Ing1 functions in DNA demethylation by directing Gadd45a to H3K4me3

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Active DNA demethylation regulates epigenetic gene activation in numerous processes, but how the target site specificity of DNA demethylation is determined and what factors are involved are still poorly understood. Here we show that the tumor suppressor inhibitor of growth protein 1 (Ing1) is required for targeting active DNA demethylation. Ing1 functions by recruiting the regulator of DNA demethylation growth arrest and DNA damage protein 45a (Gadd45a) to histone H3 trimethylated at Lys 4 (H3K4me3). We show that reduced H3K4 methylation impairs recruitment of Gadd45a/Ing1 and gene-specific DNA demethylation. Our results indicate that histone methylation directs DNA demethylation.

[**Keywords:** p33^{Ing1b}; Gadd45a; Ing1; DNA demethylation; H3K4me3; chromatin]

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DNA methylation at 5-methylcytosine (5mC) of CpGs is a common epigenetic mark in metazoa, which plays important roles in regulating gene expression, genomic imprinting, X-chromosome inactivation, genomic instability, embryonic development, and cancer (Costello and Plass 2001; Jones and Takai 2001; Ehrlich 2005). It has become clear that DNA methylation is reversible by enzymatic “active” DNA demethylation, with examples in plants, animal development, cancer, and immune cells (for review, see Niehrs 2009; Zhu 2009; Schär and Fritsch 2011). However, the molecular mechanisms underlying active demethylation are only beginning to be understood.

In *Arabidopsis*, active DNA demethylation involves DNA base excision repair of methylated cytosines, mediated by the Demeter/ROS1 DNA glycosylase family (Choi et al. 2002; Gong et al. 2002; Gehring et al. 2006). In animal cells, various DNA repair-based mechanisms seem to mediate active demethylation involving base excision repair (Kangaspeska et al. 2008; Metivier et al. 2008; Rai et al. 2008; Bhutani et al. 2010; Cortazar et al. 2011; Cortellino et al. 2011) and/or methyl oxidation (Kriaucionis and Heintz 2009; Tahiliani et al. 2009; Guo et al. 2011; He et al. 2011). We showed that growth arrest and DNA damage protein 45a (Gadd45a) mediates active DNA demethylation (Barreto et al. 2007). *Gadd45a* is member of a small gene family of stress response genes encoding 18-kDa acidic histone fold proteins (Zhan et al.

1994). Gadd45 proteins are multifunctional and regulate a range of cellular processes, including DNA repair, proliferation, apoptosis, and differentiation (Fornace et al. 1988; Kastan et al. 1992; Smith et al. 1996; Carrier et al. 1999; Harkin et al. 1999; Hollander and Fornace 2002; Zhan 2005). Gadd45a-mediated demethylation involves recruitment of the nucleotide excision repair (NER) (Barreto et al. 2007; Schmitz et al. 2009; Le May et al. 2010; Schäfer et al. 2010) and/or base excision repair (Rai et al. 2008; Cortellino et al. 2011) machineries (for review, see Niehrs and Schäfer 2012).

Gadd45a does not appear to affect global DNA demethylation (Jin et al. 2008; Engel et al. 2009; Schäfer et al. 2010). Instead, Gadd45 proteins function in gene-specific demethylation of target genes such as *rDNA* in HEK293T cells (Schmitz et al. 2009); *Bdnf IX* and *Fgf-1B* during neurogenesis in the mouse brain (Ma et al. 2009); *Bdnf IX* in psychosis in the human brain (Gavin et al. 2011); *reelin* and *Bdnf IX* upon activation of glutamate receptors (Matriciano et al. 2011); *S100* during epidermal differentiation (Sen et al. 2010); *CD11a* and *CD70* in T cells (Li et al. 2010); *RARβ2* following retinoic acid stimulation (Le May et al. 2010); *osterix*, *Dlx5*, *Runx2*, and *BGP* during bone differentiation (Zhang et al. 2011); and plasmid DNA in *Xenopus* oocytes and zebrafish embryos (Barreto et al. 2007; Rai et al. 2008). A common theme of these studies is that demethylation by Gadd45 is a highly selective process: Not only is it gene specific, but within a given gene, it typically affects distinct mCpGs, often in the promoter region. This specificity highlights a set of general, unresolved key questions in DNA demethylation: What determines the target site specificity of DNA

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demethylation? Is there a relationship between site-specific DNA demethylation and the epigenetic landscape? What may be the cofactors involved?

Gadd45a is a nuclear protein but does not bind significantly to naked DNA (Carrier et al. 1999; Sytnikova et al. 2011), yet chromatin immunoprecipitation (ChIP) shows specific recruitment to demethylation sites (Barreto et al. 2007; Ma et al. 2009; Schmitz et al. 2009; Le May et al. 2010; Gavin et al. 2011), suggesting the involvement of cofactors. Gadd45a physically interacts with a number of DNA/chromatin-binding proteins, including nuclear hormone receptors (Yi et al. 2000), core histones (Carrier et al. 1999), PCNA (Smith et al. 1994), Taf12 (Schmitz et al. 2009), and p33^{ING1b} (Cheung et al. 2001). In this study, we focused on p33^{ING1b} (ING1b [inhibitor of growth 1b]) because it shares various properties with Gadd45a. ING1b is member of the ING (ING1 to ING5) family of tumor suppressors, most of which contain a conserved plant homeodomain (PHD) finger motif (Garkavtsev et al. 1996; Campos et al. 2004; for review, see Soliman and Riabowol 2007). Like Gadd45a, ING1 is induced by stress or UV irradiation, inhibits cell growth by interacting with p21^{waf1}, and promotes NER (Cheung et al. 2001). Furthermore, *Ing1* mouse mutants are radiation-sensitive and tumor-prone, like *Gadd45a* mutants (Hollander et al. 1999; Kichina et al. 2006). ING proteins are found in complex with histone acetyl transferases (HATs) (Vieyra et al. 2002) and histone deacetylases (HDACs) (Skowrya et al. 2001; Kuzmichev et al. 2002). Important for this study, ING1 contains a PHD finger domain that specifically binds to histone H3 trimethylated at Lys 4 (H3K4me3) (Shi et al. 2006; Wysocka et al. 2006), suggesting that ING1 can read and translate the histone code (Shi et al. 2006; Pena et al. 2008).

We discovered that Ing1 is a novel factor required for gene-specific DNA demethylation. Gain- and loss-of-function experiments indicate that Ing1 functions during Gadd45a-mediated DNA demethylation. ChIP experiments and manipulation of cellular H3K4me3 levels show that ING1b recruits GADD45a to the promoter of the cancer/testis antigen *melanoma antigen gene B2* (*MAGEB2*). Genome-wide gain- and loss-of-function experiments identified additional target genes regulated synergistically by Gadd45a and Ing1 in a H3K4me3-dependent fashion. Thus, this study describes a novel factor essential for DNA demethylation and indicates that histone methylation is required to direct DNA demethylation.

Results

GADD45a requires ING1b for DNA demethylation

We first analyzed whether GADD45a and ING1b biochemically interact. In coimmunoprecipitation (co-IP), endogenous GADD45a bound ING1 in HEK293T lysates (Fig. 1A). Using in vitro pull-downs, we mapped the GADD45a-binding domain to lie between the PHD and the partial bromodomain, a region without distinctive protein motifs (Supplemental Fig. S1A). We next established

that Gadd45a and ING1b also functionally interact by performing luciferase assays using an in vitro methylated, SV40-driven reporter plasmid. Gadd45a reactivates such methylation silenced reporters by DNA demethylation (Barreto et al. 2007; Schmitz et al. 2009; Hu et al. 2010). In cotransfection experiments, *ING1b* strongly enhanced reporter activation by *Xenopus Gadd45a* in a dose-dependent manner (Fig. 1B), and this was accompanied by DNA demethylation of the reporter (Fig. 1C). The ING1b domains required for Gadd45a functional interaction mapped to the C terminus, including the PHD domain (Fig. 1D). Also, a W235A point mutation, which disrupts the PHD domain (Pena et al. 2008), was inhibitory (Supplemental Fig. S1B). In loss-of-function experiments, si*ING1b* abolished reporter gene activation (Fig. 1E; controlled in Supplemental Fig. S1C), while siRNA knockdown of another *ING1* splice variant, p47^{ING1a} (*ING1a*), lacking the N-terminal PIP and partial bromodomain enhanced the reporter activation by Gadd45a (Fig. 1E).

Not only human ING1b, but also both mouse Ing1 proteins (p31^{Ing1a/c} and p37^{Ing1b}) (Zeremski et al. 1999; Kichina et al. 2006) synergized with mouse Gadd45a in reporter reactivation (Supplemental Fig. S1D,E). Furthermore, another reporter, *Oct4TK-GFP*, was synergistically activated by *ING1b* and human *GADD45a* (Fig. 1F). Bisulfite sequencing of the *Oct4TK-GFP* reporter confirmed site-specific DNA demethylation after *GADD45a* transfection, which was potentiated by *ING1b* (Fig. 1G). This effect was direct, as both proteins bound the reporter plasmid in ChIP experiments (Fig. 1H,I). Notably, GADD45a binding was significantly increased by *ING1b* expression (Fig. 1I), suggesting that ING1b enhances GADD45a recruitment to target reporters. Importantly, reporter binding of both ING1b and GADD45a was fully dependent on the ING1b-PHD domain, suggesting a targeting to chromatin, and, indeed, the reporter plasmid was chromatinized (Supplemental Fig. S1F,G).

These results indicate that (1) Gadd45a and ING1b biochemically and functionally interact in DNA demethylation; (2) ING1b—and in particular its chromatin-binding PHD domain—is required for the Gadd45a-mediated reactivation of methylation silenced reporters; and (3) the proteins directly bind to the demethylation substrate in a PHD domain-dependent manner.

ING1b targets GADD45a to MAGEB2 demethylation sites

By a candidate approach, we identified *MAGEB2* as an endogenous demethylation target of GADD45a and ING1b. The expression of the gene family of cancer/testis antigens is high in testis and silenced in somatic tissues but is activated due to DNA demethylation specifically in many invasive cancers (De Smet et al. 1996, 1999). *MAGE* genes are promising targets for anti-tumor immunotherapy (Sang et al. 2011).

MAGEB2 expression was inhibited by *GADD45a* or *ING1b* siRNAs in RKO cells (Fig. 2A; controlled in Supplemental Fig. S2A), and this was accompanied by

Figure 1. ING1b binds GADD45a and is required for DNA demethylation of methylated reporter plasmids. (A) Co-IP and Western blot analysis of GADD45a and ING1 in HEK293T cells. Input shows 5% of lysates used for immunoprecipitation. (B) Luciferase reporter assays of HEK293T cells transiently transfected with empty vector (Co) or *Xenopus Gadd45a* (xG45a) and human *ING1b* along with in vitro methylated *SV40* reporter plasmid. (C) DNA methylation analysis by 454 bisulfite sequencing of the luciferase reporter plasmid treated as in B. (D,E) Luciferase reporter assays as in B using *ING1b* deletion mutants (D) (see scheme in Supplemental Fig. S1A) or the indicated siRNAs (E). (F,G) HEK293T cells transiently transfected with empty vector (Co), human HA-GADD45a, and myc-ING1b along with in vitro methylated *Oct4TK-GFP* reporter plasmid were analyzed for relative GFP expression monitored at the indicated time points after transfection by qPCR (F) or DNA methylation of three CpGs (-299, -10, +93) of the *Oct4TK-GFP* regulatory region by 454 bisulfite sequencing (G). (H,I) ChIP of myc-ING1b (H) or HA-GADD45a (I) of two regions of the *Oct4TK-GFP* reporter spanning CpG -299 or the ATG (see scheme in G) using HEK293T chromatin (harvested 48 h after transfection) as described in F and G. ChIP using IgG served as negative control. (ATG) Translation start site. Bar charts represent the mean of independent experiments ($n = 3$) with error bars as $\pm$ standard deviation (SD).

promoter hypermethylation (Fig. 2B). Specifically, the *MAGEB2* promoter was generally hypermethylated except for three adjacent CpGs located at -70, -65, and -57, which were methylated only between 30% and 60%, (Supplemental Fig. S2B). *GADD45a* or *ING1b* siRNAs led to hypermethylation of these three CpGs (Fig. 2B). Consistent with a direct effect, transfected epitope-tagged GADD45a bound the *MAGEB2* locus, with the highest enrichment at the transcription start site, as did myc-ING1b (Supplemental Fig. S2C). Sequential re-ChIP assays confirmed that ING1b and GADD45a coassociate in the same chromatin fraction (Fig. 2C). Moreover, endogenous GADD45a also bound to the *MAGEB2* 5' upstream region, and importantly, this binding was impaired by si*ING1b* (Fig. 2D). ING1b binding, however, was not affected by depletion of *GADD45a* (Supplemental Fig. S2D). We conclude that (1) ING1b and GADD45a maintain hypomethylation of the *MAGEB2* promoter and prevent gene silencing and (2) ING1b targets GADD45a to specific demethylation sites of *MAGEB2*.

ING1b targets *GADD45a* to H3K4me3 via its PHD domain

A hallmark of ING1b is its ability to read and translate H3K4me3 histone marks via its PHD domain. This suggests that Gadd45a-mediated DNA demethylation is targeted to H3K4me3. Consistent with this, ING1b Δ PHD and ING1b^{W235A} failed to synergize with Gadd45a in transcriptional activation of the methylated luciferase reporter (Fig. 1D; Supplemental Fig. S1B) and likewise impaired recruitment of Gadd45a to the *Oct4TK-GFP* reporter and its DNA demethylation (Fig. 1I; data not shown).

H3K4me3 is commonly associated with the promoter regions of actively transcribed genes (Santos-Rosa et al. 2002; Schneider et al. 2004). As expected, H3K4me3 showed maximal occupancy around the transcription start site of *MAGEB2* (Fig. 2E), overlapping with endogenous GADD45a binding. Sequential re-ChIP assays confirmed that GADD45a and ING1b both bind to chromatin occupied by H3K4me3 (Fig. 2F,G).

We performed pull-down assays using biotinylated histone peptides to confirm that Gadd45a can be recruited

Figure 2. ING1b targets GADD45a to the *MAGEB2* demethylation site. (A,B) RKO cells were transiently transfected for 48 h with control (Co), *ING1b*-specific, or *GADD45a*-specific siRNA. (A) Relative *MAGEB2* expression analysis. (B) Bisulfite sequencing analysis of genomic DNA of CpGs at -70 , -65 , and -57 relative to the *MAGEB2* transcription start site. Each line represents one clone. White and black circles indicate unmethylated and methylated CpG, respectively (for complete *MAGEB2* promoter methylation profile, see Supplemental Fig. S2A). (Left) The diagram indicates the analyzed CpG at -57 and sites a–d used for ChIP analysis. (TSS) Transcription start sites; (ATG) translation start sites. (C) Re-ChIP of RKO cells transfected with *SMAD4* (Co) or *HA-GADD45a* and *myc-ING1b*. A second ChIP was performed on the precipitates of the first ChIP using the indicated antibodies. Material of the first and the second ChIP was used for PCR using primers specific for site a of *MAGEB2* (see scheme in B) and for an intergenic control region (intergenic). (D) Endogenous GADD45a ChIP of *MAGEB2* regulatory regions marked in the scheme in B in RKO cells treated as in A and B. (Co) qPCR of an intergenic control region. IgG ChIP served as negative control (data not shown). Level of significance (*) is indicated versus siControl for each region. (E) H3K4me3 ChIP of *MAGEB2* regulatory regions marked in the scheme in B in untreated RKO cells. (F,G) H3K4me3 Re-ChIP of RKO cells transfected with *SMAD4* (Co) and *HA-GADD45a* (HA-G45a) (F) or *myc-ING1b* (G) as described in C. (H) Western blot analysis of in vitro pull-downs from nuclear extracts of HEK293T cells transiently transfected with the indicated constructs using biotinylated

histone peptides carrying the indicated modifications. Bar charts represent the mean of independent experiments ($n = 3$) with error bars as $\pm$ SD.

to H3K4me3 via ING1b. GADD45a alone did not bind to any of the tested histone peptides. However, when co-expressed with ING1b, GADD45a bound to H3K4me2 and H3K4me3. Expectedly, ING1b alone also bound specifically to H3K4me3 and, with decreasing affinity, di- and monomethylated H3K4 (Fig. 2H; Supplemental Fig. S3A; Pena et al. 2008). Deletion of the PHD domain (ING1b Δ PHD) or W235A point mutation abolished binding to H3K4me3 and recruitment of Gadd45a (Fig. 2H, Supplemental Fig. S3B). Toward acetylated peptides (H3K9ac and H3K14ac), ING1b had no affinity, even though it contains a partial bromodomain (Fig. 2H). These results indicate that ING1b can recruit GADD45a to H3K4me3 via its PHD domain.

Gadd45a and *Ing1* mediate *Mageb2* demethylation

We next analyzed *Mageb2* regulation genetically in primary mouse embryonic fibroblasts (MEFs), which we generated from mice single mutant (Hollander et al. 1999;

Kichina et al. 2006) or double mutant (double knockout [DKO]) (this study) for *Gadd45a* and *Ing1*. *Gadd45a* and *Ing1* are UV-inducible stress response genes (Fornace et al. 1988; Hollander et al. 1999; Cheung et al. 2001). Consistent with this, *Gadd45a* was up-regulated in UV-irradiated wild-type MEFs, as were *Ing1a* and *Ing1c*, two of the three highly related alternative transcripts encoded by the mouse *Ing1* locus (Fig. 3A; Supplemental Fig. S4A). UV irradiation also strongly induced expression of *Mageb1–3* (gene expression average of the three highly similar mouse homologs to the single human *MAGEB2*) (Fig. 3B; De Backer et al. 1995). Importantly, *Mageb1–3* induction was reduced in *Gadd45a* and *Ing1* single-mutant MEFs and was almost abolished in DKO MEFs (Fig. 3C). UV irradiation also induced 25% *Mageb2* promoter demethylation (Fig. 3D,E, CpG sites a and b; schematic in Supplemental Fig. S4B). This demethylation was reduced in *Gadd45a* and *Ing1* single-mutant MEFs and was abolished in DKO MEFs (Fig. 3E). These results indicate that Gadd45a and Ing1 are required for

Figure 3. Gadd45a and Ing1 are required for UV-induced *Mageb2* demethylation. (A,B) Gene expression kinetics in MEFs irradiated at t_0 with 40 J/m². Relative expression was monitored by qPCR. (C) *Mageb1-3* expression in MEFs irradiated with 20 or 40 J/m² (UV+, ++). Relative expression was monitored in the indicated mutant MEFs 16 h after treatment. (D,E) *Mageb2* promoter methylation in MEFs of the experiments shown in B and C (UV++) monitored by methylation-sensitive qPCR. The *Mageb2* sites (a and b) analyzed are indicated in Supplemental Figure S4B. (F) “ChIP-Chop” assay to determine the DNA methylation state of H3K4me3-occupied *Mageb2* site a. UV-irradiated MEFs (40 J/m²) were harvested at the indicated time points. After H3K4me3 ChIP, the immunoprecipitated DNA was subjected to methylation-sensitive restriction digest followed by PCR. (G,H) MEFs were transiently transfected with the indicated siRNAs. After 36 h, cells were UV-irradiated with 40 J/m² and analyzed after 16 h. (G) H3K4me3 ChIP analysis of *Mageb2* sites indicated in the scheme in Supplemental Figure S4B. (H) Relative expression analysis of *Mageb1-3*. Bar charts represent the mean of independent experiments ($n = 3$) with error bars as $\pm$ SD.

UV-induced activation and demethylation of *Mageb2*. Of note, the basal methylation level of *Mageb2* increased from 80% to 100% in DKO MEFs (Supplemental Fig. S4C), suggesting that *Gadd45a* and *Ing1* not only mediate UV-induced demethylation, but also protect the promoter from hypermethylation in steady state.

H3K4me3 is required for Gadd45a/Ing1-induced *Mageb2* demethylation

We addressed whether *Gadd45a*/*Ing1*-mediated *Mageb2* demethylation is targeted by H3K4me3 in MEFs. Despite its low expression level, uninduced *Mageb2* was substantially bound by H3K4me3, which was unaltered by UV treatment (Supplemental Fig. S5A). However, H3K4me3 chromatin was selectively demethylated: H3K4me3-bound *Mageb2* had 20% reduced DNA methylation compared with input (60% vs. 80%) (Fig. 3F vs. D) and was preferentially demethylated following UV treatment (20% vs. 60%) (Fig. 3F vs. D). Consistent with H3K4me3 targeting demethylation, siRNA knockdown of *Wdr5*, an essential component of the MLL H3K4 methyltransferase (Milne et al. 2002; Wysocka et al. 2005), reduced global and *Mageb2* promoter-associated H3K4me3 levels (Fig. 3G; Supplemental Fig. S5B) and, importantly, also impaired basal as well as UV-induced *Mageb1-3* expression (Fig. 3H). Similarly, siWdr5 strongly impaired both *Mageb1-3* expression and demethylation when triggered

by *GADD45a*/*ING1b* overexpression instead of UV irradiation (Fig. 4A,B).

ChIP analysis showed that *GADD45a* and *ING1b* bound with a similar profile to the predicted *Mageb2* transcription start site, encompassing the demethylation sites (Fig. 4C,D). Importantly, siWdr5 impaired the binding of both proteins to *Mageb2* (Fig. 4C,D). We conclude that WDR5-mediated H3K4 methylation is essential for *GADD45a*/*ING1* to bind to the *Mageb2* locus and promote gene activation by DNA demethylation.

ING1b and *GADD45a* cooperate in gene activation and demethylation

The above data support an interaction of *Gadd45a* and *Ing1* in demethylation of reporter constructs and *MAGEB2*. To extend these findings, we performed genome-wide microarray profiling to analyze global changes in gene expression in HEK293T cells after expression of *ING1b* and *GADD45a*. If *ING1b* targets *GADD45a* in active DNA demethylation, then expression of both proteins should (1) activate rather than inhibit gene expression and (2) require the *ING1b*-PHD domain for synergism. Indeed, in the set of genes that showed a significant response only when *GADD45a* and *ING1b* were cotransfected, the majority were up-regulated (85 genes up-regulated vs. 24 genes down-regulated) (Fig. 5A; Supplemental Table S1). Individually, *GADD45a* and *ING1b* had only a modest

Figure 4. GADD45a/ING1b-induced demethylation of *Mageb2* requires H3K4me3. (A) Relative *Mageb1-3* expression in MEFs transiently transfected with the indicated siRNAs and retransfected with control DNA, *HA-GADD45a*, and *myc-ING1b*. (B) Methylation analysis of *Mageb2* using genomic DNA of MEFs from the experiment described in A. Methylation-sensitive qPCR assessing the sites a, a', and b (see scheme in Supplemental Fig. S4B) and *Gapdh*. (C,D) ChIP analysis of *Mageb2* using chromatin of MEFs from the experiment in A and B. (C) HA-GADD45a ChIP. (D) myc-ING1b ChIP. Analyzed sites are indicated (see scheme Supplemental Fig. S4B). Bar charts represent the mean of independent experiments ($n = 3$) with error bars as $\pm$ SD.

impact on gene expression. Notably, GADD45a/ING1b cooperation was fully dependent on ING1b's PHD domain, as the deletion mutant did not cooperate with GADD45a in gene induction (Fig. 5B).

Examples of genes whose expression was synergistically induced by GADD45a and ING1b are *DHRS2*, *FTCD*, *ARC*, *TCEAL7*, *VFG*, and *TAF7L*, as validated by quantitative PCR (qPCR) (Fig. 5C). We exemplarily confirmed DNA demethylation of *DHRS2* and *TAF7L* regulatory regions by GADD45a and ING1b, which once again was site-specific, affecting distinct CpGs (Fig. 5D,E). These results corroborate that ING1b and GADD45a synergistically act in gene activation and DNA demethylation.

H3K4me3 directs GADD45a/ING1b to target sites

We confirmed that recruitment of GADD45a/ING1b to some of the newly identified target genes involved H3K4me3. First, DNA demethylation and transcriptional activation of *DHRS2* and *TAF7L* required the ING1b-PHD domain (Supplemental Fig. S6A–C). Second, in ChIP experiments, GADD45a and ING1b bound to the promoters of *DHRS2*, *TCEAL7*, and *TAF7L* in a ING1b-PHD domain-dependent manner (Fig. 6A,B; Supplemental Fig. S6D). Third, siWDR5, which reduced H3K4me3 levels (Supplemental Fig. S6E), inhibited recruitment of transfected

ING1b and GADD45a (Fig. 6A,B). Importantly, endogenous ING1b/GADD45a also bound to *DHRS2*, *TCEAL7*, and *TAF7L* regulatory regions, and this binding was reduced by siWDR5 as well (Fig. 6C–E). Moreover, while siING1b reduced GADD45a recruitment to *DHRS2*, *TCEAL7*, and *TAF7L*, siGADD45a did not affect endogenous ING1b binding to these target genes (Fig. 6D,E). We conclude that WDR5-mediated H3K4 methylation is essential for GADD45a/ING1b to bind to target promoters, confer demethylation, and promote gene activation.

Ing1 and *Gadd45a* genetically interact in regulating gene expression

To further corroborate the requirement of Ing1 and H3K4me3 in targeting Gadd45a-mediated demethylation, we carried out microarray expression profiling in MEFs deficient in *Ing1*, *Gadd45a*, or both (Supplemental Fig. S7A). The majority of affected genes showed differential expression only in DKO MEFs (135 genes up-regulated, and 145 genes down-regulated) (Fig. 7A; Supplemental Table S2). Differential expression of 12 genes was confirmed by qPCR and in two independent MEF lines (Fig. 7B; data not shown). These results corroborate that *Gadd45a* and *Ing1* genetically interact in regulating gene expression.

Figure 5. ING1b and GADD45a cooperate in gene activation and demethylation in gain of function. (A) Venn diagram showing differentially expressed genes (more than twofold) in HEK293T cells upon GADD45a (gray), ING1b (blue), or combined GADD45a/ING1b (red) expression compared with control transfected cells. (B) Statistics of differentially expressed genes as in A, including samples expressing ING1bΔPHD or a combination of GADD45a/ING1bΔPHD. (C) qPCR expression analysis of the indicated target genes in transiently transfected HEK293T cells. Bars show the mean of independent experiments ($n = 8$) with error bars as $\pm$ SD. (D,E) DNA methylation analysis of the regulatory regions of *DHRS2* (D) and *TAF7L* (E) in HEK293T cells treated as in C assessed by 454 bisulfite sequencing.

Direct DNA demethylation target genes of Gadd45a/Ing1 should be hypermethylated in DKO and transcriptionally silenced. Gene class enrichment analysis of the DKO down-regulated genes revealed an overrepresentation of extracellular matrix components and proteins involved in immune response and development (Supplemental Table S3). For example, expression of the chemokine *Cxcl1* and the homeobox gene *Hoxd8* was strongly reduced in DKOs (Fig. 7B), and the majority of CpGs in their promoter region was hypermethylated, while few showed hypomethylation (Fig. 7C,D). Conversely, *Cxcl1* and *Hoxd8* were activated and bound at their promoters by transfected GADD45a and ING1b (Fig. 7E–G). Both recruitment to *Cxcl1* and *Hoxd8* and activation by GADD45a and ING1b were inhibited by siWdr5 (Fig. 7E–H). siWdr5 also impaired basal expression of *Cxcl1* and *Hoxd8* (Supplemental Fig. S7B).

These results suggest that (1) endogenous Gadd45a/Ing1 prevent silencing of *Cxcl1* and *Hoxd8* by demethylating their regulatory regions and (2) Wdr5-mediated H3K4me3 is required for this process. The evidence from both gain- and loss-of-function experiments therefore supports that the chromatin reader Ing1 recruits Gadd45a to DNA demethylation sites in a H3K4me3-dependent manner.

Discussion

The proteins regulating DNA demethylation are mostly unknown. Here we demonstrate that the tumor suppressor Ing1 is a novel factor involved in DNA demethylation. A major question concerning DNA demethylation is how the targeting to specific loci is regulated because it is essential for understanding how DNA methylation patterns are established, maintained, and remodeled. One main conclusion of this study is that Ing1 serves as a cofactor that targets Gadd45a to specific loci by its ability to bind methylated H3K4 via its PHD domain. Thereby, it acts as an adapter between chromatin and Gadd45a, which in turn can recruit DNA repair enzymes capable of replacing 5mC (Barreto et al. 2007; Rai et al. 2008; Schmitz et al. 2009; Le May et al. 2010; Cortellino et al. 2011).

Our study therefore suggests a model (Supplemental Fig. S8) whereby histone methylation functions as one determinant for active DNA demethylation. The model fits the observation that the activating histone marks H3K4me2 and H3K4me3 are preferentially deposited around transcription start sites, similar to the distribution of Gadd45a and Ing1 at *Mageb2* and other gene promoters (Barreto et al. 2007; Schmitz et al. 2009; Le May et al. 2010;

Figure 6. H3K4me3 targets GADD45a/ING1b. (A,B) myc-ING1b ChIP (A) (wild type or mutant) or HA-GADD45a ChIP (B) of the regulatory regions of selected target genes, including the negative control region *GAPDH*. HEK293T cells were transiently transfected with the indicated siRNAs followed by DNA transfection of the indicated constructs. (C–E) H3K4me3 (C), endogenous ING1 (D), and endogenous GADD45a (E) ChIP of the regulatory regions of selected target genes, including the negative control region *GAPDH* and an intergenic control region (Co; C) in HEK293T cells. ChIP using IgG served as negative control (data not shown). Bar graphs represent the mean of independent experiments ($n = 3$) with error bars as $\pm$ SD.

Cortellino et al. 2011). Also consistent with this, H3K4me2 and H3K4me3 are strongly anti-correlated with 5mC (Okitsu and Hsieh 2007; Weber et al. 2007). However, while H3K4me3 is necessary, it is clearly not sufficient as a determinant for demethylation. H3K4me3 is present at many more loci than are regulated by DNA methylation. This was also observed for the *Mageb2* promoter, which shows a broad H3K4me3 distribution, including at sites that are not demethylated (cf. Fig. 4B and Supplemental Fig. S5A). Indeed, another determinant for DNA demethylation appears to be low-level transcription (D'Alessio et al. 2007; Le May et al. 2010), and in the case of *rDNA* promoter demethylation, this involves Gadd45a recruitment via Taf12 to RNA polymerase I (Schmitz et al. 2009). Another class of Gadd45-binding cofactors that may target demethylation are nuclear hormone receptors (Ito et al. 2007; Le May et al. 2010; Cortellino et al. 2011; Tian et al. 2011; Zhang et al. 2011). A further determinant may be the presence of 5mC. For example, in *Xenopus* oocytes, Gadd45 engages excision repair on methylated but not unmethylated plasmid DNA, even though it binds to both (Barreto et al. 2007).

The fact that H3K4me3 regulates DNA demethylation is in line with the reciprocal relationship between DNA

methylation and histone modifications. For example, methylated DNA is bound by methyl-CpG-binding proteins that recruit histone methyltransferases to set repressive histone marks such as H3K9me (Fujita et al. 1999). Conversely, knockdown of the polycomb gene *EZH2* or the repressive histone methyltransferases *DIM-5* or *kryptonite* disrupt the DNA methylation pattern (Jackson et al. 2002; Tamaru and Selker 2003; Vire et al. 2006).

Finally, the results suggest that *Gadd45a* and *Ing1* not only mediate UV-induced demethylation of *Mageb2*, but also protect its promoter from hypermethylation in steady state. Thus, even for steady-state methylation, there is an equilibrium, consistent with the notion that 5mC can undergo rapid turnover (Kangaspeska et al. 2008; Metivier et al. 2008).

The unanticipated role of the tumor suppressor Ing1 in DNA demethylation raises many new questions: What determines the site-specific demethylation of distinct mCpGs within a promoter covered by H3K4me3? Which signaling cascades regulate Gadd45a/Ing1-mediated demethylation? How does their function in DNA demethylation relate to the biological processes in which Gadd45a/Ing1 have been implicated, such as autoimmunity, aging,

Figure 7. Ing1 and Gadd45a genetically interact in regulating gene expression. (A) Venn diagram showing differentially expressed genes (>1.5-fold) in *Gadd45a*^{-/-}, *Ing1*^{-/-}, or DKO MEFs compared with wild-type (WT) MEFs. (B) qPCR expression analysis of *Cxcl1* and *Hoxd8* in MEFs of the indicated genotype. Bars represent the mean of independent experiments ($n = 4$) with error bars as $\pm$ SD. (C,D) DNA methylation analysis of the regulatory region of *Cxcl1* (C) and *Hoxd8* (D) in MEFs of the indicated genotype assessed by 454 bisulfite sequencing. (E) qPCR expression analysis of the indicated genes in wild-type MEFs transiently transfected with the indicated siRNAs and retransfected with the indicated genes. (F–H) ChIP for myc-ING1b (F), HA-GADD45a (G), and H3K4me3 (H) of the regulatory regions of *Cxcl1* and *Hoxd8*, including either the negative control region *Gapdh* (F,G) or an intergenic control region (Co; H) in wild-type MEFs treated as described in E. Bars represent the mean of independent experiments ($n = 3$) with error bars as $\pm$ SD.

and cancer? These questions can now be addressed in mutant mice.

Materials and methods

Tissue culture, transfection, and siRNA treatment

HEK293T, RKO, and MEF cells were grown at 37°C in 10% CO₂ in Dulbecco's modified Eagle's medium (DMEM), 10% fetal calf serum, 2 mM L-glutamine, 100 U/mL penicillin, and 100 μ g/mL streptomycin. MEFs were generated as described (Nagy et al. 2003) from embryonic day 15.5 embryos (Hollander et al. 1999; Kichina et al. 2006). Plasmid DNA was transfected using Fugene 6 (Fig. 1B–G) or X-tremeGENE 9 (Figs. 1H, 5, 6) (both from Roche) for HEK293T cells and Lipofectamine in combination with Plus reagent (Invitrogen) for RKO cells. Cells were transfected with 25 nM *GADD45a*- or *ING1a*-specific Dharmacon siRNA Smart pools or 40 nM *WDR5*-specific siRNA (Stealth, Invitrogen) using Lipofectamine 2000 (Invitrogen). For depletion of *ING1b*, a self-designed Dharmacon siRNA was used: sense 5'-GGGAGAUC GACGCGAAUAUU-3'. For Figures 1E; 2, A, B, and D; and 6, A and B, HEK293T cells were incubated for 24 h with siRNA prior to plasmid DNA transfection. Similarly, MEFs were transfected with siRNA 24 h before plasmid DNA electroporation for

Figures 4 and 7, E–H. MEF electroporation was performed according to the manufacturers' instructions using 6 μ g of DNA and 10⁶ cells using the 100- μ L Tip Neon Transfection system (Invitrogen; 1250 V, 20 msec). For UV irradiation, MEF cells were washed with Hank's buffered salt solution, placed without liquid in a Stratalinker (Stratagene), and irradiated with 20 or 40 J/m² as indicated.

Expression constructs and antibodies

p33^{ING1b} wild-type and p47^{ING1a} cDNA were subcloned into pCS2-myc vector and used throughout the study. *ING1b* deletion constructs were generated by PCR. The W235A point mutation was introduced using the QuikChange II site-directed mutagenesis kit (Stratagene). The corresponding primers are listed in the Supplemental Material. *Xenopus* Flag-*Gadd45a*-encoding and human HA-*Gadd45a*-encoding expression plasmids have been described (Barreto et al. 2007). Antibodies used were mouse α - α -tubulin (Sigma); mouse α -Flag M2 and rabbit (Western blot and ChIP) or mouse (immunoprecipitation) α -Gadd45a and goat α -ING1 (all from Santa Cruz Biotechnology); rabbit α -H3, α -H3K4me2, α -H3K4me3, mouse α -myc, rat α -HA, and IgG control (all from Abcam); or rabbit α -H3K4me1 (Upstate Biotechnology).

Dual-luciferase reporter assay

Dual-luciferase reporter assays (Promega) were performed as described (Barreto et al. 2007). In brief, HEK293T cells were transfected with 20 ng of M.SssI in vitro methylated reporter and 5 ng of effector, and for Figure 1B with 1, 5, or 10 ng of *ING1b* in a 96-well plate format. Results are shown as the mean of triplicates $\pm$ standard deviation (SD).

RT-qPCR

Total RNA was isolated using a Qiagen RNeasy minikit with on-column DNase digest (Qiagen). First strand cDNA was generated using SuperScriptII reverse transcriptase (Invitrogen). Real-time PCR was performed in technical duplicates using Roche LightCycler480 probes master and primers in combination with predesigned monocolour hydrolysis probes of the Roche Universal Probe Library (UPL) (see the Supplemental Material). For quantification, Roche LC480 quantification software module was used. All values were normalized to the level of a housekeeping gene: *GAPDH* for human cells, or *G6pd* for MEF cells. Mean values shown were calculated using the mean value of the qPCR replicates from independent experiments ($n \geq 3$).

Microarray expression analysis

HEK293T cells were transiently transfected in duplicates as described above and harvested 48 h after transfection. Male MEF cells were analyzed in duplicates derived from one MEF line per genotype. Total RNA was isolated using the RNeasy minikit with on-column DNase digest (Qiagen).

Microarray sample processing, hybridization, and scanning were performed on HumanHT-12 v4 and MouseWG-6 v2 expression BeadChips (Illumina) following the manufacturer's instructions. Data were processed, quality-controlled, and filtered with the GenomeStudio version 2011.1 software (Illumina) using the following parameters: background subtraction, cubic spline normalization, and Illumina custom error model with false discovery rate (FDR) multiple testing correction. Scored as differentially expressed were genes with FDR-adjusted differential *P*-values <0.1 and >1.5 (for the MEF experiment) or >2.0 (for the HEK293T experiment) thresholds for expression fold changes, which were calculated after flooring the normalized array signal intensities to 15 (equivalent to probe detection *P*-value of 0.1) in order to minimize the effect of random fluctuations in background noise. For the calculation of the number of differentially expressed genes in each sample, the most recent available manifest annotation files from Illumina were used, and some unassigned mouse array probes were further annotated using information from the MGI database (<http://www.informatics.jax.org/batch>). Downstream analysis of the filtered gene lists was carried out using DAVID (<http://david.abcc.ncifcrf.gov>), ToppGene (<http://toppgene.cchmc.org>), BioVenn (<http://www.cmbi.ru.nl/cdd/biovenn>), and Venny (<http://bioinfopg.cnb.csic.es/tools/venny>). The microarray data have been deposited in the NCBI's Gene Expression Omnibus (GEO) under accession number GSE43319 (<http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE43319>). For qPCR validation, the original samples were used as well as at least two additional experiments for MEF cells using two additional male lines per genotype.

Analysis of DNA methylation

For bisulfite sequencing, the regions of interest were PCR-amplified from bisulfite-converted DNA (EpiTect kit, Qiagen) using Accuprime Taq polymerase (Invitrogen). For plasmid DNA

bisulfite sequencing, the M.SssI- (Fig. 1C, *pGL3-Luc*) or HpaII (Fig. 1G, *Oct4TK-GFP*) in vitro methylated plasmid was recovered as described (Barreto et al. 2007) and linearized prior to bisulfite conversion (EpiTect kit, Qiagen).

Bisulfite sequencing of *MAGEB2* was performed as described (Mund et al. 2005). The PCR products were purified by agarose gel electrophoresis, subcloned into TA-cloning vector pCR2.1 (Invitrogen), and sequenced. Bisulfite treatment efficiency and specificity were controlled using an unmethylated or in vitro methylated plasmid carrying the *MAGEB2* promoter amplicon (data not shown).

For Figures 1C; 5, D and E; and 7, C and D, the PCR amplicons were ligated to Roche Multiplex Identifier (MID) adaptors, mixed, and analyzed using the Roche 454 GS Junior system. Only amplicons with >100 reads per sample were considered reliable.

For *Oct4TK-GFP* 454 sequencing, the primers were fused to Roche Titanium Adaptor sequences for amplification. The four samples of Figure 1G were analyzed in four lanes of the Roche GS FLX system and multiplexed for the three time points using Roche multiplex identifiers.

For methylation-sensitive PCR (MS-PCR), genomic DNA was digested with PvuII (control) or MS restriction enzyme. MS enzymes were HpaII for mouse *Mageb2* site b, HhaI for mouse *Mageb2* sites a and a' (see scheme in Supplemental Fig. S4B), and HpyCH4IV for mouse *Mageb4*. Methylation in percent was determined by qPCR from the ratio of digested versus control digested DNA for the respective CpGs. In ChIP-Chop assays, immunoprecipitated chromatin was subjected to the same procedure. The control promoter mouse *Gapdh* contained both a HpaII and a HhaI restriction site and was always unmethylated. *Gapdh* methylation based on HhaI restriction is shown in Figures 3, D–F, and 4B. All methylation analysis primers are listed in the Supplemental Material.

Co-IP and ChIP

For co-IP, HEK293T cells were lysed in 150 mM NaCl, 20 mM Tris (pH 7.5), 2 mM EDTA, 10% glycerol, 1% Triton X-100, and 1 mM β -mercaptoethanol containing Complete Mini Protease Inhibitor Cocktail (Roche). The lysates were syringe and needle-homogenized and cleared by centrifugation and by incubation with mixed protein A/G agarose beads (50% each; Roche) for 30 min at 4°C. Lysates were directly used for immunoprecipitation without the freezing step ($\sim 4 \times 10^6$ cells per immunoprecipitation). After antibody incubation overnight at 4°C on a rotating wheel, protein A/G agarose beads were added for 1 h, 30 min. Beads were washed four times using the immunoprecipitation lysis buffer and eluted with 4 $\times$ Laemmli buffer containing 10% β -mercaptoethanol. For Western blot analysis, one-third of the eluate was loaded along with 5% of the input lysate.

RKO ChIP was performed essentially as described (Wells et al. 2002) using a mixture of protein A/G agarose beads (Roche). For re-ChIP, chromatin was eluted with 50 mM Tris (pH 8.0), 1% SDS, and 10 mM EDTA. Ninety percent of the eluate was subjected to a second immunoprecipitation after a 20-fold dilution in ChIP dilution buffer (1.1% Triton, 1.2 mM EDTA, 16.7 mM Tris-Cl at pH 8.2, 167 mM NaCl).

Chromatin isolation and ChIP of HEK293T and MEF cells was also performed as described (Wells et al. 2002) with some modifications. For immunoprecipitation, a mixture of protein A/G Dynabeads (Invitrogen) was incubated with 20 μ g of chromatin (tagged proteins) or 120 μ g of chromatin (endogenous proteins, freshly used without the freezing step). After incubation with the chromatin, the beads were washed five times using the above-described co-IP buffer. After elution with 50

mM Tris (pH 8.0), 1% SDS, and 10 mM EDTA overnight at 65°C, proteinase K digestion for 2 h at 56°C, and purification via QIAquick PCR purification columns (Qiagen), samples were used for qPCR.

Mean values shown in the graphs were calculated using the mean value of the qPCR replicates from independent experiments ($n \geq 3$).

In vitro pull-down assay

For histone peptide-binding assays, biotinylated histone peptides were used (Upstate Biotechnology). Briefly, 0.2 μ g of peptide was incubated with 15 μ L of streptavidin Dynabeads (Dyna) in binding buffer (50 mM Tris-HCl at pH 8.0, 150 mM NaCl, 0.2% Triton X-100, 1 mM dithiothreitol [DTT], EDTA-free Protease inhibitor cocktail [Roche]) for 45 min at 4°C. After two washes, beads were added to 100 μ g of nuclear protein extracts expressing the indicated tagged proteins. After 3 h of incubation at 4°C in binding buffer, beads were washed five times using wash buffer (20 mM HEPES at pH 7.9, 250 mM KCl, 0.2% Triton X-100, 1 mM DTT, protease inhibitors) and subjected to Western blot analysis.

Statistical analyses

Data are expressed as mean values $\pm$ SD of independent biological replicates ($n \geq 3$). Two-tailed, unpaired Student's *t*-test was used to calculate the level of significance. A *P*-value of <0.05 was considered significant. (*) $P < 0.05$; (**) $P < 0.01$; (***) $P < 0.001$.

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