Proteomic screening identifies calreticulin as a miR-27a direct target repressing MHC class I cell surface exposure in colorectal cancer

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Impairment of the immune response and aberrant expression of microRNAs are emerging hallmarks of tumour initiation/progression, in addition to driver gene mutations and epigenetic modifications. We performed a preliminary survey of independent adenoma and colorectal cancer (CRC) miRNA data sets and, among the most dysregulated miRNAs, we selected miR-27a and disclosed that it is already upregulated in adenoma and further increases during the evolution to adenocarcinoma. To identify novel genes and pathways regulated by this miRNA, we employed a differential 2DE-DIGE proteome analysis. We showed that miR-27a modulates a group of proteins involved in MHC class I cell surface exposure and, mechanistically, demonstrated that calreticulin is a miR-27a direct target responsible for most downstream effects in epistasis experiments. In vitro miR-27a affected cell proliferation and angiogenesis; mouse xenografts of human CRC cell lines expressing different miR-27a levels confirmed the protein variations and recapitulated the cell growth and apoptosis effects. In vivo miR-27a inversely correlated with MHC class I molecules and calreticulin expression, CD8+ T cells infiltration and cytotoxic activity (LAMP-1 exposure and perforin release). Tumours with high miR-27a, low calreticulin and CD8+ T cells' infiltration were associated with distant metastasis and poor prognosis. Our data demonstrate that miR-27a acts as an oncomiRNA, represses MHC class I expression through calreticulin downregulation and affects tumour progression. These results may pave the way for better diagnosis, patient stratification and novel therapeutic approaches.

Cell Death and Disease (2016) 7, e2120; doi:10.1038/cddis.2016.28; published online 25 February 2016

Colorectal cancer (CRC) is the third major cause of cancer worldwide. In addition to gene mutations and epigenetic modifications, impairment of the immune response and dysregulation of microRNAs are emerging hallmarks of tumour initiation/progression. Tumour cells elicit a native and adaptive immune response mediated by different cell types aimed to eradicate the tumour. Antigen processing and presentation by major histocompatibility complex (MHC) class I molecules is a critical event to mount a specific antitumour response. Class I antigen peptides are originated from proteasome-mediated degradation of intracellular proteins and transported to the endoplasmic reticulum through the transporters associated with antigen-processing proteins (TAP1 and 2). Here they are loaded onto the peptide-loading complex (PLC) formed by GRP78, calnexin and tapasin on which calreticulin and ERp57 are recruited to interact with MHC class I molecules. Once loaded with ‘optimal’ antigens, these latter molecules dissociate from the PLC and translocate to the cell surface where they are recognized by cells of the immune system, contributing to immune surveillance.

Defects of MHC class I antigen presentation occur at high frequency in solid tumours and is a feature of tumour immune evasion that renders cancer cells invisible to cytotoxic T cell. A selective loss or reduced level of MHC class I is generally associated with disease progression and reduced patient survival. Specifically, downregulation of MHC class I is detected in >74% of colorectal tumours and associated with a significantly shorter mean disease-specific survival compared with MHC-class-I-high-expressing tumours. miRNAs dysregulation is a common feature of human malignancies as they act as oncomiRNAs or tumour-suppressor miRNAs. The contribution of miRNAs to antitumour immune response is still undefined and is a topic of current investigation.
under intense investigation. In this study, we performed a preliminary survey of independent adenoma and CRC miRNAs, we selected miR-27a and showed that it is already upregulated in adenoma and further increases during the evolution to adenocarcinoma. Subsequently, using a proteomic approach, we identified a series of new proteins modulated by miR-27a that are involved in MHC class I cell surface exposure. In gain- and loss-of-function experiments, we provide evidence that miR-27a impairs this pathway directly targeting calreticulin. Furthermore, miR-27a greatly affects cell proliferation and angiogenesis in vitro and in mouse xenografts where these results are recapitulated. Consistently, miR-27a is overexpressed in a large proportion of human sporadic CRCs, inversely correlates with MHC class I molecules and calreticulin and CD8+ T cells’ infiltration and activity. The combination of high miR-27a/low calreticulin is associated with distant metastasis and worse outcome.

Results

miR-27a is upregulated in human adenoma and CRC. We preliminarily analysed a publicly available adenoma data set for miRNA expression profile (E-MTAB-813): the heatmap reports the list of those that are upregulated or downregulated. We also analysed two independent CRC miRNA data sets (GSE35602 microarray and TCGA_miRNA-seq) and, upon intersecting the results as shown in the Venn diagram, identified three common upregulated miRNAs (Figures 1A–C). We selected miR-27a for further analysis and assessed its expression in our series of adenomas (n = 32) and sporadic CRCs (n = 80) by quantitative RT-PCR analysis. miR-27a was already elevated in about 60% of adenomas and further increased during tumour progression (stages I–II, n = 48; stages III–IV, n = 32), suggesting that its aberrant expression is an early event in colon tumourigenesis (Figure 1D). These data were confirmed by in situ hybridization that showed high levels in adenomas that increased in differentiated and even more in undifferentiated tumour samples (Figure 1E).

Identification of novel genes and pathways regulated by miR-27a by differential proteome analysis. To identify novel proteins and pathways modulated by miR-27a, we employed a proteomic approach. We first assessed its expression in a series of CRC-derived cell lines and selected HCT116 among the overexpressing ones (Figure 2A). A plasmid vector carrying a short hairpin anti-miR-27a RNA (shRNA) and the GFP (green fluorescence protein) cassette was stably transfected into HCT116 cells (CTRL); a cell clone, hereafter defined miR27a_KD, was chosen for further studies (Figure 2B). The efficacy of the silencing was established by assessing diminution of miR-27a and increase in validated targets (PPARG, ZBTB10 and FBXW7) (Figure 2c and Supplementary Figure S1A). We also transfected HCT116 cells with a plasmid carrying the miR-27a mimic and, among the overexpressing clones, we selected one hereafter named miR27a_OE. Extracts from CTRL and miR27a_KD cells were analysed by comparative proteomic 2DE-DIGE showing distinct expression profiles and a consistent experimental reproducibility (Supplementary Figure S1B, Supplementary Table S1). Out of the 51 differentially expressed spots, 27 were identified by LC-MS/MS, correlated with the corresponding spots and classified into 9 functional classes according to their biological activities (Figures 2d and e, Supplementary Table S2). Quantification of the selected spots is shown in a three-dimensional view along with the corresponding standard abundance in the two different cell lines (Figure 2f). Validation by western blotting analysis revealed an increase of calreticulin (CRT), tapasin (TAPBP), GRP78, ERP57 and annexin A1 (ANXA1) and reduction of TRAP1 in miR27a_KD with respect to CTRL cells; the inverse results were obtained in miR27a_OE (Figure 2g).

miR-27a downmodulates MHC class I cell surface exposure. To recognize the pathways in which the identified proteins are involved, we queried the Ingenuity Pathway Analysis (IPA) algorithm: antigen presentation, immunological, and inflammatory pathways were the most enriched ones (Figure 3a, Supplementary Figure S1C). Accordingly, we assessed cell-surface exposure of MHC class I molecules in three different cell lines (HCT116, HT29 and RKO) and their derivative clones with either a silenced or overexpressed miR-27a. In flow cytometry, miR27a_KD cells displayed more MHC class I proteins on the surface than the parental CTRL cells, whereas the miR27a_OE cells displayed less proteins (Figure 3b). Similarly, in immunofluorescence staining, the specific antibody recognized a low amount of MHC class I proteins on the membrane of non-permeabilized HCT116 CTRL cells that remarkably increased on the surface of miR27a_KD, whereas diminished in miR27a_OE cells (Figure 3c). The specificity of the staining was validated by counterstaining the nuclei with DAPI: by merging, the two stainings remained separate as they mark different subcellular compartment. To have a more quantitative assessment of the proteins exposed on the cell surface, we set up a procedure to selectively isolate plasma membrane proteins and evaluate those included. The miR27a_KD cells membrane fraction contained at least four times more MHC class I molecules than CTRL and even more than miR27a_OE cells. That the bands corresponded to real membrane proteins was confirmed by challenging the same fraction with E-cadherin, an integral membrane protein, as a positive control, and with β-Actin, a cytosolic protein as a negative control (Figure 3d). All together these data demonstrate that the surface expression of MHC class I molecules is downregulated by miR-27a.

miR-27a directly targets calreticulin affecting MHC class I exposure. To identify a direct target of miR-27a among the differentially expressed proteins, we acquired several algorithms and predicted calreticulin as a putative target owing to a conserved seed recognition sequence in the 3’UTR of the corresponding mRNA (Figure 4a). A miScript Target Protector (TP) was designed against this recognition sequence to selectively prevent the binding of miR-27a to the corresponding mRNA, without interfering with the action of the miRNA on other targets. Upon TP transfection, calreticulin increased in
miR-27a represses MHC class I surface exposure

T Colangelo et al

HCT116 and miR27a_OE more than miR27a_KD cells (Figure 4b). Interestingly, also the cognate mRNA increased, likely owing to a stabilization effect, contributing to the protein elevation detected in HCT116 and miR27a_OE cells (Figure 4c). Transfection of three independent siRNAs against calreticulin mRNA in HCT116, miR27a_KD and miR27a_OE cells successfully silenced the protein, with siRNA#3 as the most efficient (Figure 4d). To determine the impact of the miR-27a/calreticulin axis on MHC class I molecules surface exposure, we transfected calreticulin TP and siRNAs in the three cell lines. In flow cytometry, TP produced no significant variations, whereas the siRNAs,
especially siRNA#3, increased MHC class I cell membrane display by 50% in miR27a_KD and 25–30% in CTRL and miR27a_OE cells (Figure 4e), in line with the total content of the isolated plasma membrane fraction (Figure 4f). Calreticulin, thus, is a direct target of miR-27a and mediates the effects on MHC class I exposure.

Mouse xenografts recapitulate miR-27a effects on the proteomic profile and cell growth. To investigate the modulation of miR-27a on the proteins identified by 2DE-DIGE in vitro, we generated mouse xenografts.
injection of a miR-27a mimic into HT29 cell-derived masses were 450% larger than those from the parental cells (Figure 5a). The specificity and efficacy of miR-27a inhibition or overexpression was verified by qRT-PCR on total RNAs extracted from the two different types of tumours. In accordance with the size, Ki67 positivity was stronger in the high miR27a-expressing tumours than the lower ones, supporting a role of this miRNA in cell proliferation (Supplementary Figures S3A and B). By contrast, apoptosis was greatly reduced in the same tumours, whereas large areas of apoptosis were detected in those expressing low miR-27a by a terminal deoxynucleotidyl transferase (TdT)-mediated dUTP nick end-labelling test (TUNEL) assay (Figures 5b and c).

Calreticulin, ERp57, GRP78/BiP, Annexin1 and Tapaxin were downregulated in western blotting analysis of extracts from HCT116 tumours, whereas they were all upregulated in those with a silenced miR-27a. TRAP1 exhibited an inverse behaviour, in agreement with the 2DE-DIGE results in vitro (Figures 5d and e). The analysis of the same markers in HT29 tumours produced an opposite scenario, in line with the lower expression of miR-27a that was reversed upon injection of the corresponding mimic. No variations for all the indicated proteins were observed in the western blots of extracts from tumours injected with scrambled control with respect to HCT116 or HT29-derived tumours (data not shown).

MHC class I expression was monitored by immuno-histochemistry (IHC). The signal detected on tissue sections from miR-27a antisense-injected tumours was stronger than from the scrambled injected or parental cell tumours. At a higher magnification, the staining was localized at the cell membrane, consistent with stimulation of MHC class I cell surface translocation upon miR-27a silencing. A more quantitative western blotting analysis on extracts from the same tumours confirmed in a mouse model that miR-27a silencing was associated with an overall increase of MHC class I proteins (Figures 5b, d and e). Consistently, calreticulin displayed a weak and mainly cytosolic positivity on sections of HCT116 tumours by IHC. The staining was, instead, remarkably localized as ‘patches’ on cell membranes in tumours injected with a miR-27a antisense (Figure 5b). Western blotting analysis of extracts from the same tissues exhibited an overall increase of calreticulin only in those masses with a reduced miR-27a (Figures 5d and e).

Collectively, mouse xenografts confirmed that miR-27a affects cell growth and apoptosis also in CRC and clearly

Figure 3  MHC class I cell surface exposure is downregulated by miR-27a in CRC cells. (a) The drawing shows the most enriched networks generated from the list of differentially expressed (DE) proteins (red elements = upregulated proteins; green elements = downregulated proteins) after miR-27a silencing using IPA. (b) Flow cytometry analysis reveals different MHC class I molecules cell surface exposure in HCT116, RKO, HT29 and their derivative clones miR27a_KD or miR27a_OE. (c) Immunofluorescence staining using an antibody against human MHC class I molecules in HCT116 CTRL and their derivative clones miR27a_KD or miR27a_OE (scale bar, 50 μm). (d) Enrichment of MHC class I molecules in the isolated plasma membrane fraction. Positivity for E-cadherin, a membrane protein, and negativity for β-Actin, a cytosolic protein, proved that the identified proteins were truly integral membrane components. The relative fold change, obtained by densitometric analysis and normalization to E-cadherin, is reported below the bands. *P ≤ 0.05; **P ≤ 0.01 (two-tailed Student’s t-test). Data are representative of three independent experiments and error bars represent S.D. of technical replicates (mean ± S.D.)
showed that it negatively modulates a specific set of proteins identified in vitro that specifically contribute to MHC class I expression. These results definitively demonstrate that miR-27a is a CRC tumour-inducing factor acting as an oncomiRNA.

miR-27a expression in CRC inversely correlates with MHC class I and calreticulin expression and with CD3+ and CD8+ T cells’ infiltration/activation. To correlate the data obtained in vitro and in mouse xenografts with human CRCs, we performed quantitative western blotting analysis of some representative samples: high miR-27a-expressing CRCs displayed low MHC class I molecules and calreticulin (Figures 6a and b). Accordingly, IHC of tissue microarrays showed that high miR-27a-expressing tumours frequently displayed a weak or absent membrane staining for MHC class I molecules and calreticulin; the staining was stronger in low miR-27a-expressing tumours (Figure 6c). Furthermore, miR-27a inversely correlated with CD3+ and CD8+ T cells’ infiltration and perforin positivity whose relative abundance was determined (Figures 6c and d). Perforin and LAMP-1 are two membrane proteins used as surrogate markers of CD8+ T cells’ cytotoxic activity.23 Thus CD8+/perforin+ and CD8+/LAMP-1+ double-positive cells, detected by immunofluorescence on CRC specimens, were higher in low miR-27a-expressing tumours (Figures 7A and B). Altogether, these results suggest that miR-27a could impair T cells’ infiltration, activation, proliferation and degranulation.24 Consistently, Kaplan and Meier analysis of patients’ survival showed that low calreticulin expression (P < 0.001) and CD3+ and CD8+ low infiltrates (P < 0.001), taken alone, were significantly associated with a shorter overall survival, whereas high miR-27a showed only a trend (P = 0.104; Supplementary Figures S4A and B). The low calreticulin/high miR-27a association (n = 26) was the one with the worst outcome when these characteristics were combined; hazard ratios analysis of all possible associations identified calreticulin as a dominant variable that was even more discriminant when coupled with high miR-27a expression. When the CD8+ infiltrates were associated with miR-27a levels, the combination low CD8+/high miR-27a had the worse prognosis; hazard ratios analysis of all possible associations highlighted the presence of CD8+ infiltrates as a dominant variable (Figures 7C and D). High miR-27a/low calreticulin was also associated with the development of liver metastasis and CD3+/CD8+ T cells’ infiltrates were reduced in metastases.
compared with matched primary tumours (Supplementary Figures S4C–E).

The biological relevance of these data was confirmed by two publicly available data sets.15,16 miR-27a expression remarkably increased with tumour staging and inversely correlated with patients’ overall survival, consistent with the results of our data set (Supplementary Figure S5A). Interestingly, whereas calreticulin mRNA was elevated in all data sets, the corresponding protein was reduced, a discrepancy explained by the posttranscriptional control mediated by miR-27a reported here (Supplementary Figures S6A and B; Figures 6a–c). miR-27a inversely associated also with CD3+ and CD8+ T cells’ mRNAs from the early tumour stages and correlated with poor prognosis, supporting the results of our series (Supplementary
miR-27a represses MHC class I surface exposure

T Colangelo et al

Although further studies are required to provide mechanistic insight into the link between miR-27a and MHC class I antigen presentation and, ultimately, CD8+ T cells’ recognition and activation, our present data indicate that the miR-27a/calreticulin axis regulates MHC class I cell surface expression. miR-27a upregulation occurs from the very early phases of colorectal tumourigenesis and persists throughout the progression accounting for a more aggressive development. In line, CRCs expressing the combination high miR-27a/low calreticulin are associated with reduced CD3+/CD8+ T cells’ infiltrates and cytotoxic activity, a more aggressive behaviour, metastatic spreading and worse outcome. The emerging scenario is that tumour and neighbouring cells, especially infiltrating immune cells, establish a specific microenvironment through an intricate network of crosstalks. miR-27a is crucial in immune cells, as it inhibits DCs maturation and T cells’ proliferation and activation, whereas induces M2b and M2c macrophage subtypes maturation.

Figure 6  miR-27a upregulation in CRC samples correlates with reduced MHC class I and calreticulin expression and CD3+ and CD8+ T cells infiltration. (a) Western blotting analysis of CRT and MHC class I molecules in a representative group of normal tissues (n=5) and CRC samples (n=9) classified according to miR-27a expression; β-Actin was used as a loading control. (b) The box plots report CRT and MHC class I levels in normal versus tumour tissues (upper) and in tumour tissues according to miR-27a levels (lower). (c) IHC analysis of MHC class I, CRT, CD3+ and CD8+ T cells infiltrates in paraffin-embedded samples of normal and CRC specimens classified according to miR-27a levels (scale bar, 50 μm). (d) Box plots report CD3+ and CD8+ T cells infiltration related to miR-27a levels. P-values were calculated by paired t-test in panels (b) and (d).

Discussion

Unveiling the full range of a microRNA’s functions is a major task as they are involved in a vast array of biological processes. In addition, they simultaneously target many genes acting in different pathways, the interactions of which generate a network that can be different in distinct contexts and cell types. By a 2DE-DIGE proteomic approach, we identify a series of proteins modulated by miR-27a implicated in MHC class I expression. Specifically, miR-27a represses MHC class I surface exposure directly targeting calreticulin, a protein involved in the quality control of the assembly of this multi-subunit complex contributing to its stability and retrieval of suboptimally assembled MHC class I molecules. Mechanistically, calreticulin is a major downstream effector of miR-27a in repressing MHC class I surface exposure, a pivotal event in eliciting an efficient immune response and tumour eradication. Defects in this process are a common means for cancer cells to evade T cells’ recognition. The in vivo data support this notion: high miR-27a-expressing tumours inversely correlate with MHC class I expression in our CRC series.

Figures S6C–E). Collectively, miR-27a acts as an oncomiRNA from the early phases of colon tumourigenesis, impairs MHC class I and calreticulin expression, correlates with CD3+/CD8+ infiltration, development of distant metastases and poorer outcome likely affecting the host antitumour immune response in vivo.
miR-27a represses MHC class I surface exposure
T Colangelo et al

**Materials and Methods**

**Cell culture.** Human colon cancer cell lines HCT116, HT29, CaCo-2, LoVo, RKO and SW480 were purchased from American Type Culture Collection (ATCC, Rockville, MD, USA). All cell lines were maintained in Dulbecco’s modified Eagle’s or RPMI 1640 medium supplemented with 10% fetal bovine serum, 2 mM L-glutamine, penicillin and streptomycin. Cells were cultured in a humidified 37 °C incubator at 5% CO₂. Human umbilical cord endothelial cells were purchased from Lonza, Allendale, NJ, USA. Cell lines and human umbilical cord endothelial cells were maintained in EGM BulletKit medium (Lonza, Allendale, NJ, USA).

**Western blotting analysis.** Protein extracts from cell lines and tissues were prepared and analysed as previously reported. Antibodies to calreticulin (ab2907), E-cadherin (BD 610405) from BD Biosciences, San Jose, CA, USA; ANXA1 (sc-7273), TRAP1 (sc-9134), GRP78 (sc-13968), anti-mouse (sc-2031) and anti-rabbit (sc-2004) were from Santa Cruz Biotechnology (Dallas, TX, USA); PPARγ (sc-7130), TAPBP (ab14098), ERP57 (ab13506) and MHC class I (ab70328) were from Abcam (Cambridge, MA, USA; PPARγ (sc-7273), TRAP1 (sc-9134), GRP78 (sc-13968), anti-mouse (sc-2031) and anti-rabbit (sc-2004) were from Santa Cruz Biotechnology (Dallas, TX, USA); ANXA1 (sc-7273), TRAP1 (sc-9134), GRP78 (sc-13968), anti-mouse (sc-2031) and anti-rabbit (sc-2004) were from Santa Cruz Biotechnology (Dallas, TX, USA). To analyse surface proteins, we used an extraction method based on a published procedure. Positivity for calreticulin was assessed by immunofluorescence analysis of double-stained CD8 + (red) and LAMP1 + (green) or CD8 + and perforin+ cells (scale bars: 20 μm in panels a and b, and 5 μm in panels b and d). The histograms on the right show the percentage of double-positive CD8+/LAMP1+ and CD8+/perforin+ cells in tumour tissues according to miR-27a levels **P≤0.01** (two-tailed Student’s t-test).

**Oligonucleotides and plasmids transfection.** Synthetic miR-27a mimic (Syn-hsa-miR-27a), miR-27a inhibitor (anti-hsa-miR-27a) or the appropriate scrambled controls (AllStar or mirScript Inhibitor-Negative Control) were purchased from Qiagen (Hilden, Germany). The miR-27a-antisense (MZIP27a-P A-1), the pre-miR-27a expression constructs (PMIRH27a-onlyPA-1) and scrambled control miRNAs (MZIP000-PA-1; PMIRH000PA-1) plasmids (System Biosciences, Rockville, MD, USA). All cell lines were maintained in Dulbecco’s modified Eagle’s or RPMI 1640 medium supplemented with 10% fetal bovine serum, 2 mM L-glutamine, penicillin and streptomycin. Cells were cultured in a humidified 37 °C incubator at 5% CO₂. Human umbilical cord endothelial cells were purchased from ATCC and maintained in EGM BulletKit medium (Lonza, Allendale, NJ, USA).

**mRNA/miRNA extraction and qRT-PCR analysis.** Total RNA was extracted from cells and tissues using TRizol (Thermo Fisher) and treated with DNase I. microRNAs were extracted using the Qiagen miRNaseasy Mini Kit (Qiagen) according to the manufacturer’s protocol. Assessment of RNA purity and quantity was performed as described. The sequences of the specific primers are reported in Supplementary Table S3.

**Figure 7** miR-27a is inversely associated with CD8+ T cell infiltration and activation affecting tumour aggressiveness and patients’ survival. (A) IHC analysis of miR-27a, CD8+ T cells and perforin in paraffin-embedded samples of CRC specimens classified according to different tumour stages (stages I–II versus III–IV) (scale bar, 50 μm). The histograms below report the quantification of CD8/perforin staining expressed as the mean number of positive cells in high-power fields (HPFs). **(B)** Immunofluorescence analysis of double-stained CD8+ (red) and LAMP1+ (green) or CD8+ and perforin+ cells (scale bars: 20 μm in panels a and b, and 5 μm in panels b and d). The histograms on the right show the percentage of double-positive CD8+/LAMP1+ and CD8+/perforin+ cells in tumour tissues according to miR-27a levels **P≤0.01** (two-tailed Student’s t-test).

**Immuno-fluorescence analysis of double-stained CD8+ (red) and LAMP1+ (green) or CD8+ and perforin+ cells.** The histograms below report the quantification of CD8/perforin staining expressed as the mean number of positive cells in high-power fields (HPFs). **(C)** Kaplan–Meier survival analysis of CRC patients on the basis of (C) CRT/miR-27a and (D) CD8+/miR-27a combinations. Log-rank test, P = 0.012, P = 0.096. Hazard ratios analyses are reported in the boxes on the right side.

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**Immuno-fluorescence analysis of double-stained CD8+ (red) and LAMP1+ (green) or CD8+ and perforin+ cells.** The histograms below report the quantification of CD8/perforin staining expressed as the mean number of positive cells in high-power fields (HPFs). **(C)** Kaplan–Meier survival analysis of CRC patients on the basis of (C) CRT/miR-27a and (D) CD8+/miR-27a combinations. Log-rank test, P = 0.012, P = 0.096. Hazard ratios analyses are reported in the boxes on the right side.

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2DE DIGE analysis. Differential proteome analysis on HCT116 CTRL and HCT116 miR27a_KD was performed as previously described by Milone et al. The proteomic experiments included: (a) protein preparation and labelling with DIGE dyes; (b) isoelectric focussing (IEF); (c) image acquisition, analysis and processing; and (d) protein identification using LC-MS/MS. The experimental design using the three-dye approach is illustrated in Supplementary Table S1.

Animal experiments. For xenograft generation, 20 × 10^6 CRC-derived cells (HCT116 or HT29) were subcutaneously transplanted into the flank of 20 female athymic nude mice (6–8 weeks old; Charles River, Lecco, Italy). Mice were maintained according to the United Kingdom Coordinating Committee on Cancer Research guidelines, and tumour volumes, calculated as (tumour length × width^2)/2, were monitored twice a week by caliper measurement. Two weeks after transplantation, when tumours reached the volume of 200 mm^3, mice were grouped (N = 5/group) and intratumorally injected every 7 days for 4 times with anti-miR-27a (4 nmol/mm^3) for HCT116 or with miR-27a mimic (2 nmol/mm^3) for HT29 xenograft models. In both cases, the appropriate scrambled RNAs were used (indicated as anti-miR-Ctrl and miR-Ctrl, respectively). At day 36, tumour masses were measured, excised, and further analysed: qRT-PCR was performed on RNA from xenografts to establish the efficiency of miR-27a inhibition/overexpression. This experiment was carried out in duplicate. No adverse or toxic effects were observed. All animal experiments were reviewed and approved by the Ethics Commission at Menarini Ricerche, according to the guidelines of the European Directive (2010/63/UE).

Immunofluorescence and IHC. Immunofluorescence staining was carried out on non-permeabilized HCT116 and derivative clones. Cells were plated on coverslips, fixed in para-formaldehyde (4% in PBS) at room temperature for 10 min, blocked in bovine serum albumin (3% in PBS) for 30 min before incubation with specific antibodies to MHC class I (1:100 dilution) for 1 h at room temperature. Subsequently, anti-mouse IgG-R secondary antibody (sc-2092, 1:1000 dilution) (Santa Cruz Biotechnology) was incubated for 1 h at room temperature. Coverslips were washed with PBS, stained with DAPI and, after three more washes in cold PBS, mounted in mowiol 48–88 (Merck-Millipore, Darmstadt, Germany) on glass slides. For double-immunofluorescence staining, after an initial block with 10% normal serum in PBS, tissue sections were incubated overnight at 4 °C with primary antibodies specific for CD8^+ (Clone C8/144B; Dako) and perforin (Diagnostic Biosystem) were used to identify differentially expressed miRNAs having a fold change \(N=5\)–48; stages III–IV, \(n=32\) were included in this study. Each sample was matched with the adjacent apparently normal mucosa (\(n=80\)) removed during the same surgery. Patients’ familial history and tumour classification have been reported. Patients were followed up for a median of 89.79 months or until death. All patients gave informed consent for sample collection, and study protocols were approved by the Institutional Review Board of the Fatebenefratelli Hospital in accordance with the ethical guidelines of the Declaration of Helsinki.

Clinical samples. Paraffin-embedded and liquid-nitrogen–frozen specimens from adenoma (\(n=32\)) and primary sporadic CRCs (\(n=80\)) (stages I–II, \(n=48\); stages III–IV, \(n=32\)) were included in this study. Each sample was matched with the adjacent apparently normal mucosa (\(n=80\)) removed during the same surgery. Patients’ familial history and tumour classification have been reported. Patients were followed up for a median of 89.79 months or until death. All patients gave informed consent for sample collection, and study protocols were approved by the Institutional Review Board of the Fatebenefratelli Hospital in accordance with the ethical guidelines of the Declaration of Helsinki.

Independent data sets’ analysis. The following independent, publicly available adenoma and CRC data sets, deposited in the Gene Expression Omnibus (GEO): as adenoma E-MTAB-8131 (GEOs) GSE35602 series (http://www.ncbi.nlm.nih.gov/geo), three data sets consisting of 21 patients while the data set GSE35602 counts on 59 RNA samples separately extracted from stroma and epithelium of 13 CRC tissues and four normal tissues. mRNAs expression analysis was performed on Agilent-014850 Whole Human Genome Microarray 4x44K (Agilent Technologies, Santa Clara, CA, USA); for miRNA analysis, Agilent-019118 Human miRNA Microarray 2.0 G4470B was used. Robust multichip average normalization was performed using GeneSpring 11.5 (Agilent Technologies). The information from this data set was used to identify differentially expressed miRNAs having a fold change \(\geq 2\) and \(P<0.05\), as determined by Welch–Hest statistical analysis. We performed Volcano plot analysis to visualize differential expression. TCGA COAD data set consists of 224 colorectal tumours and normal pairs. Normalized Level 3 data were used for our analysis. IPA (Ingenuity Systems, (http://www.ingenuity.com website) was used for gene set enrichment analysis and gene network analysis.

Statistics. All statistical analyses were made using Statistical Package from Social Science (SPSS; version 16.0) for Windows (SPSS Inc., Chicago, IL, USA) and R/Bioconductor (Seattle, WA, USA). Association between miRNA expression and tumour stage was assessed using Fisher exact test or Pearson \(\chi^2\) test (where indicated). The Kaplan–Meier method was used to estimate survival; log-rank test was used to test differences between the survival curves; the hazard ratios were calculated by combining the variables at 95% confidence interval to correlate the chance of events. Data are reported as means ± S.D., and mean values were compared using Student’s t-test or Mann–Whitney test. Results were considered statistically significant when \(P \leq 0.05\) was obtained.

Conflict of Interest

The authors declare no conflict of interest.

Acknowledgements. This work was partially supported by a grant from AIR (Associazione Italiana per lo studio dei linfomi e plasmacitomi) to VC, by grants from the Fondazione Italiana Sclerosi Multiples (FISM) no. 2012/R11, EU Ideas Programme, ERC-SIG ‘mentORingTregs’ no. 310496, FISR MERIT grant no. RBNE08HWLZ_015 and a CNR Medicina Personalizzata grant to GM and by a grant from Juvenile Diabetes Research Foundation (JDRF) no. 1-PNF-2015-115-5-B to
MG. MS is supported by the Phd Program in Medicina Molecolare e Biotecnologie Mediche, Università degli Studi di Napoli ‘Federico II’. We thank Professor Marco E Bianchi for generous gift of reagents, critical reading of the manuscript and discussion and to Dr. Angelo Andruilli for critical discussions and suggestions. We would like to thank Dr. Caterina Vicentini and Dr. Vincenza Guzzardo (DIMED) and Mariarosaria Montagna (DBBBM) for technical assistance.

Author contributions
TC, GP and PZ performed most of the experiments; LM, LS, AF and CV performed additional experiments and obtained more data; BP, MRM and AB contributed with the bioinformatics analysis and statistics; VC and LS designed the bulk of the experiments, supervised the entire study and wrote the manuscript along with TC.

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