Synergistic activity of everolimus and 5-aza-2’-deoxycytidine in medullary thyroid carcinoma cell lines

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Short running title: everolimus and AZA in medullary thyroid cancer

List of abbreviations:
AZA: 5-aza-2′-deoxycytidine
BDNF: brain derived neurotrophic factor
CI: combination index
DEGs: differentially expressed genes
DRI: dose reduction index
FDR: false discovery rate
MTC: medullary thyroid cancer
NGF: nerve growth factor
NT-3: neurotrophins-3
NT-4/5: neurotrophins-4/5
NT-6: neurotrophins-6
PARP: poly(ADP-ribose) polymerase
PF: potentiation factor
PI: propidium iodide
SAM: significance Analysis of Microarray
SEM: structural equation modeling
SPIA: Signaling Pathway Impact Analysis

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Abstract

Medullary thyroid cancer (MTC) is a tumor highly resistant to chemo- and radiotherapy. Drug resistance can be induced by epigenetic changes such as aberrant DNA methylation. To overcome drug resistance, we explored a promising approach based on the use of 5-aza-2'-deoxycytidine (AZA), a demethylating agent, in combination with the mTOR inhibitor everolimus in MTC cells (MZ-CRC-1 and TT). This combined treatment showed a strong synergistic antiproliferative activity through the induction of apoptosis. The effect of everolimus and/or AZA on genome-wide expression profiling was evaluated by Illumina BeadChip in MZ-CRC-1 cells. An innovative bioinformatic pipeline identified four potential molecular pathways implicated in the synergy between AZA and everolimus: PI3K-Akt signaling, the neurotrophin pathway, ECM-receptor interaction and focal adhesion. Among these, the neurotrophin signaling pathway was most directly involved in apoptosis, through the overexpression of NGFR and Bax genes. The increased expression of genes involved in the NGFR-MAPK10-TP53-Bax/Bcl2 pathway during incubation with AZA plus everolimus was validated by western blotting in MZ-CRC-1 cells. Interestingly, addition of a neutralizing anti-NGFR antibody abolished the synergistic cytotoxic activity between AZA and everolimus. These results open a new therapeutic scenario for MTC and potentially other neuroendocrine tumors, where therapy with mTOR inhibitors is currently approved.
**Key words:** Medullary thyroid cancer, 5-aza-2′-deoxycytidine, mTOR, everolimus, neurotrophin pathway, NGFR

1. Introduction

Medullary thyroid cancer (MTC) is a neuroendocrine tumor originating from parafollicular C cells and it results highly resistant to chemo- and radiotherapy (Vitale et al., 2001). The hyper-activation of the PI3K/Akt/mTOR cascade has a relevant role in the pathogenesis and progression of MTC. In fact, most of the pro-oncogenic effects of RET and RAS mutations are modulated by the activation of PI3K/Akt/mTOR pathway (Lyra et al., 2014; Manfredi et al., 2015). Few preliminary studies showed promising antitumor effects of mTOR inhibitors, such as everolimus, in MTC (Druce et al., 2012; Faggiano et al., 2012; Heilmann et al., 2016; Lim et al., 2013). A recent phase II study, assessing efficacy and tolerability of everolimus on tumor progression in patients with advanced thyroid cancer, showed stable disease in 5 out of 7 (71%) patients with MTC and a low toxicity profile for everolimus (Schneider et al., 2015).

Although everolimus has been reported to inhibit cell proliferation and angiogenesis in several human tumors, long-term treatment with mTOR inhibitors can be frustrated by the induction of resistance. Moreover, the anticancer effect of everolimus may be limited because of the intrinsic lack of inhibition on mTORC2 and the activation of survival pathways in cancer cells (Chan and Kulke, 2014). On this basis, anticancer treatment with everolimus as monotherapy may not be optimal, supporting the rationale for a combined treatment approach with other targeted agents.

Epigenetic alterations, including dysregulated protein acetylation and DNA methylation, affect gene expression, and contribute to tumorigenesis in several tumors (Hervouet et al., 2013; Walenkamp et al., 2014), including MTC (Vitale et al., 2016). PI3K/Akt/mTOR pathway is frequently deregulated in several malignancies through epigenetic alterations.
Epigenetic alterations appear to be also implicated in the development of resistance to mTOR inhibitors (Bihani et al., 2015; Juengel et al., 2012). In fact, histone deacetylase inhibitors are able to re-sensitize renal carcinoma cells to mTOR inhibitors (Bihani et al., 2015; Juengel et al., 2012; Juengel et al., 2014). However, most of the epigenetic mechanisms of resistance to mTOR inhibitors, particularly those concerning aberrant DNA methylation, remain poorly defined.

In the present manuscript we have explored a novel approach to potentiate the antitumor activity of everolimus, based on the use of 5-aza-2'-deoxycytidine (AZA), a well established demethylating agent, in MTC.

2. Materials and methods

2.1 Cell lines and reagents

Everolimus and AZA were kindly supplied by Novartis Pharma (Basel, Switzerland) and Sigma-Aldrich (St. Louis, USA), respectively. Both drugs were dissolved in DMSO and stored at -20°C. Human MTC cell lines (MZ-CRC-1 and TT) were provided from Prof. Lips (Utrecht, The Netherlands). Both cell lines were cultured at 37°C and 5% CO₂ in F-12 with Kaighn’s Modification medium supplemented with 10% foetal bovine serum, 2 nM glutamine and 10^5 U/l penicillin-streptomycin. HEK-293 (ATCC) human embryonic kidney cell line was cultured at 37°C and 5% CO₂ in DMEM high glucose medium, without sodium pyruvate, supplemented with 10% foetal bovine serum, 2 nM glutamine and 10^5 U/l penicillin-streptomycin.

2.2 Cell viability assessment

MTC cells were plated in 96-well plates (3x10⁴ cells per well) and treated with various concentrations of compounds (range 10⁻¹-10² nM for everolimus, 10⁻³-10⁵ nM for AZA). Cells growing in drug free medium and vehicle treated were used as control. After 3 days, medium
and drugs were replaced. After 6 days viability of cells was measured by the MTT assay (3-(4,5-dimethylthiazol-2-yl)-2.5-diphenyltetrazolium), as previously described (Vitale et al., 2012). Under the same conditions the in vitro toxicity of everolimus and AZA at synergistic concentrations was examined through MTT assay in HEK-293 cells.

2.3 Drug combination studies

MTC cells were seeded in 96-well plates with medium without and with everolimus and/or AZA at different concentrations. After 3 days, medium and compounds were refreshed. Cell viability was evaluated after 6 days of treatment using MTT assay. Three combinations were tested for each schedule: equiactive doses of the two agents (IC₅₀), higher relative doses of everolimus (IC₇₅ of everolimus/IC₂₅ of AZA) and higher relative doses of AZA (IC₂₅ of everolimus/IC₇₅ of AZA). Assessment of synergistic interaction between drugs was performed with CalcuSyn software (Biosoft, Ferguson, MO). Combination index (CI) values of <1, 1, and >1 are suggestive of synergy, additivity and antagonism, respectively (Chou et al., 1994; Chou and Talalay, 1984). We also evaluated: the dose reduction index (DRI), providing the magnitude of how much the dose of each drug in a synergistic combination may be reduced at a given effect level compared with the doses of each drug alone; the potentiation factor (PF), defined as the ratio of the IC₅₀ of either everolimus and AZA alone to the IC₅₀ of everolimus and AZA in combination.

2.4 Cell cycle analysis

Cells were plated in 6-well plates (3x10⁵ cells/well) in duplicate with medium without (control group) and with everolimus and/or AZA. After 3 days, medium and drugs were refreshed. After 6 days, cells were collected, stained with propidium iodide (PI) (Sigma-Aldrich, St. Louis, MO, USA), and analyzed using a FACScalibur flow cytometer (Becton Dickinson, Belgium) and CellQuest Pro Software, as previously described (Vitale et al., 2007).
2.5 Flow cytometric analysis of apoptosis

Cells were plated in 6-well plates and incubated with everolimus and/or AZA, as reported in the section of cell cycle analysis. After 6 days of treatment, cells were collected and stained with Annexin V-FITC (BD Pharmingen, San Diego, USA) and PI, and all samples were analyzed with FACScalibur, as previously reported (Vitale et al., 2006).

2.6 Western blot

MZ-CRC-1 cells were plated in 6-well plates and incubated with everolimus and/or AZA, as previously reported. After 6 days of treatment, cells were scraped, washed in PBS and resuspended in RIPA lysis buffer. Lysates were centrifuged, and protein analysis was conducted on supernatant. Cell extracts (50 μg/lane) were separated on NuPage 4-12% Bis-Tris Gels (Life Technologies) and transferred with iBlot System (Life technologies). Membranes were blocked with 5% milk TBS-T and incubated with specific antibodies overnight at 4 °C: anti-caspase-3 and PARP, anti-cleaved caspase and PARP, anti-Bcl2, anti-Bax, anti-mTOR, anti-phospho-mTOR (Ser2448), anti-4E-BP1, anti-phospho-4E-BP1 (Ser65), anti-MAPK10, anti-phospho-MAPK8/9/10 (Thr183/Tyr185), anti-phospho-p53 (Ser15) (Cell Signaling Technology, Beverly, MA) and anti-NGFR (Sigma-Aldrich, St. Louis, MO, USA). Blots were detected by the Luminata Forte ECL kit (Millipore) after incubation with HRP conjugated mouse and rabbit secondary antibodies (dilution 1:1,000 for anti-MAPK10, anti-phosphoMAPK8/9/10 and anti-phospho-p53; 1:5,000 for anti-caspase-3; and 1:10,000 for other antibodies) and then exposed to X-ray film. Bands of interest were quantified through ImageJ software version 1.47. Results were normalized against the level of actin expression in each sample and the intensities of the bands were expressed as arbitrary units when compared to those of the untreated cells.
2.7 Gene expression profiling

MZ-CRC-1 cells were seeded in 6-well plates and treated with everolimus and/or AZA, as previously reported. After 6 days, total RNA was extracted using Trizol Reagent (Life Technologies) and purified by RNeasy Mini Kit (Qiagen, Hilden, Germany). Amplification Grade DNase I (Life Technologies) was used to eliminate residual genomic DNA from RNA samples. Samples with an A260/A280 ratio falling in the range 1.8-2.1 were used for experiments.

The Illumina TotalPrep RNA Amplification kit (Ambion, Foster City, USA) was employed using 200 ng of total RNA as starting material. Labelled cRNA (750 ng) was hybridized to human HT-12 v3 BeadChip arrays (Illumina, San Diego, USA) according to the manufacture's recommendation. Fluorescent images were obtained with a BeadArray reader and processed with the BeadScan software.

2.8 Microarray gene expression analysis

Significance Analysis of Microarray (SAM) was employed to identify differentially expressed genes (DEGs) by comparing the expression value of each gene between every group of treatment vs the untreated control, as previously described (Tusher et al., 2001). All the delta values were selected in order to provide a false discovery rate (FDR) <0.05. Once determined the DEGs, the next step was to discover which biological pathways were associated to the list of DEGs through Signaling Pathway Impact Analysis (SPIA) (Tarca et al., 2009), an algorithm of third generation that takes in consideration not only the identity of the genes but also the topology of the pathway. These aspects were reported by the probability values: pPERT (reflecting the amount of perturbation measured in each pathway) and pNDE (probability of obtaining a number of DEGs on the given pathway at least as large as the observed one). These two types of evidence, pPERT and pNDE, were finally combined.
into one global probability value, pGFdr, which was used to rank the pathways and to test the research hypothesis that the pathway was significantly involved in the condition under the study. To refine the SPIA analysis, a downstream analysis based on the structural equation modeling (SEM) was performed (Pepe and Do, 2015; Pepe and Grassi, 2014). The method consists in the understanding how the DEGs, source of perturbation, propagated the perturbation in the biological network composed by the significant pathways. The module was obtained by the detection and fusion in a unique model of all directed shortest paths that put in communication the DEGs present in the significant pathways. This allowed: 1) to overcome the limits of the classical pathway analysis that considers the pathways as separate entities; 2) to detect the biological network where the drug modules are searched.

2.9 DNA preparation and Infinium Methylation 450K array

MZ-CRC-1 cells were plated in 6-well plates and treated with everolimus and/or AZA, as previously reported. After 6 days, total DNA was extracted using the DNeasy Blood & Tissue Kit (Qiagen, Germany). Quality control and quantification of DNA were performed before and after bisulphite conversion. DNA was quantified with NanoDrop (NanoDrop Products Thermo Scientific Wilmington, DE) and by fluorometric reading (Quant-iT™ PicoGreen® dsDNA Assay Kit); quality was assessed by visualisation of genomic DNA on 1% agarose gel electrophoresis. Only not fragmented DNA samples and with a concentration higher than 50 ng/μl were subsequently processed. The genomic DNA was treated with sodium bisulfite using the EZ DNA Methylation Kit ™ (Zymo Research); the technique requires only 500 ng of input DNA. Four microliters of bisulfite-converted DNA were used for hybridization on Infinium HumanMethylation 450K BeadChip, according to Illumina’s standard protocol. Data were acquired through Illumina HiScan SQ scanner. Image intensities were extracted using GenomeStudio software v2010.3. The methylation score for each CpG site was represented as β values according to the fluorescent intensity ratio between methylated and unmethylated DNA.
unmethylated probes. β values may range between 0 (completely unmethylated) and 1 (completely methylated). Illumina Methylation 450K raw data were analysed using the RnBeads analysis software package (Assenov et al., 2014). Sites overlapping SNPs were firstly removed from the analysis as well as probes on sex chromosomes. Probes and samples of highest impurity were removed from the dataset using the Greedycut algorithm. We have considered every β value to be unreliable when its corresponding detection p-value was not below the threshold (T=0.05). The background was subtracted using the methylumi package (method "noob"). The signal intensity values were normalized using the SWAN normalization method, as implemented in the minfi package (Aryee et al., 2014). The “Shapiro.test” function provided in the R package “stats” was applied to test normality among variables. The “kruskal.test” function provided in the R package “stats” was used to test differences among treated and untreated groups for all non-parametric data. DunnTest provided in the R package “dunn.test” has been applied as post-hoc test.

2.10 Statistical analysis

All experiments were performed at least 3 times. Statistical differences among groups were first evaluated by the ANOVA test, followed by post-hoc test (Newman-Keuls). A P value < 0.05 was considered significant. The values reported in the figures represent the mean ± standard error of the mean. For statistical analysis GraphPad Prism 5.0 was used.

3. Results

3.1 Pharmacological combination between everolimus and AZA on cell proliferation

In MZ-CRC-1 cells the combination of everolimus and AZA was highly synergistic when either the two drugs were used with lower concentrations of everolimus (IC\(_{25}\) everolimus:IC\(_{75}\) AZA) or with equitoxic concentrations (IC\(_{50}\) everolimus:IC\(_{50}\) AZA) (table 1, figure 1). The DRI\(_{50}\) (DRI calculated for 50% cell survival) was 8.8 for everolimus and 101 for AZA (when
drugs were used at IC\textsubscript{25} everolimus:IC\textsubscript{75} AZA), 4.3 for everolimus and 590 for AZA (at equitoxic concentrations) (table 1).

In TT cells a synergistic activity was similarly detected when lower concentrations of everolimus were adopted (IC\textsubscript{25} everolimus:IC\textsubscript{75} AZA), while antagonistic effects were observed with relative higher concentrations of everolimus (IC\textsubscript{75} everolimus:IC\textsubscript{25} AZA) (table 1, figure 1).

The optimal results (lowest CI values) were obtained when the two drugs were used at lower doses of everolimus in MZ-CRC-1 cells. Therefore, we have performed subsequent experiments using 2.1 nM (IC\textsubscript{25}) of everolimus and/or 1.4x10\textsuperscript{2} nM (IC\textsubscript{75}) of AZA in MZ-CRC-1 cells.

### 3.2 In vitro toxicity assessment

The effect of drug combination on the cell viability of HEK-293 cell line, derived from human embryonic kidney cells, was evaluated through MTT assay to roughly predict the toxicity profile of both everolimus and AZA in a non-cancer cell model. Combined treatment with everolimus and AZA, at concentrations showing synergistic antiproliferative activity in MZ-CRC-1 cells, inhibited viability of HEK-293 cells (-25%) to a lesser extent than that detected in MTC cells (-54%) (figure 2A). Moreover, we did not observe any change in morphology of HEK-293 cells after treatment with everolimus and AZA compared to untreated control (figure 2B).

### 3.3 Effects of everolimus and AZA on cell cycle.

AZA alone slightly, but significantly decreased cell population in G\textsubscript{0}/G\textsubscript{1} phase (-6% vs untreated control, p<0.05) and increased cell number in G\textsubscript{2}/M phase (+23% vs untreated control, p<0.01), suggesting a cell cycle arrest in G\textsubscript{2}/M phase (figure 3). Everolimus alone
significantly decreased cells percentage in both S (-24% vs untreated control, \(p<0.05\)) and G2/M phase (-15% vs untreated control, \(p<0.05\)). This effect was comparable to that observed during the synergistic combination of everolimus plus AZA (S phase: -27% vs untreated control, G2/M phase: -14% vs untreated control, both \(p<0.05\), figure 3).

3.4 Effects of everolimus and AZA on apoptosis

Everolimus did not change the percentages of early apoptotic and late apoptotic/necrotic cells compared with the untreated control. AZA induced a not statistically significant increase in the population of late apoptotic/necrotic cells. Interestingly, the AZA/everolimus combination significantly increased the percentage of early apoptotic (+35% vs control, \(p<0.05\)) and late apoptotic/necrotic (+90% vs control, \(p<0.001\)) cells (figure 4A).

Western blot analysis of key executioners of apoptosis, such as caspase-3 and poly(ADP-ribose) polymerase (PARP) further confirmed these data (figure 4 B, C). After 6 days of treatment, the AZA/everolimus combination significantly increased caspase-3 degradation, resulting in increased expression of cleaved caspase-3 and decreased expression of full caspase-3. A moderate increase in cleaved caspase-3 has been detected after AZA alone. Similarly, the activity of PARP was moderately stimulated after 6 days of treatment with AZA, and its combination with everolimus enhanced again this effect. No significant changes in caspase-3 activity and PARP cleavage were detected after the incubation of MZ-CRC-1 cells with everolimus alone.

3.5 Effects of everolimus and AZA on mTOR activation

No detectable impact on total mTOR and 4E-BP1 protein levels has been observed during incubation with everolimus and/or AZA in MZ-CRC-1 cells (figure 5). Everolimus alone significantly reduced both mTOR and 4E-BP1 phosphorylation. A similar effect has been
observed after everolimus in combination with AZA, while AZA alone did not significantly affect this process (figure 5).

3.6 Gene Expression analysis

To clarify the molecular bases for synergy between these two compounds, gene-expression profiles with or without everolimus and/or AZA treatment were evaluated in MZ-CRC-1 cells. Following treatment with everolimus or AZA alone, 72 DEGs (supplementary table 1) and 16 DEGS (supplementary table 2) were identified, respectively. On the other hand, 74 DEGs were identified following incubation with everolimus and AZA (supplementary table 3).

On each list of DEGs, a pathway analysis was performed with SPIA, which calculated the connections of pathways by considering their topology and the expression levels of their genes. Only for the comparisons between everolimus vs control and AZA/everolimus combination vs control significant pathways were detected (table 2 and 3). The significant pathways modulated after everolimus treatment (table 2) were PI3K-Akt signaling pathway and complement and coagulation cascades. SPIA analysis revealed four significant pathways in the group treated with the synergistic AZA/everolimus combination (table 3): PI3K-Akt signaling, neurotrophin signaling pathway, ECM-receptor interaction and focal adhesion.

The module network obtained by the fusion of the four significant pathways in the group treated with the AZA/everolimus combination is described in figure 6. Edges represent interactions between two genes. The list of genes in the module is reported in supplementary table 4. All the DEGs reported in figure 6 were overexpressed compared to the untreated control. The SEM multi-group analysis was performed comparing the covariance matrices implied by the model for the treatment data against the control data. The likelihood ratio test

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had a p-value of 0.002, confirming that the module was differentially regulated between the groups. The perturbed connections identified in this module are reported in the table 4.

Considering the perturbed edges (table 4), the links between the NGFR and BAX resulted significant and intriguing because involved in the process of apoptosis (figure 6). NGFR was overexpressed during exposure to the AZA/everolimus combination compared to untreated control. In the same condition, the activation of RAC1 (mediated by NGFR) and the indirect interaction RAC1/MAPK10 resulted to be stimulated (table 4). It has been previously reported that the activation of MAPK10 (JNK3), mediated by NGFR/RAC1, can induce apoptosis through the increased expression of TP53 and BAX (Aloyz et al., 1998; Kenchappa et al., 2010). Although, in our model the path TP53→BAX was not significantly perturbed during incubation with the AZA/everolimus combination, an increase in BAX expression was detected during synergistic treatment.

The same procedure was applied for the two pathways (table 2) previously identified during everolimus treatment. The directed shortest paths between the DEGs in the significant KEGG pathways generated the module reported in supplementary figure 1. The list of genes in the module is described in supplementary table 5. SEM was used to verify if globally the strength of connections was statistically different between control and everolimus in the assessed modules, but significant differences were not detected (p-value: 0.1144). Therefore, additional analyses were not performed.

An enrichment analysis based on Disease Ontology terms was performed on the genes selected in the module related to the treatment with “everolimus plus AZA” (figure 7). Twenty-four out of 25 identified terms were tumors.
3.7 Validation of gene expression data

From the final model of perturbed biological pathways during concomitant incubation with AZA/everolimus combination (figure 6), NGFR-MAPK10-TP53-Bax/Bcl-2 pathway was selected for further analysis on the basis of its strict interaction and modulation of apoptosis, and assessed by western blot (figure 8). Indeed, in our system apoptosis was the cell death mechanism responsible for the synergistic cytotoxic activity of everolimus plus AZA.

A significant increase in NGFR expression has been observed exclusively after 6 days of treatment with the synergistic combination of everolimus plus AZA in MZ-CRC-1 cells compared to untreated cells (figure 8). Everolimus and AZA alone and in combination increased the protein expression of MAPK10 and phosphorylated MAPK8/9/10, these effects were more pronounced with the combined treatment. In addition, a significant increase in phosphorylated p53 was observed only after incubation with everolimus and AZA combination (figure 8). The pro-apoptotic and anti-apoptotic Bax and Bcl-2, respectively, represent the key proteins involved in mitochondrial pathway of apoptosis. Western blot analysis showed that only the synergistic combination of everolimus plus AZA significantly increased Bax with a concomitant decrease in Bcl2 expression, while there was no significant change in the expression of these proteins after the exposure of MZ-CRC-1 cells to everolimus or AZA alone (figure 8).

To verify that the synergistic antitumor activity observed during AZA/everolimus co-incubation was indeed due to the increased expression in NGFR, we assessed the effects of both drugs alone or in combination on MZ-CRC-1 cells proliferation and protein expression of Bax and Bcl-2 after 6 days of treatment, in the presence or absence of a neutralizing antibody raised against NGFR (figure 9). The concomitant incubation with blocking NGFR antibody counteracted the cytotoxic activity of everolimus and AZA alone, and induced a moderate stimulation in cell proliferation after treatment with AZA/everolimus combination.
(+20% vs control, p<0.01, figure 9A). In a parallel set of experiments performed without the incubation with neutralizing antibody against NGFR, the more potent antiproliferative activity after incubation with everolimus plus AZA has been confirmed compared to single drugs (figure 9B). During concomitant incubation with neutralizing antibody against NGFR, everolimus plus AZA induced a significant decrease in Bax and increase in Bcl-2 expression, while these effects were mild with each drug alone (figure 9C, D).

3.8 Effects of everolimus and AZA on DNA methylation profile

Methylation levels of genes identified in the perturbed everolimus plus AZA module (figure 6 and supplementary table 4) were extrapolated from genome-wide methylation analysis using the Infinium HumanMethylation 450K BeadChip array performed in MZ-CRC-1 with or without treatment with everolimus and/or AZA. The array covers 769 probes for these selected 31 genes. Methylation levels for these genes resulted significantly different among groups (p=3.6x10^{-5}) (supplementary figure 2A). A significant reduction in DNA methylation levels has been observed after incubation with AZA alone, similar to that observed after AZA plus everolimus compared to untreated control. The incubation with everolimus alone did not significantly modify DNA methylation levels. These data were confirmed through the density plot showing methylation level distributions of untreated and treated samples (supplementary figure 2B). A simile trend has been observed for NGFR and MAPK10, both methylated genes involved in the NGFR-MAPK10-TP53-Bax/Bcl-2 pathway (supplementary figure 2C).

4. Discussion

Aberrant promoter methylation of tumor suppressor genes plays a relevant role in the initiation and progression of cancer. In addition, it is being increasingly recognized that aberrant DNA methylation seems to be involved in the development of drug resistance through transcriptional suppression of genes implicated in drug metabolism, apoptosis, cell
cycle control, and other biological processes (Hervouet et al., 2013). Taking into consideration that DNA methylation is a reversible event unlike genetic mutations, the use of demethylating agents has been recently proposed as a new therapeutic strategy in cancer. Several reports demonstrated the capability of AZA to overcome resistance to chemotherapeutic and biological agents in tumors (Oronsky et al., 2014; Vijayaraghavalu et al., 2013; Zhang et al., 2009). This was encouraging, given that AZA is clinically used for patients with acute myeloid leukemia and myelodysplastic syndromes.

In the current study, combination analysis, based on the Chou-Talalay's method, definitely demonstrated that AZA combined with everolimus was more effective in inhibiting cell proliferation than each agent alone in MTC cells. This cytotoxic activity was highly synergistic through a potent induction of apoptosis in MZ-CRC-1 cells, which was the cell line most resistant to everolimus alone. Whereas, in TT cells combined treatment exhibited both synergistic or antagonistic effects on cell proliferation inhibition, depending on the concentration of the drugs. In MZ-CRC-1 cells, while everolimus or AZA alone were barely effective in apoptosis induction, the AZA/everolimus combination doubled the fraction of MTC apoptotic cells compared to untreated controls. Cell-cycle analysis showed that everolimus decreased the percentages of MZ-CRC-1 cells in G2/M and S phase, but the combination with AZA did not potentiate this effect. Interestingly, the concentrations of AZA and everolimus adopted \textit{in vitro} for the synergistic antiproliferative activity can be readily achieved \textit{in vivo} with acceptable risk-benefit (O'Donnell et al., 2008; van Groeningen et al., 1986). In addition, the \textit{in vitro} toxicity of this combination appeared to be moderate. The cytotoxic activity of everolimus and AZA was lower in HEK-293 cells, a human embryonic kidney normal cell line, compared to that observed in MZ-CRC-1 cells. This is very promising in terms of safety and tolerability for future clinical trials.
Synergistic antiproliferative activity between AZA and everolimus was not related to a direct effect on mTOR activation. In fact AZA was unable to potentiate the inhibitory activity of everolimus on both mTOR and 4E-BP1 phosphorylation in MZ-CRC-1 cells.

Gene expression analysis revealed potential molecular mechanisms implicated in the synergy of AZA and everolimus in MZ-CRC-1 cells. We adopted an innovative bioinformatic pipeline based on SPIA and SEM, recently validated by one of us (Pepe and Grassi, 2014), that allowed to investigate pathway modules, considering not only deregulated genes but also the connections between the perturbed ones. Several key regulatory genes modulated by the combined treatment of AZA plus everolimus were identified by this approach: PI3K-Akt signaling, neurotrophin signaling pathway, ECM-receptor interaction and focal adhesion. Interestingly, all these pathways have a critical role in the regulation of both proliferation and migration/invasion of tumor cells. In addition, a Disease Ontology enrichment analysis, performed on these selected genes, identified 25 biological terms. Twenty-four out of 25 identified terms were tumors, including endocrine gland cancer. The only non-cancer item was “autosomal dominant disease”. These data further support the pivotal roles of these pathways in the development and progression of MTC. Indeed, MTC is a neuroendocrine tumor and MZ-CRC-1 cells harbor the Multiple Endocrine Neoplasia type 2B RET-M918T mutation, transmitted in this disease as an autosomal dominant trait (Santoro et al., 1995).

A perturbed pathway module, associated to the combined treatment of everolimus plus AZA, was generated by the detection and fusion of all shortest paths that put in communication the DEGs (figure 6). In this network, the “neurotrophin signaling pathway” appeared to exert a direct influence on the apoptotic machinery through the overexpression of NGFR and the activation of MAPK10-TP53-Bax pathway. Neurotrophins are a family of proteins involved

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in differentiation, plasticity and survival of neurons and modulating several functions of the neuroendocrine-immune system (Fiore et al., 2009). This family includes nerve growth factor (NGF), brain derived neurotrophic factor (BDNF), neurotrophins-3 (NT-3), neurotrophins-4/5 (NT-4/5) and neurotrophins-6 (NT-6). Biological activity of these factors is mediated through activation of Trk tyrosine kinase receptors (TrkA, TrkB, and TrkC) and NGFR (also known as p75 neurotrophin receptor, p75NTR) (Skaper, 2012). In our study, gene expression analysis confirmed the induction of apoptosis through the upregulation of Bax. Bax is a pro-apoptotic protein antagonized by the antiapoptotic Bcl-2. The ratio of Bax to Bcl-2 expression represents a cell death switch, which determine the susceptibility of cells to an apoptotic stimulus. Indeed, a selective decrease in Bax/Bcl-2 expression represents a common mechanism of drug resistance in tumor cells (Indran et al., 2011). We have found a selective increase in the expression of Bax/Bcl-2 ratio, validated through WB analysis, exclusively after treatment with AZA and everolimus. This effect seems to be associated with the upregulation of NGFR, which has been described to be a tumor-suppressor gene in several tumors (Dimaras and Gallie, 2008; Jin et al., 2007; Khwaja et al., 2004; Khwaja et al., 2006; Kuchler et al., 2011; Kuner and Hertel, 1998; Wang et al., 2014; Yang et al., 2015; Yuanlong et al., 2008), and consecutive activation of MAPK10 and p53.

Although the specific role of NGFR in the tumorigenesis and progression of MTC is unknown, neurotrophin signaling pathway appears to be involved in both preneoplastic thyroid C cell hyperplasia and MTC progression through Trk receptors (McGregor et al., 1999). In fact, a cross-talk between the signal pathways mediated by the Ret protooncogene and Trk receptors has been described (Esposito et al., 2008; Peterson and Bogenmann, 2004). The concomitant incubation with blocking NGFR antibody counteracted the cytotoxic activity of everolimus and AZA alone, and induced a moderate stimulation in cell
proliferation and a decrease in Bax/Bcl-2 ratio expression after treatment with AZA plus everolimus compared to control. This effect could be explained by the block of the pro-apoptotic pathway mediated by NGFR, while the survival pathways remained activated. These data confirmed that the NGFR overexpression plays a main role in the synergistic cytotoxic activity of these two compounds. Although the mechanism by which the expression of NGFR significantly increased during incubation with everolimus and AZA remains unclear, in colorectal cancer it has been recently reported that NGFR expression was silenced by promoter methylation and that the overexpression of this gene significantly inhibited cell proliferation, invasion and stimulated cell apoptosis (Yang et al., 2015). In our model the overexpression of NGFR appears not to be related to a direct epigenetic effect on NGFR gene. In fact, during incubation of MZ-CRC-1 cells with AZA, we found a significant decrease in DNA methylation levels of NGFR, comparable to that observed after everolimus plus AZA, while NGFR expression increased exclusively after combined treatment.

Intriguingly, the synergistic cytotoxic activity of everolimus/AZA combination was more relevant in MZ-CRC-1 (harboring RET-M918T mutation) than in TT (harboring RET-C634W mutation) cells (Table 1). We cannot exclude that the type of RET genetic alteration may have a role in the different antitumor activity observed in both cell lines. In fact, Gild et al. (2013) demonstrated that oncogenic RET regulates mTOR activity in MZ-CRC-1 and TT cell lines. Combined incubation with RET and mTOR inhibitors (AST487 and INK128, respectively) at low concentrations cooperated to inhibit mTOR signaling and cell growth, through induction of apoptosis, in both MTC cell lines. In addition, a recent study found that methylation profiles relate closely to RET mutational status in MTC, and the most distinctive methylome was observed for RET-M918T positive tumors (Mancikova et al., 2017).

In conclusion, we described for the first time a synergistic cytotoxic activity combining AZA with everolimus in MTC. This effect occurred through the overactivation of NGFR-
MAPK10-TP53-Bax/Bcl-2 pathway and the induction of apoptosis. These data provide a new therapeutic scenario in MTC and probably in other neuroendocrine tumors, where therapy with everolimus is currently approved.

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Table 1: Combination index (CI), dose reduction index (DRI) and potentiation factor (PF), according to the different cytotoxic ratio of everolimus (EV) and 5-aza-2’-deoxycytidine (AZA) combination in MZ-CRC-1 and TT cell lines after 6 days of treatment.

1CI<sub>50</sub> was calculated for 50% cell survival (ED<sub>50</sub>) by isobologram analyses performed with CalcuSyn software.

2DRI was calculated in case of synergism. It represents the order of magnitude (fold) of dose reduction obtained for ED<sub>50</sub> effect in combination setting as compared to each drug alone.

3PF was calculated in case of synergism as the ratio between the IC<sub>50</sub> of either everolimus or AZA alone and the IC<sub>50</sub> of everolimus or AZA in combination setting.

| Cell line | Cytotoxic ratio | ED<sub>50</sub> (nM) | CI<sub>50</sub> | DRI<sub>50</sub> | PF | Interpretation |
|-----------|----------------|----------------------|---------------|----------------|-----|----------------|
|           |                | EV       | AZA       | EV           | AZA | EV       | AZA       |               |
| MZ-CRC-1  | EV-AZA (50:50) | 12.4     | 35.6      | 0.2          | 5.9x10<sup>2</sup> | 1.8     | 1.1x10<sup>3</sup> | Strong synergism |
|           | EV-AZA (25-75) | 2.1      | 1.4x10<sup>2</sup> | 0.1          | 1.0x10<sup>2</sup> | 10.4    | 2.8x10<sup>2</sup> | Strong synergism |
|           | EV-AZA (75-25) | 1.3x10<sup>2</sup> | 3.2x10<sup>1</sup> | 2.3          | -               | -       | -               | Antagonism       |
| TT        | EV-AZA (50:50) | 9.2      | 3.4x10<sup>1</sup> | 1.3          | -               | -       | -               | Antagonism       |
|           | EV-AZA (25-75) | 1.2      | 8.0       | 0.7          | 6.4x10<sup>2</sup> | 6.1     | 11              | Synergism        |
|           | EV-AZA (75-25) | 91.1     | 1.2x10<sup>3</sup> | 17           | -               | -       | -               | Antagonism       |
Table 2. Significant pathways associated to treatment with everolimus by SPIA in MZ-CRC-1 cells

pPERT: probability of observing a total accumulated perturbation of the pathway more extreme than expected by chance; pNDE: probability of obtaining a number of DEGs in the given pathway at least as large as the observed one; pGFdr: false discovery rate for the global p-value from the combination of pPERT and pNDE.

| Pathway Name                                | pPERT  | pNDE   | pGFdr  |
|---------------------------------------------|--------|--------|--------|
| PI3K-Akt signaling pathway                  | 0.066  | 0.0013 | 0.036  |
| Complement and coagulation cascades         | 0.493  | 0.0005 | 0.048  |

Table 3. Significant pathways associated to treatment with everolimus plus AZA by SPIA in MZ-CRC-1 cells

pPERT: probability of observing a total accumulated perturbation of the pathway more extreme than expected by chance; pNDE: probability of obtaining a number of DEGs in the given pathway at least as large as the observed one; pGFdr: false discovery rate for the global p-value from the combination of pPERT and pNDE.

| Pathway Name                                | pPERT  | pNDE   | pGFdr  |
|---------------------------------------------|--------|--------|--------|
| PI3K-Akt signaling pathway                  | 0.102  | 0.0002 | 0.0178 |
| Neurotrophin signaling pathway              | 0.689  | 0.0002 | 0.0308 |
| ECM-receptor interaction                    | 0.115  | 0.0013 | 0.0308 |
| Focal Adhesion                              | 0.199  | 0.0018 | 0.0415 |
Table 4. Perturbed connections in the everolimus plus AZA module compared to the untreated control. The estimation of the strength connection and p-value for “Control”, “everolimus plus AZA” and the “difference between everolimus plus AZA vs control” are reported. We have only included the connections where a significant statistical P value has been observed for the “difference between everolimus plus AZA vs control”.

| Path          | Type of process | Control |       |       |       | Everolimus + AZA |       |       |       | Difference |       |
|---------------|-----------------|---------|-------|-------|-------|------------------|-------|-------|-------|------------|-------|
|               |                 | Estimate | P value |       |       | Estimate | P value |       |       | Estimate | P value |
| MDM2→TP53     | Inhibition      | 0.8271   | 0.0001 | -0.9864 | 0.0022 | -1.8135 | 0.0000 |
| NGFR→ARHGDIB  | Binding/association | 0.8935   | 0.0001 | 2.7208 | 0.0001 | 1.8273 | 0.0125 |
| EPHA2→IRS1    | Activation      | -0.5077  | 0.0079 | 0.3521 | 0.2699 | 0.8598 | 0.0208 |
| PIP5K1C→VCL   | Indirect        | -0.6464  | 0.0237 | 0.3072 | 0.0001 | 0.9536 | 0.0013 |
| VCL→ACTN1     | Binding/association | -0.0985  | 0.8827 | 2.4853 | 0.0002 | 2.5839 | 0.0062 |
| ITGA1→PTK2    | Binding         | -0.3282  | 0.0206 | 0.1908 | 0.2790 | 0.5190 | 0.0218 |
| NGFR→RAC1     | Activation      | -0.9042  | 0.0180 | 0.5223 | 0.0155 | 1.4265 | 0.0012 |
| RAC1→MAPK10   | Indirect        | 0.0499   | 0.6359 | 0.5799 | 0.0001 | 0.5300 | 0.0033 |
| NGFR→CDC42    | Activation      | 0.2736   | 0.3068 | -1.0567 | 0.0001 | -1.3303 | 0.0004 |
| PAK1→MAP2K1   | Phosphorylation | 0.4008   | 0.0018 | -0.0669 | 0.0998 | -0.4678 | 0.0005 |
FIGURE LEGENDS

**Figure 1.** Evaluation of synergism between everolimus (EV) and 5-aza-2’deoxyctydine (AZA) in medullary thyroid cancer cells by isobologram analysis. These experiments were performed with MTT assay. Combination index (CI)/fractional effect curves were elaborated with the dedicated software CalcuSyn (developed by Chou and Talalay) as described in Materials and methods. Curves show the CI vs the fraction of medullary thyroid carcinoma cells MZ-CRC-1 (A-C) and TT (D-F) affected by the EV-AZA combinations at 50:50 (A/D), 25:75 (B/E), 75:25 (C/F) cytotoxic ratios. CI represent the assessment of synergy induced by drug interaction. In detail, CI values of <1, 1, and >1 indicate synergy, additivity, and antagonism, respectively. Each point (represented in graph by x-mark) is the mean of at least 4 different replicates. The statistical significance of each point was evaluated with ANOVA and the derived p values were always less than 0.01.

**Figure 2.** In vitro toxicity profile of everolimus (EV) and 5-aza-2’deoxyctydine (AZA). Cells were treated for 6 days with EV (2.1 nM) and/or AZA (1.4x10^2 nM). (A) The cytotoxicity was measured by MTT cell viability assay in HEK-293 and MZ-CRC-1 cell lines. (B) The morphology of HEK-293 cells was determined after 6 days of exposure to drug free medium (CTR) and EV plus AZA. Images were captured using a phase-contrast microscopy at x 20 magnification.***p<0.001.

**Figure 3:** Cell cycle distribution, detected by FACS analysis, in propidium iodide stained MZ-CRC-1 cells after 6 days of treatment with 2.1 nM everolimus (EV) and/or 1.4x10^2 nM 5-aza-2’deoxyctydine (AZA). Control values have been set to 100%. *p<0.05  ** p<0.01

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Figure 4: Effects of 2.1 nM everolimus (EV) and/or 1.4x10^2 nM 5-aza-2’deoxyctydine (AZA) on apoptosis in MZ-CRC-1 cells after 6 days of incubation. (A) Flow cytometry with Annexin V and propidium iodide. Values of early apoptotic cells and late apoptotic/necrotic cells were expressed as percentage compared with the untreated control. (B) Representative western blot analysis of apoptotic markers (full and cleaved forms of caspase 3 and PARP). Actin was used as a loading control. (C) Quantification of western blot analysis from at least three independent experiments. *p<0.05; ** p<0.01; *** p<0.001

Figure 5: Effects of 2.1 nM everolimus (EV) and/or 1.4x10^2 nM 5-aza-2’deoxyctydine (AZA) on protein expression of total and phosphorylated mTOR and 4E-BP1 in MZ-CRC-1 cells after 6 days of incubation. (A) Representative western blot analysis. Actin was used as a loading control. (B) Quantification of western blot analysis from at least three independent experiments. *p<0.05; ** p<0.01

Figure 6. Perturbed pathway module in the group treated with everolimus and 5-aza-2’deoxyctydine. The green nodes are the DEGs, the yellow nodes represent the not DEGs microarray genes that connect the DEGs. Edges represent interactions between two genes.

Figure 7. Disease ontology enrichment analysis for the genes selected in the significant perturbed module, in the group treated with everolimus and 5-aza-2’deoxyctydine.

Figure 8. (A) Representative western blot analysis of NGFR, MAPK10, phosphorylated MAPK8/9/10, phosphorylated p53, Bax and Bel-2 performed in MZ-CRC-1 cells without (CTR) or after incubation with 2.1 nM everolimus (EV) and/or 1.4x10^2 nM 5-aza-
2′deoxycytidine (AZA). Actin was used as a loading control. (B) Quantification of western blot analysis from at least three independent experiments. *p<0.05; ** p<0.01; *** p<0.001

Figure 9: Effect of 2.1 nM everolimus (EV) and/or 1.4x10^2 nM 5-aza-2′deoxycytidine (AZA) on cell proliferation in MZ-CRC-1 cells (A, B) and protein expression of Bax and Bcl-2 (C, D) after 6 days of treatment, in the presence (A, C, D) or absence (B) of neutralizing antibody against NGFR (1:250). (C) Representative western blot analysis. Actin was used as a loading control. (D) Quantification of western blot analysis from at least three independent experiments. *p<0.05; ** p<0.01; *** p<0.001
