

Epigenetic overlap in autism-spectrum neurodevelopmental disorders: *MECP2* deficiency causes reduced expression of *UBE3A* and *GABRB3*

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Autism is a common neurodevelopmental disorder of complex genetic etiology. Rett syndrome, an X-linked dominant disorder caused by *MECP2* mutations, and Angelman syndrome, an imprinted disorder caused by maternal 15q11–q13 or *UBE3A* deficiency, have phenotypic and genetic overlap with autism. *MECP2* encodes methyl-CpG-binding protein 2 that acts as a transcriptional repressor for methylated gene constructs but is surprisingly not required for maintaining imprinted gene expression. Here, we test the hypothesis that *MECP2* deficiency may affect the level of expression of *UBE3A* and neighboring autism candidate gene *GABRB3* without necessarily affecting imprinted expression. Multiple quantitative methods were used including automated quantitation of immunofluorescence and *in situ* hybridization by laser scanning cytometry on tissue microarrays, immunoblot and TaqMan PCR. The results demonstrated significant defects in *UBE3A*/E6AP expression in two different *Mecp2* deficient mouse strains and human Rett, Angelman and autism brains compared with controls. Although no difference was observed in the allelic expression of several imprinted transcripts in *Mecp2*-null brain, *Ube3a* sense expression was significantly reduced, consistent with the decrease in protein. A non-imprinted gene from 15q11–q13, *GABRB3*, encoding the $\beta 3$ subunit of the GABA_A receptor, also showed significantly reduced expression in multiple Rett, Angelman and autism brain samples, and *Mecp2* deficient mice by quantitative immunoblot. These results suggest an overlapping pathway of gene dysregulation within 15q11–q13 in Rett, Angelman and autism and implicate MeCP2 in the regulation of *UBE3A* and *GABRB3* expressions in the postnatal mammalian brain.

INTRODUCTION

Autism is a complex genetic neurodevelopmental disorder characterized by severe impairments in social interaction, communication and behavioral patterns that are restrictive and stereotypical (1). Rett syndrome (RTT; OMIM 312750) is an X-linked dominant neurodevelopmental disorder caused by mutations in *MECP2*, which encodes methyl-CpG-binding protein 2 (MeCP2) (2). Angelman syndrome (AS; OMIM 105830) is an imprinted disorder caused by maternal deficiency of chromosome 15q11–q13, methylation defects or maternal mutation of *UBE3A* encoding the ubiquitin ligase UBE3A/E6-AP (3). RTT and AS share overlapping clinical features with autism including developmental delay, language impairment, seizures and stereotypic behaviors (4). Furthermore, clinical assessments of social behavior have demonstrated a high frequency of autism in patients with

RTT (5) and AS (6). A *Mecp2* mutant mouse model of RTT also shows abnormalities in social interactions (7). In addition, *MECP2* mutations have been found in a few patients diagnosed with AS (8) and autism (9,10) and 15q11–q13 duplications are present in ~1% of autism cases (11), suggesting overlap in the pathogenesis of these distinct genetic syndromes. *GABRB3*, encoding the GABA_A receptor β subunit, is located within the 15q11–q13 (12) and shows association with autism (13,14) but is biallelically expressed (15). We have previously demonstrated MeCP2 expression defects in autism and AS brain samples by a quantitative immunofluorescence approach (16). In contrast, expression level of proteins encoded by genes within 15q11–q13 in RTT, AS and autism have not been well characterized.

MeCP2 binds to methylated CpG sites (17), represses transcription of methylated constructs (18), and associates with chromatin modifying factors histone deacetylase (19,20) and

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DNA methyltransferase (21,22). *Mecp2*-null (23,24) or mutant (25) mice recapitulate the phenotype of RTT and demonstrate that MeCP2 is essential for postnatal mammalian brain development. Although MeCP2 was predicted to be a global transcriptional repressor, a paucity of MeCP2 target genes have been identified in *Mecp2*-null mouse brain (26), RTT brain (27) or cell lines (28) by gene expression profiling. The gene encoding brain derived neurotrophic factor has been identified as a target of MeCP2 transcriptional repression where a low basal level of transcription prior to neuronal stimulation was increased in *Mecp2*-null neurons (29,30). The imprinted gene *H19* also showed increased expression in *Mecp2*-null cells (21), but it was not previously determined whether the increase was due to de-repression of the methylated paternal allele. Because of the limited number of MeCP2 target genes, the role of MeCP2 in the pathogenesis of RTT and other autism-spectrum disorders remains elusive.

Although *MECP2* mutation does not affect imprinted expression of several genes within 15q11–q13 (31), we hypothesized that MeCP2 may regulate the expression level of genes in this region without necessarily affecting allele-specific expression. Here, we demonstrate that *Mecp2* deficiency results in significant reduction of UBE3A/*Ube3a* and GABRB3/*Gabrb3* expressions in mouse cerebrum without apparent alterations in allele-specific expression. Furthermore, significant reduction of UBE3A and GABRB3 expressions was observed in AS, RTT and autism human cerebral samples compared with controls. These results demonstrate overlapping epigenetic defects in these phenotypically similar but genetically distinct neurodevelopmental disorders and implicate MeCP2 in the regulation of gene expression within 15q11–q13.

RESULTS

Reduced UBE3A expression in *Mecp2* deficient mice

As a non-cell-autonomous dominant effect of *Mecp2* mutation in the wild-type (wt)-expressing cells of mosaic *Mecp2*^{-/+} females has been described (32), we performed single cell quantitative analyses of mutant (mt)-expressing (MeCP2-negative) and wt-expressing (MeCP2-positive) cells for UBE3A immunofluorescence on a tissue microarray by laser scanning cytometry (LSC). This approach demonstrated a reproducible 1.5–2-fold decrease in UBE3A immunoreactivity in 10 *Mecp2*^{-/+} and two *Mecp2*^{-/-} cerebral samples from two different *Mecp2*-null strains (23,24) compared with wt controls (Fig. 1A and B). Both mt-expressing cells from *Mecp2*^{-/+} and *Mecp2*^{-/-} (blue histograms) and wt-expressing cells from *Mecp2*^{-/+} cerebrum (red histograms) showed significantly lower UBE3A expression compared with wt-expressing cells from *Mecp2*^{+/+} cerebrum (orange histograms). To demonstrate that the defects observed in *Mecp2*^{-/+} and *Mecp2*^{-/-} brain samples were actually due to expression differences in UBE3A, not immunofluorescence detection, immunoblot analyses were performed (Fig. 1C). Reduced UBE3A/E6-AP expression was observed by immunoblot for both *Mecp2*^{-/+} and *Mecp2*^{-/-} brain samples from both *Mecp2*-null strains compared with controls but with lower significance than the LSC results. The precise sampling of single cells within the

cerebral cortex in the LSC analysis compared with the whole brain analysis by immunoblot may explain the greater sensitivity of the LSC approach. These results demonstrate that UBE3A protein expression in the cerebral cortex is reduced by both cell-autonomous and non-cell-autonomous effects of *Mecp2* mutation.

Ube3a expression level is lower in *Mecp2* deficient mouse brain without alterations in imprinted gene expression

As the maternal expression of *Ube3a* in postnatal neurons is correlated with the paternal expression of an antisense *Ube3a* transcript from the imprinting control region (ICR) of the *Snrpn/Snrnf* promoter (33), we examined the role of MeCP2 on the imprinting status of this locus. Chromatin immunoprecipitation (ChIP) demonstrated that MeCP2 was bound to the promoter of *Snrpn* and positive control *U2af1-rs1* (34) in mouse cerebrum (Fig. 2A). In contrast, neither the *Ube3a* nor *Gabrb3* promoter was found to be associated with MeCP2 at a detectable level, consistent with a lack of methylation. Allele-specific analyses of (*Mecp2*^{-/+} × PWK)F1 progeny demonstrated the expected preferential maternal expression of *Ube3a* sense transcript, paternal expression of *Ube3a* antisense and *Snrpn* and biallelic expression of *Gabrb3*, regardless of *Mecp2* genotype (Fig. 2B). These results demonstrated that reduced expression of UBE3A/*Ube3a* in *Mecp2* deficient brain was not directly due to alterations in allele-specific expression and confirm similar results from human brain (31). Two additional imprinted genes from other loci (*Rasgrf1* and *H19*) were also tested for allele-specific expression changes in *Mecp2*-null mice with similar results (Fig. 2B). These results further demonstrate that *Mecp2* is not required for maintenance of imprinted gene expression.

To determine whether *Ube3a* transcript levels were also reduced, TaqMan PCR was performed on RNA samples obtained from *Mecp2*^{-/-} and *Mecp2*^{+/+} mice. The results showed a reduction in *Ube3a* (sense) and *Gabrb3* transcripts, but not in *Snrpn*, in *Mecp2* deficient brain (Fig. 2C). Fluorescence *in situ* hybridization with single stranded riboprobes was then performed on the tissue microarray in order to separately analyze the sense and antisense transcripts of *Ube3a* and confirm the reduction of the *Ube3a* sense transcript by another method. Figure 2D shows the results demonstrating a significantly lower expression of *Ube3a* sense but not *Ube3a* antisense transcripts. These combined results suggest that the decrease in UBE3A protein expression in *Mecp2* deficient brain is due to decreased expression of the *Ube3a* sense transcript without a significant change in *Snrpn* or the *Ube3a* antisense transcript or the maternal-biased expression of *Ube3a*.

UBE3A expression is deficient in RTT, AS and autism cerebral samples

To determine whether significant reductions in UBE3A expression were also observed in human brain samples, a postmortem brain tissue microarray described previously containing human cerebral samples (16) was analyzed by immunofluorescence and LSC. The histograms in Figure 3A demonstrate significant reductions in UBE3A expression of

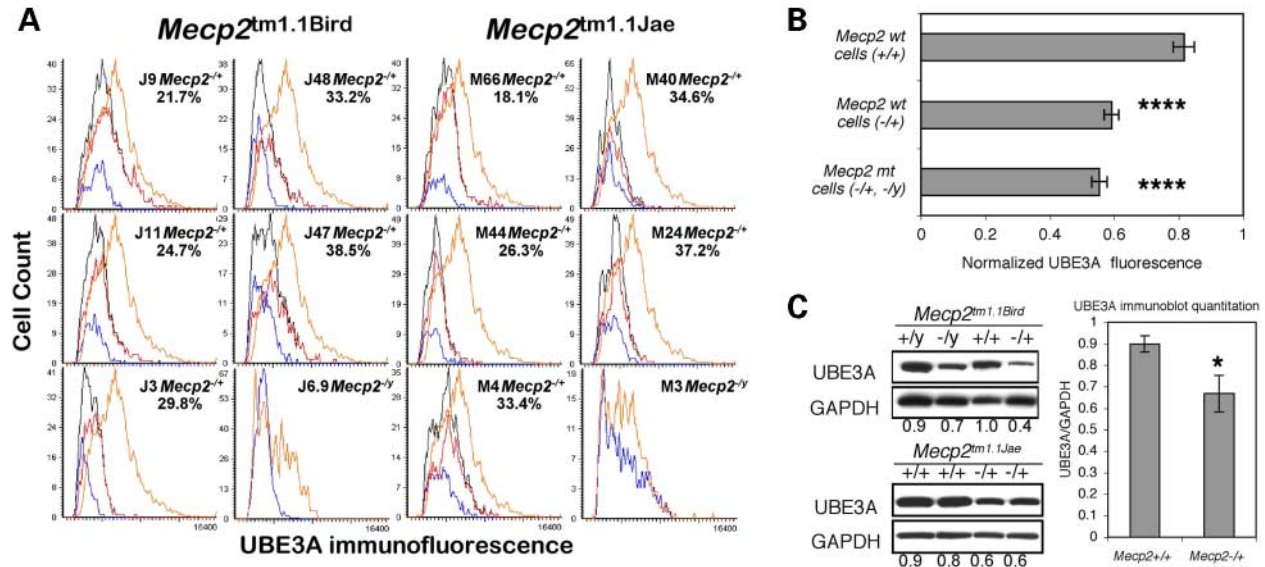


Figure 1. UBE3A expression in *Mecp2*-null mouse brain. (A) A multiple tissue microarray containing wt and knock-out mouse cerebral cortex samples (16) was stained for immunofluorescence using a C-terminal reactive anti-MeCP2 and an anti-UBE3A antibody. The MeCP2-negative cells (*Mecp2* wt-expressing, blue histograms, percentage shown) were separately gated from MeCP2-positive cells (*Mecp2* wt-expressing, red histograms) and the total population (black histograms) and compared with age-matched wt control female or male samples (orange histograms). (B) The graph shows combined LSC results (mean ± SEM) from four replicate slides using two different anti-UBE3A antibodies. (C) Protein extracts from whole adult mouse brain were probed on an immunoblot with anti-UBE3A or anti-GAPDH. A representative image shows lower expression in brain of UBE3A in both hemizygous male (-/y) and heterozygous female (-/+) brain compared with wild-type (wt, +/y and +/+) littermate controls for both *Mecp2*^{tm1.1Bird} and *Mecp2*^{tm1.1Jae} strains. The graph demonstrates combined GAPDH-normalized results (mean ± SEM, quantitation as previously described (32), example ratios shown below each lane) from eight different -/+ and six different +/+ brain samples from *Mecp2*^{tm1.1Bird} and *Mecp2*^{tm1.1Jae} strains. **P* < 0.05, *****P* < 0.0001 by *t*-test.

RTT, AS and autism samples (black) compared with age-matched controls (gray). Ages of each sample are shown, but no significant differences were observed in the level of UBE3A expression with increasing age of the controls (data not shown), unlike the striking increase with age observed for MeCP2 (16,35). Mean UBE3A fluorescence was normalized to control histone H1 expression and graphed (Fig. 3B), demonstrating significant UBE3A reductions in RTT and autism samples. The reduced expression of UBE3A in the AS 15q11-q13 deletion sample was expected and consistent with a low level of paternal *UBE3A* expression in the cerebral cortex (36). The LSC results were confirmed by immunoblot (Fig. 3C shows a representative blot). Furthermore, reduced UBE3A expression in AUT1174 was also shown by immunoblot in a recent report (37).

GABRB3 expression defects are observed in *Mecp2* deficient mouse and RTT, AS and autism human brain

GABRB3 has been strongly implicated in autism by multiple association studies (14,38,39), but alterations in expression level have not been previously demonstrated. Because *Mecp2* deficiency affected not only *Ube3a* expression but also the biallelically expressed gene, *Gabrb3* (Fig. 2C), we further investigated its protein product, the gamma aminobutyric acid (GABA) receptor $\beta 3$ subunit, for defects in *Mecp2*/MeCP2 deficient brain. Immunoblot analyses demonstrated reduced-GABRB3 expression in *Mecp2*^{-/+} and *Mecp2*^{-/-} mouse brain compared with controls (Fig. 4A), suggesting

an additional target from the 15q11-q13 locus may also be dysregulated in RTT, AS and autism.

As a transmembrane protein, GABRB3 was less amenable to LSC analysis than nuclear or cytoplasmic proteins because individual cells are identified by nuclear contouring. Immunoblot analyses were, therefore, performed on frozen frontal cerebral cortex, Brodmann area 9 (BA9), from a total of three RTT (with *MECP2* mutations), two AS (with 15q11-q13 deletions), nine autism (no detectable *MECP2* mutations or 15q11-q13 duplications) and 11 different age-matched normal controls. Two representative immunoblots are shown in Figure 4B. Table 1 shows the combined data of normalized GABRB3/GAPDH ratios for all individual samples. When analyzed as combined groups, RTT, AS and autism samples showed significantly lower GABRB3 expression compared with controls. The autism samples showed the highest significant decrease in GABRB3 as a group (*P* < 0.0001) despite higher variability between samples, with some showing extremely low expression (i.e., 797, 1174, 967 and 5173, with 5–10% of control expression) and others showing little change (i.e., 3924 and 5342, Fig. 4B and Table 1). The ages of samples are shown in Table 1, but reveal no significant change in GABRB3 expression with age of the control samples tested. The average GABRB3 expression for each patient category is, therefore, also represented as a group for statistical comparisons. Although GABRB3 expression deficiencies are not universal in all autism brain samples, the defects are more significant than in RTT and AS and occur at an unexpectedly high frequency (5/9 or 56%). The specificity of GABRB3 defects to AS, RTT and autism was tested by analyzing three

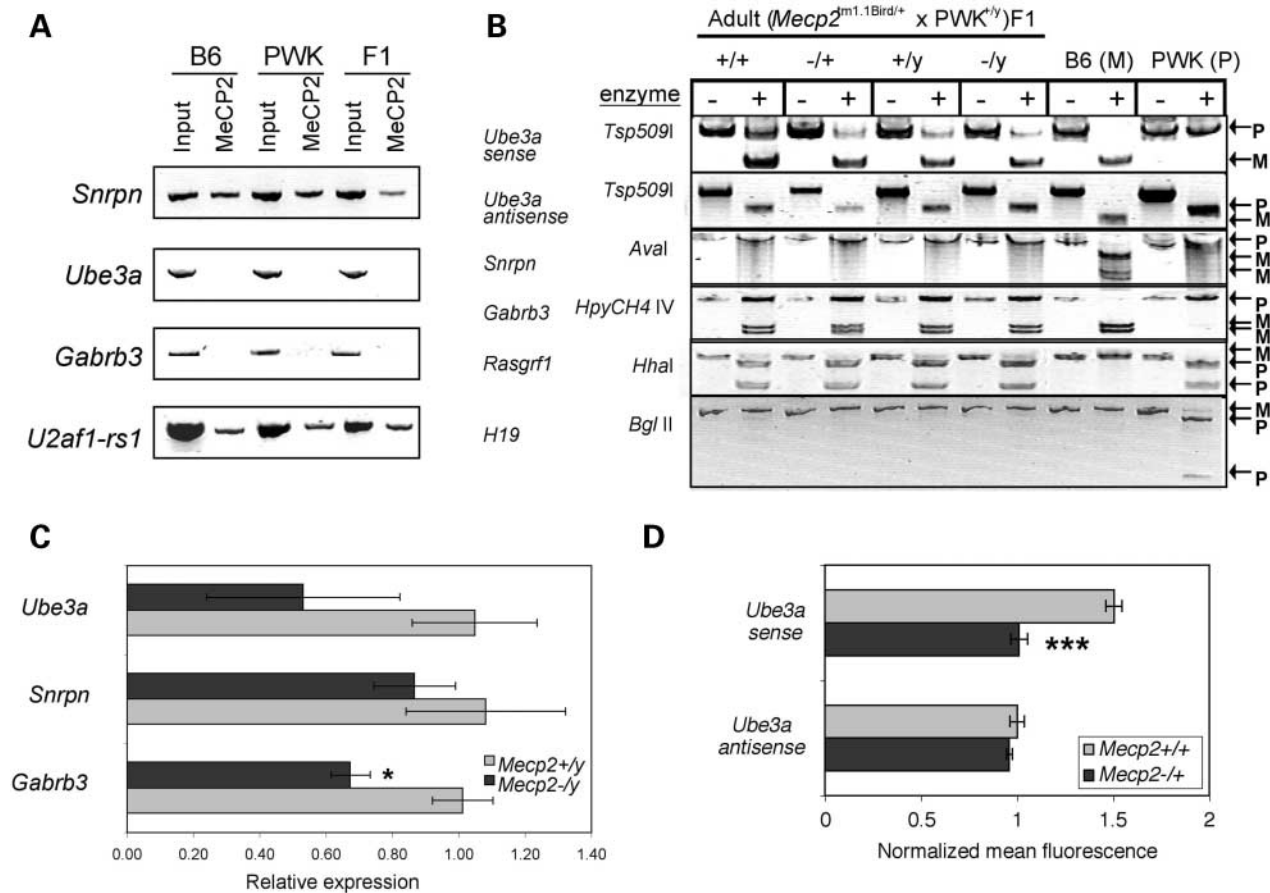


Figure 2. Imprinting and transcriptional analyses in mouse brain. (A) Chromatin from adult mouse cerebrium samples [C57B6, PWK or (B6 × PWK)F1] was isolated for ChIP. Anti-MeCP2 (C-terminal) was used to immunoprecipitate DNA fragments from 'Input' control. *Ube3a* and *Gabrb3* promoters were not detected in the anti-MeCP2 precipitated chromatin, in contrast to the *Snrpn* promoter sequences that showed association with MeCP2. *U2af1-rs1* was a positive control for a promoter previously demonstrated to bind MeCP2 in brain (34). (B) *Mecp2*^{tm1.1Bird/+} (B6) females were crossed with wt PWK males to obtain F1 mice heterozygous for single nucleotide polymorphisms in the coding regions of several imprinted genes. RT-PCR followed by restriction enzyme digestion (+) was performed on RNA from adult brain samples of all *Mecp2* genotypes as well as parental B6 and PWK samples (P, paternal; M, maternal). As reported previously for wt (B6 × PWK)F1 brain (67), *Ube3a* sense exhibited preferential maternal expression, *Ube3a* antisense and *Snrpn* were exclusively paternal and *Gabrb3* was biallelic, with no significant effect of *Mecp2* genotype on these imprints. In addition, *Rasgrf1* showed preferential paternal expression (68), whereas *H19* showed exclusive maternal expression (70) in all samples. Results are representative of adult and neonatal F1 brain samples. (C) TaqMan PCR was used to quantitatively determine expression levels of *Ube3a*, *Snrpn* and *Gabrb3* in 10-week-old *Mecp2*^{-/-} and *Mecp2*^{+/-} brain RNA. Primers spanned intron/exon boundaries and are specific to the *Ube3a* sense transcript. Results shown are for the average ± SEM of four experimental replicates of two mice per genotype. Although *Ube3a* and *Gabrb3* showed consistently lower expression in *Mecp2*-deficient brain compared with controls, *Snrpn* was not significantly changed. (D) Sense and antisense transcripts of *Ube3a* were detected by fluorescence *in situ* hybridization using single stranded riboprobes and quantitated by LSC as described previously (16). Results (mean ± SEM of 3 wt and 10 *Mecp2*^{-/+} samples) were normalized to control β-actin probe. ****P* < 0.001 by *t*-test.

Down syndrome (DS) brain samples with trisomy 21. Table 1 shows no significant defects in GABRB3 expression in DS compared with controls. These results, therefore, demonstrate that reduced-GABRB3 expression is a common characteristic of RTT and autism brain.

DISCUSSION

Uncovering the genetic basis of autism has been challenging because of its complex etiology. Although the heritability of autism in families is high, multiple gene loci and environmental influences are expected (1). Even for the autism-spectrum disorders with known genetic causes, the molecular

pathogenesis is not well understood. Because of the overlap in phenotype among RTT, AS and autism (4), we tested the hypothesis that gene expression in the 15q11–q13 region may be altered in all three syndromes. We show for the first time that the AS gene, *UBE3A/Ube3a*, is significantly reduced in RTT and *Mecp2*-null mice, demonstrating that *MECP2/Mecp2* deficiency causes reduced expression of *Ube3a/UBE3A*. In addition, we demonstrate *UBE3A* deficiency in autism brain samples, consistent with a recent report (37). Furthermore, we also show for the first time deficiencies in the expression of the GABA receptor β3 subunit gene (*GABRB3*) in RTT and autism brain and *Mecp2*-null mice.

Maternal duplications in 15q11–q13 have been observed in ~1% of autism cases as the most common cytogenetic

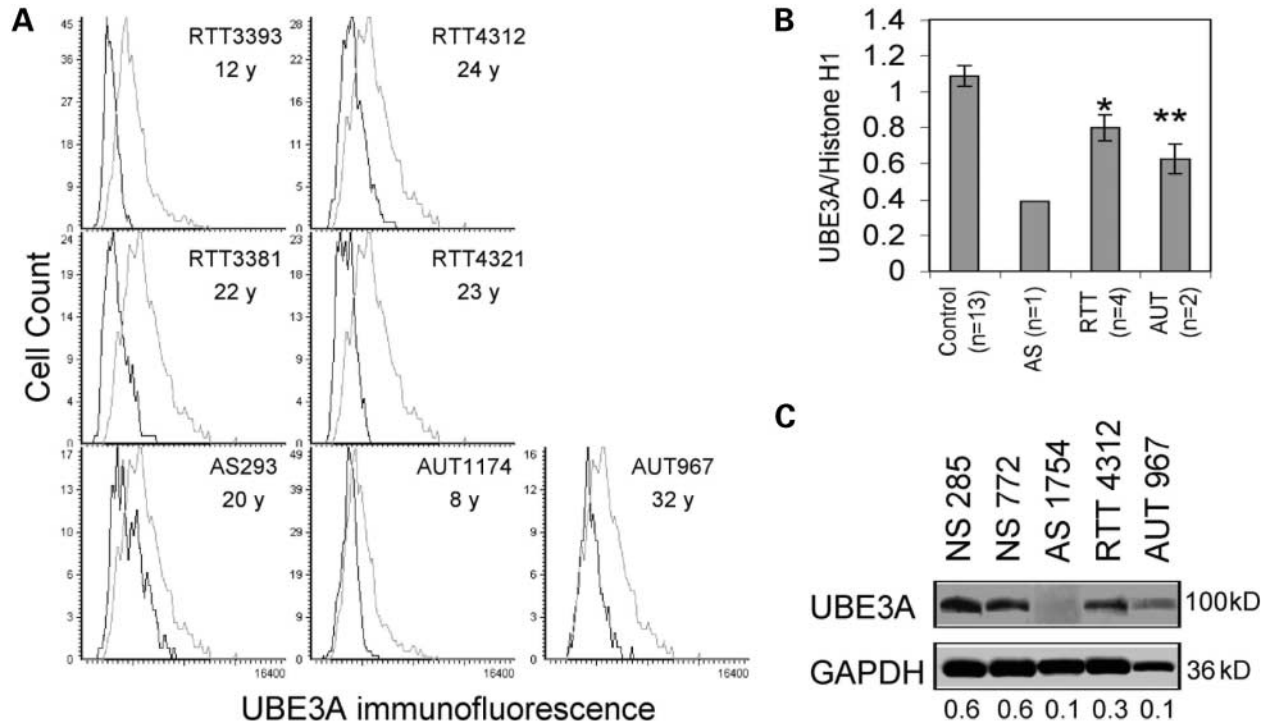


Figure 3. Quantitation of UBE3A expression in Rett, Angelman and autism brain samples. (A) The human tissue microarray previously described (16) containing frontal cortex layers III–V from Brodman area (BA9) was analyzed for UBE3A expression by LSC. Histograms show lower expression of patient samples (black) from Rett (RTT), Angelman (AS) and autism (AUT) compared with age-matched controls (gray). (B) UBE3A immunofluorescence values were normalized to those of control histone H1 and the results of four replicate slides using two different anti-UBE3A antibodies (Affinity Bioreagents and BD Biosciences) were averaged (mean \pm SEM) for each brain sample shown in (A) (*n*, number of different samples). (C) Immunoblot analysis of UBE3A expression (anti-UBE3A, Affinity, recognizing all known isoforms) confirmed decreased expression in AS, RTT and AUT compared with controls (UBE3A/GAPDH ratios are shown). **P* < 0.05, ***P* < 0.01 by *t*-test.

abnormality in autism (11). Genome-wide linkage studies have identified the 15q11–q13 locus in some but not all scans (40–43). Furthermore, higher resolution scans of linkage within 15q11–q13 in autism families have identified significant linkage to *UBE3A* and *GABRB3* in some studies but not others (14,44–48). While the linkage of autism to 15q11–q13 may be somewhat disputed at the genetic level, here we show that expression level defects of two genes within 15q11–q13 are common to RTT, AS and autism. The reduction of UBE3A and GABRB3 in AS cerebrum was expected because of the maternal 15q11–q13 deletion in these samples. Interestingly, neither *Ube3a* deficient nor uniparental disomy mice showed evidence for decreased *Gabrb3* expression (49,50), but the lack of AS brain samples with uniparental disomy or *UBE3A* mutations prevented our confirmation of this finding in human samples.

Our results are consistent with a recent report showing decreased UBE3A in autism brain (37), but extend the observation to demonstrate GABRB3 deficiency as a common defect in autism. The GABA_A receptor genes are attractive candidate genes for autism as several studies have demonstrated linkage to one or more of these genes (13,14,48). Autoradiography has demonstrated reduced GABA_A binding in autism brain (51) and elevated circulating GABA levels have been found in autistic children (52–54). Although our sample size was small owing to the requirement for post-mortem human brain tissue, the frequency of UBE3A and

GABRB3 expression defects within autism brain samples was much higher than for previously described genetic defects. The defects in UBE3A and GABRB3 expressions were unlikely to be due to degradation in postmortem tissue as they were not observed in multiple normal and DS controls and because the results were all normalized to housekeeping gene controls. These results suggest that reduced expression of *UBE3A* and *GABRB3* are correlated and may be a downstream consequence of multiple different genetic or epigenetic etiologies. The phenotypic consequence of reduced expression of these genes is expected to be less severe than gene mutation (*UBE3A* mutations in AS) or nullisomy. Both *Ube3a*^{−/+} and *Gabrb3*^{+/-}, ^{−/−} mice have learning and motor defects and inducible seizures, similar to AS and some cases of autism (55,56). While our autism brain sample population seems to be somewhat enriched for associated seizure disorders (Supplementary Material, Table S1), significantly reduced-GABRB3 expression was observed in brain samples from individuals with and without seizures.

We also identify the X-linked gene, *MECP2*, as one genetic cause of UBE3A and GABRB3 expression defects in the mammalian cerebrum. We have used two *Mecp2* deficient RTT mouse strains (23,24) to demonstrate that UBE3A/*Ube3a* and GABRB3/*Gabrb3* are significantly decreased in adult mouse brain compared with wt controls. Furthermore, we show significant defects in expression of UBE3A and GABRB3 in human RTT brain samples with *MECP2*

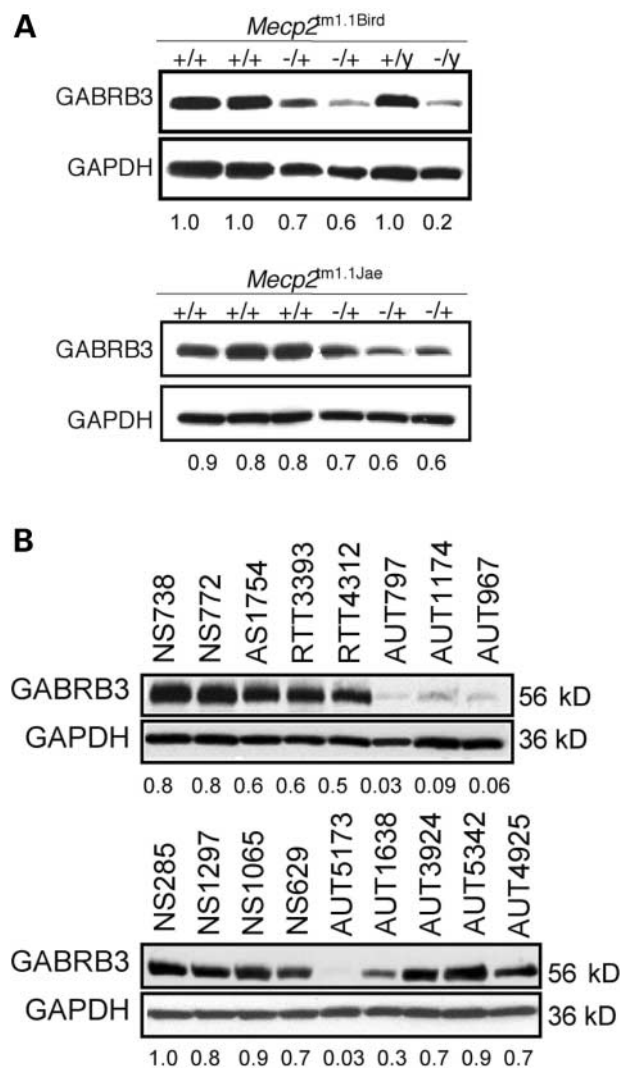


Figure 4. Analysis of GABRB3 expression in mouse and human brain. (A) Immunoblot analysis of GABRB3 expression (anti-GABRB3, Affinity Bioreagents, recognizing all known isoforms) shows lower expression in *Mecp2^{tm1.1Bird}* and *Mecp2^{tm1.1Jae}* adult mouse brain compared with controls. Combined GAPDH-normalized results from eight different *Mecp2^{tm1.1Bird}* and six different *Mecp2^{tm1.1Jae}* adult brain samples from *Mecp2^{tm1.1Bird}* and *Mecp2^{tm1.1Jae}* strains were significantly lower than wild-type controls ($P < 0.01$). (B) Representative immunoblot analyses of GABRB3 and GAPDH in AS, RTT, autism and control (NS) cerebral samples with GABRB3/GAPDH ratios shown. Replicate immunoblot results for all samples and significance tests are shown in Table 1. Additional information about postmortem brain samples used in these studies is provided online (Supplementary Material, Table S1).

mutations. Our results are consistent with *GABRB3* showing reduced expression in RTT brain by genome-wide expression profiling in one report (27). The relatively subtle changes in expression of UBE3A and GABRB3 observed in this study may explain why these genes were not identified in other genome-wide profiling studies of *MECP2/Mecp2* mutant tissues (26,28). The observation of down-regulation as well as up-regulation of MeCP2 target genes has also been observed in expression profiling studies, arguing against a single role for MeCP2 as a transcriptional repressor.

Table 1. Significant differences in GABRB3 expression in RTT, AS and autism brain by quantitative immunoblot

Category	Sample no.	Age (years)	No. of replicates	GABRB3/GAPDH (mean \pm SEM)
Control	629	7	6	0.75 \pm 0.016
	738	8	5	0.64 \pm 0.064
	3835	9	1	0.78
	1297	15	1	0.84
	1065	15	1	0.87
	602	27	5	0.69 \pm 0.047
	285	31	5	0.80 \pm 0.081
	1104	35	1	0.57
	772	36	1	0.79
	4192	46	2	0.75 \pm 0.025
Control average	4503	56	2	0.52 \pm 0.001
				0.73 \pm 0.032
Rett	3393	12	2	0.61 \pm 0.001 ^{a*}
	4312	23	3	0.53 \pm 0.017 ^{a**}
	5020	24	2	0.43 \pm 0.093
Rett average				0.52 \pm 0.031 ^{b**}
Angelman	1754	4	2	0.51 \pm 0.037 ^{a***}
	1494	43	2	0.13 \pm 0.078 ^{a***}
AS average				0.32 \pm 0.120 ^{b***}
Autism	797	9	3	0.17 \pm 0.103 ^{a***}
	1174	7	3	0.25 \pm 0.099 ^{a***}
	967	32	2	0.04 \pm 0.011 ^{a*}
	5000	27	3	0.55 \pm 0.117
	5342	11	2	0.79 \pm 0.085
	5173	30	2	0.05 \pm 0.014 ^{a*}
	4925	9	2	0.56 \pm 0.097
	3924	16	2	0.69 \pm 0.004
	1638	20	2	0.31 \pm 0.001 ^{a*}
				0.39 \pm 0.060 ^{b*}
Autism average				
Down	1267	10	3	0.76 \pm 0.046
	707	23	3	0.80 \pm 0.005
	1753	24	3	0.71 \pm 0.027
Down average				0.76 \pm 0.028

^aReplicate results from each individual sample were compared with the three closest age-matched controls by *t*-test.

^bTotal sample replicates were combined from each category and compared with total control sample replicates by *t*-test.

* $P < 0.01$.

** $P < 0.05$.

*** $P < 0.001$.

**** $P < 0.0001$.

As we have previously demonstrated *MECP2* expression defects in the same autism brain samples (16), perhaps *MECP2* expression defects could be responsible for the lower UBE3A and GABRB3 expressions in these autistic individuals without characterized genetic defects. Alternatively, defects in expression of all three genes may be a downstream consequence of abnormal brain development. As the *Mecp2^{308/y}* RTT mouse model has been demonstrated to show some autistic behaviors, including forepaw stereopathies and social avoidance (7,25), the role of MeCP2 in the pathogenesis of at least some cases of autism appears likely (4).

In this report, we also begin to uncover a rather complex and unexpected mechanism by which MeCP2 regulates expression of UBE3A and GABRB3. Although MeCP2 binds to methylated DNA and is a predicted transcriptional

repressor (17,18), it is not required for maintaining parental imprints (31). Here, we demonstrate that MeCP2 binds to the ICR in the *Snrpn* promoter, yet it is not required for silencing the maternal allele of *Snrpn* and *Ube3a* antisense or the paternal allele of the *Ube3a* sense transcript. Furthermore, expression level of *Gabrb3* is affected even though this gene is biallelically expressed. We also demonstrate the maintenance of exclusive maternal expression of *H19* in *Mecp2*-null mice despite its description as a 'bona fide' MeCP2 target gene (21). These results suggest that the effect of MeCP2 binding to the ICR is more complex than simply repressing imprinted genes in *cis* and may instead involve long-range chromatin interactions. Perhaps chromatin loop formation such has been recently demonstrated in the imprinted *Igf2/H19* region (57) is adversely affected by *Mecp2* deficiency, as MeCP2 has been previously described to have matrix binding activity (58). The ~1 Mb region from *UBE3A* to *GABRB3* also includes multiple small nucleolar RNA genes (33,59,60) and *ATP10A* (61,62), and two other GABA_A receptors (GABRA5 and GABRG3) located telomeric of *GABRB3* (63) that also could be dysregulated by MeCP2 deficiency. In addition to the three CpG islands investigated here for association with MeCP2, there are 13 other CpG islands in this region that remain to be investigated as potential MeCP2 binding sites (UCSC Genome Browser, May 2004 assembly). In addition to potential long-range chromatin organization in *cis*, 15q11–q13 homologous *trans* interactions have been demonstrated in human lymphocytes (64) and brain (Thatcher *et al.*, submitted for publication) and mouse ES cells (65) that may be important in regulating the expression level of multiple genes. Although autistic individuals with 15q11–q13 duplications have been predicted to express higher levels of *UBE3A* and *GABRB3* compared with controls, lack of proper biparental homologous pairing could instead result in lower expression of these genes similar to the autism brain samples in our study.

In conclusion, in this report, we have made an important connection by uncovering a molecular brain alteration common to three different autism-spectrum neurodevelopmental disorders of different genetic etiologies. The expression defects in *UBE3A* and *GABRB3* observed in RTT and autism brain are likely to be consequences of abnormal neuronal function due to deficiencies in *MECP2* or other genetic or environmental causes. Further understanding of how MeCP2 regulates expression of these and other genes within 15q11–q13 may help in the identification of genetic causes of autism and aid in the design of therapies for all three disorders.

MATERIALS AND METHODS

Immunofluorescence analysis of tissue microarrays

Construction of tissue microarrays used in this study have been previously described for *Mecp2*-null mouse (32) and human neurodevelopmental disorders (16). Briefly, for the mouse array, triplicate 600 µm cores of cerebral cortex were removed from five *Mecp2*^{tm1.1Bird/+}, five *Mecp2*^{tm1.1Jae/+}, and three *Mecp2*^{+/+} controls (P20 w–P27 w); one *Mecp2*^{tm1.1Bird/y}, and one *Mecp2*^{+/+} littermates (P10 w); one *Mecp2*^{tm1.1Jae/y} and one *Mecp2*^{+/+} littermate controls (P10 w).

The human tissue microarray contains triplicate frontal cerebral cortex (BA9, layers III–V) obtained frozen <30 h post-mortem. 5 µm section was processed for immunofluorescence as described previously for anti-MeCP2 (C-terminal, Affinity Bioreagents) and anti-histone H1 (Upstate Biotechnology) antibodies (66). For mouse tissues, combined staining was performed with anti-UBE3A antibody (BD Biosciences and monoclonal gift from Howley, Harvard Medical School) and a cascade-blue labeled goat-anti-mouse IgG (Molecular Probes). For human tissues, an anti-UBE3A antibody (Affinity Bioreagents) was used. LSC analysis of MeCP2 negative and positive populations was performed as described previously (32).

In situ hybridization analysis of tissue microarrays

Sense and antisense transcripts of *Ube3a* were detected by fluorescence *in situ* hybridization using single stranded riboprobes from a 796 bp cloned fragment spanning exons 6–10 (primers, 5'-tcacatgatgaagctacgaa-3' and 5'-ttctttgcttgaa tattccgg-3') labeled with biotin (Lofstrand Laboratories). A control β-actin probe digoxigenin labeled (Roche) was simultaneously hybridized at 55°C. Hybridization conditions, fluorescent detection and quantitation by LSC were performed as described previously (16).

Immunoblot analyses

A total of 10 µg of protein extract from whole adult mouse brain was analyzed per lane by immunoblot as described previously (32) using anti-UBE3A (BD Biosciences), anti-GABRB3 (Affinity Bioreagents) or anti-GAPDH (Advanced Immunochemical) antibodies at recommended concentrations. Protein extracts were isolated from frozen human postmortem cerebral cortex (BA9) from samples previously described (16) and additional samples (Harvard Brain Bank and University of Maryland Brain and Tissue Bank). Similar to mouse samples, 10 µg of human protein was analyzed per lane with the same reagents except the substitution of a human-specific anti-UBE3A antibody (Affinity Bioreagents). Additional human cerebral (BA9) samples were obtained from autism, Down and age-matched control samples from the Autism Tissue Program for immunoblot analyses. Quantitation was performed using Nucleotech Gel Expert version 2.0.

Allelic expression of imprinted genes in interspecies hybrids

Mecp2^{tm1.1Bird/+} (C57B6) female mice (23) were mated to wt PWK males and the resulting offsprings were genotyped for *Mecp2* as described previously (32). RNA was isolated from adult (*Mecp2*^{tm1.1Bird/+} × PWK)F1 brain samples from all four genotypes as well as parental B6 and PWK mice. RT-PCR followed by restriction enzyme digestion was performed as described previously for *Ube3a*, *Snrpn*, *Gabrb3* (67) and *Rasgrfl* (68). *H19* primers (5'-GAACCACTAACAC TACCTGCC-3' and 5'-GGAAGTCTTCCAGACTAGG-3', 58°C with 30 cycles) amplified a 585 bp fragment that was digested with *Bgl*II.

Chromatin immunoprecipitation

Chromatin from adult mouse cerebrum samples [C57B6, PWK or (B6 × PWK)F1] was isolated for ChIP as described previously (34,69). Anti-MeCP2 (C-terminal) was used to immunoprecipitate DNA fragments from 'Input' control as described previously for primers spanning 5' CpG islands for *Snrpn* and positive control *U2af1-rs1*(34), *Ube3a* (5'-CCCTTCTGCTTCTCTTCGGAGT-3', 5'-CAGAAGCAG CACACGAATAAA-3', 58°C anneal, 37 cycles) and *Gabrb3* (5'-CCTCAGAGCCACCCGTAC-3', 5'-GTCTAGGACCCC GCGACA-3, 60°C anneal, 40 cycles).

TaqMan PCR

Each PCR contained 400 nM of each primer, 80 nM of the TaqMan probe and commercially available PCR mastermix (TaqMan Universal PCR Mastermix, Applied Biosystems) containing 10 mM Tris-HCl (pH 8.3), 50 mM KCl, 5 mM MgCl₂, 2.5 mM deoxynucleotide triphosphates, 0.625 U AmpliTaq Gold DNA polymerase per reaction, 0.25 U AmpErase UNG per reaction and 5 µl of the diluted cDNA sample in a final volume of 25 µl. *Ube3a* forward primer spanned exons 7–8 (5'-TCCAGATATTGGTATGTTTCA TATG-3'), reverse primer (5'-GGGAAAATGGACATCCA GTATACAA-3') and probe (5'-CTGATTGGCATAGTCCT GGGTCTGGC-3') were from exon 8 (NM_173010). *Snrpn* forward primer spanned exons 3–4 (5'-TCAGGAAGAT CAAGCCAAAGAATG-3'), reverse primer (5'-AAGTTCT CCCCACGTAGCAAGAC-3') and probe (5'-ACAGCCAG AACGTGAAGAAAAACGGGT-3') were from exon 4 (NM_013670). *Gabrb3* forward primer spanned exons 6–7 (5'-CCGTCTGGTCTCCAGGAATGTT-3'), reverse primer (5'-CCGATATTTCTCTTCAACCGAAAA-3') and probe (5'-TTCCGCCACAGGTGCCTATCCTCGAC-3') were from exon 7. The samples were placed in 96 well plates and amplified in an automated fluorometer (ABI PRISM 7700 Sequence Detection System, Applied Biosystems). Amplification conditions were 2 min at 50°C, 10 min at 95°C, 40 cycles of 15 s at 95°C and 60 s at 60°C. Final quantitation was done using the comparative CT method (Applied Biosystems) and is reported as relative transcription or the *n*-fold difference relative to a calibrator cDNA (*Mecp2*^{+/y} of each pair) following normalization to *HPRT1* housekeeping control gene.

SUPPLEMENTARY MATERIAL

Supplementary Material is available at HMG Online.

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REFERENCES

1. Volkmar, F.R. and Pauls, D. (2003) Autism. *Lancet*, **362**, 1133–1141.
2. Amir, R.E., Van den Veyver, I.B., Wan, M., Tran, C.Q., Francke, U. and Zoghbi, H.Y. (1999) Rett syndrome is caused by mutations in X-linked *MECP2*, encoding methyl-CpG-binding protein 2. *Nat. Genet.*, **23**, 185–188.
3. Lalande, M. (1996) Parental imprinting and human disease. *Annu. Rev. Genet.*, **30**, 173–195.
4. Zoghbi, H.Y. (2003) Postnatal neurodevelopmental disorders: meeting at the synapse? *Science*, **302**, 826–830.
5. Mount, R.H., Charman, T., Hastings, R.P., Reilly, S. and Cass, H. (2003) Features of autism in Rett syndrome and severe mental retardation. *J. Autism Dev. Disord.*, **33**, 435–442.
6. Peters, S.U., Beaudet, A.L., Madduri, N. and Bacino, C.A. (2004) Autism in Angelman syndrome: implications for autism research. *Clin. Genet.*, **66**, 530–536.
7. Moretti, P., Bouwknecht, J.A., Teague, R., Paylor, R. and Zoghbi, H.Y. (2005) Abnormalities of social interactions and home cage behavior in a mouse model of Rett syndrome. *Hum. Mol. Genet.*, **14**, 205–220.
8. Watson, P., Black, G., Ramsden, S., Barrow, M., Super, M., Kerr, B. and Clayton-Smith, J. (2001) Angelman syndrome phenotype associated with mutations in *MECP2*, a gene encoding a methyl CpG binding protein. *J. Med. Genet.*, **38**, 224–228.
9. Lam, C., Yeung, W., Ko, C., Poon, P., Tong, S., Chan, K., Lo, I., Chan, L., Hui, J., Wong, V. *et al.* (2000) Spectrum of mutations in the *MECP2* gene in patients with infantile autism and Rett syndrome. *J. Med. Genet.*, **37**, E41.
10. Beyer, K.S., Blasi, F., Bacchelli, E., Klauck, S.M., Maestrini, E., Poustka, A. and International Molecular Genetic Study of Autism (2002) Mutation analysis of the coding sequence of the *MECP2* gene in infantile autism. *Hum. Genet.*, **111**, 305–309.
11. Schroer, R.J., Phelan, M.C., Michaelis, R.C., Crawford, E.C., Skinner, S.A., Cuccaro, M., Simensen, R.J., Bishop, J., Skinner, C., Fender, D. *et al.* (1998) Autism and maternally derived aberrations of chromosome 15q. *Am. J. Med. Genet.*, **76**, 327–336.
12. Wagstaff, J., Chaillet, J.R. and Lalande, M. (1991) The GABAA receptor beta 3 subunit gene: characterization of a human cDNA from chromosome 15q11q13 and mapping to a region of conserved synteny on mouse chromosome 7. *Genomics*, **11**, 1071–1078.
13. Cook, E.H., Courchesne, R.Y., Cox, N.J., Lord, C., Gonen, D., Guter, S.J., Lincoln, A., Nix, K., Haas, R., Leventhal, B.L. *et al.* (1998) Linkage-disequilibrium mapping of autistic disorder, with 15q11–q13 markers. *Am. J. Hum. Genet.*, **62**, 1077–1083.
14. Buxbaum, J.D., Silverman, J.M., Smith, C.J., Greenberg, D.A., Kilifarski, M., Reichert, J., Cook, E.H., Fang, Y., Song, C.Y. and Vitale, R. (2002) Association between a GABRB3 polymorphism and autism. *Mol. Psychiat.*, **7**, 311–316.
15. Nicholls, R.D., Gottlieb, W., Russell, L.B., Davda, M., Horsthemke, B. and Rinchik, E.M. (1993) Evaluation of potential models for imprinted and nonimprinted components of human chromosome 15q11–q13 syndromes by fine-structure homology mapping in the mouse. *Proc. Natl Acad. Sci. USA*, **90**, 2050–2054.
16. Samaco, R.C., Nagarajan, R.P., Braunschweig, D. and LaSalle, J.M. (2004) Multiple pathways regulate MeCP2 expression in normal brain development and exhibit defects in autism-spectrum disorders. *Hum. Mol. Genet.*, **13**, 629–639.
17. Lewis, J.D., Meehan, R.R., Henzel, W.J., Maurer-Fogy, I., Jeppesen, P., Klein, F. and Bird, A. (1992) Purification, sequence, and cellular localization of a novel chromosomal protein that binds to methylated DNA. *Cell*, **69**, 905–914.
18. Nan, X., Campoy, F.J. and Bird, A. (1997) MeCP2 is a transcriptional repressor with abundant binding sites in genomic chromatin. *Cell*, **88**, 471–481.
19. Nan, X., Ng, H.H., Johnson, C.A., Laherty, C.D., Turner, B.M., Eisenman, R.N. and Bird, A. (1998) Transcriptional repression by the methyl-CpG-binding protein MeCP2 involves a histone deacetylase complex. *Nature*, **393**, 386–389.

20. Jones, P.L., Veenstra, G.J., Wade, P.A., Vermaak, D., Kass, S.U., Landsberger, N., Strouboulis, J. and Wolffe, A.P. (1998) Methylated DNA and MeCP2 recruit histone deacetylase to repress transcription. *Nat. Genet.*, **19**, 187–191.
21. Fuks, F., Hurd, P.J., Wolf, D., Nan, X., Bird, A.P. and Kouzarides, T. (2003) The methyl-CpG-binding protein MeCP2 links DNA methylation to histone methylation. *J. Biol. Chem.*, **278**, 4035–4040.
22. Kimura, H. and Shiota, K. (2003) Methyl-CpG-binding protein, MeCP2, is a target molecule for maintenance DNA methyltransferase, Dnmt1. *J. Biol. Chem.*, **278**, 4806–4812.
23. Guy, J., Hendrich, B., Holmes, M., Martin, J.E. and Bird, A. (2001) A mouse Mecp2-null mutation causes neurological symptoms that mimic Rett syndrome. *Nat. Genet.*, **27**, 322–326.
24. Chen, R.Z., Akbarian, S., Tudor, M. and Jaenisch, R. (2001) Deficiency of methyl-CpG binding protein-2 in CNS neurons results in a Rett-like phenotype in mice. *Nat. Genet.*, **27**, 327–331.
25. Shahbazian, M., Young, J., Yuva-Paylor, L., Spencer, C., Antalffy, B., Noebels, J., Armstrong, D., Paylor, R. and Zoghbi, H. (2002) Mice with truncated MeCP2 recapitulate many Rett syndrome features and display hyperacetylation of histone H3. *Neuron*, **35**, 243–254.
26. Tudor, M., Akbarian, S., Chen, R.Z. and Jaenisch, R. (2002) Transcriptional profiling of a mouse model for Rett syndrome reveals subtle transcriptional changes in the brain. *Proc. Natl Acad. Sci. USA*, **99**, 15536–15541.
27. Colantuoni, C., Jeon, O.H., Hyder, K., Chenchik, A., Khimani, A.H., Narayanan, V., Hoffman, E.P., Kaufmann, W.E., Naidu, S. and Pevsner, J. (2001) Gene expression profiling in postmortem Rett syndrome brain: differential gene expression and patient classification. *Neurobiol. Dis.*, **8**, 847–865.
28. Traynor, J., Agarwal, P., Lazzaroni, L. and Francke, U. (2002) Gene expression patterns vary in clonal cell cultures from Rett syndrome females with eight different MECP2 mutations. *BMC Med. Genet.*, **3**, 12.
29. Chen, W.G., Chang, Q., Lin, Y., Meissner, A., West, A.E., Griffith, E.C., Jaenisch, R. and Greenberg, M.E. (2003) Derepression of BDNF transcription involves calcium-dependent phosphorylation of MeCP2. *Science*, **302**, 885–889.
30. Martinowich, K., Hattori, D., Wu, H., Fouse, S., He, F., Hu, Y., Fan, G. and Sun, Y.E. (2003) DNA methylation-related chromatin remodeling in activity-dependent BDNF gene regulation. *Science*, **302**, 890–893.
31. Balmer, D., Arredondo, J., Samaco, R.C. and LaSalle, J.M. (2002) MECP2 mutations in Rett syndrome adversely affect lymphocyte growth, but do not affect imprinted gene expression in blood or brain. *Hum. Genet.*, **110**, 545–552.
32. Braunschweig, D., Simcox, T., Samaco, R.C. and LaSalle, J.M. (2004) X-chromosome inactivation ratios affect wild-type MeCP2 expression within mosaic Rett syndrome and Mecp2^{-/+} mouse brain. *Hum. Mol. Genet.*, **13**, 1275–1286.
33. Runte, M., Huttenhofer, A., Gross, S., Kieffmann, M., Horsthemke, B. and Buiting, K. (2001) The IC-SNURF-SNRPN transcript serves as a host for multiple small nucleolar RNA species and as an antisense RNA for UBE3A. *Hum. Mol. Genet.*, **10**, 2687–2700.
34. Gregory, R.I., Randall, T.E., Johnson, C.A., Khosla, S., Hatada, I., O'Neill, L.P., Turner, B.M. and Feil, R. (2001) DNA methylation is linked to deacetylation of histone H3, but not H4, on the imprinted genes Snrpn and U2af1-rs1. *Mol. Cell. Biol.*, **21**, 5426–5436.
35. Balmer, D., Goldstine, J., Rao, Y.M. and LaSalle, J.M. (2003) Elevated methyl-CpG-binding protein 2 expression is acquired during postnatal human brain development and is correlated with alternative polyadenylation. *J. Mol. Med.*, **81**, 61–68.
36. Rougeulle, C., Glat, H. and Lalande, M. (1997) The Angelman syndrome candidate gene, UBE3A/E6-AP, is imprinted in brain (letter). *Nat. Genet.*, **17**, 14–15.
37. Jiang, Y.H., Sahoo, T., Michaelis, R.C., Bercovich, D., Bressler, J., Kashork, C.D., Liu, Q., Shaffer, L.G., Schroer, R.J., Stockton, D.W. et al. (2004) A mixed epigenetic/genetic model for oligogenic inheritance of autism with a limited role for UBE3A. *Am. J. Med. Genet.*, **131A**, 1–10.
38. Cook, E.H., Jr, Courchesne, R.Y., Cox, N.J., Lord, C., Gonen, D., Guter, S.J., Lincoln, A., Nix, K., Haas, R., Leventhal, B.L. et al. (1998) Linkage-disequilibrium mapping of autistic disorder, with 15q11–q13 markers. *Am. J. Hum. Genet.*, **62**, 1077–1083.
39. McCauley, J.L., Olson, L.M., Delahanty, R., Amin, T., Nurmi, E.L., Organ, E.L., Jacobs, M.M., Folstein, S.E., Haines, J.L. and Sutcliffe, J.S. (2004) A linkage disequilibrium map of the 1-Mb 15q12 GABA(A) receptor subunit cluster and association to autism. *Am. J. Med. Genet.*, **131B**, 51–59.
40. IMGSOA Consortium (1998) A full genome screen for autism with evidence for linkage to a region on chromosome 7q. *Hum. Mol. Genet.*, **7**, 571–578.
41. Philippe, A., Martinez, M., Guilloud-Bataille, M., Gillberg, C., Rastam, M., Sponheim, E., Coleman, M., Zappella, M., Aschauer, H., Van Maldergem, L. et al. (1999) Genome-wide scan for autism susceptibility genes. Paris Autism Research International Sibpair Study. *Hum. Mol. Genet.*, **8**, 805–812.
42. IMGSOA Consortium (2001) A genomewide screen for autism: strong evidence for linkage to chromosomes 2q, 7q, and 16p. *Am. J. Hum. Genet.*, **69**, 570–581.
43. Yonan, A.L., Alarcon, M., Cheng, R., Magnusson, P.K., Spence, S.J., Palmer, A.A., Grunn, A., Juo, S.H., Terwilliger, J.D., Liu, J. et al. (2003) A genomewide screen of 345 families for autism-susceptibility loci. *Am. J. Hum. Genet.*, **73**, 886–897.
44. Salmon, B., Hallmayer, J., Rogers, T., Kalaydjieva, L., Petersen, P.B., Nicholas, P., Pingree, C., McMahon, W., Spiker, D., Lotspeich, L. et al. (1999) Absence of linkage and linkage disequilibrium to chromosome 15q11–q13 markers in 139 multiplex families with autism. *Am. J. Med. Genet.*, **88**, 551–556.
45. Shao, Y., Cuccaro, M.L., Hauser, E.R., Raiford, K.L., Menold, M.M., Wolpert, C.M., Ravan, S.A., Elston, L., Decena, K., Donnelly, S.L. et al. (2003) Fine mapping of autistic disorder to chromosome 15q11–q13 by use of phenotypic subtypes. *Am. J. Hum. Genet.*, **72**, 539–548.
46. Nurmi, E.L., Amin, T., Olson, L.M., Jacobs, M.M., McCauley, J.L., Lam, A.Y., Organ, E.L., Folstein, S.E., Haines, J.L. and Sutcliffe, J.S. (2003) Dense linkage disequilibrium mapping in the 15q11–q13 maternal expression domain yields evidence for association in autism. *Mol. Psychiatr.*, **8**, 624–634, 570.
47. Veenstra-VanderWeele, J., Gonen, D., Leventhal, B.L. and Cook, E.H., Jr (1999) Mutation screening of the UBE3A/E6-AP gene in autistic disorder. *Mol. Psychiatr.*, **4**, 64–67.
48. Nurmi, E.L., Bradford, Y., Chen, Y.-h., Hall, J., Arnone, B., Gardiner, M.B., Hutcheson, H.B., Gilbert, J.R., Pericak-Vance, M.A., Copeland-Yates, S.A. et al. (2001) Linkage disequilibrium at the Angelman syndrome gene UBE3A in autism families. *Genomics*, **77**, 105–113.
49. Sinkkonen, S.T., Homanics, G.E. and Korpi, E.R. (2003) Mouse models of Angelman syndrome, a neurodevelopmental disorder, display different brain regional GABA(A) receptor alterations. *Neurosci. Lett.*, **340**, 205–208.
50. Buettner, V.L., Longmate, J.A., Barish, M.E., Mann, J.R. and Singer-Sam, J. (2004) Analysis of imprinting in mice with uniparental duplication of proximal chromosomes 7 and 15 by use of a custom oligonucleotide microarray. *Mamm. Genome*, **15**, 199–209.
51. Blatt, G.J., Fitzgerald, C.M., Guptill, J.T., Booker, A.B., Kemper, T.L. and Bauman, M.L. (2001) Density and distribution of hippocampal neurotransmitter receptors in autism: an autoradiographic study. *J. Autism Dev. Disord.*, **31**, 537–543.
52. Moreno-Fuenmayor, H., Borjas, L., Arrieta, A., Valera, V. and Socorro-Candanoza, L. (1996) Plasma excitatory amino acids in autism. *Invest. Clin.*, **37**, 113–128.
53. Dhossche, D., Applegate, H., Abraham, A., Maertens, P., Bland, L., Bencsath, A. and Martinez, J. (2002) Elevated plasma gamma-aminobutyric acid (GABA) levels in autistic youngsters: stimulus for a GABA hypothesis of autism. *Med. Sci. Monit.*, **8**, PR1–6.
54. Aldred, S., Moore, K.M., Fitzgerald, M. and Waring, R.H. (2003) Plasma amino acid levels in children with autism and their families. *J. Autism Dev. Disord.*, **33**, 93–97.
55. Jiang, Y.H., Armstrong, D., Albrecht, U., Atkins, C.M., Noebels, J.L., Eichele, G., Sweatt, J.D. and Beaudet, A.L. (1998) Mutation of the Angelman ubiquitin ligase in mice causes increased cytoplasmic p53 and deficits of contextual learning and long-term potentiation. *Neuron*, **21**, 799–811.
56. DeLorey, T.M., Handforth, A., Anagnostaras, S.G., Homanics, G.E., Minassian, B.A., Asatourian, A., Fanselow, M.S., Delgado-Escueta, A., Ellison, G.D. and Olsen, R.W. (1998) Mice lacking the beta3 subunit of the GABAA receptor have the epilepsy phenotype and many of the behavioral characteristics of Angelman syndrome. *J. Neurosci.*, **18**, 8505–8514.
57. Murrell, A., Heeson, S. and Reik, W. (2004) Interaction between differentially methylated regions partitions the imprinted genes Igf2 and H19 into parent-specific chromatin loops. *Nat. Genet.*, **36**, 889–893.
58. Weitzel, J.M., Buhrmester, H. and Stratling, W.H. (1997) Chicken MAR-binding protein ARBP is homologous to rat methyl-CpG-binding protein MeCP2. *Mol. Cell. Biol.*, **17**, 5656–5666.

59. Cavaille, J., Buiting, K., Kieffmann, M., Lalande, M., Brannan, C.I., Horsthemke, B., Bachellerie, J.P., Brosius, J. and Huttenhofer, A. (2000) Identification of brain-specific and imprinted small nucleolar RNA genes exhibiting an unusual genomic organization. *Proc. Natl Acad. Sci. USA*, **97**, 14311–14316.
60. de los Santos, T., Schweizer, J., Rees, C.A. and Francke, U. (2000) Small evolutionarily conserved RNA, resembling C/D box small nucleolar RNA, is transcribed from PWCR1, a novel imprinted gene in the Prader–Willi deletion region, which is highly expressed in brain. *Am. J. Hum. Genet.*, **67**, 1067–1082.
61. Meguro, M., Kashiwagi, A., Mitsuya, K., Nakao, M., Kondo, I., Saitoh, S. and Oshimura, M. (2001) A novel maternally expressed gene, *ATP10C*, encodes a putative aminophospholipid translocase associated with Angelman syndrome. *Nat. Genet.*, **28**, 19–20.
62. Herzing, L.B.K., Kim, S.-J., Cook, E.H. and Ledbetter, D.H. (2001) The human aminophospholipid-transporting ATPase gene *ATP10C* maps adjacent to UBE3A and exhibits similar imprinted expression. *Am. J. Hum. Genet.*, **68**, 1501–1505.
63. Sinnett, D., Wagstaff, J., Glatt, K., Woolf, E., Kirkness, E.J. and Lalande, M. (1993) High-resolution mapping of the gamma-aminobutyric acid receptor subunit beta 3 and alpha 5 gene cluster on chromosome 15q11–q13, and localization of breakpoints in two Angelman syndrome patients. *Am. J. Hum. Genet.*, **52**, 1216–1229.
64. LaSalle, J. and Lalande, M. (1996) Homologous association of oppositely imprinted chromosomal domains. *Science*, **272**, 725–728.
65. Tsai, T.F., Bressler, J., Jiang, Y.H. and Beaudet, A.L. (2003) Disruption of the genomic imprint in trans with homologous recombination at *Snrpn* in ES cells. *Genesis*, **37**, 151–161.
66. LaSalle, J., Goldstine, J., Balmer, D. and Greco, C. (2001) Quantitative localization of heterologous methyl-CpG-binding protein 2 (MeCP2) expression phenotypes in normal and Rett syndrome brain by laser scanning cytometry. *Hum. Mol. Genet.*, **10**, 1729–1740.
67. Yamasaki, K., Joh, K., Ohta, T., Masuzaki, H., Ishimaru, T., Mukai, T., Niikawa, N., Ogawa, M., Wagstaff, J. and Kishino, T. (2003) Neurons but not glial cells show reciprocal imprinting of sense and antisense transcripts of *Ube3a*. *Hum. Mol. Genet.*, **12**, 837–847.
68. Yoon, B.J., Herman, H., Sikora, A., Smith, L.T., Plass, C. and Soloway, P.D. (2002) Regulation of DNA methylation of *Rasgrf1*. *Nat. Genet.*, **30**, 92–96.
69. Yasui, D., Miyano, M., Cai, S., Varga-Weisz, P. and Kohwi-Shigematsu, T. (2002) SATB1 targets chromatin remodelling to regulate genes over long distances. *Nature*, **419**, 641–645.
70. Yang, Y., Hu, J.F., Ulaner, G.A., Li, T., Yao, X., Vu, T.H. and Hoffman, A.R. (2003) Epigenetic regulation of *Igf2/H19* imprinting at CTCF insulator binding sites. *J. Cell. Biochem.*, **90**, 1038–1055.