Overexpression of S100A4 in human cancer cell lines resistant to methotrexate

Nuría Mencía†1, Elisabet Selga†1, Isabel Rico1, M Cristina de Almagro1, Xenia Villalobos1, Sara Ramirez1, Jaume Adan2, Jose L Hernández2, Véronique Noé1 and Carlos J Ciudad*1

Abstract

Background: Methotrexate is a chemotherapeutic drug that is used in therapy of a wide variety of cancers. The efficiency of treatment with this drug is compromised by the appearance of resistance. Combination treatments of MTX with other drugs that could modulate the expression of genes involved in MTX resistance would be an adequate strategy to prevent the development of this resistance.

Methods: The differential expression pattern between sensitive and MTX-resistant cells was determined by whole human genome microarrays and analyzed with the GeneSpring GX software package. A global comparison of all the studied cell lines was performed in order to find out differentially expressed genes in the majority of the MTX-resistant cells. S100A4 mRNA and protein levels were determined by RT-Real-Time PCR and Western blot, respectively. Functional validations of S100A4 were performed either by transfection of an expression vector for S100A4 or a siRNA against S100A4. Transfection of an expression vector encoding for β-catenin was used to inquire for the possible transcriptional regulation of S100A4 through the Wnt pathway.

Results: S100A4 is overexpressed in five out of the seven MTX-resistant cell lines studied. Ectopic overexpression of this gene in HT29 sensitive cells augmented both the intracellular and extracellular S100A4 protein levels and caused desensitization toward MTX. siRNA against S100A4 decreased the levels of this protein and caused a chemosensitization in combined treatments with MTX. β-catenin overexpression experiments support a possible involvement of the Wnt signaling pathway in S100A4 transcriptional regulation in HT29 cells.

Conclusions: S100A4 is overexpressed in many MTX-resistant cells. S100A4 overexpression decreases the sensitivity of HT29 colon cancer human cells to MTX, whereas its knockdown causes chemosensitization toward MTX. Both approaches highlight a role for S100A4 in MTX resistance.

Background

Methotrexate (MTX) is a classical drug that is used for the treatment of a wide variety of cancers, both alone and in combination with other chemotherapeutic agents [1,2]. Drug resistance is usually observed upon treatment with MTX, thus compromising its effectiveness. Combination treatments of MTX with other drugs that could modulate the expression of genes involved in MTX resistance would be an adequate strategy to prevent the development of resistance. With this premise, we searched for genes differentially expressed in common among seven MTX-resistant cell lines representative of five different types of cancer. In that work we identified and validated DHFR as the only gene overexpressed in common among all the studied cell lines [3]. We also identified and validated other genes that, aside of DHFR, played an important role in MTX resistance (AKR1C1, DKK1) [4] or that indirectly contributed to the resistance phenotype (MSH3, SSBP2, ZFYVE16) [5].

In the present report, we identified genes differentially expressed in at least 4 out of the seven cell lines. Among the genes that fulfilled this requisite, we found some genes that we had previously studied as modulators of MTX resistance, and S100A4, a gene overexpressed in five out of the seven cell lines studied.
S100A4 is a member of the S100 calcium binding protein family, which is composed of more than 20 members. Their name was given because they are soluble in 100% saturated ammonium sulfate [6]. As many of the S100 family members, S100A4 is a symmetric homodimer characterized by the presence of two calcium binding sites of the EF-hand type (helix-loop-helix) [7] that enable S100 proteins to respond to calcium stimulus induced by cell signaling.

S100A4 has been described to be involved in a wide variety of intra- and extracellular processes, such as protein phosphorylation, dynamics of cytoskeleton components or Ca2+ homeostasis, which are regulated through interaction of S100A4 with its target proteins [8,9]. Overexpression of S100A4 has been associated with tumor malignancy [10] as well as to metastasis [11], angiogenesis [12] and chemoresistance [13].

In this work, we searched for genes differentially expressed in common among cell lines resistant to MTX. We identified S100A4 as a gene overexpressed in five out of the seven MTX-resistant cell lines studied, which had been previously associated with chemotherapy resistance. Functional validations using either an expression vector encoding for S100A4 or siRNA against its RNA show a role for S100A4 in MTX resistance.

Methods

Cell Lines

Cell lines representative of five types of human cancer were used: HT29 and Caco-2 of colon cancer, MCF-7 and MDA-MB-468 of breast cancer, MIA PaCa-2 of pancreatic cancer, K562 of erythroleukemia, and Saos-2 of osteosarcoma [3]. Resistant cells were obtained in the laboratory upon incubation with stepwise concentrations of MTX (Lederle) as previously described [4]. HT29, Caco-2 and K562 resistant cells were able to grow in 10-5M MTX, which corresponds to a 1000-fold increase in resistance with respect to their respective sensitive cells; MIA PaCa-2, Saos-2, MCF-7 and MDA-MB-468 were resistant to 10-4M MTX, which represents an increase in resistance of 100 times with respect to their sensitive counterparts.

Cell Culture

Human cell lines were routinely grown in Ham’s F12 medium supplemented with 7% fetal bovine serum (FBS, both from Gibco) at 37°C in a 5% CO2 humidified atmosphere. Resistant cells were routinely grown in selective DHFR medium (GHT medium, Gibco) lacking glycine, hypoxanthine and thymidine, the final products of DHFR activity. This medium was supplemented with 7% dialyzed fetal bovine serum (GIBCO).

Global microarray data analyses of cell lines resistant to MTX

A global comparison of all cell lines was performed using GeneSpring GX v 7.3.1 (Agilent Technologies), using the latest gene annotations available (March 2009), in order to find differentially expressed genes in the majority of the resistant cell lines. The triplicate samples for each condition, sensitive and resistant, in each of the seven cell lines (42 samples in total) were imported into one single experiment. Normalization was applied in two steps: i) "per Chip normalization" by which each measurement was divided by the 50th percentile of all measurements in its array; and ii) "per Gene normalization" by which the samples of each cell line were normalized against the median of the respective sensitive cells (control). The expression of each gene was calculated as the ratio of the value obtained for each condition relative to the control condition after normalization of the data. Then, data were filtered using the control strength, a control value calculated using the Cross-Gene Error Model on replicates [14] and based on average base/proportional value. Measurements with higher control strength are relatively more precise than measurements with lower control strength. Genes that did not reach this value were discarded. Additional filtering was performed to determine the differentially expressed genes. We selected the genes that displayed a p-value corrected by false discovery rate (Benjamini and Hochberg FDR) of less than 0.05. A list was generated with the genes that passed those filters in at least four out of the seven resistant cell lines.

RT-Real-Time PCR

mRNA levels were determined by RT-Real-time PCR. Total RNA was extracted from cells using Ultraspec™ RNA reagent (Biotec) following the recommendations of the manufacturer. Complementary DNA was synthesized in a total volume of 20 μl by mixing 500 ng of total RNA, 125 ng of random hexamers (Roche), in the presence of 75 mM KCl, 3 mM MgCl2, 10 mM dithiothreitol, 20 units of RNasin (Promega), 0.5 mM dNTPs (Appli-Chem), 200 units of M-MLV reverse transcriptase (Invitrogen) and 50 mM Tris-HCl buffer, pH 8.3. The reaction mixture was incubated at 37°C for 60 min. The cDNA product was used for subsequent Real-time PCR amplification using an ABI Prism 7000 Sequence Detection System (Applied Biosystems) with 25 ng of the cDNA mixture and the assays-on-demand from Applied Biosystems (Hs00243202_m1 for S100A4 and Hs00356991_m1 for APRT).

Gene copy number determination

Genomic DNA from either sensitive or resistant cells was obtained with the Wizard™ Genomic DNA Purification Kit (Promega) following the manufacturer’s recommen-
Preparation of total extracts for Western blotting

Total extracts from cells were used to assay S100A4 protein levels. Cells were scrapped in 700 μl of ice-cold PBS and centrifuged for 10 min at 2,000 rpm on a microfuge. The supernatant was discarded and cells were resuspended in 50 μl of RIPA buffer (50 mM Tris-HCl, 150 mM NaCl, 5 mM EDTA, 1% Igepal CA-630 (Sigma), 0.5 mM PMSF, Protease inhibitor cocktail from Sigma, 1 mM NaF; pH 7.4). Cells were incubated in ice for 30 min with vortexing every 10 minutes and then centrifuged at 15,000 × g at 4°C for 10 min. Five μl of the extract were used to determine protein concentration by the Bradford assay (Bio-Rad). The extracts were frozen in liquid N₂ and stored at -80°C. Different amounts of total extracts were resolved on SDS 15%-polyacrylamide gels. After protein resolution, the gels were cut and stored at -80°C.

ELISA assay

Cells were transfected with pCMV-S100A4 as described above and cell culture media was collected 72 h after transfection. Sandwich ELISA was performed in MaxiSorb plates (NUNC), coated with a mouse anti-S100A4 antibody O/N at 4°C. Blocking was performed with 1% BSA for 1 h at 37°C. Incubation with the samples was performed for 2.5 h at 37°C, followed by incubation with a rabbit anti-S100A4 antibody. Finally, incubation with a secondary goat anti-rabbit horseradish peroxidase-conjugated antibody was performed for 1 h at 37°C. Reading of absorbance upon TMB addition was performed in a Multiskan Ascent (Thermo) plate-reading spectrophotometer at 630 nm. Background signal was determined in parallel and was used to correct the absorbance values. All analyses were performed at least in duplicate for each sample in each experiment.

Transfection of siRNAs against S100A4 RNA

HT29 cells were plated in 1 ml of -GHT medium. Transfection was performed eighteen hours later with a siRNA designed against S100A4 RNA (siS100A4) or with a siS100A4-4 MIS bearing 4 mismatches with respect to siS100A4 (underlined in the sequence below).

siS100A4: 5'-CAGGGACAACGAGGTGGAC-3'
siS100A4-4 MIS: 5'-CACCTCAACGAGGTGGAC-3'  

Cells were lipofected with the siRNAs using Lipofectamine™2000 (Invitrogen) in accordance to the manufacturer’s instructions. MTX (10⁻7M) was added 48 hours after siRNA treatment and MTT assays were performed as described previously [15] after 5 days from the treatment with the siRNA. S100A4 knockdown was monitored by determining its mRNA levels after 48 h and its protein levels after 72 h upon siRNA transfection, respectively.

The siRNAs were designed using the software iRNAi v2.1. Among the possible alternatives, sequences rich in A/T on the 3’ of the target were chosen. Then, BLAST resources in NCBI [16] were used to assess the degree of specificity of the sequence recognition for these siRNAs. We selected the siRNA that reported the target gene as the only mRNA hit. To further assess siRNA specificity for S100A4, we determined the mRNA levels of Enolase 2, Topoisomerase II, Clusterin and UGT1A7 upon incubation with siS100A4 (off-target effects) by Real-Time PCR using SYBR Green dye. See Additional file 1: Primers for off-target effects determination.

Transfection of an expression vector encoding for β-catenin

Cells were seeded into 6-well plates. Eighteen hours later, transfection with the expression plasmid for β-catenin (pcDNA3-β-catenin) kindly provided by Dr. Duñach, Universitat Autònoma de Barcelona, Spain) was performed as described for pCMV-S100A4. Cells were collected 48 h after transfection for RNA determination. S100A4 mRNA levels were monitored by RT-Real-Time PCR. The empty vector was transfected in parallel with pcDNA3-β-catenin and was used as negative control.

Transfection of an expression vector encoding for S100A4

Cells were seeded into 6-well plates in 1 ml of HAM’s F12 selective medium. Eighteen hours later, cells were transfected with the expression plasmid for S100A4 (pCMV6-XL5-S100A4, OriGene Technologies; abbreviated in the manuscript as pCMV6-S100A4), using either Lipofectamine™2000 (Invitrogen) or Fugene (Roche) following the manufacturer’s specifications. MTX (10⁻5M) was added 18 hours after transfection with the expression plasmid for β-catenin (pcDNA3-β-catenin) kindly provided by Dr. Duñach, Universitat Autònoma de Barcelona, Spain) was used as negative control. We used 25 ng of DNA and the following primers for Real-Time PCR amplification: For S100A4: 5’-CTTCTGGG-GCTGCTTATAT-3’ and 5’-ACTGGGCCT-TCTGT-TTTTCTATC-3’; and for APRT: 5’-CGGGAAC-CCTGGTCTTTGCCC-3’ and 5’-GCCTCGGG-GGCT-CAATCTCAC-3’.

Preparation of total extracts for Western blotting

Total extracts from cells were used to assay S100A4 protein levels. Cells were scrapped in 700 μl of ice-cold PBS and centrifuged for 10 min at 2,000 rpm on a microfuge. The supernatant was discarded and cells were resuspended in 50 μl of RIPA buffer (50 mM Tris-HCl, 150 mM NaCl, 5 mM EDTA, 1% Igepal CA-630 (Sigma), 0.5 mM PMSF, Protease inhibitor cocktail from Sigma, 1 mM NaF; pH 7.4). Cells were incubated in ice for 30 min with vortexing every 10 minutes and then centrifuged at 15,000 × g at 4°C for 10 min. Five μl of the extract were used to determine protein concentration by the Bradford assay (Bio-Rad). The extracts were frozen in liquid N₂ and stored at -80°C. Different amounts of total extracts were resolved on SDS 15%-polyacrylamide gels. After protein transference using a semidry electroblotter, PVDF membranes (Immobilon P, Millipore) were incubated with an antibody against S100A4 (DAKO), and detection was accomplished by secondary horseradish peroxidase-conjugated antibody and enhanced chemiluminescence, as recommended by the manufacturer (Amersham). To normalize the results, blots were re-probed with antibodies against either Actin or GAPDH (Sigma).

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ELISA assay

Cells were transfected with pCMV-S100A4 as described above and cell culture media was collected 72 h after transfection. Sandwich ELISA was performed in MaxiSorb plates (NUNC), coated with a mouse anti-S100A4 antibody O/N at 4°C. Blocking was performed with 1% BSA for 1 h at 37°C. Incubation with the samples was performed for 2.5 h at 37°C, followed by incubation with a rabbit anti-S100A4 antibody. Finally, incubation with a secondary goat anti-rabbit horseradish peroxidase-conjugated antibody was performed for 1 h at 37°C. Reading of absorbance upon TMB addition was performed in a Multiskan Ascent (Thermo) plate-reading spectrophotometer at 630 nm. Background signal was determined in parallel and was used to correct the absorbance values. All analyses were performed at least in duplicate for each sample in each experiment.

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HT29 cells were plated in 1 ml of -GHT medium. Transfection was performed eighteen hours later with a siRNA designed against S100A4 RNA (siS100A4) or with a siS100A4-4 MIS bearing 4 mismatches with respect to siS100A4 (underlined in the sequence below).

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Transfection of an expression vector encoding for β-catenin

Cells were seeded into 6-well plates. Eighteen hours later, transfection with the expression plasmid for β-catenin (pcDNA3-β-catenin) kindly provided by Dr. Duñach, Universitat Autònoma de Barcelona, Spain) was performed as described for pCMV-S100A4. Cells were collected 48 h after transfection for RNA determination. S100A4 mRNA levels were monitored by RT-Real-Time PCR. The empty vector was transfected in parallel with pcDNA3-β-catenin and was used as negative control.
**Results**

**S100A4 is overexpressed but not amplified in human cells resistant to methotrexate**

In previous studies, we used the HG U133 PLUS 2.0 microarrays from Affymetrix as a tool to analyze the differential gene expression between sensitive and MTX-resistant cells derived from different human cell lines representative of colon cancer (HT29 and CaCo-2), breast cancer (MCF-7 and MDA-MB-468), pancreatic cancer (MIA PaCa-2), erythroblastic leukemia (K562) and osteosarcoma (Saos-2) (GEOS series accession number GSE16648), which also contains the current data). Now we performed a global analysis of the seven cell lines to find out genes differentially expressed in at least four out of the seven cell lines. The list of these genes is provided in Table 1. Among them, S100A4 was a gene differentially expressed in five out of the seven cell lines and previously reported to have a role in chemotherapy resistance [13]. We validated S100A4 overexpression in HT29, MCF-7, MIA PaCa-2, K562 and Saos-2 resistant cell lines at the levels of mRNA and protein, and determined that S100A4 upregulation in the resistant cells was not due to changes in its gene copy number (Table 2).

**Ectopic overexpression of S100A4 desensitizes HT29 cells toward MTX**

HT29 MTX-resistant cells displayed the highest S100A4 expression values, considering both the mRNA and protein levels (Table 2). Thus, HT29 cells were selected for further studies using an expression vector for S100A4 (pCMV-S100A4). Transfection with 250 ng of this vector produced a 8-fold increase in the levels of S100A4 RNA after 48 hours (Figure 1A). The intracellular levels of S100A4 protein were also increased, reaching almost 5 fold when transfecting 1 μg of the expression plasmid (Figure 1B). The overexpression of S100A4 was associated to a decrease in the sensitivity toward MTX (Figure 1C).

**S100A4 is secreted by HT29 cells transfected with pCMV-S100A4**

Given that it had been described that S100A4 has extracellular functions [12,17], we explored S100A4 protein levels in the culture media of pCMV-S100A4 transfected cells. The results from the ELISA assays showed increased amounts of S100A4 protein in media from transfected cells (Figure 1D), thus indicating that a fraction of the protein produced upon vector transfection was secreted outside the cells.

**Knocking down S100A4 with siRNA chemosensitizes HT29 toward MTX**

We used iRNA technology to study the role of S100A4 in MTX resistance. Treatment of sensitive HT29 cells with increasing concentrations (10-100 nM) of the siRNA against S100A4 (siS100A4) showed a progressive decrease in its mRNA levels (Figure 2A), causing an 80% reduction at 100 nM. This treatment also caused a vast decrease in S100A4 protein levels (Figure 2B) and increased the sensitivity of HT29 cells toward MTX by about 50% (Figure 2C). To further assess siRNA specificity for S100A4, we determined the mRNA levels of other cellular genes upon incubation with siS100A4 (off-target effects). These experiments did not show any significant variation of the mRNA levels for Enolase 2, Topoisomerase II, Clusterin and UGT1A7 (Table 3)

Treatment with 100 nM siS100A4 was also performed in HT29 MTX-resistant cells. This approach led to a reduction of 75% in the levels of S100A4 RNA (Figure 3A) but had no effect on cell viability (Figure 3B). A 4-mismatch siRNA against S100A4 (siS100A4-4 MIS) was used as negative control, and did not cause any significant effects on S100A4 mRNA, protein levels or cell viability, in neither of the two cell lines.

**S100A4 is transcriptionally regulated by the Wnt pathway in HT29 cells**

It had been previously described that S100A4 was a target of the Wnt signaling pathway and a functional TCF binding site has been identified in its promoter sequence [18]. To investigate whether S100A4 expression was regulated through this pathway in HT29 cells, we overexpressed β-catenin in HT29 sensitive or resistant cells and quantified S100A4 mRNA levels 48 h after transfection. A 2-fold increase in S100A4 mRNA levels was observed upon transfection of 1 μg of pcDNA3–β-catenin in HT29 sensitive cells (Figure 4A) whereas no changes were obtained when transfection was performed in HT29 resistant cells (Figure 4B).

**Discussion**

The principal aim of this work was to find out genes differentially expressed in cell lines resistant to MTX representative of five different types of human cancer. In a previous report [3], we determined and compared the patterns of differential gene expression associated to MTX resistance in seven human cancer cell lines. In that work, we established that the only differentially expressed gene in common among all the cell lines studied was DHFR. In the present work, we identified genes differentially expressed in at least four out of the seven cell lines.
Table 1: Genes differentially expressed in at least four out of the seven MTX-resistant cell lines studied.

| Genbank  | Gene Name      | Description                                                                 | HT29 | Caco-2 | MCF-7 | MDA-MB-468 | MIA | PaCa-2 | KS62 | SaOs-2 |
|----------|----------------|------------------------------------------------------------------------------|------|--------|-------|------------|-----|--------|------|--------|
| U05598   | AKR1C2         | aldo-keto reductase family 1, member C2                                      | 4.6  | 10.2   | 0.03  | 31.1       | 0.02| 5.0    | NS   |         |
| AB018580 | AKR1C3         | aldo-keto reductase family 1, member C3                                      | 1.8  | 4.6    | 0.8   | 73.3       | 0.7 | 12.0   | NS   |         |
| NM_000691| ALDH3A1        | aldehyde dehydrogenase 3 family, member A1                                   | 2.9  | 3.8    | 6.6   | 11.9       | 0.4 | NS     | NS   | NS     |
| AL136912 | ATG10          | ATG10 autophagy related 10 homolog                                          | 8.7  | NS     | NS    | 1.5        | 2.1 | 10.1   | 2.0  |         |
| U36190   | CRIP2          | cysteine-rich protein 2                                                      | 2.7  | 1.3    | 1.9   | NS         | 2.2 | NS     | 0.3  |         |
| NM_001321| CSRP2          | cysteine and glycine-rich protein 2                                          | 2.0  | 0.3    | 2.7   | NS         | 2.0 | 2.9    | 0.5  |         |
| AU144855 | CYP1B1         | cytochrome P450, family 1, subfamily B, polypeptide 1                        | 3.1  | 2.3    | 0.5   | NS         | 4.2 | NS     | 0.4  |         |
| AI144299 | DHFR           | dihydrofolate reductase                                                     | 7.3  | 50.2   | 52.8  | 1.8        | 16.9| 17.8   | 8.9  |         |
| AA578546 | DHFRL1         | dihydrofolate reductase-like 1                                              | 7.0  | 10.8   | 5.5   | NS         | 3.0 | 17.5   | 2.1  |         |
| NM_012242| DKK1           | dikkopf homolog 1 (Xenopus laevis)                                          | 4.3  | 2.5    | 2.1   | NS         | 0.7 | NS     | 19.1 |         |
| AI745624 | ELL2           | elongation factor, RNA polymerase II, 2                                     | NS   | 1.7    | 0.4   | 2.6        | 8.0 | 2.3    | 2.7  |         |
| AF052094 | EPAS1          | endothelial PAS domain protein 1                                            | 2.8  | NS     | 2.4   | NS         | 6.4 | 0.4    | 0.1  |         |
| AU144565 | EPB41L4A       | erythrocyte membrane protein band 4.1 like 4A                              | 1.6  | 1.3    | 1.7   | 1.4        | 2.3 | 4.4    | 4.7  |         |
| AW246673 | FAM46A         | family with sequence similarity 46, member A                                | NS   | 2.6    | NS    | NS         | 3.9 | 2.5    | 1.5  |         |
| NM_000142| FGFR3          | fibroblast growth factor receptor 3                                         | NS   | 4.6    | 3.1   | 0.4        | 1.5 | 0.4    | 3.4  |         |
| NM_018071| FLJ10357       | hypothetical protein FLJ10357                                               | NS   | 3.8    | NS    | NS         | 2.1 | 3.2    | 3.5  |         |
| BE550452 | HOMER1         | homer homolog 1 (Drosophila)                                                | NS   | NS     | 27.9  | NS         | 2.4 | 13.8   | NS   |         |
| AF243527 | KLK5           | kallikrein-related peptidase 5                                             | 2.3  | NS     | 2.2   | NS         | 3.5 | NS     | 4.6  |         |
| AK022625 | LOC92270       | V-type proton ATPase subunit S1-like protein                                | 54.0 | 2.3    | NS    | NS         | 3.4 | 25.0   | NS   |         |
| BC006471 | MLLT11         | myeloid/lymphoid or mixed-lineage leukemia                                   | 2.2  | 2.6    | NS    | NS         | 1.8 | 0.08   | 1.7  |         |
| NM_002439| MSH3           | mutS homolog 3 (E. coli)                                                    | 6.3  | 8.2    | 3.8   | NS         | 13.2| 35.9   | 1.5  |         |
| NM_006117| PECI           | peroxisomal D3, D2-enoyl-CoA isomerase                                      | 7.5  | 2.6    | 1.9   | NS         | NS | NS     | 1.8  |         |
studied. Among the genes that fulfilled this requisite, we found some genes that we had previously validated and reported to be directly associated with MTX resistance in HT29 MTX-resistant cells (DHFR, AKR1C1 and DKK1) or suggested to contribute to the phenotype in this cell line (MSH3, SSBP2 and ZFYVE16) [3-5]. We also found UGT1A6, a gene that seems to play a role in MTX resistance in MCF-7 and in MDA-MB-468 cells (data not shown). Among the other genes in the list, we selected, for expression, genomic and functional validation, S100A4, a gene overexpressed in five out of these seven cell lines that was the only one previously reported to be associated with chemoresistance [13]. This gene has also been related with malignancy, metastasis, and tumor progression.

The human S100A4 is a member of the S100 family of EF-hand Ca²⁺-binding proteins. The multigenic family of S100 proteins is expressed in vertebrates exclusively and has been described to play intracellular and extracellular regulatory activities on protein phosphorylation, on the dynamics of cytoskeleton components or on Ca²⁺ homeostasis [8,9]. S100A4 in particular has been described to have a function in cell cycle progression, in cell motility and as a modulator of intercellular adhesion and of the...
invasive properties of cells [9,19] (see Sherbet et al. for a review [20]). Several cancers, including colon cancer, breast cancer and osteosarcoma, are known to produce S100A4 [10], and particularly, HT29 cells have been described to express a substantial amount of S100A4 [21]. Overexpression of S100A4 is more frequently found in cancer cells than in normal colonic mucosa, as well as more in liver metastasis than in primary tumors [21]. A correlation between S100A4 expression levels and the invasive potential of HT29 cells has been suggested [21]. S100A4 gene overexpression has been related to the malignant potential of some tumors [10] and has been closely associated with metastasis in several human cancers, including colorectal [21], breast [22,23], ovary [24], thyroid [25], pancreatic [26], lung [27,28], esophageal [29], prostate [30] and gastric cancer [31,32].

We tried to mimic the overexpression of S100A4 in the resistant cells by transfecting HT29 sensitive cells with an expression vector encoding for S100A4. We tested that S100A4 mRNA and intracellular protein levels were significantly increased after transfection. Importantly, we could observe a reversion of the cytotoxicity caused by MTX upon transfection with S100A4 expression vector. It is worth mentioning that a moderate S100A4 overexpression has been described in a colon cancer cell line resistant to doxorubicin [33]. S100P, another member of the S100 family, has also been related to doxorubicin resistance in colon cancer [33]. Moreover, S100A4 expression has been detected in patients that had received adjuvant 5-fluorouracil, and quantitation was performed upon densitometry of the images. Values represent the mean (in fold change relative to each respective sensitive cells) of three experiments ± SE; N/D, non-determined.

### Table 2: Validation of S100A4 overexpression and copy number determination in the different cell lines.

| Cell Line | Expression |
|-----------|------------|
|           | Microarray |          |
|           | Signal Sensitive | Signal Resistant | Ratio | RT-PCR Validation | Copy-number | Protein |
| HT29      | 1876       | 6862      | 3.7   | 6.2 ± 0.2         | 1.4 ± 0.1   | 22.5 ± 1.0 |
| MCF-7     | 49         | 213       | 4.3   | 6.2 ± 0.3         | 1.5 ± 0.1   | 10.8 ± 2.2 |
| MiaPaCa-2 | 3304       | 7925      | 2.4   | 4.7 ± 0.6         | 1.2 ± 0.1   | 24.4 ± 8.1 |
| K562      | 147        | 249       | 1.5   | 3.6 ± 0.2         | 1.6 ± 0.1   | 4.0 ± 1.0  |
| Saos-2    | 137        | 345       | 2.5   | 2.2 ± 0.1         | 2.16 ± 0.07 | 2 ± 0.4   |
| CaCo-2    | 4515       | 2769      | 0.6   | N/D                | N/D         | N/D       |
| MDA-MB-468| 317        | 235       | 0.7   | N/D                | N/D         | N/D       |

The table shows the S100A4 expression levels obtained in the microarray experiments for all the cell lines studied. The level of expression is presented as the mean of the signal value in sensitive and resistant cells, as well as the ratio between them (fold change). Gene overexpression was validated in five of the seven cell lines by RT-Real-Time PCR. S100A4 gene copy number was determined by Real-Time PCR. Protein levels were determined by Western blot using a specific antibody, and quantitation was performed upon densitometry of the images. Values represent the mean (in fold change relative to each respective sensitive cells) of three experiments ± SE; N/D, non-determined.
Figure 1 Effects on S100A4 expression and MTX sensitivity upon pCMV-S100A4 transfection of HT29 cells. A) mRNA levels of S100A4 determined by RT-Real-Time PCR 48 h after treatment of HT29 cells (30,000) with 250 ng of the expression vector for S100A4 (pCMV-S100A4). B) A representative image of the intracellular protein levels of S100A4 determined by Western Blotting 72 h after ectopic transfection with its expression vector is shown in the upper panel, and the quantification of the blots is shown in the lower panel. Purified S100A4 protein was used as a reference marker (Abnova; first lane). An additional panel showing endogenous S100A4 protein levels in HT29 sensitive (S) and resistant (R) cells is also provided. C) Effects of S100A4 overexpression on cell viability. HT29 cells (100,000) were treated with 1 μg of pCMV-S100A4 and 5 × 10⁻⁸ M MTX was added 48 h later. Cell viability was assessed by the MTT assay six days after MTX treatment. D) Extracellular S100A4 protein levels quantified by ELISA 72 h after S100A4 overexpression upon pCMV-S100A4 transfection. The expression and viability results are expressed as percentages referred to the untreated cells. Values are the mean of three independent experiments ± SE. *p < 0.05, **p < 0.01, *** p < 0.001.
ity of osteocarcinoma cells to undergo apoptosis [36] and reverses the metastatic potential of osteosarcoma [37] and lung carcinoma cells [28]. These reports together with our results using either the expression vector or siRNA technology, provide evidence that S100A4 acts as a pro-survival factor that contributes, together with DHFR, AKR1C1 and DKK1 (see above), to chemoresistance in HT29 MTX-resistant cells. S100A4 could be considered as biomarker of resistance in the sense that it is overexpressed in the majority of the resistant cell lines analyzed, and its determination is easy to perform. In addition it could be considered as a drug target in the category of resistance modulator using inhibitors of S100A4 as an adjuvant in MTX treatment.

In an attempt to determine the mechanism for the overexpression of S100A4 in HT29 resistant cells, we determined S100A4 copy number in both HT29 cell lines. Although it has been previously described that the chromosomal region where the gene is located (1q21) is amplified in breast cancer [38,39] and in tumor tissue from osteosarcoma patients [40], we did not observe copy number changes for S100A4 in any of the cell lines studied. Other mechanisms have been proposed that attempt to explain the mechanism for S100A4 overexpression: i) by hypomethylation, since increased S100A4 expression levels have been associated with decreased methylation of the first intron of this gene in cell lines and carcinomas of pancreatic origin [26] and of the second intron in colon cancer cell lines [41]; and ii) through the Wnt signaling pathway, given that it has been shown that S100A4 is a target of this pathway in colon cancer and a functional TCF binding site has been identified in its promoter sequence [18]. In this sense, we had previously proposed the activation of the Wnt/β-catenin pathway to be an important step in MTX resistance in HT29 colon cancer cells [5]. In this cell line, E-cadherin is lost at the chromosome level and underexpressed, thus allowing β-catenin to play its function in gene transcription. In this direction, an inverse correlation has been established between the expression levels of S100A4 and E-Cadherin [42] and has been associated with poor differentiation of cancer cells [31]. Moreover, transfection of an E-Cadherin expression vector has been reported to cause a decrease in S100A4 expression levels [43]. To further elucidate if the Wnt signaling pathway could be responsible for the increased expression levels of S100A4 in the resistant cells, we overexpressed β-catenin and determined

Table 3: Assessment of siS100A4 specificity.

| Gene        | S100A4 | Enolase 2 | Topoisomerase II | Clusterin | UGT1A7 |
|-------------|--------|-----------|------------------|-----------|--------|
| Control     | 100    | 100       | 100              | 100       | 100    |
| siRNA       | 18.6 ± 6 | 135.8 ± 21.6 | 127.5 ± 6.4     | 133.9 ± 14.4 | 159.5 ± 28.4 |
| S100A4      |        |           |                  |           |        |

HT29 cells were transfected with 100 nM siS100A4. Forty-eight hours later, the mRNA levels of S100A4 and four unrelated genes were determined by RT-Real-Time PCR. Data represent the mean of three experiments ± SE.

Figure 2 Effects on S100A4 expression and MTX sensitivity upon siS100A4 transfection of HT29 cells. A) HT29 cells (30,000) were transfected with siS100A4 as described in Methods. Total RNA was extracted after 48 h and S100A4 mRNA levels were determined by RT-Real-Time PCR. B) S100A4 protein levels were determined by Western Blotting 72 h after transfection, using specific antibodies against S100A4 and Actin to normalize the results. C) Chemosensitization assays toward methotrexate: cells were treated with siS100A4 for 48 h and then incubated with MTX. Cell viability was determined 3 days after MTX treatment. The expression and viability results are expressed as percentages referred to the untreated cells. Values are the mean of three independent experiments ± SE. A representative image of Western Blots is presented. *p < 0.05, ** p < 0.01, *** p < 0.001.
S100A4 mRNA levels. The resistant cells, which already have the Wnt/β-catenin pathway highly activated, do not respond to β-catenin overexpression. However, the results obtained in HT29 sensitive cells showing a two-fold increase in S100A4 mRNA expression upon β-catenin transfection give support to a Wnt/β-catenin pathway-mediated S100A4 transcription. Another target of the Wnt signaling pathway that had been previously related with MTX resistance is P-glycoprotein [44,45]. However, no changes in its expression levels were evidenced by our microarray results in HT29 cells (data not shown).

Although many biological functions have been attributed to S100A4, the exact molecular mechanisms by
which S100A4 exerts these functions have not been fully elucidated [46]. Extracellular S100A4 has been related to metastasis and angiogenesis [17,47,48] and has been reported to activate NF-κB signaling. S100A4-mediated activation of NF-κB has been proposed to repress the expression of BNIP3, a pro-apoptotic member of the Bcl-2 family, leading to decreased apoptosis [13] and increased resistance to chemotherapy [49]. Our ELISA experiments demonstrated that S100A4 ectopic overexpression in the sensitive cells leads to the secretion of a fraction of the synthesized protein. Secretion of S100A4 by tumor cells has been previously demonstrated in vitro [12]. One can hypothesize that secreted S100A4 would contribute to resistance through BNIIP3 regulation by the NF-κB pathway in HT29 cells. However, our microarray results show no changes in BNIP3 expression levels in HT29 resistant cells (data not shown), thus suggesting that this gene does not play a role in chemoresistance in this cell line.

S100A4 has been reported to interact with p53, thus interfering in p53 activation and DNA binding capacity [50]. Mutations in p53 are frequently found in colorectal cancer [51], and HT29 cells bear mutations in this gene. These mutations reduce p53-dependent responses, thus allowing the cells to continue to proliferate and therefore give rise to drug resistance [52,53]. To further establish MTX resistance, interaction between p53 and S100A4 would be necessary to reduce p53 remaining activity. The significant 2-fold decrease in p53 expression levels in HT29 resistant cells shown in our microarray results, in accordance with others [54], would lead to increased proliferation and reduced apoptosis, both of which would contribute to MTX resistance.

In summary, our results show a role for S100A4 in MTX resistance. Its overexpression in HT29 MTX-resistant cells is not due to changes in gene copy-number but to a transcriptional regulation, probably through the Wnt pathway. Cellular knockdown of S100A4 leads to chemosensitization toward MTX and ectopic gene overexpression desensitizes the cells toward this chemotherapeutic agent.

Conclusions
S100A4 was identified and confirmed as a gene overexpressed in five out of the seven MTX-resistant cell lines studied. S100A4 ectopic overexpression in HT29 cells lead to a desensitization toward MTX. Conversely, transfection experiments using interference RNA produced a chemosensitization to MTX. The results obtained in this report establish a relation between S100A4 and resistance to MTX.

Additional material

Additional file 1 Primers for off-target effects determination. The word table shows the sequences of the primers used to assess the off-target effects of transfected siS100A4 by determining the mRNA levels of Enolase 2, Topoisomerase II, Clusterin and UGT1A7 by RT-Real-Time PCR. APRT mRNA was used to normalize the results.

Abbreviations
MTX: methotrexate; DHFR: dihydrofolate reductase; S100A4: S100 calcium binding protein A4.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
NM, ES, CA, XV, IR, JA, SR, and JLH performed the experimental work. ES and CJC analyzed the data. VN and CJC designed the study, helped with data interpretation and supervised the experimental work. All authors wrote and approved the final manuscript.

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Author Details
1Department of Biochemistry and Molecular Biology, School of Pharmacy, University of Barcelona, Diagonal Avenue 643, Barcelona, Spain and 2Leitat Technological Center (Biomed Division), Helix building PCB, Barcelona, Spain

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