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Differential roles for MBD2 and MBD3 at methylated CpG
islands, active promoters and binding to exon
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Differential roles for MBD2 and MBD3 at methylated CpG islands, active promoters and binding to exon sequences

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ABSTRACT

The heterogeneous collection of nucleosome remodelling and deacetylation (NuRD) complexes can be grouped into the MBD2- or MBD3-containing complexes MBD2–NuRD and MBD3–NuRD. MBD2 is known to bind to methylated CpG sequences in vitro in contrast to MBD3. Although functional differences have been described, a direct comparison of MBD2 and MBD3 in respect to genome-wide binding and function has been lacking. Here, we show that MBD2–NuRD, in contrast to MBD3–NuRD, converts open chromatin with euchromatic histone modifications into tightly compacted chromatin with repressive histone marks. Genome-wide, a strong enrichment for MBD2 at methylated CpG sequences is found, whereas CpGs bound by MBD3 are devoid of methylation. MBD2-bound genes are generally lower expressed as compared with MBD3-bound genes. When depleting cells for MBD2, the MBD2-bound genes increase their activity, whereas MBD2 plus MBD3-bound genes reduce their activity. Most strikingly, MBD3 is enriched at active promoters, whereas MBD2 is bound at methylated promoters and enriched at exon sequences of active genes.

INTRODUCTION

The nucleosome remodelling and deacetylation (NuRD) complex harbours a multi-functional and highly conserved combination of chromatin-modifying activities. Through the MBD2 and MBD3 proteins, with their methyl-CpG-binding domains (MBD), the NuRD complex combines reading of DNA methylation marks with modifying histones [for recent reviews see (1,2)]. Through the combinatorial assembly of similar, paralogous versions of histone deacetylases, nucleosome-remodelling ATPases, metastasis-associated (MTA) factors and others, biological specificity of the NuRD complex is achieved during development, oncogenesis and cancer progression.

Specificity mediated by or associated with MBD2 and MBD3 has been documented on several levels. MBD2-knockout mice are viable and fertile with subtle defects only, whereas MBD3-knockout mice are embryonic lethal (3). On the molecular level, both proteins differ in respect to binding to methylated DNA. MBD2 is able to bind DNA with a 5-methylcytosine (5 mC) modification (4,5), in contrast to MBD3, which binds to 5-hydroxymethylcytosine (5 hmC), but not to 5 mC (4–7). Purification and analysis of NuRD complexes revealed that MBD2 and MBD3 are components of mutually exclusive NuRD complexes, MBD2–NuRD and MBD3–NuRD (8). Despite these and other differences, both factors have been shown to interact with GATAD2A/p66α and GATAD2B/p66β within the NuRD complex (9–11), as well as with CSB (Cockayne Syndrome Protein B) (12), HIC1 (Hypermethylated in Cancer 1) (13) and DOC-1 (Deleted in Oral Cancer 1) (14).

Testing of Hela cell promoter regions revealed preferential MBD2 binding near transcriptional start sites (TSS) (15). Genome-wide binding analysis of NuRD complexes in ES cells, which are primarily MBD3–NuRD complexes, revealed a broad binding near promoters, but with a gap at the TSS (16). In contrast, ChIPseq analysis of MBD3 in ES cells revealed a sharp occupancy at the TSS (7). In addition to these published differences for MBD3–NuRD binding distribution, a direct comparison between MBD2 and MBD3 on the whole genome level is missing.

Because of the increasing information for both functional differences as well as similarities, we wanted to know whether MBD2 and MBD3 differ in respect to chromatin modification and in genome-wide binding. Functional tests revealed a dramatic difference in that MBD2–NuRD, but not MBD3–NuRD, transformed...
euchromatin into repressed chromatin. Analysis of the genome-wide binding pattern of MBD2 and MBD3 within the same cell type showed a preference for MBD2 to be bound at methylated CpG islands and inactive promoters, whereas MBD3 was found at unmethylated CpG islands and active promoters. Most strikingly, exon sequences of active genes were enriched for MBD2 binding.

MATERIALS AND METHODS

Antibodies

Immunostaining was done using commercial antibodies recognizing Mi2 (Santa Cruz, sc-11378), RbAp46 (Santa Cruz, sc-8272), HDAC1 (Santa Cruz, sc-9397), MBD2 (Santa Cruz, sc-9397) and MBD3 (Santa Cruz, sc-9402). For ChIP H3K9ac (Abcam, 4441), H3K9me3 (Abcam, 8898), GFP (polyclonal rabbit antiserum was raised against full length GFP), MBD2a/b (Sigma Aldrich, M-7318), MBD3 (Bethyl Laboratories A302-528A), normal rabbit control IgG (Abcam, ab46540) antibodies and in addition V5-agarose (Sigma Aldrich, A7345) was used. GFP, MBD2a/b and MBD3 antibodies were used for Western Blot as well.

Luciferase assay

A10 cells were co-transfected with 1 μg of either GFP-LacI, GFP-LacI-VP16, GFP-LacI-MBD2a, GFP-LacI-MBD2b or GFP-LacI-MBD3 and βGAL-vector using TurboFect (Fermentas) in six-well plates. Cells were harvested 48 h after transfection and lysed with 300 μl lysis-buffer (25 mM Tris/HCl pH 7.5, 8 mM MgCl2, 1 mM EDTA, 1% Triton X 100, 15% glycerin and freshly added 1 mM DTT) per well. Luciferase reporter activity was determined and corrected for βGAL expression.

Data deposition

The microarray and the ChIPseq data from this publication have been submitted to the GEO database as entry GSE41010.

Cell lines and transfection, Immunofluorescence analysis, Western Blot, Chromatin immunoprecipitation and ChIPseq analysis and bioinformatics analyses

See Supplementary online material.

RESULTS

The NuRD complex can be assembled on LacO repeats

To study possible MBD2- or MBD3-induced changes in chromatin compaction, we used the LacO/LacI system that uses 100s of LacO repeats integrated into single genomic loci (17). In many cases, genomic integration of the repeat cluster is found in heterochromatic regions. For such a situation, we used the F42B8 cell clone of U2OS cells, with a repeat cluster integrated close to the centromere (18). As a model for a euchromatic array, we used the RREB1 cell clone of CHO cells (19). In a first set of experiments, we wanted to know whether an MBD2 or MBD3 fusion with the DNA-binding domain of the Lac repressor LacI can be targeted to a LacO array, and whether such a bound protein recruits additional components of the NuRD complex. For this, we started with the heterochromatic array of the F42B8 cells, as the compact structure of such an array can be easily visualized and analysed by cytological methods. As a requirement for these experiments, we analysed the heterochromatic nature of the array to assure that it is lacking NuRD components. The array can be easily identified by expression of a GFP-Lac fusion (Supplementary Figure S1) and was found to be devoid of NuRD components. In contrast to the expression of GFP-LacI, both MBD fusions, GFP-LacI-MBD2b and GFP-LacI-MBD3, resulted in positive staining by immunofluorescence with antibodies against the NuRD components Mi2, HDAC1 and RbAp46 (Figure 1A). Thus, these NuRD factors were recruited by MBD2 or MBD3 targeted to the LacO array.

For MBD2, two isoforms are known, MBD2b and MBD2a, that contains an N-terminal extension. Within the assays described in this manuscript, there was no difference seen between using GFP-LacI-MBD2a or GFP-LacI-MBD2b (Supplementary Figure S2). Recruitment of GATAD2B was detected by co-expressing Cherry-GATAD2B (Figure 1A). When using GATAD2B fused to GFP-LacI (GFP-LacI-GATAD2B), the NuRD components Mi2, HDAC1 and RbAp46 could be similarly recruited to the LacO array (Supplementary Figure S3). Furthermore, GFP-LacI-GATAD2B also recruited MBD2 and MBD3 to the array. This might potentially be a source for generating mixed complexes, i.e. MBD2 recruits GATAD2B, which in turn might recruit MBD3. To test for this potential problem, we co-expressed GFP-LacI-MBD2 with Cherry-MBD3 or GFP-LacI-MBD3 with Cherry-MBD2. These combinations clearly showed that binding of neither of the MBD factors resulted in recruitment of the paralogous factor to the array and that only homogeneous NuRD complexes were generated on the array (Supplementary Figure S4). In addition, we targeted HDAC1 to the LacO array by the use of GFP-LacI-HDAC1, but this did not result in assembly of the NuRD complex (Supplementary Figure S5). The results of these experiments are summarized in Figure 1B and demonstrate that the LacO array system is suitable to study NuRD. Furthermore, MBD2, MBD3 or GATAD2B can be used as starting component to assemble the NuRD complex. When targeting MBD2 to the DNA, only MBD2–NuRD is assembled, as is the case for MBD3, which assembles solely MBD3–NuRD.

In contrast to MBD3, MBD2 converts euchromatin into repressed chromatin

To study the repressive function of NuRD, we used the above LacO/LacI targeting system within a euchromatic background. This was provided by the RREB1 cell clone (19). NuRD complex assembly was similarly possible with the heterochromatin array of the F42B8 cells above (Supplementary Figure S6). After transfection of GFP-LacI, the expanded array can be visualized. Upon expression of GFP-LacI-MBD2, the expanded array
dramatically shrinks to a small spot similarly as seen for the heterochromatin array of the F42B8 cells. This array compression is observed when targeting MBD2a or MBD2b to the LacO array (Figure 2), but is not seen when GFP-LacI-MBD3 is expressed. Similarly, the array remains expanded when expressing GFP-LacI-HDAC1, a factor that is not able to assemble the NuRD complex on the LacO array (Figure 1B). If array binding, and therefore compaction, was indeed caused by the DNA-bound MBD2–NuRD complex, recruitment should be dependent on the LacI DNA-binding domain. Indeed, the deleted DNA-binding domain in construct GFP-LacI-MBD2b resulted in a nuclear GFP staining not localized to the LacO array (Supplementary Figure S7). We wanted to test whether the compaction of the chromatin array is marked by the chromatin modification H3K9me3, which is specific for inactive and repressed chromatin. Therefore, we purified chromatin for immune-precipitation with antibodies specific for H3K9me3 or H3K9ac. The ratio of these precipitations is a mark for active or for repressed chromatin. In contrast to cells transfected with GFP-LacI, expression of GFP-LacI-MBD2a resulted in a strong increase of the repressive mark and a similarly strong decrease of the active mark. In contrast, expression of GFP-LacI-MBD3 induced a lower decrease in H3K9ac and a lower increase in H3K9me3 (Supplementary Figure S8). This low, but reproducible, effect of MBD3 in the absence of chromatin compaction may indicate that these modification changes are not sufficient to compact chromatin as is the case with MBD2.

To study the relationship between the chromatin status and gene activity, we generated HeLa cell clones containing an integrated luciferase reporter gene. The luciferase gene is controlled by seven copies of the LacO sequence, such that any LacI fusion protein (Figure 3A) can be tested for its regulatory capacity. We selected a single clone (A10), from several different cell clones based on the robust ground state of luciferase activity, which was expected to be further increased by an activator or to be reduced by a repressor. Indeed, transient control transfection with an expression plasmid coding for a GFP-LacI-VP16 fusion resulted in a 8-fold increase of luciferase activity (Figure 3C). Expression of either MBD2a or MBD2b led to a ∼10-fold reduction of luciferase activity, in contrast to the GFP-LacI control or to GFP-LacI-MBD3. Clearly, MBD2 mediates strong chromatin compaction and transcriptional repression, whereas MBD3 does not.
Therefore, we determined the chromatin modification at the reporter gene with antibodies against H3K9me and against H3K9ac. The ratio of methylated K9 versus acetylated K9 is an indication for repressed chromatin. GFP-LacI-MBD2a and GFP-LacI-MBD2b expression resulted in an increased repression ratio in contrast to GFP-LacI-MBD3, which did not change the repression ratio as compared with the GFP-LacI control expression (Figure 3E and F).

**Both MBD2 or MBD3 bind to similar sequence classes**

Because of the functional differences of MBD2 and MBD3, we wanted to know whether both factors are targeted to factor-specific sites throughout the genome. Such an analysis has not yet been performed within a single cell type. The quality and specificity of available antibodies directed against these two proteins is a limiting factor in carrying out chromatin precipitation experiments. Therefore, we generated MBD versions fused to the V5 domain. This domain has been shown to be a universal and optimal tag for ChIPchip or ChIPseq analysis (20).

Expression of V5, V5-MBD2a and V5-MBD2b in HeLa cells did not alter cell proliferation. Western-blot detection of the tagged MBD factors revealed similar amounts of proteins expressed from the different constructs (Supplementary Figure S9A). Therefore, we generated chromatin from HeLa cells after expression of these three constructs and purified V5 bound DNA for ChIPseq. We identified regions of significant binding for MBD2 and MBD3 by comparing precipitations from cells transfected with V5-MBD2 and V5-MBD3 with those

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**Figure 3.** In contrast to GFP-LacI-MBD3, GFP-LacI-MBD2a or GFP-LacI-MBD2b repress the euchromatic LacO promoter. (A) Illustration of the reporter-gene including seven LacI-binding sites (LacO-repeats). Reporter construct was used to generate stable reporter cell clones in HeLa cells. (B) Western Blot analysis of the indicated GFP-LacI-fusion proteins expressed in the stable A10 clone. GAPDH antibody was used as loading control. (C) The A10 clone was transiently transfected with either GFP-LacI, GFP-LacI-VP16, GFP-LacI-MBD2a, GFP-LacI-MBD2b or GFP-LacI-MBD3. Gene activity is expressed as fold change relative to the untransfected A10 clone. (D) Binding analysis by ChIP-qPCR was performed after transfection of the indicated constructs using a GFP antibody and expressed relative to untransfected cells. (E, F) GFP-LacI-MBD2a and GFP-LacI-MBD2b change the H3K9me3/H3K9ac ratio after binding to the reporter. ChIP-qPCR was performed with antibodies recognizing H3K9ac (E) and H3K9me3 (F) and expressed relative to input. Error bars indicate the standard error of three individual replicates.
from cells transfected with the vector expressing the V5 tag only. We used PeakRanger to call peaks with a $P$-value $<10^{-5}$ at an FDR $<5\%$. Using this approach, we were able to detect $\sim8200$ MBD2- and 490 MBD3-binding regions. Despite this difference in the number of binding sites, both factors were found to bind to similar sequence classes. The peak distribution indicates highly specific binding to transcriptional start sites including upstream sequences (Figure 4A and B). Furthermore, exon sequences are highly enriched including transcriptional end sites. In contrast, intron sequences are underrepresented as compared with the total genomic sequence distribution (Figure 4A and B). In contrast to sequences associated with genes, intergenic sequences are almost depleted from either MBD2 or MBD3 binding (Figure 4B).

Binding specificity was verified at positive and negative sites (Supplementary Figure S9B). To confirm that MBD2- or MBD3-binding sites are represented by the V5-tagged MBD proteins, we used the host HeLa cells without transfection and immune-precipitated chromatin with antibodies directed against MBD2 or MBD3. Although the specificity and efficiency of these antibodies was low, we could confirm MBD2 and MBD3 binding even without expression of the tagged protein (Supplementary Figure S10).

Binding to the transcriptional start sites was highly position specific with a sharp peak at the TSS (Figure 4C). Because of the lower number of MBD3-bound sites, the binding peak at the TSS was less pronounced. No specific enrichment of promoter DNA could be detected in the ChiPseq data from cells transfected with the empty V5 vector.

Because similar sequence classes are bound by both MBD factors, we tested for a possible overlap of binding sites. We determined the cumulative binding profiles for MBD2 and MBD3 across MBD2 and MBD3 peak regions. As expected MBD2 and MBD3 binding can be detected across the respective peak regions (Figure 4D). Interestingly, we are able to detect strong MBD2 binding even across MBD3 peak regions (Figure 4D, central panel; for genome-browser examples see Supplementary Figure S11), although the analysis of overlapping peaks suggested only moderate overlap between MBD2 and MBD3 peak ranges (Figure 5A). In contrast, we cannot detect much MBD3 binding across MBD2 peak regions (Figure 4D, right panel).

Thus, overall binding of MBD2 or MBD3 is targeted to similar sequence classes and MBD3 sites are often marked by robust MBD2 binding.

MBD2 sites are highly enriched for methylated DNA, whereas MBD3 sites are devoid of methylation

Despite the similarity of the MBD domains of MBD2 and MBD3, it has been shown that only MBD2 is able to bind to methylated CpG islands in vitro (4). The genome-wide methylation pattern of HeLa cells is known [GSE40699 (22)]. CpG dinucleotides are underrepresented in the genome and occur in two classes, dispersed throughout the genome and clustered within CpG islands (23). Both MBD2- and MBD3-binding sites are highly enriched for CpG islands (Figure 5A). Seventy-nine per cent of the MBD2 peaks are overlapping with CpG islands and in case of MBD3, 69% are overlapping. The MBD2 overlap with CpG islands involves a sub-fraction (22%) of the CpG islands only. Taking into account that a sub-fraction of CpG islands in HeLa cells is methylated, we determined binding of both MBD factors to methylated CpG islands. For MBD2, a high percentage of the MBD2-bound CpG islands were methylated, whereas for MBD3, <10% of the bound CpG islands were methylated (Figure 5B).

Thus, although both factors bind to similar sequence elements such as TSS and CpG islands, their binding differs dramatically in respect to CpG methylation.

MBD2 is associated with repressed genes and MBD3 with activated genes

Because we have seen the striking difference between MBD2 and MBD3 in mediating reporter-gene repression and chromatin compaction, we compared MBD2 and MBD3 promoter binding in respect to gene activity. Indeed, MBD2 is bound to promoters of genes with low transcriptional activity, whereas MBD3-bound promoters show a 4-fold higher gene activity, as determined from the mean values (Figure 6A). When comparing these differences with the activity of all genes in HeLa cells, MBD2 was found to be associated with genes with lower transcriptional activity as compared with all genes. In contrast, gene activity of MBD3-associated genes was higher than the mean activity of all genes (Figure 6A).

To gain insights into the MBD2 or MBD3 function in the context of the endogenous gene repertoire, we depleted HeLa cells by RNAi against either MBD protein or as a negative control with control siRNA, determined the respective expression profiles on the Affymetrix GeneChip® Gene 1.0 ST Array platform and verified expression changes at individual genes (Supplementary Figures S12 and S13). In almost all cases, when well-studied repressors have been depleted, the expression profiles of target genes not only increased, but also a substantial number of genes decreased their expression. Similarly, depletion of well-characterized activators resulted in decreased as well as in increased gene activity (e.g. data compiled at http://www.ncbi.nlm.nih.gov/geo/). One of several reasons for this is an indirect effect on genes that do not harbour a binding site for the respective factor. Therefore, we focussed our view on 1080 genes with MBD2 binding within the promoter region (Figure 6B). The majority of these genes show a mild change in gene activity towards activation. In comparing MBD2 depletion with MBD3 depletion of MBD2-bound genes, an increase in gene activity of >75% is seen after MBD2 RNAi (red curve in Figure 6C shifted to the right). Because the expression changes are rather low, we wanted to determine the significance of this observation. Therefore, we determined the expression change of all genes, irrespective of a MBD2-binding site (Figure 6C) and found that 50% of these are either induced or repressed. This difference between MBD2-bound genes and all genes was significant,
with a $P$ value of $7 \times 10^{-76}$. Expression changes after MBD3 depletion of MBD2-bound genes resulted in equal gene numbers repressed or induced. Similarly, we analysed genes with MBD2 binding to exon sequences (see below) and found that these genes responded with a slight shift towards activation upon MBD2 depletion (Figure 6C green line). Similar results were achieved, when we used the gene set enrichment analysis (GSEA; Supplementary Figure S14). Thus, again MBD2 functions primarily as a repressor at MBD2-bound genes.

Because MBD3 is found at non-methylated DNA (see above), we wanted to test for a possible functional difference of both MBD factors by analysing gene expression changes of 271 MBD3-bound genes upon MBD3 depletion. Small changes were found, resulting in both slightly increased or decreased activity of genes (Figure 6D). The binding analysis of MBD3-bound promoters shows a significant number of cases with MBD2 binding as well (heatmap in Figure 6D). In contrast to the whole genome binding analysis, where we applied a stringent peak-calling algorithm (Macs) with less stringent settings. This revealed ~140 of MBD3-bound promoters to be significantly associated with MBD2 peaks as well. Therefore, we asked for the effect of MBD2 depletion on the activity of these genes. For these, we found expression changes towards repression (red curve in Figure 6E shifted to the left). The difference in expression change after MBD2 depletion at MBD2 plus MBD3-bound genes and expression change of all genes after MBD3 depletion (black curve in Figure 6E) is significant ($P < 6 \times 10^{-5}$).

Therefore, we can conclude that MBD2 primarily causes gene repression, whereas MBD2 in the context of MBD3 promoter binding might be involved in gene activation.

**In contrast to MBD2, MBD3 is associated with chromatin modification characteristic for active chromatin**

To identify additional chromatin features MBD2 and MBD3 might be associated with, we compared the binding sites with known features of histone modification in HeLa cells. First, we identified that MBD3-bound sites are generally associated with open chromatin as indicated...
compared with the average island methylation (Wilcoxon signed-rank test) whereas MBD3-associated islands are devoid of methylation as defined from (21). MBD2-associated islands show high methylation, islands associated with MBD2, MBD3 or of all CpG islands (island definition)

**Figure 5.** MBD2 sites are highly enriched for methylated DNA, whereas MBD3 sites are devoid of DNA methylation. (A) Venn diagram shows overlap of MBD2- and MBD3-binding regions with CpG islands. (B) Average relative (percentage) CpG methylation level within CpG islands associated with MBD2, MBD3 or of all CpG islands [island definition from (21)]. MBD2-associated islands show high methylation, whereas MBD3-associated islands are devoid of methylation as compared with the average island methylation [Wilcoxon signed-rank test: \( P = 0 \) (MBD2) and \( P = 3.5 \times 10^{-7} \) (MBD3)].

by the precise peaks in the FAIRE (Formaldehyde-Assisted Isolation of Regulatory Elements) (24) and DNAseI profiles. In contrast, MBD2 sites are not in an open conformation.

Additionally, we found for MBD3 a correlation with active promoter marks H3K4me2, H3K4me3 and H3K9ac at MBD3-binding sites, which is in line with MBD3 binding to promoters of actively transcribed genes (Figure 6A) and to non-methylated CpG islands (Figure 5B), which often are found at active promoters. Indeed, MBD3 binding clearly correlated with these active promoter marks (Figure 7) with the characteristic ‘dip’ of the respective marks, since transcriptional start sites and other regulatory sites are depleted from nucleosomes. In general, sites occupied by both MBD factors (grey line in Figure 7) show similar features as the MBD3 sites. This binding is in stark contrast to MBD2, which does not show any correlation with these marks. A similar difference for the two MBD factors is seen with polymerase II binding at MBD3 sites, but not at MBD2-bound regions. Contrary, testing specifically for marks associated with silent chromatin, such as H3K9me3, we do not find a specific enrichment at MBD2 sites.

Because we have seen for both, MBD2 and MBD3, an enrichment over gene and exon sequences (Figure 4A and B), we wanted to compare the binding profiles with chromatin marks characteristic for the gene body and for transcriptional elongation, such as H3K36me3 and H3K79me2 (25–27). These showed a preference for MBD3 binding, but with a wide gap at the MBD3-binding site (Figure 7). This pattern is best explained by MBD3 binding to promoters, whereas the H3K36me3 and the H3K79me2 signals are known to mark the gene body with H3K79me2 peaking within the first kb of the transcribed region (28). Because these plots are compiling genes in both orientations, the increase for these two marks is seen on both sides of the MBD3-binding site.

We can conclude that chromatin modifications associated with MBD2 or MBD3 binding is in line with the LacO array studies, with reporter gene activity and with the response of endogenous gene activity upon depletion of either MBD factor.

### MBD2 marks specifically exon sequences

Overall sequence class enrichment for MBD2 or MBD3 binding revealed that, among others, exon sequences were targeted (see Figure 4A and B). Therefore, we compiled the MBD2- and MBD3-binding sites at gene regions to test for enriched exon sequences in comparison with intron sequences. An instructive method is to plot binding sites over a meta-gene. Therefore we plotted the average MBD2/3 occupancy across 5′-upstream, first, internal and last exons as well as 3′-downstream regions according to their position relative to all RefSeq transcripts. (Figure 8A). With this analysis, a clear correlation for MBD2 with exon sequences was evident, whereas MBD3 was primarily detected at the promoter region and overlapping into the first exon. Control ChIPseq using the V5 tag resulted in only a marginal preference for the promoter region. The last exon, which is often not translated to a varying extend, depicts a decline of MBD2 binding from the 5′end towards the transcriptional end site. Such a finding is in line with results on DNA methylation that promoter regions of inactive genes may be methylated, whereas methylation of active genes is found in gene bodies (29,30). To verify the MBD2-binding pattern over the meta-gene and to test for a dependency on transcriptional activity, we grouped the MBD2-binding sites into those binding to genes with low or high transcriptional activity. Again, the preference for exons is evident with a stronger enrichment at the promoter and first exon at genes with low or no transcriptional activity, whereas the active genes show reduced binding to the promoter and first exon, but increased binding to internal exons (Figure 8B).

Thus, we can conclude that MBD2 shows an exon specificity within active genes.

### DISCUSSION

MBD2 and MBD3 have been analysed extensively for possible functional differences and similarities. Striking differences are embryonic lethality for MBD3-knockout
Figure 6. MBD2 binding to promoters is associated with gene repression and combined MBD2/MBD3 binding with gene activation. (A) MBD3-bound genes are significantly higher expressed than MBD2-bound genes. Gene expression in HeLa cells as measured by Affymetrix Gene 1.0 ST arrays [Wilcoxon signed rank test: \( P = 2 \times 10^{-28} \) (MBD2) and \( P = 8 \times 10^{-33} \) (MBD3)]. (B) Analysis of the expression level of 271 MBD3-bound genes sorted by fold change after MBD2 knockdown on all genes (black) (Wilcoxon signed rank test: \( P = 6 \times 10^{-5} \)). (D) MBD3-knockdown results in gene repression. Analysis of the expression level of 271 MBD3-bound genes sorted by fold change after MBD3 knockdown on all genes (black) (Wilcoxon signed rank test: \( P = 6 \times 10^{-5} \)).
our observation that depletion of MBD2 results in changes of gene expression characterized by a release from repression. A more direct repressive role by MBD2 is evident from our binding site analysis that shows a preference for MBD2 to be associated with genes of low activity as compared with the mean activity of all genes. Therefore, we focussed on these genes bound by MBD2 and could find gene activation upon MBD2 depletion, supporting the repressive role of MBD2 bound to methylated CpGs.

For MBD3, a repressive function has been postulated and has also been documented in a few cases. For example, PML-RARα binds and recruits MBD3–NuRD to target genes, thereby facilitating Polycomb binding and H3K27 methylation (37). Recently, deacetylation of H3K27 by MBD3 could be shown, which subsequently causes H3K27 trimethylation (16). Nevertheless, when depleting mouse ES cells from MBD3, a majority of responsive genes are down-regulated (16), indicating that a potential activation function is associated with MBD3. This is in line with our HeLa cell analysis showing that genome-wide binding of MBD3 is associated with genes showing a significantly higher expression as compared with the mean value of the expression of all genes. This non-repressive activity of MBD3 is further supported by our finding that MBD3 neither causes chromatin compaction nor does it repress reporter gene activity. In general, we find MBD3 binding to non-methylated CpGs, i.e. active promoters, which often show MBD2 binding as well. This is the small fraction of MBD2 sites that are not methylated. In these cases, depletion of MBD3 activates as well represses gene activity, whereas depletion of MBD2 causes a small down-regulation of the associated genes. A dual role of a repressor complex in gene repression as well as in gene activation is often triggered by the neighbouring chromatin features. In case of MBD3, repression may require synergizing repressive factors, which may be absent in TSS regions bound by MBD3.

Overall, both MBD factors bind similar sequence classes, which might support earlier results on MBD2 recruiting MBD3 complexes (3), but the differences arguing for separate complexes are prevailing. First, there is an enrichment of promoter sequences, with MBD2 binding to methylated and MBD3 binding to unmethylated promoters. Previously, the analysis of MBD3 or of MBD3–NuRD revealed differing results concerning binding to promoters in ES cells (7,16,38,39). As discussed previously (40), distribution of the NuRD complex was identified in broad regions with a depletion at the TSS, or specifically binding at the TSS. Here, we show that within a single cell type, both MBD2 and MBD3 are found at the TSS tailing into the first exon.
Most striking is the binding to gene bodies and exon sequences. In general, we found MBD2 and MBD3 binding at exon sequences. However the absolute number of cases with exon binding by MBD2 was much higher and therefore we analysed these in detail. Primarily internal (translated) exons are enriched for MBD2 binding. The terminal exon with varying length of the open reading frame shows MBD2 binding enriched within the 3′-region. Previously, CpG methylation has been seen to be enriched in the exons of active genes (41–43). This enrichment is further specified by the fact that non-coding exons are hypomethylated, whereas internal and last exons are more frequently methylated (29,44). This methylation pattern reflects the MBD2 distribution within the gene bodies, with declining MBD2 binding towards the 3′-end of the last exon, which usually is not translated. Furthermore, we find a correlation of MBD2 binding to internal exons and gene activity. It has been suggested that this methylation pattern may foreshadow the content of mature mRNA (41). A functional connection between DNA methylation and splicing has been recently demonstrated at the CD45 gene. In this case, methylation inhibits binding of CTCF, which influences the splicing reaction (45). Here we are extending this idea that MBD2 binding may affect splicing, although formally, the reverse may be true as well in that the transcribing polymerase/splicing complex may determine CpG methylation and MBD2 binding.

Thus, what has been generally called the NuRD repressor complex turns out to be associated with a diverse set of functions, such as repression, activation and potentially connecting chromatin with splicing.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online: Supplementary Figures 1–14, Supplementary Methods and Supplementary References [21,46–54]

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