Abstract. A previous case report described an adrenal incidentaloma initially misdiagnosed as adrenocortical carcinoma (ACC), which was treated with mitotane. The final diagnosis was metastatic melanoma of unknown primary origin. However, the patient developed rapid disease progression after mitotane withdrawal, suggesting a protective role for mitotane in a non-adrenal-derived tumor. The aim of the present study was to determine the biological response of primary melanoma cells obtained from that patient, and that of other established melanoma and ACC cell lines, to mitotane treatment using a proliferation assay, flow cytometry, quantitative PCR and microarrays. Although mitotane inhibited the proliferation of both ACC and melanoma cells, its role in melanoma treatment appears to be limited. Flow cytometry analysis and transcriptomic studies indicated that the ACC cell line was highly responsive to mitotane treatment, while the primary melanoma cells showed a moderate response in vitro. Mitotane modified the activity of several key biological processes, including ‘mitotic nuclear division’, ‘DNA repair’, ‘angiogenesis’ and ‘negative regulation of ERK1 and ERK2 cascade’. Mitotane administration led to elevated levels of DNA double-strand breaks, necrosis and apoptosis. The present study provides a comprehensive insight into the biological response of mitotane-treated cells at the molecular level. Notably, the present findings offer new knowledge on the effects of mitotane on ACC and melanoma cells.

Introduction

Adrenocortical tumors are relatively common, with a prevalence of 3-10% in the general population (1). These tumors can be categorized into adrenocortical adenoma and adrenocortical carcinoma (ACC), which is rare, with a reported incidence of only 2 cases per 1,000,000 individuals per year (2). ACC is most common in women (55-60% of cases) in their 4th or 5th decade of life. However, ACC can affect patients at any age, including children (3). Geographical factors also appear to serve a role, with a higher incidence in certain regions of the world, such as in southern and southwestern Brazil (4). The 5-year survival rate of individuals with adrenocortical carcinoma is 50% (5).

The main drug currently approved for the treatment of ACC is mitotane (2,4'-dichlorodiphenyl)dichloroethane, 1-(2-chlorophenyl)-1-(4-chlorophenyl)-2,2-dichloroethane). The recommended therapeutic window for plasma mitotane levels is 14-20 mg/l (~50 µM) (6,7). However, the efficacy of this drug is limited due to its low pharmacokinetic properties and dose-limiting toxicity (8-10). Mitotane is an insecticide-derivative lipophilic drug that accumulates in lipoproteins, and dyslipidemia has been observed in certain mitotane-treated patients with ACC (11). Mitotane raises the concentration of cortisol-binding globulin hormone, steroid-binding globulin and thyroxine-binding globulin, and it may also impair pituitary gland function by reducing the secretion of thyroid stimulating hormone, thus leading to hypothyroidism (12). The action of mitotane on adrenal steroidogenesis has been associated with the inhibition of a number of mitochondrial cytochrome P450-dependent enzymes, including cholesterol side-chain cleavage, 11β-hydroxylase and 18β-hydroxylase, as...
well as P450-independent enzymes, such as 3β-hydroxysteroid id-dehydrogenase (13). Mitotane contributes to the induction of respiratory chain impairment, leading to decreased aspartate and increased glutamate content (14). This drug has also been shown to inhibit the expression of the voltage-dependent anion channel, a protein anchored to the outer mitochondrial membrane (15,16).

A recent report described an unusual case of a patient admitted to the Department of Endocrinology, Metabolism and Internal Medicine at Poznan University of Medical Sciences (Poznan, Poland) in September 2017 (17). The patient was initially diagnosed with ACC and treated with mitotane therapy. However, due to the unusual course of the disease, an experienced pathologist re-analyzed all the tissue samples, and noticed that all had the same immunophenotype and morphology. As a result, the initial diagnosis of ACC was changed to metastatic melanoma of unknown primary origin. The most notable aspect of that case was the rapid disease progression after mitotane withdrawal, which suggested that mitotane may play a protective and stabilizing role in non-adrenal-derived tumors (17).

Based on the aforementioned case report, the present study aimed to compare the biological response of ACC and melanoma cells after mitotane treatment. The cell proliferation rate was determined, and cytometric analysis of key processes involved in the response to cytostatic treatment [including mitochondrial membrane potential, DNA double-strand breaks (DSBs), necrosis, apoptosis and cell cycle] was performed, alongside gene expression profiling.

Materials and methods

Cell culture of metastatic melanoma in the left adrenal gland of unknown primary origin. To establish a primary culture of metastatic melanoma, a sample from a specimen for which several immunohistochemical analyses targeting melanoma markers were previously performed, as described in detail in a previous publication (17), was used.

Adrenal metastases of melanoma tumor samples obtained during surgery, were cut into several small pieces, and the fragments were further dissociated enzymatically in 25 ml DMEM F12 (Thermo Fisher Scientific, Inc.) containing 0.1% type I collagenase (cat. no. 17018029; Thermo Fisher Scientific, Inc.) for 45 min at 37°C in a water bath with intermittent mixing. After digestion, the mixture was filtered through a 70-µM sieve. Next, the tissue was centrifuged at 4°C and 300 x g for 7 min. The cell pellet was resuspended in the DMEM F12 containing 0.5% penicillin-streptomycin (cat. no. P4333; Merck KGaA) and 10% fetal bovine serum (FBS; HyClone; Cytiva). The primary cell line was cultured at 37°C in the presence of 5% CO₂ in a humidified atmosphere until reaching 70% confluence. The cells exhibited key melanoma markers (18,19): CSPG4, FN1, TYRP1, MCAM and SPP1 (selected by transcriptomic studies using microarrays) (Fig. S1).

Cell culture of commercially available cells lines. The HAC15 cell line (cat. no. CRL-3301TM; American Type Culture Collection) was cultured in a defined medium consisting of DMEM/F12 without phenol red (Thermo Fisher Scientific, Inc.), 10% Cosmic Calf Serum (HyClone; Cytiva), 1% insulin-transferrin-selenium + Premix (cat. no. 25-800-CR; Corning, Inc.) and 1% penicillin-streptomycin. Importantly, the HAC15 cell line was authenticated using STR analysis by the supplier.

The metastatic human melanoma WM266-4 cell line (Rockland Immunochemicals, Inc.) was cultured in medium consisting of high-glucose DMEM (Thermo Fisher Scientific, Inc.), 10% FBS (HyClone; Cytiva) and 1% P/S (MilliporeSigma). The wild-type TTP53 status of both commercial cell lines was previously confirmed (20,21).

HAC15, WM266-4 and primary melanoma cell treatment. These cells (2x10⁶) were incubated with mitotane (50 µM) for 24 h (37°C in the presence of 5% CO₂ in a humidified atmosphere) and then subjected to further analyses. Based on the opinion of clinicians (HK, Department of Endocrinology, Metabolism and Internal Medicine, Poznan University of Medical Sciences, Poznan, Poland; MK, General and Transplantation Surgery, Poznan University of Medical Sciences, Poznan, Poland; TW, General, Endocrinological and Gastroenterological Surgery, Poznan University of Medical Sciences, Poznan, Poland; and MareR, Department of Endocrinology, Metabolism and Internal Medicine, Poznan University of Medical Sciences, Poznan, Poland), a time point (24 h) that is both clinically and biologically important was selected for this study. Specifically, at this time point, the greatest changes in cell proliferation were observed, which should reflect dynamic changes in gene expression (22,23). Therefore, material collected after 24 h of mitotane treatment was selected for further microarray analysis.

Real-time cell analyzer (RTCA)-based cell proliferation assay. To verify the effect of a wide range of concentrations of mitotane (10, 20, 40, 60 and 80 µM) on the proliferation rate of the evaluated cells (n=6/group; 1,000 cells per well), an RTCA electrical impedance-based cell proliferation assay (xCELLigence RTCA; Roche Diagnostics GmbH) was used. The RTCA system detects fluctuations in electrical impedance on the integrated sensory electrodes, which are located at the bottom of the 16-hole slide plates of the chamber (E-Plate 16), which are covered by dividing cells. Electrical impedance is determined at 15-min intervals during the culture period (~40 h after mitotane delivery). The main RTCA parameter is the cell index (CI), which refers to the relative change in electrical impedance depending on the rate of proliferation or apoptosis of the cultured cells. The CI values were normalized to obtained a normalized cell index (NCI), which was calculated for each time point according to the following formula: NCI=CI time point/CI mitotane delivery.

The characteristic parameters of cell proliferation, including doubling time and dose-dependent rate of decrease, and the half-maximal response to mitotane, were calculated. Doubling time is the time (in h) required to double NCI (positive values) or to reduce it by half (negative values) based on the curve-fit from the first 24 h after mitotane administration. All calculations, such as R² (regression line fit), rate of decrease and slope were performed using RTCA software v.1.2.1.1002 according to the manufacturer’s protocol (Roche Diagnostics GmbH) and then visualized in the ggplot2 package of the R programming language (https://cran.r-project.org).
org/web/packages/ggplot2/index.html). The effect of mitotane on HAC15, WM266-4 and primary melanoma cell proliferation was evaluated in two independent experiments.

Flow cytometry analysis of phosphorylated at serine 139 version of histone H2AX (γH2AX) and cleaved poly (ADP-ribose) polymerase 1 (PARP-1). Cells (HAC15, WM266-4 and primary melanoma) were stained for γH2AX with Alexa Fluor® 647 Mouse anti-H2AX (cat. no. 560447; BD Biosciences) and cleaved PARP-1 with PE Mouse anti-Cleaved PARP (Asp214) (cat. no. 562253; BD Biosciences) antibodies according to the manufacturer's instructions. Briefly, 1x10^6 untreated and mitotane-treated cells were fixed and permeabilized with Cytotox/Cytperm™ Fixation/Permeabilization Solution (BD Biosciences) for 30 min at room temperature. Additional permeabilization and fixation as recommended by the manufacturer’s protocol were then performed. The fixed cells were once washed with 1 ml Perm/Wash Buffer (BD Biosciences) at room temperature for 5 min and stained with an appropriate antibody (5 µl/assay in 20 µl BD Perm/Wash Buffer) for 20 min at room temperature. Cells were then resuspended in 1 ml PBS and analyzed with a flow cytometer (CytoFlex; Beckman Coulter, Inc.). Fluorescence intensity in arbitrary units was represented in histograms, and the mean fluorescence intensity was calculated. Data were analyzed using FlowJo software v10 (FlowJo LLC). Untreated cells served as a control. The mean fluorescence intensity from three experiments was normalized to 1.

Flow cytometry analysis of necrosis and the cell cycle. Cell cycle analysis was performed using a propidium iodide (cat. no. P1304MP; Thermo Fisher Scientific, Inc.). Briefly, treated cells (1x10^6) were fixed in cold (-20°C) 70% ethanol by adding this dropwise to the cell suspension while vortexing. Next, the cells were rinsed with PBS and incubated with a mixture of 10 µl propidium iodide (1 mg/ml), 188 µl PBS and 2 µl RNAse A (10 mg/ml; Thermo Fisher Scientific, Inc.). Cells were incubated at 37°C for 30 min, and then rinsed and resuspended in 200 µl PBS. The fluorescence intensity of the sample was determined with a blue laser (488 nm) and detection filters 610/20 nm bandpass for PI. The procedure to determine necrosis was similar to that used for cell cycle analysis, with the following exceptions: i) RNAse was not added to the unfixed cells and ii) incubation with propidium iodide was performed at 4°C for 30 min.

Flow cytometry analysis of mitochondrial membrane potential. Mitochondrial membrane potential analysis was performed using JC-1 solution (cat. no. T3168; Thermo Fisher Scientific, Inc.) at a final concentration of 100 µM. The resuspended cells (1x10^6) were stained for 20 min at 37°C. At low concentrations (low mitochondrial membrane potential), JC-1 is predominantly a monomer that yields green fluorescence with emission of 530 nm. In turn, at high concentrations (high mitochondrial membrane potential), the dye aggregates yielding a red to orange colored emission (590 nm). This correlates to the FL-2 and FL-1 channels (24). The results were depicted as mean fluorescence that was subsequently normalized (the value of the control was set as 1).

Reverse transcription-quantitative PCR (RT-qPCR). Total RNA was extracted from cells with Direct-zol™ RNA MiniPrep columns (Zymo Research Corp.). Total RNA (1 µg per 20 µl reaction volume) was reverse transcribed using an iScript cDNA Synthesis Kit (Bio-Rad Laboratories, Inc.). qPCR (Initial denaturation at 95°C for 5 min; 44 cycles of denaturation at 95°C for 10 sec, amplification at 60°C for 30 sec and elongation at 72°C for 1 sec) was performed using the LightCycler® 480 Probes Master (Roche Diagnostics) and the appropriate probe for each primer (5'-3'): TTP53 forward, 5'-CTTTCCACGAGTTGACA-3' and reverse, 5'-TCTTCCATGCGACGACCC-3'; BRCA2 forward, 5'-CCTTGCAGCC TTACACCTTCT-3' and reverse, 5'-GCAGGGCGAGTA TGTTTACG-3'; RAD51 forward, 5'-ATACACTAACAG GTGGTAGAC-3' and reverse, 5'-CCCCCTTCTCTTTCTTCA GA-3'; XRCC4 forward, 5'-TGGTGAACTGAGAAAAGC ATTG-3' and reverse, 5'-TGAAGGAAACAGTCTGAGTA GA-3'; PRKDC forward, 5'-AGAGGCTGGGAGCATCAC T-3' and reverse, 5'-CACCAAGGGCTCAAAACAACA-3'; BAX forward, 5'-ATGTTTCATCTGCGCAACTCTC-3' and reverse, 5'-ATCAGTTCCGGCAGCTTCC-3'; BCL-2 forward, 5'-GCACCTGACACCTTGGAA-3' and reverse, 5'-AGC CAGGAGAATCAAACAG-3'; CDK2 forward, 5'-TGG GGAATATGGAAATAAT-3' and reverse, 5'-CAGGAC TGCTGTTGGGACATA-3'; and CDK4 forward, 5'-AACCTC TGATTGACGCTACAGTG-3' and reverse, 5'-GGTGGG GATAGTTGAGCAG-3'.

cDNA samples were analyzed for the following genes of interest: BAX, BCL-2, CDK2, CDK4, TTP53, RAD51 recombinase (RAD51), BRCA2 DNA repair associated (BRCA2), X-ray repair cross complementing 4 (XRCC4) and protein kinase DNA-activated catalytic subunit (PRKDC), and for the reference gene GAPDH (cat. no. 05-190-541-001; Roche Diagnostics). The expression level for each target gene was calculated with the 2^(-ΔΔCq) method (25). The reaction was performed in triplicate for each gene of interest. qPCR analysis was performed using a LightCycler® 96 (Roche Diagnostics).

Microarray expression analysis. Microarray analysis was conducted as previously described (26-28). The isolated RNA (50-300 ng/µl and not degraded) from melanoma and ACC cells was pooled into two samples per group, and the following groups were established: i) Untreated primary melanoma cell line; ii) mitotane-treated primary melanoma cell line; iii) untreated WM266-4 cell line; iv) mitotane-treated WM266-4 cell line; v) untreated HAC15 cell line; and vi) mitotane-treated HAC15 cell line.

Cells were treated with mitotane (final concentration, 50 µM) for 24 h. Transcription in vitro, cDNA labelling and Affymetrix fragmentation for further hybridization were performed with an Affymetrix GeneChip IVT Express Kit (Affymetrix; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. The biotin-labelled fragments were hybridized with Affymetrix Gene Chip Human Genome U219 microarrays (Affymetrix; Thermo Fisher Scientific, Inc.) together with control cDNA and oligo B2 (Affymetrix; Thermo Fisher Scientific, Inc.). Hybridization was performed with an AccuBlock™ Digital Dry Bath Hybridization Oven (Labnet International, Inc.) at 45°C for 16 h. The microarrays were then washed 3x at room temp. for 5' and stained using Affymetrix
GeneAtlas™ Fluidics Station (Affymetrix; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. The array strips were scanned using the imaging station of the GeneAtlas™ system (Affymetrix; Thermo Fisher Scientific, Inc.). Preliminary analysis of the scanned chips was performed using GeneAtlas™ operating software (Affymetrix; Thermo Fisher Scientific, Inc.). The quality of the gene expression data was verified using the quality control criteria established by the software. The obtained CEL files were imported for downstream data analysis.

**Microarray data analysis.** All analyses were performed using the Bioconductor repository (https://bioconductor.org) with the relevant Bioconductor libraries as an extension of the statistical R programming language. The robust multiarray average normalization algorithm implemented in the ‘Affy’ Bioconductor library was used for normalization, background correction and calculation of the expression values of all examined genes (29). Biological annotation was obtained from the Bioconductor ‘oligo’ library, where an annotated data frame object was merged with a normalized data set to obtain a complete gene data table (30). Differential expression and statistical assessment were determined by applying the linear models for microarray data implemented in the ‘limma’ library (31). The selection criteria for significant changes in gene expression were based on absolute fold-change >1.5 and P-value with false discovery rate (FDR) correction (adjusted P-value) of 0.05. The result of this selection was presented as a volcano plot, showing the total number of upregulated and downregulated genes affected by mitotane. The top 10 upregulated and downregulated genes are presented in tables. The raw data files were also deposited in the Gene Expression Omnibus (GEO) repository at the National Center for Biotechnology Information (http://www.ncbi.nlm.nih.gov/geo/) under the GEO accession number GSE186870.

**Assignment of differentially expressed genes to relevant Gene Ontology (GO) biological process (BP) terms.** ENTREZ ids with fold-change values of differentially expressed genes (DEGs) were subjected to GO enrichment analysis using the Database for Annotation, Visualization and Integrated Discovery (DAVID) bioinformatics tool (32). Gene symbols of DEGs were uploaded to DAVID using the ‘RDAVIDWebService’ Bioconductor library (33), where DEGs were assigned to relevant GO terms, with subsequent selection of significantly enriched GO terms from the GO-BP direct database. The P-values of the selected GO terms were corrected using Benjamini-Hochberg procedure and were described as adjusted P-values (34). Relevant GO ontological groups with adjusted P<0.05 and N/group >5 were visualized using a bubble plot. Detailed analysis of genes belonging to selected ontological groups, with their fold-change in expression, were presented as Circos plots using the ‘GOplot’ library (35).

**Gene Set Enrichment Analysis (GSEA).** GSEA was used to determine enrichment or depletion in gene expression between two biological groups within gene sets defined *a priori* (GO terms, pathways). The Kolmogorov-Smirnov statistical test was used to identify significantly enriched or depleted groups of genes (36). GSEA was conducted using the ‘fgsea’ library (37). The normalized fold-change values of all the genes in the microarray were log2-transformed. A predefined gene set from the GO-BP database (from the Molecular Signatures Database) was selected (38). Genes belonging to the selected set were ranked according to the difference in their expression level using the signal-to-noise ratio with a 10,000-fold permutation. The enrichment score was calculated for each selected gene set (39). These scores were normalized by their gene set size, and false-positives were corrected according to their FDR. Gene sets with an adjusted P<0.1 were exported to Cytoscape v.3.7.2 to generate links between significantly enriched processes in the form of an enrichment map (40). Enriched terms were clustered and annotated using the AutoAnnotate v.1.3.2 Cytoscape plugin (41).

**Identification of common mitotane-regulated genes in all experimental models.** To identify common mitotane-regulated genes, DEGs in ≥1 cell type were selected according to the aforementioned cut-off criteria, with the additional requirement that their fold-change values in other comparable groups were >1.2. This selection was carried out in the R statistical programming language. The expression values for the gene sets were transformed into z-scores, hierarchically clustered and visualized as heat maps using the heatmap library (42). To determine the BPs regulated by mitotane, enrichment analysis was performed in the relevant ontological groups from the DAVID Bioinformatics Resources 6.8 tool with the GOTE RM BP DIRECT database (https://david.ncifcrf.gov/tools.jsp). The procedure was performed as aforementioned. Relevant GO ontological groups with n/group >3 were visualized using a bubble plot. Interactions between individual genes and relevant GO terms were evaluated using Cytoscape v.3.7.2.

**Clinically significant mitotane-affected genes regulated in the studied cells.** Clinical descriptions with RNA sequencing data from 94 cases of ACC and 470 cases of skin cutaneous melanoma (SKCM) were downloaded from the public The Cancer Genome Atlas (TCGA) database using the FireBrowse server (http://gdac.broadinstitute.org/) (43). Next, the ‘voom’ algorithm from the ‘limma’ package was used for data normalization (31). Normalized data for common mitotane-regulated genes were extracted from the whole dataset. The obtained expression set was divided into two separate populations using median values as the cut-off point: Genes with expression above and below the median value were assigned to the high or low expression group, respectively. The hazard ratio (HR) value of the selected genes was calculated using Gene Expression Profiling Interactive Analysis 2 software (44). Genes with a statistically significant HR in ACC and SKCM underwent Kaplan-Meier survival analysis. Survival plots with log-rank P-value estimates were performed using the ‘survival’ R library (45) based on mortality events described in the clinical records.

**Statistical analysis.** The statistical analysis of microarray data was described in detail in previous subsections. All other data were statistically analyzed using GraphPad software v.5.0 (GraphPad Software, Inc.). All experiments were performed ≥3 times. Results are presented as the mean ± standard
deviation. Differences between two groups were analyzed using an unpaired Student's t-test (treated cells were compared with the control). P<0.05 was considered to indicate a statistically significant difference.

Results

Mitotane administered in a wide concentration range (10–80 µM) inhibits the proliferation of primary melanoma cells, and of the WM226-4 melanoma and HAC15 ACC cell lines. The effect of a wide concentration range (10–80 µM) of mitotane on the proliferation of primary melanoma, WM266-4 and HAC15 cells was determined by measuring the impedance of proliferating cells. As shown in Fig. 1A, mitotane inhibited primary melanoma, WM226-4 and HAC15 cell proliferation in a dose-dependent manner. The proliferation growth curve profile based on data collected from the first 24 h of culture after mitotane administration showed that mitotane increased the doubling time of primary melanoma cells at each of the evaluated doses (Fig. 1B). The doubling time increased as the dose increased, confirming the dose-specific effect of mitotane on the cultured cell population. HAC15 cells continued to proliferate at mitotane doses ranging from 10 to 80 µM, but required a much longer time to double the cell population compared with that of the controls. A linear regression model was then used to calculate the proliferation decrease rate. The proliferation profile of primary melanoma cells decreased the most with increasing doses (decrease rate, 97.9%), followed by that of the melanoma cell line (63.6%). A relatively small dose-dependent decrease in cell proliferation was observed in HAC15 cells (decrease rate, 16.1%) (Fig. 1C).

Mitotane at a dose of 50 µM leads to necrosis and cell cycle arrest in G1 phase in the cell lines evaluated. First, it was confirmed that the mitochondrial membrane potential decreased in both HAC15 and WM266-4 cells after 24 h of incubation with mitotane (~0.64 fold-change (P<0.05) and ~0.76 fold-change (P<0.001), respectively) (Fig. 2A). The slight changes observed in the primary melanoma cell line (~0.94 fold-change) were not statistically significant.

DSBs were determined after mitotane administration (50 µM for 24 h), and the results revealed that DSBs were most abundant in the treated HAC15 cell line (Fig. 2B). A ~40% increase in DSBs was observed in treated HAC15 cells compared with that in control cells. In the WM266-4 cell line, a statistically significant increase of almost 10% in DSBs was also observed (mean fluorescence values). By contrast, primary melanoma cells did not show changes in the formation of DSBs after 24 h of exposure to chemotherapy.

Next, the expression of the key genes involved in DNA repair as well as DNA damage response (DDR) was analyzed, including TTP53, RAD51 [homologous recombination (HR)], BRCA2 (HR), X-ray repair cross complementing 4 (XRCC4) [non-homologous end joining (NHEJ)] and protein kinase DNA-activated catalytic subunit (PRKDC) (NHEJ). However, no significant changes in TTP53 expression in any of the cells evaluated were observed. Notably, after mitotane treatment, the expression of the RAD51 gene was significantly higher in the HAC15 cell line (~1.85 fold-change) but lower in primary melanoma cells, suggesting that these cells may have distinct DDR mechanisms. BRCA2 gene expression was significantly decreased only in primary melanoma cells (P<0.05). The expression of the XRCC4 gene was significantly reduced in HAC15 cells (~0.56 fold-change). No significant changes in the expression of PRKDC were observed in any of the treated cells.

Next, the type of cell death that was most likely to be activated after mitotane treatment was determined. The results indicate that it is probable that all the cells evaluated underwent necrosis more frequently than apoptosis (based on propidium iodide and cPARP staining, respectively). In the HAC15 cell line, a ~60% increase in the number of cells undergoing necrosis was observed. The WM266-4 cell line also exhibited an elevated level of necrosis (~44% of the mean fluorescence; P<0.01). A higher level of necrosis was observed in the primary melanoma cells (~72%) compared with that observed in the WM226-4 cell line (P<0.05) (Fig. 2C). The only cell line to demonstrate an elevated level (16%) of apoptosis was HAC15 (P<0.05) (Fig. 2D). Similarly, the primary melanoma cells appeared to be less sensitive to mitotane, as evidenced by the absence of apoptosis. To confirm these results, the expression of the pro-/-antiapoptotic genes BAX and BCL-2 was investigated. In the HAC15 cell line, these genes showed an elevated expression level in treated cells (~1.5 fold-change for BAX and ~2.16 fold-change for BCL-2) compared with that of the controls. WM266-4 cells were characterized by greater expression of the BCL-2 gene (~1.61 fold-change; P<0.05). Importantly, no significant changes were observed in BAX or BCL-2 gene expression in the primary melanoma cells.

Cell cycle arrest at G1 occurred after mitotane administration in all the examined cell types, with ≤10% more cells present in the G1 phase of the cell cycle. Consequently, a smaller percentage of cells were detected in the S and G2 phases of the cell cycle. The largest changes (P<0.001) were observed in the WM266-4 cell line (Fig. 2E). These results were confirmed by RT-qPCR analysis, which was employed to analyze the expression of genes involved in the cell cycle (CDK2 and CDK4). An elevated level of CDK4 gene expression (~3.8 fold-change) was observed in the HAC15 cells, which is characteristic of the G1 phase. The expression of the CDK2 gene was also found to be significantly reduced in the HAC15 and WM266-4 cell lines, with 0.4 ~0.5 fold-change, respectively. Since CDK2 is responsible for G1/S and S/G2 transitions (46), the present findings suggest that mitotane treatment reduces the number of cells in the S and G2 phases of the cell cycle. No changes were observed in the primary melanoma cells in terms of the expression of CDK2 or CDK4. However, cell cycle analysis by flow cytometry indicated arrest of these cells in the G1 phase of the cell cycle (with a ~10% increase; Fig. 2E).

Mitotane at a dose of 50 µM significantly modulates the transcriptomic profile of primary melanoma, WM226-4 and HAC15 cells. A transcriptome study was carried out after 24 h of cell incubation with mitotane and the results of the treated cells were compared with those of the untreated (control) group. The overall transcriptome profiles are shown as volcano plots in Fig. 3A. The following DEG selection criteria were used: |Fold-change| (absolute value) >1.5 and P-value with FDR correction <0.05. According to the accepted cut-off criteria, in primary melanoma cells,
mitotane induced a significant decrease in the expression of 21 genes, while it stimulated the expression of 17 genes. In the WM266-4 cell line, 47 genes were upregulated and 229 genes downregulated. The strongest effect of mitotane on transcriptome modulation was observed in HAC15 cells, where mitotane stimulated and inhibited the expression of 466 and 411 genes, respectively.

The 10 genes with the highest and lowest fold-change values are presented in a tabular format in Fig. 3B. In primary melanoma cells, this group of genes includes aldehyde dehydrogenase 1 family member L2 (fold-change=2.42) and mitochondrial MT-RNR2-like 2 gene (fold-change=-3.25). In the WM266-4 cell line, mitotane most strongly influenced the expression of the following genes: Matrix metalloproteinase 1 (fold-change=3.25), zinc finger and BTB domain containing 20 (fold-change=-3.06) and ST8 α-N-acetyl-neuraminide α-2,8-sialyltransferase 6 (fold-change=-3.06) (belonging to the protein family that synthesizes sialyl glycoconjugates), which are involved in multidrug resistance in cancer cells (47). In the ACC HAC15
Figure 2. Effect of mitotane on mitochondrial membrane potential. (A) The black and red colors reflect the changes shown in the chart below, where the black color corresponds to the untreated control and the red color indicates changes in red fluorescence after mitotane treatment. In the WM266-4 and HAC15 cell lines, significant changes in red fluorescence were observed (P<0.001 and P<0.05, respectively). By contrast, primary melanoma cells showed no significant changes in fluorescence intensity. (B) The most significant change in DSBs (as reflected by H2AX expression) was observed in the HAC15 cell line (~40% increase; P<0.01). The WM266-4 cell line also showed a notable increase in DSBs formation (~10%; P<0.05). Primary melanoma cells showed no significant change in the number of DSBs. Analysis of the expression of genes involved in DNA damage response showed that the HAC15 cell line may promote homologous recombination (RAD51, ~1.85 fold-change). Primary melanoma cells showed decreased gene expression of BRCA2 (~0.6 fold-change) and RAD51 (~0.2 fold-change). No significant changes in gene expression of TP53 or PRKDC were observed in any of the cell types evaluated. (C) In all the investigated cell types, an elevated level of necrosis was observed: ~72% (primary melanoma), ~44% (WM266-4) and ~60% (HAC15). (D) Apoptosis was analyzed via flow cytometry detecting cleaved PARP-1. The HAC15 cell line presented significantly higher levels of pro-apoptotic markers (BAX) and antiapoptotic markers (BCL-2) compared with those of other cell types. Specifically, BAX and BCL-2 gene expression was significantly increased in HAC15 cells (~1.5 and ~2.16 fold-change, respectively). WM266-4 cell line also showed an elevated level of BCL-2 gene expression (~1.61 fold-change; P<0.05). (E) All the investigated cell types (primary melanoma, WM266-4 and HAC15) showed a higher percentage of cells in the G1 phase of the cell cycle, as follows: ~33.72 to ~38.53% in primary melanoma cells; from ~71.22 to ~81.15% in WM266-4 cells; and from ~38.95 to ~46.40% in HAC15 cells. The CDK2 and CDK4 genes showed, respectively, decreased and increased expression levels. The most notable changes were observed in the HAC15 and WM266-4 cell lines [CDK2, ~0.4 fold-change (HAC15) and ~0.5 fold-change (WM266-4); and CDK4, ~3.8 fold-change (HAC15)]. Primary melanoma cells did not show a significant change in the expression of these genes. *P<0.05, **P<0.01, ***P<0.001. DSB, DNA double-strand breaks; RAD51, RAD51 recombinase; BRCA2, breast cancer type 2 susceptibility protein; PRKDC, protein kinase, DNA-activated, catalytic subunit; PM, primary melanoma.
cell line, mitotane significantly stimulated the expression of growth differentiation factor 15 (fold-change=27.38), a major secretory protein induced by mitochondrial dysfunction (48), and DNA damage inducible transcript 3 (DDIT3; fold-change=8.07), an endoplasmic reticulum stress-induced apoptosis factor (49). In HAC15 cells, mitotane most potently inhibited the expression of genes involved in adrenal steroidogenesis, such as low-density lipoprotein receptor (fold-change=-4.97).

Mitotane at a dose of 50 µM exerts a significant effect on genes involved in the cell division in both WM266-4 and HAC15 cells. This analysis was performed separately for the previously obtained DEGs from individual comparisons using the DAVID bioinformatics tools. The results of this analysis are shown as bubble plots in Fig. 4A, which contain only those ontological groups that met the following criteria: Adjusted P<0.05 and ≥5 genes in the group. In primary melanoma cells, none of the ontological groups met these criteria; however, in these cells,
only a limited number of DEGs (21 upregulated and 17 downregulated) were used for the analysis. In the WM266-4 cell line, mitotane inhibited genes belonging to ontological groups associated with cell proliferation such as ‘GO:0007067-mitotic nuclear division’, (n=17; P=7.83x10^-6); ‘GO:0007062-sister chromatid cohesion’ (n=12; P=7.83x10^-6); ‘GO:0051301-cell division’ (n=19; P=3.11x10^-4) and ‘GO:0007069-chromosome segregation’ (n=8; P=0.002). In HAC15 cells, decreased expression was also observed in genes belonging to proliferation-related ontology groups, including: ‘GO:0008203-cholesterol metabolic process’ (n=9; P=4.14x10^-4), ‘GO:0051301-cell division’ (n=24; P=9.01x10^-5) and ‘GO:0007067-mitotic nuclear division’ (n=19; P=3.11x10^-4). In the HAC15 cell line, mitotane also stimulated the expression of genes belonging to the following ontology groups, which are key for the present study due to the fact that they are directly involved in the processes associated with the action of cytostatics: ‘GO:0007059-intrinsic apoptotic signaling pathway in response to endoplasmic reticulum stress’ (n=9; P=0.0003),
‘GO:0034976~response to endoplasmic reticulum stress’ (n=10; P=0.009) and ‘GO:0070373~negative regulation of ERK1 and ERK2 cascade’ (n=8; P=0.04). A detailed analysis of genes assigned to several ontology groups is shown in two circos plots in Fig. 4B. Several of these genes exhibited expression changes above the cut-off threshold in the two comparison groups, including DDIT3 (primary melanoma cells: Fold-change=2.22; HAC15 cells: Fold-change=8.1); kinesin family member 14 (KIF14; WM266-4 cells: Fold-change=1.83; HAC15 cells: Fold-change=1.64); cyclin A2 (CCNA2; WM266-4 cells: Fold-change=1.84; HAC15 cells: Fold-change=1.62); cyclin dependent kinase 1 (CDK1; WM266-4 cells: Fold-change=1.76; HAC15 cells: Fold-change=1.51) and centromere protein I (CENPI; WM266-4 cells: Fold-change=1.63; HAC15 cells: Fold-change=1.74).

Only a relatively limited number of genes involved in cell proliferation and apoptosis were regulated in a similar manner by mitotane (regardless of the cell type). Consequently, an alternative bioinformatic evaluation of transcriptome modulation was performed by using GSEA. This approach was based on full transcriptomic profile analysis regardless of the predefined cut-off criteria (fold-change >2, P<0.05). In this approach, genes pre-ranked by logarithmic fold-change values were employed to determine enrichment [positive normalized enrichment score (NES)] or depletion (negative NES) in the GO-BP database after mitotane treatment.

In primary melanoma cells (Fig. 5A), mitotane treatment led, inter alia, to the enrichment of genes involved in the regulation of mitochondrial gene expression (‘mitochondrial gene expression’, NES=1.96; ‘mitochondrial translocation’, NES=2.02), as well as genes responsible for electron transport (‘respiratory electron transport’, NES=2; ATP synthesis coupled electron transport, NES=1.98).

The largest number of significantly regulated ontology groups obtained with GSEA analysis was observed in the WM266-4 cell line; thus, only the names of ontology term clusters obtained by using the AutoAnnotate Cytoscape plugin are shown in Fig. 5B. The highest absolute NES values were as follows: ‘Recombinational repair’ (NES=-2.73); ‘double strand break repair’ (NES=-2.65), ‘centromere complex assembly’ (NES=-2.64) and ‘mitotic sister chromatid segregation’ (NES=-2.63).

Despite using a different methodological approach in HAC15 cells, the GSEA analysis revealed relatively similar groups to those shown after the analysis of ontological groups (DAVID), with enrichment of genes belonging to apoptosis-related ontological groups, including ‘intrinsic apoptotic signaling pathway in response to endoplasmic reticulum stress’ (NES=2.1) and ‘positive regulation of response to endoplasmic reticulum stress’ (NES=2.19).

In HAC15 cells, genes belonging to proliferation and steroidogenesis-related ontology terms were depleted. The following groups had the highest absolute NES values: ‘Anaphase promoting complex dependent catabolic process’ (NES=-2.45), ‘sterol biosynthetic process’ (NES=-2.24), ‘nucleosome assembly’ (NES=-2.2), ‘regulation of cholesterol biosynthetic process’ (NES=-2.18) and ‘mitotic sister chromatid segregation’ (NES=-2.09) (Fig. 5C).
Since the GO analyses performed with DAVID and GSEA indicated differences between cell types, similar gene regulation under less restrictive selection conditions was evaluated. Genes for which the fold-change value for ≥1 group was above the cut-off value (|fold-change|>1.5), and the fold-change for the other groups had the same direction (positive or negative) and was within the range of 1.2 to 1.5 were selected. Fold-change values for these genes were transformed into z-scores and presented on a heatmap in Fig. 6A.

To identify the biological role of the selected genes, another GO analysis was performed using DAVID. Interactions between individual genes and relevant GO terms were evaluated with Cytoscape v.3.7.2. The largest cluster of ontology terms involved processes associated with cell division ('sister chromatid cohesion', 'cell division', 'mitotic nuclear division', 'nuclear division' and 'mitotic cytokinesis'), where most genes were downregulated (Fig. 6B). The ontological groups included the following genes: KIF20A (primary melanoma cells: Fold-change=-1.41; WM266-4 cells: Fold-change=-1.67; HAC15 cells: Fold-change=-2.1), polo like kinase 1 (PLK1; primary melanoma cells: Fold-change=-1.32; WM266-4 cells: Fold-change=-1.45; HAC15 cells: Fold-change=-1.62), cell division cycle associated 8 (CDCA8; primary melanoma cells: Fold-change=-1.22; WM266-4 cells: fold-change=-1.42; HAC15 cells: Fold-change=-1.54), CENPI (primary melanoma cells: Fold-change=-1.24; WM266-4 cells: Fold-change=-1.63; HAC15 cells: Fold-change=-1.74), shugoshin 1 (SGO1; primary melanoma cells: Fold-change=-1.23; WM266-4 cells: Fold-change=-1.66; HAC15 cells: Fold-change=-1.24), KIF18B (primary melanoma cells: Fold-change=-1.38; WM266-4 cells: Fold-change=-1.21; HAC15 cells: Fold-change=-1.7), CCNA2 (primary melanoma cells: Fold-change=-1.23; WM266-4 cells: Fold-change=-1.83; HAC15 cells: Fold-change=-1.62), Rho/Rac guanine nucleotide exchange factor 2 (primary melanoma cells: Fold-change=-1.24; WM266-4 cells: Fold-change=-1.39; HAC15 cells: Fold-change=3.3) and cell division cycle-associated 3 (primary melanoma cells: Fold-change=-1.2; WM266-4 cells: Fold-change=-1.27; HAC15 cells: Fold-change=-2.5) (Fig. 6C).

The hypothesis that there may be a link between the expression of a potential biomarker and disease progression was verified using expression data from TCGA database (94 and 470 cases of ACC and SKCM, respectively), which were analyzed for the predictive significance of genes commonly regulated by mitotane. Fig. 7 shows genes for which elevated expression may be a negative predictor of tumour progression for both ACC and SKCM. These genes include CCNA2, CDCA3, CDCA8, PLK1, CENPI and SGO1.

Discussion

In our previous study (17), a patient was diagnosed with metastatic melanoma in the left adrenal gland of unknown primary origin and received mitotane treatment. On follow-up, there was no sign of recurrence in the adrenal bed. At the 3-month follow-up after mitotane withdrawal, multiple metastases were identified. Thus, in the present study, experiments were conducted on an in vitro model using HAC15 (adrenal...
The current study had three main aims: i) To evaluate the impact of mitotane on the proliferation rate of the investigated cells; ii) to examine the expression of a wide panel of proteins and genes involved in DDR mechanisms; and iii) to investigate the gene expression profile of the treated cells.

The HAC15 cell line constitutes a suitable model for defining the molecular mechanisms regulating aldosterone and cortisol production, and can be applied to studies on normal adrenal cell function or ACC (50). Furthermore, unlike the commonly used NCI-H295R cell line, the HAC15 cell line responds to stimulation with all major adrenal secretagogues such as angiotensin II, K⁺ and adrenocorticotropic hormone (ACTH), which lead to the stimulation of adrenal hormone biosynthesis (51). Thus, this cell line appears to closely reflect adrenal physiology and is widely used in research. Additionally, the present microarray results confirm the high expression of key adrenal steroidogenesis genes in the HAC15 line, as well as its susceptibility to ACTH and forskolin stimulation (26).

Poli et al. (52) cultured NCI-H295R cells in the presence of increasing concentrations of mitotane. Treatment with mitotane reduced cell numbers in a dose- and time-dependent manner, which was attributed to an inhibitory effect on cell viability and proliferation. The data showed that mitotane levels in the therapeutic window (30-50 µM) exerted a cytotoxic effect that was associated with the inhibition of cell proliferation. The present study has also demonstrated that mitotane reduces the proliferation rate of both ACC and melanoma cells (Fig. 1). This effect was dose-dependent (10-80 µM) and particularly noticeable in primary melanoma cells, where inhibition of proliferation was evident even at low concentrations of mitotane.

Seidel et al. (53) generated a mitotane-resistant HAC15 cell line, and found that resistant clones had the ability to maintain normal mitochondrial and nucleolar morphology during treatment. Resistance was attributed to, among other factors, altered intracellular lipid homeostasis and decreased steroid production (53). This finding is consistent with the present study result showing that mitochondrial integrity plays a crucial role in response to mitotane treatment (Fig. 2A).

Bikas et al. (54) examined the effect of mitotane on different histological subtypes of thyroid cancer, using the same concentration (50 µM) and time point (24 h) as those used in the present study. The study observed that treatment with mitotane promoted DSBs and activated the apoptotic process, while also reducing the mitochondrial membrane potential. Consistent with these reports, the present study also found that mitotane-treated adrenal cells presented higher levels of histone γH2AX expression and cleaved PARP-1 based on flow

Figure 7. Clinical significance of genes commonly regulated by mitotane in the survival of patients with cancer. (A) Expression results for ACC and from The Cancer Genome Atlas database. The expression level for all analyzed genes was divided into high (above median) and low (below median) expression. Those groups were used to calculate HRs. (B) Genes in which both HRs were statistically significant were subjected to Kaplan-Meier survival analysis. ACC, adrenocortical carcinoma; SKCM, skin cutaneous melanoma; HR, hazard ratio; CCNA2, cyclin A2; CDCA, cell division cycle associated; PLK1, polo like kinase 1; CENPI, centromere protein I; SGO1, shugoshin 1; TPM, transcripts per kilobase million.
cytometry reflecting DSBs and apoptosis (Fig. 2B and 2D) as well as necrosis (Fig. 2C). Cerquetti et al (55) found that mitotane interferes with the modulation of two DNA mismatch repair protein enzymes (DNA mismatch repair protein MSH1 and DNA mismatch repair protein MSH2), which form part of the mismatch DNA repair mechanism, which could explain the radiosensitizing properties of mitotane. In the current study, mitotane treatment promoted the expression of numerous genes involved in DSB repair, including BRCA2, XRCC4 and RAD51. Importantly, those genes were upregulated or downregulated according to the type of cell, indicating that ACC and melanoma cells appear to promote different DNA repair mechanisms at different levels.

It has been reported that combined treatment with mitotane and ionizing radiation (IR) induces accumulation of H295R and SW13 cells in G2 phase (56). Mitotane can enhance the cytotoxic effects of IR via the attenuation of DNA repair and interference in the activation of mitosis-promoting factor, which is mainly regulated by cyclin B1 degradation. This phenomenon may explain the defective activation of CDC2, which is involved in G2/M phase arrest, and probably induced by concurrent treatment with mitotane and IR (56). However, the present study found that mitotane administered alone triggered the accumulation of cells (ACC and melanoma) in the G1 phase of the cell cycle (Fig. 2E), leading to a smaller percentage of cells in the S and G2 phases. This finding was further confirmed by analyzing CDK2 and CDK4 gene expression, since an inverse correlation between CDK2 and CDK4 gene expression was observed, which is logical, considering that each gene is characteristic of a different phase of the cell cycle (namely, CDK2 of the G2/M transition and CDK4 of the G1 phase).

Volante et al (57) evaluated the expression of ribonucleotide reductase large subunit 1 (RRM1) in a cohort of patients with ACC, and evaluated the association of gene expression with clinical outcomes, finding that RRM1 gene expression was functionally associated with mitotane sensitivity, thus supporting a potential role for RRM1 determination as a novel molecular biomarker to predict response to adjuvant mitotane in patients with ACC. The present study analyzed the whole transcriptome of treated cells using microarray analysis (Figs. 3-6). The results revealed that primary melanoma, WM266-4 and HAC15 cells had distinct gene expression profiles. The largest changes in gene expression profiles after mitotane administration were observed in HAC15 cells (Fig. 3). Next, it was demonstrated that mitotane triggered changes in BPs involved in DDR; more specifically, it decreased cell division activity with mitotic nuclear division in adenocortical cells (Fig. 4A and B). Seidel et al (53) found that untreated resistant HAC15 cells showed significant upregulation of genes involved in apoptosis regulation and downregulation of pathways associated with steroid metabolism, regulation of the ERK cascade, apoptotic cell clearance and response to xenobiotics. In that study, mitotane treatment (70 μM) of control cells upregulated several pathways, including cell death and unfolded protein response. By contrast, pathways related to lipid homeostasis and transport were downregulated (53). In another study, GSEA of GO categories revealed reduced expression of 36 gene sets 48 h after mitotane treatment of NCI-H295R cells, reduced expression of 124 gene sets at 72 h of treatment, overexpression of 1 gene set at 48 h of treatment and overexpression of 21 gene sets at 72 h of treatment. Reduced expression of lipid biosynthesis, steroid biosynthesis, steroid metabolic process and several cell cycle categories (such as mitosis and M phase) were typically observed at 48 and 72 h post-treatment (58).

By pathway analysis of expression genomics data, Sbiera et al (16) revealed activation of endoplasmic reticulum stress and marked alteration of lipid-related genes caused by mitotane treatment in NCI-H295 cells. To the best of our knowledge, there are no studies on the gene expression profile of different types of cells treated with mitotane. Based on GSEA, the present study obtained similar results to those described by Sbiera et al (16). Mitotane caused ER stress, and reduced the sterol biosynthetic process, mitotic sister chromatid segregation and cell cycle G2 checkpoint, inter alia (Fig. 5). In all the cell types evaluated in the current study, common modifications in BPs were observed, such as activation of cellular amino acid biosynthetic process, and attenuation of cell division (Fig. 6A and B). Several genes, including CCNA2, CDC43, CDC48, PLK1, CENP1, SGOL1, KIF18B, KIF20A, anillin actin binding protein and NUF2 component of NDC80 kinetochore complex, are of particular interest, as their expression was attenuated in all cell types after mitotane treatment (Fig. 6C). These genes are all involved in the control of the cell cycle and division.

Regarding cancer survival, the present study focused on the role of the following genes, which are commonly regulated by mitotane: CCNA2, CDC43, CDC48, PLK1, CENP1 and SGOL1 (Fig. 7). Expression of these genes below the median levels correlates with higher survival rates in patients diagnosed with ACC and SKCM. The role of these selected genes in cancer prognosis has been reported. Liu et al (59) observed that PLK1 is a potential target for cancer therapy, as it plays multiple roles in the cell cycle, controlling mitotic entry and G2/M checkpoint, and coordinates centrosome and cell cycle. In addition, PLK1 also regulates spindle assembly and chromosome segregation, and facilitates DNA replication. Overexpression of this gene is associated with a poor prognosis in patients with cancer. Similarly, KIF18B and KIF20A mediate basic cell physiology through the regulation of the cell cycle, DNA replication, and biological DNA repair processes and pathways. This explains why KIF18B and KIF20A, both of which play an important role in cell cycle regulation, influence clinical outcomes in patients with lung adenocarcinoma (60). Furthermore, overexpression of tumor-related KIFs is correlated with worse outcomes in breast cancer and hepatocellular carcinoma; thus, KIFs may serve as prognostic biomarkers in these cancer types (61,62). The mRNA and protein levels of CENP1 are significantly increased in estrogen receptor (ER)-positive breast carcinoma, but not in ER-negative breast carcinoma. Well-established prognostic tests, such as Adjuvant! Online and the Nottingham Prognostic Index, suggest that the overexpression of CENP1 is a strong independent marker for a poor prognosis and poor survival in patients with ER-positive breast cancer (63). A previous study showed that CCNA2 was a significant prognostic indicator in ER-positive breast cancer progression and tamoxifen resistance (64). CCNA2 is overexpressed in numerous cancer types, which indicates its potential role in cancer transformation and progression. CCNA2 may also be involved in the processes of epithelial-mesenchymal transition and metastasis (65). Three members of the cell division cycle-associated gene family
(CDCA3, CDCA5 and CDCA8) are distinctly overexpressed in breast cancer tumors and cell lines. This overexpression is associated with a poor prognosis, with a low survival probability (66). Overall, decreased expression of these genes after mitotane treatment in the present study appears to be a reliable prognostic factor.

In conclusion, the present study aimed to clarify the mechanism by which mitotane, which is routinely used to treat ACC, affects other cell types. The current in vitro findings suggest that mitotane is not as effective in melanoma as in adrenal carcinoma. In the present study, mitotane had the greatest effect on a human ACC cell line, followed by a medium effect on an established metastatic human melanoma cell line. In the cell line derived from metastatic human melanoma, the response to mitotane was moderate. The three cell lines differed in terms of the intensity of their response to mitotane (primary metastatic melanoma cell line < established metastatic melanoma cell line < human adenocarcinoma cell line), and in their activation of different signaling pathways.

It is important to emphasize that the patient reported in the present study, who exhibited metastatic melanoma in the adrenal glands, responded well to mitotane therapy, and this real-world evidence is superior to the findings derived from in vitro experiments, including those performed in the current study. As a result, it is difficult to clarify the mechanism by which mitotane affects different cell types, and therefore, further studies are required. In conclusion, the present study provides a detailed description of the mechanisms that appear to be activated in response to mitotane treatment.

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Availability of data and materials

The microarray datasets generated and/or analyzed during the current study are available in the Gene Expression Omnibus repository at the National Center for Biotechnology Information (http://www.ncbi.nlm.nih.gov/geo/) under GEO accession number GSE186870. The remaining datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions

ES, HK and MR confirm the authenticity of all the raw data. ES was involved in the performing experiments and writing of the original draft; HK participated in conceptualization, methodology, writing, reviewing and editing; KJ performed experiments; AZ validated the data and took part in experiments concerning cell culture and proliferation analysis; DI conducted formal analysis of the data; AM was responsible for bioinformatic data analysis and participated in flow cytometry analyses; BS conducted experiments; ZK bioinformatic data analysis and was engaged in qPCR analysis; MK and TW were responsible for obtaining and handling primary material from the patient as well establishing the primary cell line; WMS validated the data and performed microarray analysis; MRuch was responsible for TCGA data analysis and cell culture; and MRuci participated in project supervision and took part in all experiments. All authors were involved in the preparation and modification of the figures and manuscript. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

The patient mentioned in the present study provided written informed consent. The current study was approved by the Bioethics Committee of Poznan University of Medical Sciences (Poznan, Poland; approval no. 255/15).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


