that in addition to initial response, low RPL5 expression is
also correlated with significantly longer PFS for patients treated with bortezomib both upon
diagnosis and after relapse. Unfortunately, we were unable to analyze OS data because of
cross-over of patients with progressive disease to an alternative treatment in both trials.

Although the introduction of bortezomib has greatly improved prognosis of MM, it has been
shown before that outcomes vary significantly among patient groups. Therefore, Terragna
et al. recently aimed to molecularly characterize complete response (CR) to bortezomib in
diagnostic MM. In line with our results, they found loss of a region on 1p22 to be the CNV
most significantly associated with CR after bortezomib treatment. Interestingly, the 660 kb
region identified by them includes *EVI5* but not *RPL5*. However, the size of *RPL5* and the resolution of their copy number arrays may prevent proper delineation of the boundary of the deleted area, as we also needed to perform MLPA and FISH to confirm that *RPL5* was included in one of our focal deletions. Additionally, they list another overlapping region on 1p22 as significantly associated with CR that does include *RPL5*. Terragna and colleagues did not find *RPL5* to be significantly downregulated on mRNA level in CR cases. We suspect however that this could be because of the limited number of patients they analyzed. We also only see a slight reduction in the MMRC cohort which might escape statistical cutoff in smaller cohorts.

Previously, mutations in *NRAS* have been associated with lower response rates to single-agent bortezomib treatment while high tight junction protein 1 (*TJP1*) mRNA expression has been linked to a greater likelihood of responding to bortezomib. Indeed, both *NRAS* and *TJP1* are ranked highly in our list of genes differentially expressed between responders and non-responders (Supplementary table 6). Additionally, proteomics profiling on diagnostic MM patient cells revealed that responders of bortezomib based protocols are characterized by altered expression of EIF2 signaling and by extension in refractory patients by altered expression of a subset of ribosomal proteins.

In conclusion, our genetic analyses narrow down the MDR on 1p22 to two genes: *EVI5* and *RPL5*, although the exact role of these genes in promoting MM progression remains to be determined. Survival analysis shows *EVI5* and *RPL5* expression are associated with worse survival in newly diagnosed patients. In addition, low *RPL5* expression levels are linked to initial bortezomib response in relapse patients and to survival benefit on bortezomib treatment for both diagnostic and relapse patients, leading to a complete recovery of the bad prognosis of low *RPL5* levels. We thus identify *RPL5* mRNA expression as a novel biomarker correlating with benefit from bortezomib treatment.
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AUTHORSHIP CONTRIBUTION

IH designed and performed experiments and wrote the manuscript. MVD and PS performed outcome and gene expression analyses and critically reviewed the manuscript. EDB designed research and performed experiments. LF helped with bio-informatic analyses. GM provided survival and expression data and critically reviewed the manuscript. E Geerdens performed copy number arrays. E Garelli, CM and AA performed MLPA. HL performed FISH. IW designed FISH, wrote and critically reviewed the manuscript. MD, LM and PVDB collected patient samples and critically reviewed the manuscript. KVDK designed research and performed supervision. KDK designed and performed research, supervised the entire study and wrote the manuscript.

DISCLOSURE OF CONFLICTS OF INTEREST

GM discloses employment with Takeda Pharmaceuticals International Co.
REFERENCES


12 Boyd KD, Ross FM, Walker BA, Wardell CP, Tapper WJ, Chiecchio L et al. Mapping of Chromosome 1p Deletions in Myeloma Identifies FAM46C at 1p12 and CDKN2C at 1p32.3 as


FIGURE LEGENDS

Figure 1. Genetic analysis of chromosomal region 1p22 reveals \textit{RPL5} and \textit{EVI5} as candidate tumor suppressors

(A) Ideogram of chromosome 1 with indication of the size of the 1p22 deletions detected in this study. (B) Average weighted log2 array ratio of 1p22 genes. Each dot in the graph represents a gene on 1p22, and genes are represented from telomeric (left, 85 Mbp) to centromeric (right, 95 Mbp). The region that shows the lowest average weighted log2 array ratio defining the MDR in our cohort is indicated in red and gene names are shown. (C) Focal 1p22.1 deletions in cases MM05 and MM02. Deleted areas are indicated by the grey shaded areas, the dark grey shaded area shows the overlapping deleted region (MDR). (D) FISH with probes RP11-1E09 and RP11-456E23 on bone marrow of patients MM05 and MM02. Schematic representation of the genomic region targeted by the FISH probes is shown in panel C. Note loss of one RP11-1E09 (green) signal in MM02 and loss of one RP11-456E23 (green) signal in MM05. The latter case displayed both RP11-1E09 (red) signals, because the BAC covers a larger region than the deleted sequences in MM05. (E) MLPA assay confirming deletion of exon 1-4 of the \textit{RPL5} gene in case MM02. The assay measures copy number of exons of several ribosomal protein genes. Peaks representing signals that correspond to exons of \textit{RPL5} are indicated, with deleted exons in red and non-deleted exons in black.

Figure 2. \textit{EVI5} and \textit{RPL5} show the highest incidence of mutations predicted to impair protein function and their expression levels correlate with 1p22 deletion status

(A) Mutation score for all 1p22.1 genes. Mutation score per gene was calculated by taking the mutation count for each gene and by correcting this value for gene length and predicted functional impact of the mutations. The MDR identified in Figure 1B is again indicated in red. (B) Gene expression plots of \textit{EVI5} (probe set 209717\_s\_at) and \textit{RPL5} (200937\_s\_at) in 1p22 wt versus deleted cases. The red horizontal lines indicate the average value in the group and the standard deviations. p-values were calculated using a 2-tailed Mann-Whitney test. The fold change below the plot indicates the fold downregulation in 1p22 deleted cases.
Figure 3. Low EVI5 and RPL5 expression correlates with shorter PFS and OS in newly diagnosed but not in relapse patients

(A) Kaplan-Meier curves comparing PFS (left) and OS (right) of RPL5 low and high expressing cases in the phase III HOVON-65/ GMMG-HD4 trial. (B) PFS and OS of EVI5 low and high expressing cases in the phase III HOVON-65/ GMMG-HD4 trial. ‘Low’ and ‘high’ are defined here as expression below and above median. p-values were calculated with Log-rank tests.

Figure 4. Reduced expression of RPL5 and other ribosomal proteins correlates with response to bortezomib

(A) Gene expression plots of RPL5 (probe set 200937_s_at) in bortezomib responders and non-responders in the APEX trial. The red horizontal lines indicate the average value in the group and the standard deviations. P-values were calculated using a 2-tailed Mann-Whitney test. The fold change indicates the fold downregulation in responders. (B) GSEA plot supporting downregulation of the genes in KEGG pathway ‘ribosome’ in bortezomib responders.

Figure 5. RPL5 expression levels are associated with the benefit of bortezomib on PFS

(A) Kaplan-Meier curves comparing PFS of RPL5 low (left) and high (right) expressing patients for bortezomib versus non-bortezomib arms in the HOVON-65/ GMMG-HD4 trial. (C) Kaplan-Meier curves comparing PFS of RPL5 low (left) and high (right) expressing patients for bortezomib versus dexamethasone arms in the APEX trial. Low and high expression are defined here as below and above median. All p-values were calculated with Log-rank tests.
Figure 2

A  

Mutation score

<table>
<thead>
<tr>
<th>Gene</th>
<th>Mutation Score</th>
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<tbody>
<tr>
<td>CDC7</td>
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<tr>
<td>TGFB3</td>
<td>0.29</td>
</tr>
<tr>
<td>BRDT</td>
<td>0</td>
</tr>
<tr>
<td>EPHX4</td>
<td>0</td>
</tr>
<tr>
<td>BTBD8</td>
<td>0</td>
</tr>
<tr>
<td>KIAA1107</td>
<td>0</td>
</tr>
<tr>
<td>C1orf146</td>
<td>0.21</td>
</tr>
<tr>
<td>GLMN</td>
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</tr>
<tr>
<td>RPAP2</td>
<td>0</td>
</tr>
<tr>
<td>GFI1</td>
<td>0</td>
</tr>
<tr>
<td>EVI5</td>
<td>0.74</td>
</tr>
<tr>
<td>RPL5</td>
<td>0.85</td>
</tr>
<tr>
<td>FAM69A</td>
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<tr>
<td>MTF2</td>
<td>0</td>
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<tr>
<td>TMED5</td>
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<td>CCDC18</td>
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<td>DR1</td>
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<td>FNBP1L</td>
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<td>DNTTIP2</td>
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<tr>
<td>GCLM</td>
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<tr>
<td>ABCA4</td>
<td>0.05</td>
</tr>
</tbody>
</table>

B

Log2 normalized expression

-2 -1 0 1 2

1p22 wt 1p22 del

Fold change: 0.59

EVI5

p<0.0001

Fold change: 0.71

RPL5

p<0.0001
Figure 3

A

HOVON-65/ GMMG-HD4

Median survival (Months)

- RPL5 low (n=163) 25.03
- RPL5 high (n=164) 33.31

Months

PFS (%)

OS (%)

p=0.03

p=0.05

B

HOVON-65/ GMMG-HD4

Median survival (Months)

- EVI5 low (n=164) 25.82
- EVI5 high (n=163) 33.31

Months

PFS (%)

OS (%)

p=0.04

p=0.01
Figure 4

A

**RPL5**

```
Log2 expression
```

- Bortezomib Responders
- Bortezomib Non-Responders

Fold change: 0.68

B

**KEGG_RIBOSOME**

- up in bortezomib responders
- down in bortezomib responders

FDR: 0.000
Figure 5

A

**HOVON-65/ GMMG-HD4**

**RPL5 low**

<table>
<thead>
<tr>
<th></th>
<th>Bortezomib (n=86)</th>
<th>Control (n=77)</th>
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</thead>
<tbody>
<tr>
<td>PFS (%)</td>
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<tr>
<td>Months</td>
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<tr>
<td>p</td>
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</table>

<table>
<thead>
<tr>
<th>Median PFS (Months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bortezomib</td>
</tr>
<tr>
<td>Control</td>
</tr>
</tbody>
</table>

B

**HOVON-65/ GMMG-HD4**

**RPL5 high**

<table>
<thead>
<tr>
<th></th>
<th>Bortezomib (n=83)</th>
<th>Control (n=81)</th>
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<tbody>
<tr>
<td>PFS (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Months</td>
<td></td>
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<tr>
<td>p</td>
<td>0.94</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Median PFS (Months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bortezomib</td>
</tr>
<tr>
<td>Control</td>
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</tbody>
</table>

B

**APEX**

**RPL5 low**

<table>
<thead>
<tr>
<th></th>
<th>Bortezomib (n=84)</th>
<th>Dexamethasone (n=36)</th>
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</thead>
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<tr>
<td>PFS (%)</td>
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</tr>
<tr>
<td>Days</td>
<td></td>
<td></td>
</tr>
<tr>
<td>p</td>
<td>0.01</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Median PFS (Days)</th>
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</thead>
<tbody>
<tr>
<td>Bortezomib</td>
</tr>
<tr>
<td>Dexamethasone</td>
</tr>
</tbody>
</table>

B

**APEX**

**RPL5 high**

<table>
<thead>
<tr>
<th></th>
<th>Bortezomib (n=89)</th>
<th>Dexamethasone (n=33)</th>
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<tbody>
<tr>
<td>PFS (%)</td>
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<td></td>
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<tr>
<td>Days</td>
<td></td>
<td></td>
</tr>
<tr>
<td>p</td>
<td>0.81</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Median PFS (Days)</th>
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<tbody>
<tr>
<td>Bortezomib</td>
</tr>
<tr>
<td>Dexamethasone</td>
</tr>
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</table>
Table 1. Cox regression values

Cox regression analysis of OS and PFS (Figure 3)

<table>
<thead>
<tr>
<th></th>
<th>PFS</th>
<th>OS</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Exp (B)</td>
<td>95% CI</td>
</tr>
<tr>
<td>HOVON-65/GMMG-HD4</td>
<td></td>
<td></td>
</tr>
<tr>
<td>RPL5</td>
<td>0.81</td>
<td>0.70-0.93</td>
</tr>
<tr>
<td>EVI5</td>
<td>0.89</td>
<td>0.79-1.01</td>
</tr>
<tr>
<td>APEX</td>
<td></td>
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<tr>
<td>RPL5</td>
<td>1</td>
<td>0.85-1.14</td>
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<tr>
<td>EVI5</td>
<td>0.98</td>
<td>0.85-1.12</td>
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</tbody>
</table>

Cox regression analysis on PFS in RPL5 low versus high expressing cases for bortezomib versus control treatment (Figure 5)

<table>
<thead>
<tr>
<th></th>
<th>PFS</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exp (B)</td>
</tr>
<tr>
<td>HOVON-65/GMMG-HD4</td>
<td></td>
</tr>
<tr>
<td>RPL5 low: Bz vs. CTRL</td>
<td>0.67</td>
</tr>
<tr>
<td>RPL5 high: Bz vs. CTRL</td>
<td>0.99</td>
</tr>
<tr>
<td>APEX</td>
<td></td>
</tr>
<tr>
<td>RPL5 low: Bz vs. CTRL</td>
<td>0.57</td>
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<tr>
<td>RPL5 high: Bz vs. CTRL</td>
<td>0.95</td>
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Table 2. Top 20 of probe sets with differential signal in bortezomib responders versus non-responders in the APEX trial

<table>
<thead>
<tr>
<th>Probe</th>
<th>Gene</th>
<th>log2FC</th>
<th>P adj</th>
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<tbody>
<tr>
<td>210532_s_at</td>
<td>C14orf2</td>
<td>-0.408</td>
<td>0.002</td>
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<tr>
<td>225335_at</td>
<td>ZNF496</td>
<td>-0.540</td>
<td>0.002</td>
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<tr>
<td>217988_at</td>
<td>CCNB1IP1</td>
<td>-0.590</td>
<td>0.002</td>
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<tr>
<td>229586_at</td>
<td>CHD9</td>
<td>0.424</td>
<td>0.008</td>
</tr>
<tr>
<td>224985_at</td>
<td>NRAS</td>
<td>-0.470</td>
<td>0.008</td>
</tr>
<tr>
<td>224616_at</td>
<td>DYNC1LI2</td>
<td>0.459</td>
<td>0.014</td>
</tr>
<tr>
<td>213941_x_at</td>
<td>RPS7</td>
<td>-0.340</td>
<td>0.014</td>
</tr>
<tr>
<td>200937_s_at</td>
<td>RPL5; SNORD21</td>
<td>-0.553</td>
<td>0.014</td>
</tr>
<tr>
<td>206790_s_at</td>
<td>NDUFB1</td>
<td>-0.388</td>
<td>0.019</td>
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<tr>
<td>200834_s_at</td>
<td>RPS21</td>
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<td>202232_s_at</td>
<td>EIF3M</td>
<td>-0.521</td>
<td>0.019</td>
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<tr>
<td>224841_x_at</td>
<td>GAS5; SNORD44; SNORD47; SNORD74; SNORD76; SNORD77; SNORD79; SNORD80; SNORD81</td>
<td>-0.854</td>
<td>0.019</td>
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<tr>
<td>224741_x_at</td>
<td>GAS5; SNORD44; SNORD47; SNORD74; SNORD76; SNORD77; SNORD79; SNORD80; SNORD81</td>
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<tr>
<td>221180_at</td>
<td>MAP3K19</td>
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<td>208752_x_at</td>
<td>NAP1L1</td>
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<td>0.019</td>
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<td>213846_at</td>
<td>COX7C</td>
<td>-0.394</td>
<td>0.019</td>
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<tr>
<td>238025_at</td>
<td>MLKL</td>
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<td>0.019</td>
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<td>200921_s_at</td>
<td>BTG1</td>
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<td>201094_at</td>
<td>RPS29</td>
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<td>201592_at</td>
<td>EIF3H</td>
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