

SHORT REPORT

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Specific inhibition of one DNMT1-including complex influences tumor initiation and progression

Mathilde Cheray^{1,2}, Romain Pacaud^{1,2}, Arulraj Nadaradjane^{1,2}, François M Vallette^{1,2,3} and Pierre-François Cartron^{1,2,4*}

Abstract

Background: Reactivation of silenced tumor suppressor genes by DNMT inhibitors has provided an alternative approach to cancer therapy. However, DNMT inhibitors have also been shown to induce or enhance tumorigenesis via DNA hypomethylation-induced oncogene activation and chromosomal instability. To develop more specific DNMT inhibitors for efficient cancer therapy, we compared the effects of peptides designed to specifically disrupt the interaction of DNMT1 with different proteins.

Findings: Our data indicated that the use of an unspecific DNMT inhibitor (5aza-2deoxycytidine), a DNMT1 inhibitor (procainamide) or peptides disrupting the DNMT1/PCNA, DNMT1/EZH2, DNMT1/HDAC1, DNMT1/DNMT3b and DNMT1/HP1 interactions promoted or enhanced *in vivo* tumorigenesis in a mouse glioma model. In contrast, a peptide disrupting the DNMT1/DMAP1 interaction, which per se did not affect tumor growth, sensitized cancer cells to chemotherapy/irradiation-induced cell death. Finally, our data indicated that the peptide disrupting the DNMT1/DMAP1 interaction increased the efficiency of temozolomide treatment.

Conclusion: Our data suggest that the DNMT1/DMAP1 interaction could be an effective anti-cancer target and opens a new avenue for the development of new strategies to design DNMT inhibitors.

Keywords: DNMT1, Epigenetic, DNA methylation, Glioma, GBM, Cell death

Findings

Numerous reports have identified the occurrence of global DNA hypomethylation as an oncogenic event in human tumorigenesis. Indeed, DNMT1 silencing and the disruption of the DNMT1/PCNA/UHRF1 complexes were described as two events that promote the initiation of tumorigenesis (malignant transformation from a non-tumor cell to tumor cell) via the induction of global DNA hypomethylation [1-4]. However, besides PCNA and UHRF1 proteins, DNMT1 has multiple partners of interaction, the loss of which could potentially promote the

initiation of tumorigenesis or progression via the generation of the global DNA hypomethylation phenotype.

Disruption of certain DNMT1/protein-x interactions

Among the DNMT partners, we focused our analysis on the impact of tumorigenesis on the disruption of the DNMT1/PCNA, DNMT1/HDAC1, DNMT1/HP1 β , DNMT1/EZH2, DNMT1/Sp1, DNMT1/DMAP1 and DNMT1/DNMT3b complexes (Figure 1A) [5-14].

To investigate the impact of the disruption of these interactions on tumorigenesis, we transfected an astrocyte cell line (Astro#40) and a glioma cell line (U87) with plasmid constructions encoding for peptides mimicking certain amino-acid regions implicated in the interactions existing between DNMT1 and DMAP1, DNMT3b, PCNA, EZH2, HDAC1, Sp1 and HP1 β (Figure 1A and Additional file 1: Figure S1).

* Correspondence: pierre-francois.cartron@univ-nantes.fr

¹Centre de Recherche en Cancérologie Nantes-Angers, INSERM, U892, Equipe Apoptose et progression tumorale, Equipe labellisée Ligue Nationale Contre le Cancer, 8 quai moncoussu, BP7021, 44007, Nantes, France

²Département de Recherche en Cancérologie, Université de Nantes, Faculté de Médecine, IFR26, F-4400, Nantes, France

Full list of author information is available at the end of the article

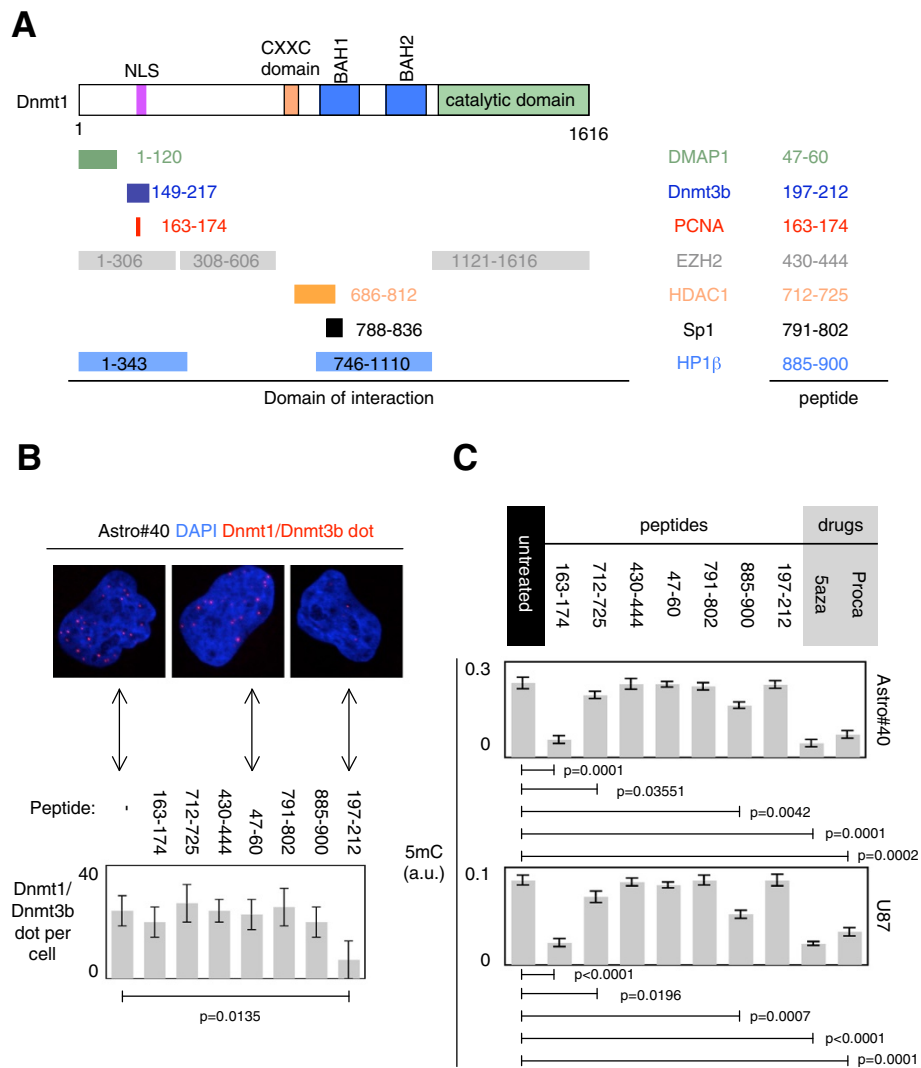


Figure 1 Specific disruption of DNMT1/protein-x interactions. (A) Representation of DNMT1 with its functional domain (nuclear localization signal: NLS, BAH1 and two domains, domain CXXC and its catalytic domain) and with amino acid domains of interaction with DMAP1, HDAC1, Sp1, HP1β, EZH2, DNMT3b and PCNA according to the UniProt website and the literature. Amino acid sequences of peptides cloned in plasmid and able to disrupt specific DNMT1/protein-x interactions are indicated in the "peptide" column; "untreated" indicates that cells were not treated with a peptide encoding by a plasmid, i.e., that cells were only treated with a plasmid encoding for the NLS of interest. (B) Only the expression of the 197–212 peptide promotes a significant decrease of the DNMT1/DNMT3b interaction in Astro#40 cells. Pictures (blue: DAPI staining; red: DNMT1/DNMT3b interaction or close proximity) illustrate the specific disruption of the DNMT1/DNMT3b interaction or close proximity induced by the expression of the 197–212 peptide, while the expression of the 47–60 peptide did not affect the DNMT1/DNMT3b interaction or close proximity. Pictures are representatives of data obtained from 100 cells in three independent experiments. The graph illustrates the average ± SD of these data. (C) Measure of the 5-methylcytosine (5 mC) level in cells (Astro#40 or U87) transfected by plasmid encoding for the indicated peptides or pre-treated with procainamide (0.5 mmol/l each 5 days for 4 weeks) or 5aza-2deoxycytidine (1 μM each 5 days for 4 weeks).

After 4 weeks of transfection/selection/amplification of cells, proximity ligation in situ assays (P-LISA) revealed that the 47–60, 197–212, 163–174, 430–444, 712–725, 791–802 and 885–900 peptides were specific to disrupting the interactions between DNMT1 and DMAP1, DNMT3b, PCNA, EZH2, HDAC1, Sp1 and HP1β, respectively. For example, we observed in Astro#40

peptide disrupted the DNMT1/DNMT3B interaction, but not the DNMT1/HP1β, DNMT1/HDAC1, DNMT1/Sp1, DNMT1/EZH2 and DNMT1/DMAP1 interactions (Figure 1B). More generally, we noted that peptides designed to specifically disrupt an interaction disrupted only the targeted and expected interactions in Astro#40 and U87 cells (Additional file 2: Figure S2).

Impact of the disruption of DNMT1/protein-x interactions on the global DNA methylation level

We then investigated the impact of these disruptions on the global level of DNA methylation, i.e., on the 5-methylcytosine level. Two DNMT inhibitors (5aza-2deoxycytidine and procainamide) were used as these drugs induced global DNA hypomethylation (Figure 1C). In agreement with our previous reports, the 163–174 plasmid was used as a peptide that induced global DNA hypomethylation [3,4] (Figure 1C). Thus, we noted that the disruption of DNMT1/HDAC1 and DNMT1/HP1 β interactions promoted a global decrease in the 5-methylcytosine level in Astro#40 and U87 cells, while the disruption of the DNMT1/EZH2, DNMT1/Sp1 and DNMT1/DMP1 interactions did not affect the 5-methylcytosine level in these cells (Figure 1C).

Impact on tumorigenesis of the disruption of DNMT1/protein-x interactions

Tumorigenic assays performed in Swiss nude mice revealed that the subcutaneous injection of Astro#40 cells treated 4 weeks with 5aza-2deoxycytidine and procainamide or transfected with plasmid encoding for the 163–174, 712–725 and 885–900 peptides generated tumor formation in 3/5, 2/5, 5/5, 2/5 and 2/5 cases, respectively (Figure 2A). Thus, we observed that gliomagenesis is induced by the loss of DNMT1/PCNA, DNMT1/HDAC1 and DNMT1/HP1 β interactions, i.e., under conditions that also induced a decrease in the global DNA methylation level. In parallel, we noted that the subcutaneous injections of U87 cells treated with 5aza-2deoxycytidine and procainamide or transfected with plasmid encoding for the 163–174, 712–725, 430–444, 885–900 and 197–212 peptides generated the formation of larger tumors as compared to untreated U87 cells (Figure 2B). These data suggest that the disruption of DNMT1/PCNA, DNMT1/EZH2, DNMT1/DNMT3B, DNMT1/HDAC1 and DNMT1/HP1 β interactions acts such as an enhancer of tumorigenesis. Thus, our work stratifies the specific disruption of certain DNMT1/protein-x interactions into two categories (Figure 2C). The first category includes the specific disruption of certain DNMT1/protein-x interactions acting as inducers and/or enhancers of gliomagenesis (DNMT1/PCNA, DNMT1/HDAC1 and DNMT1/HP1 β , DNMT1/EZH2, DNMT1/DNMT3b), i.e., acting as pro-tumorigenic-specific disruption of DNMT1/protein-x interactions. The second category includes the specific disruption of DNMT1/protein-x interactions devoid of any action on gliomagenesis. This category is called neutral-tumorigenic-specific disruption of DNMT1/protein-x interactions. Although absent in our data, we added a third category called anti-tumorigenic-specific disruption of DNMT1/protein-x interactions including the specific

disruption of DNMT1/protein-x interactions promoting the decrease in tumorigenesis (Figure 2C).

Focus on two disruptions of DNMT1/protein-x interactions devoid of action on gliomagenesis

In the absence of any observations of anti-tumorigenic-specific disruptions of DNMT1/protein-x interactions, we next analyzed whether the specific disruption of a DNMT1/protein-x interaction devoid of action on gliomagenesis could have an impact on the response to the standard GBM treatment combining temozolomide (TMZ) and irradiation. For this, we measured the percentage of cell death induced by a TMZ + irradiation treatment of U87 cells transfected with the plasmids encoding for the 47–60 and 791–802 peptides (Additional file 3: Figure S3). We noted that the 47–60 peptide, which induced the disruption of the DNMT1/DMP1 interaction, increased the percentage of TMZ + irradiation-induced cell death, while the 791–802 peptide, which induced the disruption of the DNMT1/Sp1 interaction, had no effect on the percentage of TMZ + irradiation-induced cell death (Figure 2D). In addition, our data also indicated that the 5aza-2deoxycytidine and procainamide treatments and the 163–174 peptide induced a phenotype resistant to TMZ + irradiation cell death since the percentage of TMZ-irradiation-induced cell death significantly decreased under these conditions (Figure 2D). Taken together, these data indicate that, among the considered peptides and DNMT1 inhibitors, only the specific 47-60-induced disruption of the DNMT1/DMP1 interaction increased the percentage of TMZ + irradiation-induced cell death without promoting or increasing the tumorigenesis. Thus, we conclude that the specific inhibition of the DNMT1/DMP1 interaction is a neutral-tumorigenic-specific inhibition of DNMT1/protein-x interaction harboring the capacity to sensitize cells to TMZ + irradiation-induced cell death.

Disruption of DNMT1/DMP1 interactions enhances the anti-tumor effect of TMZ treatment in mice

To investigate this point, established tumors were treated with TMZ, TMZ + 47-60^{DNMT1} and mutated 47-60^{DNMT1} (m47-60^{DNMT1}) (Additional file 4: Figure S4). Mice received 6 weeks of treatment as in treatment of human GBM, and tumor weight was analyzed 2 weeks after the end of the treatment. As illustrated in Figure 2E, we first noted that TMZ treatment reduced the tumor weight of U87-induced glioma. Second, we observed that the addition of 47-60^{DNMT1} increased the efficiency of TMZ treatment and decreased tumor growth, while m47-60^{DNMT1} had no effect on the efficiency of TMZ treatment (since this peptide does not disrupt the DNMT1/DMP1 interaction; Additional file 5: Figure S5). Thus,

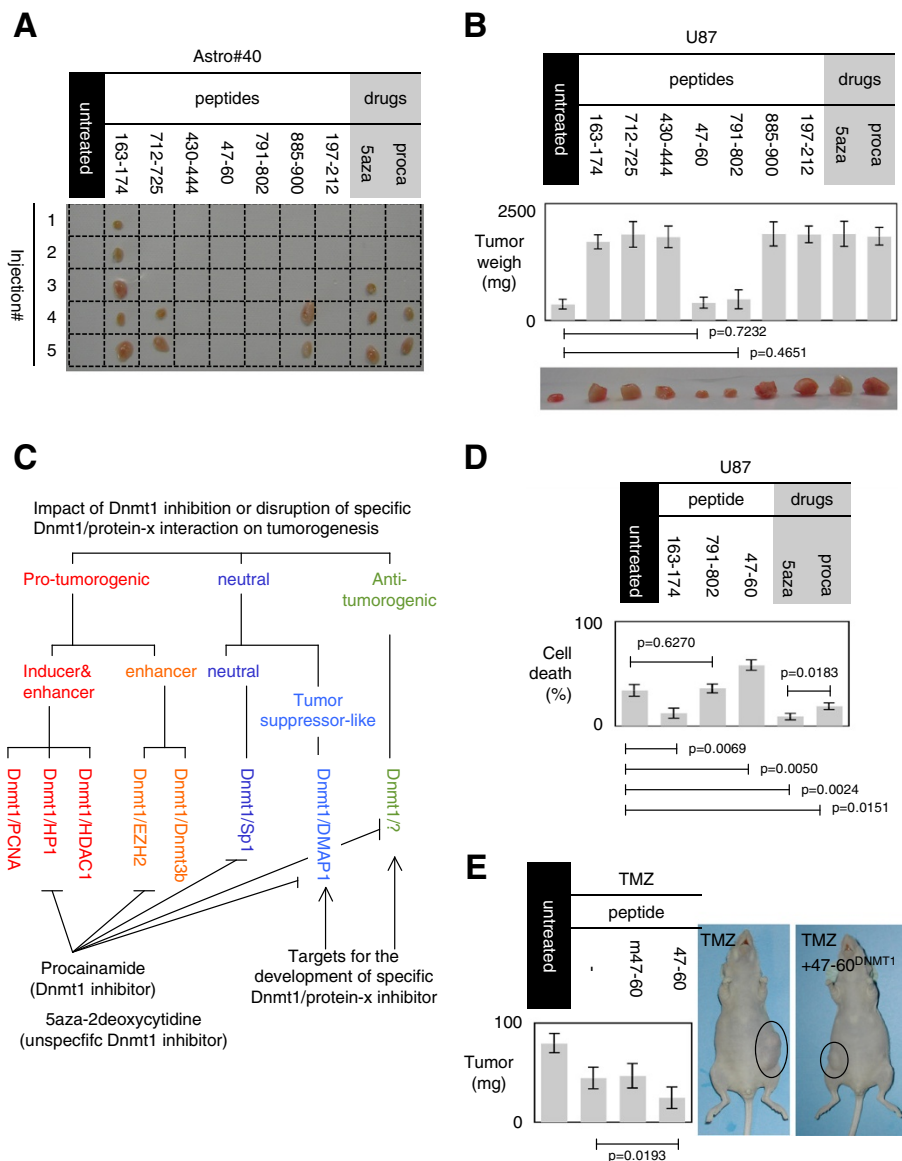


Figure 2 Impact of specific inhibition of DNMT1/protein-x interaction. (A) Tumorigenicity test of Astro#40 cells transfected with plasmids encoding for indicated peptides or pre-treated with procainamide. For each condition, five subcutaneous injections of 10^6 cells were performed in Swiss nude mice. Picture illustrates the tumor obtained (or not) 5 weeks after injections. (B) Tumorigenicity test of U87 cells transfected with plasmids encoding for indicated peptides or pre-treated with epigenetic drugs. For each condition, three subcutaneous injections of 10^6 cells were performed in Swiss nude mice. Graph illustrates the average \pm SD of these data. Picture illustrates representative tumors obtained 4 weeks after injections. (C) Effect of different DNMT inhibitor strategies. DNA methylation governs the expression of tumor suppressor genes (TSG) but also of oncogenes and DNA repeat element and retrotransposons (whose demethylation and/or expression promotes the tumorigenesis). By using an unspecific DNMT inhibitors (5aza-2deoxycytidine) or a specific DNMT1 inhibitor (procainamide), we postulate that we induced the hypomethylation not only of TSG but also of the oncogenes and retrotransposons. By reducing the effect of DNMT1 inhibition to certain specific DNMT/protein-x interactions, we still keep the opportunity to promote the expression of the TSG without promoting the hypomethylation-induced activation of retrotransposons and oncogenes. (D) Analysis of TMZ + irradiation-induced cell death in U87 cells. Percentages of cell death were evaluated by using a trypan blue stain 0.4%, and the Countess[®] Automated Cell Counter (Life Technology, France). Graph illustrates the average \pm SD of three independent experiments. (E) Impact of the addition of 47-60^{DNMT1} on TMZ treatment in an *in vivo* model of glioma. After the tumor establishment, mice were treated with TMZ and TMZ + 47-60^{DNMT1} (Additional file 4: Figure S4). Two negative controls were used: "untreated" represents a treatment with DMSO "m47-60^{DNMT1}" symbolized a co-treatment using TMZ and mutated 47-60^{DNMT1}. Graph illustrates the average \pm SD of five independent experiments.

our data identify 47-60^{DNMT1} as an enhancer of TMZ treatment.

Discussion

To summarize, our data indicate that the specific inhibition of certain DNMT1/protein-x interactions (DNMT1/PCNA interaction, for example) and the use of a specific DNMT1 inhibitor (procainamide) or unspecific DNMT1 inhibitor (5-aza-2deoxycytidine) could be used as a treatment, acting as inducers and/or enhancers of tumorigenesis. Our data indicate that the specific inhibition of the DNMT1/DNAH1 interaction acts as a tumor suppressor-like event since the disruption of the DNMT1/DNAH1 interaction increased TMZ + irradiation-induced cell death without promoting the initiation and progression of tumorigenesis. Consequently, we distinguish between among the neutral-tumorigenic-specific inhibition of DNMT1/protein-x interaction and the tumor suppressor-like neutral-tumorigenic-specific inhibition of DNMT1/protein-x interaction (Figure 2C).

Using this example, our data underline the necessity to consider the interaction partners of DNMT1 and not only the DNMT1 structure or activity to develop a DNMT1 inhibitor. In other terms, our data introduce, for the first time, the notion of protein/protein inhibition into the development of DNMT inhibitors. Indeed, without being innovative in the development of drugs or small molecules for a therapeutic application, this strategy is novel in the conception/research of DNMT inhibitors since the identification of DNMT inhibitors is, to date, based on docking-based virtual screening methods, the screening of natural products, the design and generation of derivatives of DNMT inhibitors already known, or molecular modeling of DNMT inhibitors by using crystal structure studies of DNMTs [15-19]. Thus, like ABT-737 and MI-129, two compounds targeting specific protein-protein interactions (pro-apoptotic/anti-apoptotic and p53/MDM2 interactions, respectively) have opened a new area in targeted therapy; our data argue that the targeting of certain DNMT1/protein-x interactions opens a new area in the development of targeted epigenetic therapy.

However, despite its promising character, the development of specific inhibitors of DNMT1/protein-x interactions requires the identification of tumor suppressor-like neutral-tumorigenic-specific inhibition of DNMT1/protein-x interaction or anti-tumorigenic-specific disruption of certain DNMT1/protein-x interactions. Thus, studies are ongoing in our laboratory to identify a DNMT1/protein-x-including complex promoting the methylation-induced silencing of the tumor suppressor gene without being implicated in the methylation-induced silencing of oncogenes.

Concerning the use of 5aza-2deoxycytidine and procainamide, we are aware that the tumorigenesis processes associated with the use of these drugs are obtained after a long exposure. However, 4 weeks is not a long period on the scale of the majority of the chemotherapeutic treatments that last several months. Regardless, our data provide "a warning" concerning the use of DNA demethylating agents as an anti-cancer therapy and provide proof reinforcing the necessity of using specific DNMT inhibitors or unspecific DNMT1 inhibitors with an adequate and optimized dose schedule, as already described by several publications [20,21]. Besides, several publications also argue that the use of DNMT inhibitors could promote oncogene activation [22,23]. Thus, Chik and Szyf [22] report that 5aza-2deoxycytidine activated both silenced tumor suppressor genes and pro-metastatic genes by demethylation, raising the concern that it could promote metastasis [22]. Associated with our results, these data support the idea of developing specific DNMT1, particularly developing a specific inhibitor of DNMT1/protein-x interaction.

Methods

Proximity ligation in situ assay (P-LISA)

P-LISA is a technology permitting the visualization of stable and transient interactions at endogenous protein levels directly *in situ* [20]. Briefly, two primary antibodies raised in different species recognize the target antigen or antigens of interest. Species-specific secondary antibodies, called PLA probes, each with a unique short DNA strand attached to it, bind to the primary antibodies. When the PLA probes are in close proximity (<40 nm), the DNA strands can interact through a subsequent addition of two other circle-forming DNA oligonucleotides. After ligating the two added oligonucleotides, creating a circle DNA molecule, they are amplified via rolling circle amplification. After amplification, several-hundredfold replication of the DNA circle has occurred, and labeled complementary oligonucleotide probes highlight the product. The resulting high concentration of fluorescence in each single-molecule amplification product is easily visible as a distinct bright dot when viewed with a fluorescence microscope.

Cells were cultured for 24 h on cover slips. Cells were then fixed with 4% paraformaldehyde in PBS, pH 7.4, for 15 min at room temperature. Permeabilization was performed with PBS containing 0.5% Triton X-100 for 20 min at room temperature. Blocking, staining, hybridization, ligation, amplification and detection steps were realized according to the manufacturer's instructions (Olink Bioscience). All incubations were performed in a humidity chamber. Amplification and detection steps were performed in a dark room. Fluorescence was visualized by using the Axiovert 200M microscopy system (Zeiss,

Le Pecq, France) with the ApoTome module (X63 and numerical aperture 1.4). Preparations were mounted using ProLong[®] Gold antifade reagent with DAPI (Invitrogen, France). Picture acquisition was realized in structured illumination microscopy [24]. Finally, the images were analyzed using the freeware “BlobFinder” available for download from www.cb.uu.se/~amin/BlobFinder. Thus, we obtained the number of signals per nuclei since nuclei can be automatically identified.

Plasmid construction and transfection

To express peptides in cells, we subcloned in pcDNA3.3 (Life Technology, France) the sequences encoding for the indicated peptides. In addition, the NLS sequence (PKKKRKV) was added to the sequences encoding for the indicated peptides in order to address the peptides in the nucleus. Transfections were next realized by using 2.10⁵ cells, 5 µg of plasmid and Lipofectamine[™] 2000 reagents (Life Technology, France). Selection was realized by adding 500 µg/ml of Geneticin selective antibiotic in complete medium of cell culture for 3 weeks. Next, 1 week was used to amplify the cells. Veracity of transfection was determined by PCR analyses using primers directed against pcDNA3.3 (ACGTTGTCCTGAAGCGG and CCTGATGCTCTTCGTCCA) and by the fact that each plasmid affects the DNMT1/protein-x interaction of interest.

Measure of the 5-methylcytosine level

DNA was extracted using the QiaAmp DNA mini Kit (Qiagen, France). The quantification of 5-methylcytosine is performed by using the Methylamp Global DNA methylation Quantification kit (Euromedex-Epigenetiek, France).

Tumorigenicity assay

Cultured cells were harvested by trypsinization, washed and resuspended in saline buffer. Cell suspensions were injected s.c. as 10⁶ cells in 0.2 ml volume in the flank of 7-/8-week-old nude NMRI-nu female mice (Janvier, France).

Additional files

Additional file 1: Figure S1. Representation of the pcDNA3.3 plasmid used to express the indicated peptides in cells. For each peptide, the amino acid position, sequences and corresponding cDNA sequences are indicated in the table.

Additional file 2: Figure S2. Graphs illustrate the impact of the expression of the considered peptide on the indicated DNMT1/protein-x interaction of close proximity. Graph illustrates the average ± SD obtained from 100 cells in three independent experiments. *Only the corresponding interaction was significantly decreased ($p < 0.05$, t -test) by the considered peptides in comparison with the data obtained from untreated cells. “Untreated” indicates that cells are not transfected by a plasmid encoding for a peptide or are not treated with a DNMT inhibitor.

Thus, this condition is used as a control. For each peptide, the specificity of inhibition of disruption of one considered peptide is reinforced by the use of six other peptides.

Additional file 3: Figure S3. Schematic representation of the TMZ + irradiation treatment administered to the cells.

Additional file 4: Figure S4. Schematic representation of the treatments administered to mice.

Additional file 5: Figure S5. Effect of 47-60^{DNMT1} and m47-60^{DNMT1} on the DNMT1/DMAP1 interaction. Graphs illustrate the impact of the considered peptide on the indicated DNMT1/protein-x interaction of close proximity. Graph illustrates the average ± SD obtained from 100 cells in three independent experiments. *Only the corresponding interaction was significantly decreased ($p < 0.05$, t -test) by the considered peptides in comparison with the data obtained from untreated cells.

Competing interests

The authors declare that they have no financial relationship with the organization that sponsored the research.

Authors' contributions

PFC initiated and conceived the study, undertook and interpreted analyses, and drafted the manuscript. FMV provided data interpretation and manuscript review. PFC, MC, RP and AN performed epigenetic laboratory studies. All authors read and approved the final manuscript.

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Author details

¹Centre de Recherche en Cancérologie Nantes-Angers, INSERM, U892, Equipe Apoptose et progression tumorale, Equipe labellisée Ligue Nationale Contre le Cancer, 8 quai moncoussu, BP7021, 44007, Nantes, France. ²Département de Recherche en Cancérologie, Université de Nantes, Faculté de Médecine, IFR26, F-4400, Nantes, France. ³LaBCT, Institut de Cancérologie de l'Ouest, Boulevard J Monod, 44805, Nantes, Saint Herblain Cedex, France. ⁴Institut de Recherche Thérapeutique INSERM U892 – CRCNA, Equipe 9 – Apoptose et Progression tumorale, 8 Quai Moncoussu, BP 70721, 44007, Nantes, Cedex 1, France.

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References

- Eden A, Gaudet F, Waghmare A, Jaenisch R: **Chromosomal instability and tumors promoted by DNA hypomethylation.** *Science* 2003, **300**(5618):455.
- Gaudet F, Hodgson JG, Eden A, Jackson-Grusby L, Dausman J, et al: **Induction of tumors in mice by genomic hypomethylation.** *Science* 2003, **300**(5618):489–492.
- Hervouet E, Lallier L, Debien E, Cheray M, Geairon A, et al: **Tumor induction by disruption of the Dnmt1, PCNA and UHRF1 interactions.** *Nature Precedings* 2008. <http://hdl.handle.net/10101/npre.2008.2509.1>.
- Hervouet E, Debien E, Cheray M, Hulín P, Loussouarn D, et al: **Disruption of Dnmt1/PCNA/UHRF1 interactions promotes tumorigenesis by inducing genome and gene-specific hypomethylations and chromosomal instability.** *PLoS One* 2010, **5**(6):e11333.
- Bostick M, Kim J, Estève P, Clark A, Pradhan S, et al: **UHRF1 plays a role in maintaining DNA methylation in mammalian cells.** *Science* 2007, **27**(15): 2187–2197.
- Estève P, Chin H, Pradhan S: **Human maintenance DNA (cytosine-5)-methyltransferase and p53 modulate expression of p53-repressed promoters.** *Proc Natl Acad Sci U S A* 2005, **102**(4):1000–1005.
- Estève P, Chin H, Pradhan S: **Molecular mechanisms of transactivation and doxorubicin-mediated repression of survivin gene in cancer cells.** *J Biol Chem* 2007, **282**(4):2616–2625.

8. Fuks F, Hurd P, Deplus R, Kouzarides T: **The DNA methyltransferases associate with HP1 and the SUV39H1 histone methyltransferase.** *Nucleic Acids Res* 2003, **31**(9):2305–2312.
9. Fuks F, Burgers WA, Brehm A, Hughes-Davies L, Kouzarides T: **DNA methyltransferase Dnmt1 associates with histone deacetylase activity.** *Nat Genet* 2000, **24**(1):88–91.
10. Hervouet E, Vallette FM, Cartron PF: **Dnmt1/transcription factor interactions: an alternative mechanism of DNA methylation inheritance.** *Genes & Cancer* 2010, **1**(5):434–443.
11. Margot JB, Ehrenhofer-Murray AE, Leonhardt H: **Interactions within the mammalian DNA methyltransferase family.** *BMC Mol Biol* 2003, **4**(1):7.
12. Muromoto R, Sugiyama K, Takachi A, Imoto S, Sato N, et al: **Physical and functional interactions between Daxx and DNA methyltransferase 1-associated protein, DMAP1.** *J Immunol* 2004, **172**(5):2985–2993.
13. Sharif J, Muto M, Takebayashi S, Suetake I, Iwamatsu A, et al: **The SRA protein Np95 mediates epigenetic inheritance by recruiting Dnmt1 to methylated DNA.** *Nature* 2007, **450**(7171):908–912.
14. Viré E, Brenner C, Deplus R, Blanchon L, Fraga M, et al: **The Polycomb group protein EZH2 directly controls DNA methylation.** *Nature* 2006, **439**(7078):871–874.
15. Kuck D, Singh N, Lyko F, Medina-Franco J: **Novel and selective DNA methyltransferase inhibitors: docking-based virtual screening and experimental evaluation.** *Bioorg Med Chem* 2010, **18**(2):822–829.
16. Medina-Franco J, López-Vallejo F, Kuck D, Lyko F: **Natural products as DNA methyltransferase inhibitors: a computer-aided discovery approach.** *Mol Divers* 2011, **15**(2):293–304.
17. Suzuki T, Tanaka R, Hamada S, Nakagawa H, Miyata N: **Design, synthesis, inhibitory activity, and binding mode study of novel DNA methyltransferase 1 inhibitors.** *Bioorg Med Chem Lett* 2010, **20**(3):1124–1127.
18. Yoo J, Medina-Franco J: **Inhibitors of DNA methyltransferases: insights from computational studies.** *Curr Med Chem* 2012, **19**(21):3475–3487.
19. Yoo J, Kim J, Robertson K, Medina-Franco J: **Molecular modeling of inhibitors of human DNA methyltransferase with a crystal structure: discovery of a novel DNMT1 inhibitor.** *Adv Protein Chem Struct Biol* 2012, **87**:219–247.
20. Issa J: **Optimizing therapy with methylation inhibitors in myelodysplastic syndromes: dose, duration, and patient selection.** *Nat Clin Pract Oncol* 2005, **2**(1):S24–S29.
21. Momparler R, Côté S, Eliopoulos N: **Pharmacological approach for optimization of the dose schedule of 5-Aza-2'-deoxycytidine (Decitabine) for the therapy of leukemia.** *Leukemia Suppl* 2007, **1**:S1–S6.
22. Chik F, Szyf M: **Effects of specific DNMT gene depletion on cancer cell transformation and breast cancer cell invasion; toward selective DNMT inhibitors.** *Carcinogenesis* 2010, **32**(2):224–232.
23. Liu L, Ling X, Liang H, Gao Y, Yang H, et al: **Hypomethylation mediated by decreased DNMTs involves in the activation of proto-oncogene MPL in TK6 cells treated with hydroquinone.** *Toxicol Lett* 2012, **209**(3):239–245.
24. Schaefer L, Schuster D, Schaffer J: **Structured illumination microscopy: artefact analysis and reduction utilizing a parameter optimization approach.** *J Microsc* 2004, **216**(Pt 2):165–174.

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