MiR-129-5p is required for histone deacetylase inhibitor-induced cell death in thyroid cancer cells

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Abstract

The molecular mechanism responsible for the antitumor activity of histone deacetylase inhibitors (HDACi) remains elusive. As HDACi have been described to alter miRNA expression, the aim of this study was to characterize HDACi-induced miRNAs and to determine their functional importance in the induction of cell death alone or in combination with other cancer drugs. Two HDACi, trichostatin A and vorinostat, induced miR-129-5p overexpression, histone acetylation and cell death in BCPAP, TPC-1, 8505C, and CAL62 cell lines and in primary cultures of papillary thyroid cancer (PTC) cells. In addition, miR-129-5p alone was sufficient to induce cell death and knockdown experiments showed that expression of this miRNA was required for HDACi-induced cell death. Moreover, miR-129-5p accentuated the anti-proliferative effects of other cancer drugs such as etoposide or human α-lactalbumin made lethal for tumor cells (HAMLET). Taken together, our data show that miR-129-5p is involved in the antitumor activity of HDACi and highlight a miRNA-driven cell death mechanism.

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Introduction

Mutation and epigenetic reorganization of chromatin are key events of tumor initiation and progression that lead to alteration of gene expression (Hanahan & Weinberg 2000, Jones & Baylin 2007). These epigenetic events include hypermethylation of promoters, modification of histone tails, and aberrant miRNA expression. Furthermore, histone deacetylases are overexpressed in tumors and promote carcinogenesis by blocking the transcription of tumor suppressor genes (Bolden et al. 2006). Finally, because of their efficacy in treatment of human malignancies many histone deacetylase inhibitors (HDACi) are currently being tested in clinical trials (Bolden et al. 2006, Prince et al. 2009), and have shown biological effects including tumor cell growth arrest, differentiation, and apoptosis. In thyroid cancer, several studies have reported that HDACi induce cell death through
caspase activation and B-cell CLL/lymphoma 2 (BCL2) downregulation (Greenberg et al. 2001, Catalano et al. 2005, Mitsiades et al. 2005). Moreover, HDACi were shown to sensitize tumor cells to chemotherapy, radiation, and surgery (Catalano et al. 2006, Noguchi et al. 2009). However, the molecular mechanisms responsible for the antitumor activity of HDACi remain elusive.

miRNAs are non-protein-coding small RNAs that negatively regulate the expression of hundreds of mRNA by inhibiting their translation or by promoting their degradation. miRNA has been identified as important developmental regulators and their misexpression may play a critical role in tumorigenesis and in clinical outcome (Calin & Croce 2006a,b, Esquela-Kerscher & Slack 2006). Moreover, previous studies have shown that cancer drugs including HDACi quickly modulate expression of several miRNAs (Scott et al. 2006), including miR-129-5p in bladder carcinoma cell lines (Dyrskjot et al. 2009). The precise impact of changes in miRNA expression in response to HDACi in cancer drug-associated tumor regression remains essentially unknown.

Few treatment options are currently available for poorly differentiated thyroid carcinomas and the prognosis is poor. Therefore, we have assessed the efficacy of HDACi on a panel of thyroid cancer cells ranging from well-differentiated carcinomas to aggressive, poorly differentiated tumors. We show that HDACi-induced cell death correlated with a specific increase in miR-129-5p expression, regardless of the oncogene involved and that miR-129-5p is necessary for HDACi-induced cell death. Taken together, the results suggest that miR-129-5p might be used as a biomarker for in vitro HDACi treatment efficacy.

Materials and methods
Reagents and HDACi
Etoposide, staurosporine and the most widely used HDACi trichostatin A (TSA) were provided by Sigma–Aldrich and the clinically relevant HDACi vorinostat (SAHA) by Cayman Chemical (Ann Arbor, MI, USA). PremiR and antagomiR were purchased from Applied Biosystems (Life Technologies, Carlsbad, CA, USA) (PM10195, AM10195, AM17110, and AM17010). Human α-lactalbumin made lethal for tumor cells (HAMLET) was produced from native purified human milk α-lactalbumin on an oleic acid-conditioned ion exchange matrix as described previously (Svensson et al. 2000).

Cell culture
The culture conditions for TPC-1, BCPAP, and 8505C (kind gift of Pr Santoro, Napoli, Italy) and CAL62 (kind gift of the Centre Antoine-Lacassagne (CAL), Nice, France) cells have been described previously (Gioanni et al. 1991, Ito et al. 1993, Fabien et al. 1994) and were grown in DMEM with 5% FCS, sodium pyruvate, and penicillin/streptomycin (Life Technologies). The toxicity of HDACi treatment on primary cultures of thyocytes derived from thyroid tissue of adenomas or hyperplasias or cancer cells derived from thyroid tissue of papillary thyroid carcinoma, obtained immediately after thyroidectomy, was tested. Patient consent was obtained and the institutional review board approved the project. After dissociation, primary culture cells were maintained in culture medium complemented with TSH. Contamination by fibroblast was controlled by the pathologists. An antibody against vimentin (Ventana, Paris, France; diluted 1:200; 45 min) was used to detect a potential contamination by fibroblast. No immunostaining was observed. Conversely, all cultured cells were immunostained by antibodies against HBME1 (Ventana; diluted 1:100; 45 min) and/or CK19 (Ventana; diluted 1:200; 45 min).

For all experiments, cells were grown to 70% confluence, and TSA (330 nM) or vorinostat (2.5 μM), were added for 16 h in fresh complete medium. Control cells were left untreated. The effect of HDACi on cell viability (cell cycle, 2,3-bis(2-methoxy-4-nitro-5-sulfophenyl)-2H-tetrazolium-5-carboxanilide (XTT) and morphology) was assessed by biochemical assays as well as by microarray analyses of miRNA and mRNA.

miRNA transfection
Cells were plated at 100 000 cells/well in a six-well plate and transfected with synthetic premiRs or antagomiRs using Lipofectamine RNAiMAX reagent (Life Technologies), following the manufacturer’s instructions at a final concentration of 5 and 20 nM respectively.

Western blotting
Western blotting was performed as described previously (Brest et al. 2007). The antibodies used were rb-achH4; ms-phospho-ser139-H2AX or rb-poly-ADP-ribose polymerase (PARP) (06-866, 05-636, AB16661 from Millipore, Billerica, MA, USA) and rb-GAPDH-HRP conjugate (Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA). Immunoreactive bands were revealed by enhanced chemiluminescence (Amersham-GE Healthcare, Velizy-Villacoublay, France) using Kodak film.
Isolation of RNA

Isolation of RNA from cultured cells or human samples was performed as described previously (Triboulet et al. 2007, Pottier et al. 2009).

miRNA microarrays

miRNAs of BCPAP cells treated with TSA (330 nM, 16 h) were profiled by the University of Nice Sophia-Antipolis Core facility (www.microarray.fr) using home-made arrays (Triboulet et al. 2007, Pottier et al. 2009) and deposited in NCBI Gene Expression Omnibus (GEO; http://www.ncbi.nlm.nih.gov/geo/) under series GSE19934 and platform GPL4717. Briefly, total RNA was labeled with Cy3 or Cy5 fluorescent dyes (a dye-swap method was used) using the Ulysis Alexa fluor nucleic acid labeling kit (Amersham Bioscience), and miRNA was then isolated using the mirVana miRNA isolation kit (Applied Biosystems). Three independent experiments were performed to identify miRNA up- or downregulated in cells. For each oligonucleotide microarray, TIF images containing data from each fluorescence channel were quantified with GenePix Pro 6.1 program (Molecular Devices, Sunnyvale, CA, USA). Normalization was obtained with limmaGUI Software from Bioconductor (www.bioconductor.org) (Wettenhall & Smyth 2004, Ritchie et al. 2007), according to the variance stabilization and calibration for microarray data (VSN) approach described by Huber et al. (2002).

Expression microarrays

miR-129-5p transfected TPC-1 cells were profiled using pangenomic microarrays printed with the human RNG/MRC oligonucleotide collection as described previously (Le Brigand et al. 2006) and deposited in GEO under series GSE19933 and platform GPL3241. Total RNA (2 μg) was amplified using the Amino Allyl MessageAmp aRNA kit (Ambion) according to the manufacturer’s instructions. Cy3- and Cy5-labeled aRNA were hybridized on the array for 17 h at 62 °C (a dye-swap method was used). Arrays were scanned and quantified as described earlier.

Intra- and inter-slide normalization of three independent experiments was performed by the Global Loess and the quantile methods respectively. Means of ratios from all comparisons were calculated and a B test analysis was performed by the Limma package available from Bioconductor (Gentleman et al. 2004). Differentially expressed genes were selected using a Benjamini–Hochberg correction of the P value for multiple tests, with a positive B value. Microarray data was archived in GEO under reference GSEX. Before analysis, the Sylamer algorithm was used to characterize the specificity of miR-129-5p on global mRNA gene expression. The human targets of the
differentially expressed miRNAs were predicted using public web-based prediction tools, such as PicTar, TargetScan, and miRBase Targets.

Quantitative PCR

Quantitative PCR (QPCR; sequences described in the Supplementary methods, see section on supplementary data given at the end of this article) for mRNA was carried out using the SYBR Green Master Mix from Applied Biosystems. QPCR for miRNAs was carried out using probes for miR-129-5p and RNU19-RNA, and the TaqMan miRNA kit from Applied Biosystems.

Viability assay

Cell survival was examined using the Cell Proliferation Kit II (XTT) from Roche and quantification of the DNA fragmented-subG1 population was performed by cell cycle analysis as described previously (Brest et al. 2007). All experiments were repeated at least three times, and each XTT experimental sample (5000 cells/well) was repeated at least in quadruplicate wells for each experiment. The data are average values ± S.D. of representative experiments.

Statistical analysis

For statistical analysis the SPSS16 program was used (SPSS, Chicago, IL, USA). The results were evaluated for statistical significance by the Student’s t-test or the ANOVA test. Error bars represent the S.D. of the mean. P values < 0.05 were regarded as significant.

Results

HDACi increase miR-129-5p expression in thyroid cancer cells

The molecular mechanisms responsible for the anti-tumor activity of HDACi remain poorly characterized and the relationship between the sensitivity to HDACi and miRNAs expression has not been investigated in thyroid tumor cells. To test this, we first treated the BCPAP papillary thyroid cancer cell line with the HDACi-reference TSA (330 nmol/l). The BCPAP cell line was sensitive to TSA (Fig. 1A), as shown by an increase in histone H4 acetylation. TSA-induced apoptosis, as shown by PARP cleavage (Fig. 1A), an elevated fraction of cells in subG1 (Fig. 1B), and phosphorylation of histone H2AX (Fig. 1A) that forms before the appearance of internucleosomal DNA fragments and the externalization of phosphatidylserine to the outer membrane leaflet (Rogakou et al. 2000). The decrease in viability was also observed after XTT quantification (Fig. 1C), the concentrations selected for the rest of the study were 330 nmol/l and 2.5 μmol/l for TSA and vorinostat respectively. Thus, RNA from the BCPAP cell line treated or not with TSA overnight (330 nmol/l) was analyzed using a miRNA microarray (Triboulet et al. 2007, Pottier et al. 2009). TSA-treated cells showed increased expression of several miRNAs, such as miR-129, miR-513, and miR-663 (Table 1).

Table 1 Microarray results of statistically regulated microRNA in BCPAP cells treated with trichostatin A (330 nmol/l; n=3 experiments)

<table>
<thead>
<tr>
<th>Name</th>
<th>LogFC</th>
<th>P value</th>
<th>Adj. P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mir-513 (hsa)</td>
<td>0.95</td>
<td>1.20×10^{-5}</td>
<td>1.66×10^{-2}</td>
</tr>
<tr>
<td>Mir-663 (hsa)</td>
<td>0.97</td>
<td>2.55×10^{-5}</td>
<td>1.96×10^{-2}</td>
</tr>
<tr>
<td>Mir-129 (hsa,xtr)</td>
<td>0.73</td>
<td>1.01×10^{-3}</td>
<td>8.05×10^{-2}</td>
</tr>
<tr>
<td>Mir-373 (hsa)</td>
<td>0.94</td>
<td>1.10×10^{-3}</td>
<td>8.06×10^{-2}</td>
</tr>
</tbody>
</table>

LogFC, logarithm of fold-change.

Figure 2 HDACi increases cell death and mir-129-5p expression. Malignant thyroid cells (TPC-1, BCPAP, 8505c, CAL62, and primary papillary thyroid cancer PTC n=3 cells) and non-malignant primary culture cells (adenoma n=3 and hyperplasia n=4) were treated without or with TSA (18 h, 330 nmol/l). (A) The expression of miRNA was analyzed by RT-PCR. Results represent the fold increase in miR-129-5p in treated cells vs untreated cells used as a reference. (B) HDACi-induced cytotoxicity analyzed by XTT. *, P<0.05; **, P<0.01.
with untreated cells whereas other changes in miRNA expression were not confirmed by this method.

We corroborate these results obtained for BCPAP (BRAF V600E) with another papillary thyroid cell line TPC-1 (RET/PTC1) and the anaplastic thyroid cell lines 8505c (BRAF V600E) and CAL62 (KRAS G12R) when treated with the HDACi-reference TSA (330 nmol/l). Specific TaqMan PCR confirmed that TSA treatment induced an increase in miR-129-5p (mature form) expression ranging from 7- (CAL62) to 21-fold (TPC-1) in comparison with untreated cells (Fig. 2A). The highest induction was found in papillary cell lines (BCPAP and TPC-1), which correlated with their high sensitivity to cell death (Fig. 2B). These results indicate that TSA induces apoptotic cell death in thyroid cancer cells, regardless of the cancer type and their oncogenic status.

Moreover, the increased cell death and expression of miR-129-5p was only observed in primary papillary thyroid cancer (PTC) treated with TSA while no increase was observed in primary thyroid cell cultures form adenomas or hyperplasias treated with TSA (Fig. 2). The results indicate that HDACi treatment upregulates miR-129-5p expression in tumor cells, but not in non-malignant cells.

**miR-129-5p is cytotoxic for thyroid cancer cells**

We next examined the effect of miR-129-5p overexpression on cell death. For this purpose we used TPC-1 cells that we showed to be the most sensitive and found that miR-129-5p-transfection is cytotoxic for cancer cells, as shown by increased propidium iodide staining, loss of viability (XTT), and PARP cleavage (Fig. 3A–C respectively). This result was confirmed in all the tested cell lines (Fig. 3D). Moreover, no or few cell death was observed in thyroid primary cells (data not shown) showing a specific effect for transformed cancer cells. We then analyzed the influence of limited miR-129-5p overexpression (always <25-fold increase) on transcript levels using human pan genomic arrays, as described previously (Triboulet et al. 2007, Pottier et al. 2009). Data collected from three independent biological

![Figure 3](https://example.com/fakeimage.png)
experiments on TPC-1 cells, revealed that a total of 224 transcripts were significantly modulated (102 up- and 122 downregulated, \( P < 0.01 \)) following premiR-129-5p transfection compared with the control condition (Supplementary Table 1, see section on supplementary data given at the end of this article). Analysis of this signature with Ingenuity Pathway Software (Redwood City, CA, USA) indicated a significant enrichment for ‘molecular functions’ with terms such as ‘cellular compromise, cellular growth and proliferation, cell cycle, or cell death’ (Supplementary Table 2, see section on supplementary data given at the end of this article).

Using our bioinformatic tool ‘MicroToptable’ (Pottier et al. 2009), we then looked for potential over-representation of miR-129-5p-predicted targets among the downregulated transcripts (cut off equal to 8.0 for the \( \log_2 \) (signal), to \(-1\) for the \( \log_2 \) (ratio), and to 0.01 for the adjusted \( P \) values) and isolated 42 transcripts corresponding to putative targets (Supplementary Table 3, see section on supplementary data given at the end of this article). By RT-PCR, we validated a decrease in UDP-N-acetyl-\( \alpha \)-D-galactosamine: polypeptide \( N \)-acetylgalactosaminyltransferase 1 (GALNT1; \( 2^{-2.0} \sim -4 \) fold), a gene previously described to be regulated by miR-129-5p (Dyrskjot et al. 2009). Interestingly, we also confirmed an increase in the pro-apoptotic protein programed cell death 2 (\( 2^{1.7} \sim 3.2 \) fold; Supplementary Table 3, see section on supplementary data given at the end of this article) concomitant to a decrease in its repressor BCL6 (\( -2^{-2.4} \sim -5.3 \) fold). These results were in accord with the predictive action of this miR-129-5p on BCL6 and GALNT1 using the PicTar and TargetScan algorithm. These results suggest that induction of miR-129-5p expression may participate in HDACi-induced cell death possibly through the repression of GALNT1 and BCL6.

**miR-129-5p is necessary for HDACi-induced cell death**

To examine if the induction of miR-129-5p expression was required for HDACi-induced cell death, TPC-1 cells were transfected for 24 h with specific antagomiR directed against miR-129-5p and then incubated in the presence of TSA (330 nmol/l, 24 h; Fig. 4A). AntagomiR-129-5p significantly blocked TSA-increased cell death in contrast to antagomiR-neg transfected cells.
treated with HDACi, as shown by both microscopy and the XTT viability assay (Fig. 4A–C). Furthermore, HDACi-dependent repression of GALNT1 expression was reverted by antagomiR-129-5p. GALNT1 was even overexpressed in presence of the antagomiR and TSA (Fig. 4D). However, HDACi-dependent BCL6 decreased expression was not reversed by antagomiR-129-5p showing a potential combined effect of TSA-induced microRNA on BCL6 expression (data not shown). In conclusion, silencing of miR-129-5p in thyroid cells was able to decrease HDACi-induced cell death, demonstrating that miR-129-5p expression is a key mediator of HDACi-mediated killing of tumor cells.

miR-129-5p promotes drug-induced cell death

HDACi have previously been shown to induce cell death in response to other cancer drugs such as etoposide, cisplatin, or HAMLET (Brest et al. 2007). To examine if miR-129-5p may have a similar effect, we transfected TPC-1 cells with either miR-129-5p or miR-CTL for 24 h before the addition of increasing concentrations of etoposide or HAMLET. Combination with miR-129-5p was additive with HAMLET, staurosporine, or etoposide treatment (Fig. 5). The loss of viability in response to HAMLET (0.3 mg/ml, 24 h) was 40% in miR-CTL transfected control cells, but increased to 70% in tumor cells transfected with miR-129-5p. A similar increase in cell death was obtained in cells treated with staurosporine or etoposide. In CAL62 cells, the results were similar to TPC-1 cells (data not shown). Thus, while miR-129-5p affects tumor cell viability per se, it also accentuates the effect to different cancer drugs.

Discussion

Thyroid cancers represent a broad spectrum of neoplastic disorders that include differentiated cancers with good prognosis after surgery and radioiodine therapy, but also poorly differentiated carcinomas and some medullary carcinomas that are associated with poor prognosis. These tumors are often inoperable, resistant to radiotherapy and have a high tendency to metastasize. Conventional chemotherapy is often ineffective, highlighting the need for novel therapeutic strategies. Despite the therapeutic efficacy of HDACi in several malignancies, the mechanism of action remains unclear. In this study, we show that the activity of miR-129-5p is directly related to the efficiency of HDACi as well as to substances with a related or synergistic mechanism of action. For this study, we used the most widely used HDACi TSA for which the development as a therapeutic agent has been neglected partly due to: i) the high cost of production and isolation from fermentation broth and ii) the clinically relevant HDACi vorinostat has been approved by the U.S. FDA for the treatment of cutaneous T-cell lymphoma and has been tested for treatment of patients with metastatic and/or locally advanced or locally recurrent thyroid cancer (clinical trial phase II).

Few studies have evaluated the effect of HDACi on thyroid cancer cells and no study has compared the sensitivity of HDACi on a panel of thyroid cancers harboring different oncogenes. It has been shown that vorinostat induces cell death in FRO81-2 anaplastic thyroid cancer cells (BRAF V600E) by activation of caspase and degradation of Bcl-2 family proteins. (Mitsiades et al. 2005). However, to our knowledge, no study has investigated the effect of HDACi on the overexpression of specific miRNA in thyroid cancer cells. We show here that both TSA and vorinostat have a potent effect on miR-129-5p expression in thyroid cancer cells, while cultures of non-malignant primary cells were insensitive to these drugs. In addition, miR-129-5p was strongly induced in the entire thyroid cell lines tested (Fig. 2) and in some other cell lines such as A549 (lung carcinoma) and MEL-501 (melanoma) cells (data not shown). While this work was in progress, it was shown that the HDACi 4-phenylbutyric acid restores the expression of miR-129-2 (miR-129-5p precursor) in colorectal cancers (Bandres et al. 2009), which is in agreement with our results.

We next investigated the impact of miR-129-5p expression on HDACi-induced cell death. Overexpression
of miR-129-5p alone was able to induce cell death in our model of TPC-1 thyroid cancer, confirming a recent study showing that miR-129-5p was cytotoxic in bladder carcinoma cell lines (Dyrskjot et al. 2009). A direct link between miR-129-5p and the two putative targets GALNT1 and SOX4 was documented (Dyrskjot et al. 2009). In our model, transfection with miR-129-5p, led to a strong decrease in GALNT1 (approximately fourfold; Supplementary Table 3, see section on supplementary data given at the end of this article) and BCL6 (~5.2-fold) expression, but to a modest decrease in SOX4 (~1.6-fold, data not shown) expression. Using our bioinformatic tools, we also isolated, for further functional studies, a group of 42 transcripts (including GALNT1) corresponding to putative miR-129-5p targets (Supplementary Table 1, see section on supplementary data given at the end of this article). We showed that miR-129-5p knockdown blocked HDACi-induced cell death in our model. This result showed that miR-129-5p is both sufficient and mandatory for HDACi-induced cell death and obviously through a GALNT1-dependent function.

Over the last decade, the explosion of efforts into drug discovery has led to the development of a large group of HDACi, many of which have been shown preclinically to have potent antitumor activity. Moreover, HDACi have been shown to synergize with many anti-cancer agents, including cytotoxic agents such as paclitaxel (Dowdy et al. 2006), cisplatin (Strait et al. 2005), etoposide and doxorubicin (Marchion et al. 2004), HAMLET (Brest et al. 2007), or radiotherapy (Noguchi et al. 2009). In this study, we showed that miR-129-5p enhanced the antitumor effects of staurosperine, etoposide, and HAMLET when used in combination (Fig. 5).

There is a need to better define and target HDACi for clinical use in specific patient groups. Specific biomarkers for different tumor types or mechanisms may be vital to stratify patients and tumors into subgroups that are responsive, as well as to enable target modulation to be monitored. Our findings provide new evidence for a miRNA-driven antitumor effect of HDACi and therefore we propose that miR-129-5p might be useful as a biomarker in following the HDACi-induced response in tumor cells, to define susceptible and resistant tumors.

Supplementary data
This is linked to the online version of the paper at http://dx.doi.org/10.1530/ERC-10-0257.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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