The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser2 phosphorylation of RNA polymerase II in cervical cancer

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Abstract
Cervical cancer is the fourth most frequently diagnosed and fatal gynecological cancer. 15–61% of all cases metastasize and develop chemoresistance, reducing the 5-year survival of cervical cancer patients to as low as 17%. Therefore, unraveling the mechanisms contributing to metastasis is critical in developing better-targeted therapies against it. Here, we have identified a novel mechanism where nuclear Caspase-8 directly interacts with and inhibits the activity of CDK9, thereby modulating RNAPII-mediated global transcription, including those of cell-migration- and cell-invasion-associated genes. Crucially, low Caspase-8 expression in cervical cancer patients leads to poor prognosis, higher CDK9 phosphorylation at Thr186, and increased RNAPII activity in cervical cancer cell lines and patient biopsies. Caspase-8 knock-out cells were also more resistant to the small-molecule CDK9 inhibitor BAY1251152 in both 2D- and 3D-culture conditions. Combining BAY1251152 with Cisplatin synergistically overcame chemoresistance of Caspase-8-deficient cervical cancer cells. Therefore, Caspase-8 expression could be a marker in chemoresistant cervical tumors, suggesting CDK9 inhibitor treatment for their sensitization to Cisplatin-based chemotherapy.

Keywords Caspase-8 • Non-apoptotic function • Transcription • CDK9 • P-TEFb • RNAPII
Introduction

Cervical cancer is the fourth most common gynecological cancer worldwide, accounting for 3.3% of all cancer-related deaths in 2018 [1]. Histologically, it is classified into Squamous Cell Carcinoma (SCC) (70%), adenocarcinoma (25%), and adeno-squamous carcinoma (3%). Rarer subtypes include neuro-endocrine or small-cell carcinomas. 99% of all cervical cancers are caused by persistent infections with the human papilloma virus (HPV) [2], whose early stages are usually asymptomatic. Only at later stages do symptoms arise, usually mistaken for other ailments. The development of Papanicolaou (PAP) smear screening and the HPV vaccine Gardasil has led to significant improvements in the early detection, treatment, and reduction of cervical cancer deaths, especially in developed countries. Treatment of cervical cancer involves surgery, radiotherapy, and platinum-based chemotherapy (Carboplatin and Cisplatin), usually in combination, depending on the cancer stage. Nevertheless, 15–61% of all cases develop into metastatic disease [2], or drug resistance [3]. Also, ~33% of patients develop tumor recurrence within 18–24 years of initial diagnosis, depending on the tumor stage [4]. If detected early, cervical cancer has a 5-year survival rate of 92%; 56% if it has spread to the surrounding organs and/or lymph nodes; and only 17% when metastasized to distant organs [5]. Accordingly, a better understanding of molecular mechanisms like cell-migration and cell-invasion, critical for metastasis [6], is essential for developing novel treatment strategies against cervical cancer.

Pro-caspase-8 (hereafter Caspase-8) is a member of the Caspase family of cysteine–aspartate-specific proteases. It is the principal initiator of the extrinsic apoptotic pathway, besides being involved in intrinsic apoptosis, anoikis, autophagy, necroptosis, and pyroptosis [7–11]. The evasion of apoptosis is a hallmark of cancer and is observed in multiple cancer entities [10, 12, 13]. Mutations that inhibit Caspase-8’s activity are relatively rare. Instead, lesions accumulating in cancers (loss-of-function mutations of the pro-apoptotic BAX and BAK1 genes, over-expressions of the anti-apoptotic Bcl-2 and Bcl-XL proteins, up-regulation of IAP family members, and post-translational modifications of Caspase-8) over their developmental phase, frequently interfere with the ability of Caspase-8 to execute apoptosis [10, 14–23]. Intriguingly, the down-regulation of Caspase-8’s activity has been shown to be beneficial for several cancers due to its involvement in non-apoptotic functions like cell-cycle regulation, proliferation, angiogenesis, and chemotherapy resistance [24–31]. For example, in neuroblastoma, head and neck squamous cell carcinomas, and triple-negative breast cancer, the down-regulation of Caspase-8 expression or its catalytic activity facilitates their cell-migration [10, 27–29]. However, the underlying molecular mechanisms are poorly understood.

Very limited information is available about the expression and functions of Caspase-8 in cervical cancer, although its activity decreases gradually as the malignancy progresses [21, 32]. Work in this direction has primarily attributed the enhanced metastasis or drug resistance in cervical cancer to the regulation of apoptosis or necroptosis in the presence or absence of Caspase-8 [32–37]. Therefore, understanding the non-apoptotic functions of Caspase-8 could help overcome cervical cancer metastasis and chemoresistance.

In the present study, we have demonstrated that Caspase-8 expression is frequently down-regulated in cervical cancer patients with high Tumor Mutational Burden (TMB), correlating significantly with poor prognosis. While investigating this mechanism, our interactome analysis revealed a novel interaction between Caspase-8 and CDK9. In cervical cancer cell lines, Caspase-8 negatively regulated the phosphorylation of CDK9 at Thr186 (pCDK9), thereby compromising its activation and the activity of the CDK9/CyclinT1 complex—P-TEFb (Positive Transcription Elongation Factor b) in phosphorylating the Ser2 residue at the C-terminal domain (CTD) of RNA polymerase II (RNAPII), thereby altering global transcription. The correlation between Caspase-8 expression and pCDK9 level was also observed in a cohort of cervical cancer primary materials. Knock-out and RNAi of Caspase-8 expression in cervical cancer cell lines enhanced cell-migration and cell-invasion by altering the RNAPII-mediated transcription of genes that regulated these functions. Work on cervical cancer cell lines revealed that Caspase-8, located in their nuclear fractions, could directly interact with and inhibit the phosphorylation of CDK9 at Thr186. Furthermore, our translational work with the small-molecule CDK9 inhibitor BAY1251152 [38] revealed that combining it with Cisplatin synergistically enhanced the sensitivity of Caspase-8 depleted cervical cancer cells to Cisplatin under both 2D- and 3D-cell culture conditions.

To our knowledge, this is the first-time Caspase-8 has been implicated in regulating the transcription of genes involved in the cell-migration and cell-invasion and contributing to chemoresistance and poor prognosis of cervical cancer patients. This likely occurs by altering RNAPII-mediated transcription elongation through the negative regulation of CDK9 autophosphorylation and its activity toward endogenous substrates.
Materials and methods

Cell culture

The HeLa, OVCAR-3, and OVCAR-8 cell lines were obtained from ATCC and cultured as per their instructions. The SiHa cells were cultured as described previously [39].

Antibodies, reagents, siRNAs, and plasmids

Antibodies and sources: CDK9, pCDK9, RNAPII, phospho-RNAPII, TGM2, Cyclin B1, Cyclin E1, and NUP98 (Cell Signaling Technology); CDC37, SPT5, Stomatatin, PLK1, Cyclin A1 (Santa Cruz Biotechnology); BRD4, Cyclin T1 and GFP (Abcam); Caspase-8 (Enzo Life Sciences); β-Actin, pCDK9, Vimentin, and Flag (Sigma-Aldrich).

Reagents and sources: CellTiter-Blue Cell Viability assay and Caspase-Glo 3/7 assay (Promega); BrdU kit (Roche); Thymidine, 5-ethyluridine (EU), Azide-fluor 488 (Sigma-Aldrich); Annexin V and 7AAD (BD); [γ-32P] ATP (3000Ci/mmol, Amersham Pharmacia); Trail, FasL (Enzo Life Sciences); PLA assay kit (Olink Biosciences); BAY1251152, Cisplatin, and Carboplatin (Selleckchem); BioCoat Matrigel invasion chamber (Corning); Migration chamber (Ibidi); RNeasy Plus kit (Qiagen), active GST-CDK9/Cyclin K (SRP5012, Sigma-Aldrich).

The following vectors were used: pCas9(BB)-2A-Puro (PX459) V2.0 (62988, Addgene); p3xFlag-CMV-7.1 (E7533, Sigma); pGEX-5X-3 (28–9545-55, GE Healthcare) and pEGFP-C2 (6083-1, Clontech). All siRNAs and primers were from Sigma-Aldrich.

CRISPR/Cas9-mediated stable knock-out of CASP8

As described earlier [17], the expression of the CASP8 gene was stably knocked-out in the HeLa cells by targeting two distinct regions within its Exon 1, using the (PX459) plasmid. The target sequences were: (1) GCCTGGACTACATTCCGCAAAGG and (2) GCCCTTCCGAATTAA TAGACTGG. Positive clones were confirmed through sequencing.

In vitro kinase assays

Both radioactive and non-radioactive in vitro kinase assays were performed as described previously [40].

Proximity Ligation Assay (PLA)

The proximity ligation assay was performed per the manufacturer’s protocol and as described previously [18].

Caspase-Glo 3/7 assay

The Caspase-Glo 3/7 assay was performed per the manufacturer’s protocol and as described previously [41].

Generation of spheroids and measurement of spheroid volumes

The generation and treatment of 3D spheroids were performed as described earlier [42].

The areas of the spheroids were measured using the ImageJ program, from which their radii were calculated. These radii were then used to determine the spheroid volumes. The following formula was used (\(A\) = area of spheroid; \(r\) = radius of spheroid; \(V\) = volume of spheroid):

\[
A = 4\pi r^2
\]

\[
r = \sqrt{\frac{A}{4\pi}}
\]

\[
V = \frac{4}{3}\pi r^3
\]

Phenotypic analysis

Cell-proliferation was measured using an CellTiter-Blue Cell Viability assay, as described earlier [41]. Briefly, the cells were seeded in 96-well flat-bottom plates, in 100 µl medium, for 24, 48, 72, 96, and 120 h. At each time point, 20 µl Resazurin reagent was added to each well, incubated at 37 °C for 3 h, and the fluorescence measured at 562/615 nm (Excitation/Emission).

Cell-cycle distribution was measured using PI-FACS measurement as described earlier [20].

2D cell-migration was assessed as described earlier [43]. Briefly, 0.1 × 10⁶ cells were seeded in 70 µl medium on each side of an ibidi migration chamber and allowed to settle down overnight. For each cell type, 3 chambers were seeded per well of a 6-well plate. The chambers were then removed, the wells flooded with medium, and the migration of the cells, between the two cell populations from each chamber, was analyzed using time-lapse brightfield microscopy. The reductions in the areas between the two populations at each time point were quantified.

3D cell invasion was assessed as described earlier [43]. Briefly, 5 × 10⁴ cells were seeded in Matrigel-coated invasion chambers in a serum-starved medium. The chambers were placed in 12-well plates, also filled with serum-starved medium. Following overnight culture, only the medium in the wells was replaced with a serum-containing medium to allow the cells from the chambers to migrate toward this area.
medium, through the Matrigel, for 24 h. The invaded cells were fixed and permeabilized, and their nuclei were stained with DAPI to be observed with fluorescence microscopy.

5-bromo-2'-deoxyuridine (BrdU) assay

The BrdU assay was performed as described previously [44] and per the manufacturer's protocol. Briefly, HeLa WT or KO cells were first synchronized with a double-thymidine block in a 96-well plate. Following the second thymidine treatment, the cells were released into a fresh medium for 0, 3, 6, and 9 h. 2 h before each time point (except 0 h), 10 µM BrdU-labeling solution was added to the wells. Next, the labeling solution was removed, and the cells were dried at 60 °C for 1 h. After denaturing the DNA of the dried cells with the kit-supplied FixDenat solution, 100 µl of an anti-BrdU antibody was added to the wells for 1.5 h at 25 °C. Finally, the antibody was removed, 100 µl of the kit-supplied substrate solution was added to the wells, and their absorbance was measured after 5 min at 370 nm.

5-ethynyl uridine (EU) assay

The EU assay was performed as described previously [45]. Briefly, cells were seeded onto coverslips and incubated overnight. The growth medium was then replaced with a fresh medium including 500 µM 5-ethynyl uridine (EU) and incubated for 24 h. The cells were then fixed, permeabilized, and stained with a staining solution containing 10 µM Azide-fluor 488 for 30 min. Finally, the cells were stained with DAPI before being observed with fluorescence microscopy.

Cytosol and nuclear fractionation

The cytosol and nuclear fractionation were performed, as mentioned earlier [46]. Briefly, harvested cells were incubated in Buffer A (10 mM HEPES, 10 mM KCl, 1.5 mM MgCl₂, 0.34 M Sucrose, 10% glycerol, 1 mM DTT, protease inhibitor cocktail, and 0.1% Triton X-100) for 5 min on ice and centrifuged at 1500×g. Cytosolic fraction was present in the supernatant. The nuclear fraction was obtained from the pellet by lysing with Buffer B (3 mM EDTA, 0.2 mM EGTA, 1 mM DTT, and protease inhibitor cocktail) for 10 min on ice and centrifuging at 2000×g.

RNA extraction and qRT-PCR

Total RNAs of cells were extracted with the EXTRACTME Total RNA Kit (Blirt), and their reverse transcription was performed using the GoScript Reverse Transcription Kit (Promega), as per the manufacturers’ instructions. All probes used for the quantitation of mRNA expression were obtained from Applied Biosystems: GAPDH (Hs02758991_g1), Vimentin (Hs00958111_m1), and E-Cadherin (Hs01023895_m1).

Transcriptomics analysis

RNA from non-synchronized (NS) and S/G2-phase synchronized (synch.) HeLa WT and KO cells were extracted using the RNeasy Plus kit (Qiagen) and subjected to microarray expression profiling using the HumanHT-12 v3 arrays (Illumina, San Diego, USA) at the Microarray Unit of DKFZ (Heidelberg, Germany).

Proteomics analysis

Global protein expression profiling was conducted as described before [47]. Briefly, non-synchronized (NS) and S/G2-phase synchronized (synch.) HeLa WT and KO cells were suspended in 500 µl lysis buffer [1% SDS/50 mM HEPES (pH 8.0) (with protease and phosphatase inhibitors)] and sonicated. After precipitation from the cleared lysates with acetone, the dried protein pellets were suspended in solubilization buffer (9 M urea, 20 mM HEPES, pH 8.0, 1 mM sodium orthovanadate, 2.5 mM sodium pyrophosphate, 1 mM beta-glycerophosphate), reduced with DTT and alkylated with iodoacetamide. Next, the protein samples were digested with Lys-C (Wako Chemicals) and trypsin (Promega), followed by a C18 peptide clean-up. Tandem Mass Tag (TMT) labeling was carried out according to the manufacturer’s instructions (Thermo Fisher Scientific), and the peptide mixtures were combined in a multiplexed sample, including one TMT channel containing a reference sample consisting of equal peptide amounts from each sample. After high-pH C18 RP kit pre-fractionation (Thermo Fisher Scientific), the samples were subjected to LC–MS/MS analysis on a Q Exactive HF Orbitrap mass spectrometer (Thermo Fisher Scientific) coupled to an UltiMate 3000 RSLC Nano HPLC system (Dionex). After clean-up and concentration on a pre-column (ReproSil-Pur 120 C18-AQ (5 µm); Dr. Maisch GmbH), the samples were separated on an analytical column (ReproSil-Pur 120 C18-AQ (1.9 µm); Dr. Maisch GmbH).

Interactome analysis

To identify Caspase-8 interacting proteins, the interactome analysis was performed as described earlier [48]. Essentially, Caspase-8 was immunoprecipitated (IP) from non-synchronized (NS) and S/G2-phase synchronized (synch.) HeLa WT cells and the co-immunoprecipitates (Co-IP) were loaded in a 12% SDS-PAGE gel and stained with Coomassie blue. IP from non-synchronized (NS) and S/G2-phase synchronized (synch.) HeLa KO cells were used as negative
control (NC-1). IgG control was also included as an additional negative control (NC-2). After Coomassie staining, the lanes in the gel were excised into individual fragments and prepared for liquid chromatography-mass spectrometric (LC-MS) analysis.

The analysis of the Caspase-8 co-IP from the HeLa WT cells initially revealed a list of ~6200 proteins. After ruling out any proteins which were also identified in NC-1 and NC-2, we narrowed down this list to ~4000 proteins. Using a Log2 FC (Fold Change) cut-off of ≥±0.5, ~1000 and ~850 proteins were identified in the non-synchronized (NS) and S/G2-phase synchronized (synch.) data sets, respectively.

**Statistical analysis**

For The Cancer Genome Atlas (TCGA) analysis, RNA-Seq and associated clinical data for CESC (cervical squamous cell carcinoma and endocervical adenocarcinoma) patients were obtained from the TCGA database (https://portal.gdc.cancer.gov/) and analyzed as described elsewhere [51]. Briefly, the RNA-Seq Expectation–Maximization (RSEM) normalized gene expressions for CASP8 for the CESC-TCGA patients were stratified by quartiles, where the 1st quartile represented the lowest, 2nd and 3rd quartiles, the intermediate and 4th quartile the highest expressions. Overall Survival (OS) was computed by considering the patient’s date of death or the last follow-up date. Progression-Free Survival (PFS) was considered when distant metastasis or the patient’s death occurred. Tumor Mutational Burden (TMB) was obtained from TCGA Research Network [52]. Kaplan–Meier survival curves and log-rank tests were calculated with IBM SPSS Statistics v.25 (IBM, Ehningen, Germany).

Illumina arrays were quantile normalized and analyzed for the transcriptomics analysis as previously described [53]. Briefly, groups were compared by t tests adjusting for unequal variances and correcting for multiple testing.

Raw MS data were processed using the MaxQuant software (version 1.6.5.0, MPI for Biochemistry) [54]. Using the Andromeda search engine, fragment ion mass spectra were searched against the UniProt human reference proteome supplemented with 245 frequently observed contaminants [55]. The FDR was set to 1% on a decoy database approach at both the peptide and protein levels. Further data was evaluated using the Perseus software (v.1.6.0.7, MPI for Biochemistry) [56]. Potential contaminants, hits to the decoy database, and proteins identified solely with modified peptides were removed. Proteins identified with less than 2 unique peptides were excluded from further analysis. The TMT reporter intensities within each LC/MS run were normalized for equal sample loading and individual measurements by scaling to an internal reference containing equal peptide amounts of each sample [57].

The Combination Index (CI) was calculated based on the cell-proliferation and colony-formation assay results, using the Chou–Talalay method, as described earlier [58], and the Compusyn v.1 software, according to the developers’ instructions and as described earlier [59].

For the IHC staining, the correlation between pCDK9 expression, Caspase-8 levels, and pathological factors was assessed by computing Spearman’s correlation coefficient. p < 0.05 was considered statistically significant in all testing. IBM SPSS Statistics v.25 was used for the statistical analyses.
Fig. 1 The effect of Caspase-8 expression on patient prognosis. A RNA-Seq data of cervical cancer patients with higher than median Tumor Mutational Burden (TMB) were obtained from the TCGA database (CESC-TCGA) and used to determine the correlation between high or low Caspase-8 expression with the Overall Survival (OS) and Progression-Free Survival (PFS) of 88 patients (upper panel). The tables show the mean OS and PFS of patients with low, intermediate, or high Caspase-8 expression, along with their 95% Confidence Interval (CI). B Correlation between high or low Caspase-8 expression with the OS and PFS of 88 patients with lower than median TMB based on their RNA-Seq data obtained from the CESC-TCGA database.
Results

Low Caspase-8 expression correlates with poor prognosis in cervical cancer patients

Cervical cancer possesses one of the highest Tumor Mutational Burden (TMB) of all cancers [60]. To determine the significance of Caspase-8 expression in the prognosis of cervical cancer patients, we split the patients from the CESC-TCGA database, based on their RNA-Seq data, into those with TMB lower or higher than the median. Surprisingly, patients with high non-synonymous somatic mutations and low CASP8 expression displayed significantly poorer OS (Overall Survival) and PFS (Progression-Free Survival) (Fig. 1A). In contrast, patients with TMB lower than the median, high CASP8 expression displayed significantly poorer OS and PFS (Fig. 1B).

In summary, these data highlighted that the correlation between low CASP8 expression and the prognosis of cervical cancer patients depends on their TMB status.

Knock-out of Caspase-8 alters cellular behaviors

To further explore the roles of Caspase-8 expression in the malignant behavior of cervical cancer cells that might contribute to poor patient prognosis, we used the CRISPR/Cas9 system to generate CASP8−/− HeLa and SiHa cells (Fig. 2A, Supplementary Fig. 1A). To avoid clonal variations [61], we mixed individual CASP8−/− knock-out clones—Clones K5, 7 and 8 for HeLa and Clones 7, 11 and 22 for SiHa, to form a mixed knock-out population (henceforth KO) and determined different biological traits of the resulting cells. We first determined the 2D cell-proliferation of the WT vs. KO cells over 120 h and observed that the KO HeLa cells proliferated at a significantly slower rate than the WT cells (Fig. 2B). Importantly, none of the individual knock-out clones displayed any significant differences in their cell-proliferations (Supplementary Fig. 2A), indicating that none of the individual clones significantly affected the differential proliferation of the KO population.

We next decided to determine whether the reduced proliferative activity of KO cells could be attributed to any alterations in their cell-cycle profiles. For this, HeLa and SiHa WT and KO cells were synchronized with a double-thymidine block, released into a fresh medium, and analyzed by FACS. We observed that the cell-cycle profile of synchronized KO cells was retarded primarily at the 6 h time point, which corresponded with the S/G2-phase of the cell-cycle, compared to that of WT cells, albeit more prominently in HeLa than in SiHa cells (Fig. 2C, Supplementary Fig. 1B). The expression of cell-cycle markers for G1/S (Cyclin E1), S/G2 (Cyclin A1) and G2/M (PLK1, Cyclin B1) confirmed this cell-cycle delay in the KO cells of both cell lines (Fig. 2D, Supplementary Fig. 1C). Furthermore, the significant increase in the BrdU signals, which relies on the incorporation of the thymidine analogue 5-bromo-2′-deoxyuridine (BrdU) into the nascently synthesized DNA strands [44], after 3 and 6 h release from double-thymidine synchronization, confirmed the cell-cycle delay observed in the HeLa KO cells (Fig. 2E).

Moreover, comparing the 2D migration and 3D invasion between the KO and WT cells demonstrated that KO cells migrated (Fig. 2F, Supplementary Fig. 1D) and invaded (Fig. 2G, Supplementary Fig. 1E) at significantly faster rates than their WT counterparts. Importantly, none of the individual knock-out clones displayed any significant differences in their cell-migrations (Supplementary Fig. 2B), indicating that none of the individual clones significantly affected the differential migration of the KO population.

Knocking-down Caspase-8 expression using siCasp8 in HeLa and SiHa WT cells (Supplementary Fig. 3A) significantly enhanced their invasiveness as compared to Empty Flag-Vector (EV) transfected or non-transfected (NT) cells (Supplementary Fig. 3B). Reversely, over-expressing a Flag-tagged-Casp8 (Flag-Casp8) in HeLa WT and KO cells significantly reduced invasion (except WT Flag-Casp8 vs. WT Empty Flag-Vector) and migration, as compared to their respective Empty Flag-Vector (EV) counterparts (Supplementary Figs. 3B, C).

In summary, these experiments demonstrated that the loss of Caspase-8 expression in HeLa and SiHa cells reduced cell-proliferation, which could be attributed to a delay in the cell-cycle progression at the S/G2 phase. In addition, the loss of Caspase-8 expression caused a significant enhancement of the 2D-migration and 3D-invasion of both cell lines. Importantly, we observed no significant increase in apoptosis in the KO cells, even after treating them with Trail (Supplementary Fig. 3D), further validating the veracity of our Caspase-8 knock-out.

CDK9 is an interacting partner of Caspase-8

The results of the previous experiments raised a fundamental question on how Caspase-8 alters the migration and invasion of cervical cancer cells. To elucidate this, we IPed Caspase-8 from non-synchronized (NS) exponentially growing and S/G2-phase synchronized (synch.) HeLa WT. The co-IPed proteins were subjected to an interactome analysis to identify Caspase-8 interacting proteins that might be involved in cell-migration and cell-invasion.

After performing the initial analysis of the non-synchronized (NS, ~1000 proteins) and S/G2-phase synchronized (synch., ~850 proteins) data sets, we compared the ratios of their Log2 FC ≥ ±0.5 values (up- and down-regulated) to obtain 551 common proteins between the NS and synch. data sets (Supplementary Figs. 4A, B). Finally, we selected 291
proteins for further analysis, which were present in the non-synchronized (NS) and S/G2-phase synchronized (synch.) data sets, with a Log2 FC ≥ +0.5 (up-regulated), representing the pool of potential Caspase-8 interacting proteins.
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We next used IPA to perform their network analysis to elucidate the correlation of Caspase-8 with proteins involved in these pathways. Interestingly, IPA predicted that Caspase-8 is not only directly involved with the cell-migration pathway (Fig. 3C, black arrow), whose constituent proteins, including Caspase-8, appear to form a tight cluster (Supplementary Fig. 4E, black arrow), but also with several other biological processes, including transcription (Fig. 3D). This data encouraged us to investigate potential correlations between cell-migration and transcription-associated proteins. Notably, IPA identified the kinase CDK9, which was involved in both the pathways, to interact with Caspase-8, albeit indirectly via the transcription factor STAT3 and the CDC37/HSP90 chaperone complex (Fig. 3E, red arrow).

In summary, the interactome analysis of Caspase-8 revealed that it could potentially interact with a wide variety of proteins, including CDK9, regulating different molecular pathways, including those involved in cell-migration. Surprisingly, in both analyses, IPA predicted Caspase-8 as a nuclear protein (Fig. 3C, E, black arrow), even though it is generally known to be a cytoplasmic protein [10, 17]. Based on these predictions and the fact that CDK9 is a predominantly nuclear protein [38], we next aimed to study the potential interaction between Caspase-8 and CDK9.

To achieve this goal, we first performed a ‘pull-down’ assay from the lysates of HeLa KO cells, using GST-Casp8-WT, -NT (N-terminal region/prodomain), -p18, and -p12 domains (making up its C-terminal region/catalytic domain) [10] (Fig. 4A) and observed that endogenous CDK9 interacted with the GST-Casp8-WT, specifically with the p12 domain (Fig. 4B). Since most cellular CDK9 is found either as part of the large negative regulatory complex—7SK snRNP, within the nucleus or with the CDC37/HSP90 chaperone complex, within the cytosol [38, 62], we next decided to identify whether the interaction between CDK9 and Caspase-8 occurred directly, or via any intermediate proteins, which could be a part of these complexes. For this purpose, we incubated a commercially available, recombinant, active P-TEFb (GST-CDK9/Cyclin K) of Baculovirus origin with recombinant GST-Casp8-WT of bacterial origin in a cell-free system and performed immunoprecipitations using specific antibodies. IP with an anti-CDK9 antibody immunoprecipitated GST-CDK9 and co-immunoprecipitated the GST-Casp8-WT, whereas the IP with an anti-Caspase-8 antibody immunoprecipitated GST-Casp8-WT and co-immunoprecipitated the GST-CDK9. This confirmed that the interaction between the two proteins occurs directly, without the involvement of any intermediate bridging proteins (Fig. 4C).

Reversibly, GST-CDK9 also pulled down endogenous Caspase-8 in both HeLa and SiHa WT cells, but, expectedly, not from their respective KO cells (Fig. 4D). Furthermore, endogenous CDK9 was also co-IPed in a Caspase-8 IP of HeLa WT cells, but expectedly, not using two KO clones.
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In summary, we have confirmed that CDK9 interacts with the p12 domain of Caspase-8 within the nucleus of cervical cancer cells. We cannot rule out any potential interaction between Caspase-8 and CDK9 in the cytosol. However, at least under our experimental conditions, this interaction in the cytosol was below the limit of detection of the Caspase-8 IP (Fig. 4J).

Caspase-8 inhibits the activity of CDK9

Having demonstrated the interaction between Caspase-8 and CDK9, the next question was about the effect of this interaction on each other. As CDK9 is a kinase, we first asked whether Caspase-8 may be a target for phosphorylation. Therefore, we performed an in vitro kinase assay with γ-32P ATP involving the active P-TEFb (GST-CDK9/Cyclin K) incubated with increasing amounts of GST-Casp8 or GST (Fig. 5A). Surprisingly, instead of P-TEFb phosphorylating Caspase-8, we observed a GST-Casp8 amount-dependent reduction of P-TEFb phosphorylation. Increasing amounts of GST, used as a control, did not influence the phosphorylation of P-TEFb (Fig. 5A, Supplementary Fig. 5A). Importantly, both GST-Casp-8 and GST did not affect the levels of non-phosphorylated GST-CDK9, as determined by immunoblot (Fig. 5A, lower panel).

The principal cellular function of P-TEFb is to mediate the productive transcriptional elongation by phosphorylating the Ser2 residue of RNAPII [38]. As we have demonstrated that Caspase-8 impedes the phosphorylation of P-TEFb in vitro (Fig. 5A, Supplementary Fig. 5A), we next sought to investigate whether Caspase-8’s interaction with P-TEFb would affect its kinase activity toward RNAPII. With this aim, we performed a similar kinase assay but also included the carboxy-terminal domain (CTD) of RNAPII as a His-tagged recombinant protein (His-CTD). Intriguingly, the presence of GST-Casp-8, but not GST, triggered an amount-dependent reduction in the phosphorylation of His-CTD (Fig. 5B, Supplementary Fig. 5B). These data suggested that Caspase-8’s interaction with CDK9 blocked P-TEFb’s kinase activity, which in turn affected the P-TEFb dependent phosphorylation of RNAPII.

Next, we were curious whether this inhibition of P-TEFb’s kinase activity also extended to other substrates. P-TEFb has been reported to phosphorylate the SPT5 subunit of DSIF ([DRB] Sensitivity-Inducing Factor), a negative regulator of transcription elongation [38]. As no commercial phosphospecific antibodies against SPT5 exist, we IPed SPT5 from HeLa WT cells and performed a similar kinase assay with P-TEFb. In line with the previous results, we observed a GST-Casp8, but not GST, amount-dependent reduction in the phosphorylation of the IPed SPT5 by P-TEFb (Fig. 5C,
A) Prodomain and Catalytic domain of Caspase 8.

B) Blot analysis of GST-Casp8 WT (81 KDa), CDK9 p42 (42 KDa), and GST-Casp8 p18 (44 KDa) and GST-Casp8 p12 (38 KDa).

C) GST-Casp8 and GST-CDK9 blots showing Caspase 8 (55/53 KDa) and CDK9 p42 (42 KDa).

D) IP-Casp8 and IB-Casp8 showing Caspase 8 (55/53 KDa) and CDK9 p42 (42 KDa).

E) IB-GFP showing GFP-Casp8 (81 KDa) and CDK9 p55 (55 KDa).

F) IB-CDK9 showing CDK9 p42 (42 KDa).

G) PLA showing CDK9, Casp8, and their co-localization.

H) Western blots showing Caspase 8 (55/53 KDa), pCDK9 p42 (42 KDa), CDK9 p42 (42 KDa), and β-Tubulin (55 KDa).

I) Western blots showing FlagCasp8 (58 KDa), CDC37 (50 KDa), CDK9 (42 KDa), and β-Actin (42 KDa).

J) Western blots showing Casp8 (55/53 KDa), GAPDH (37 KDa), and NUP98 (98 KDa).
The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser2 phosphorylation of RNAPII.

Our next objective was to specify whether this inhibition occurred at Thr186, the activation site of CDK9, and Ser2, CDK9’s phosphorylation site on RNAPII [38]. With this aim, we performed a non-radioactive kinase assay by first precipitating active CDK9, using a Cyclin T1 antibody from WT and two KO clones of HeLa cells, and incubating the precipitates with His-CTD and ATP. We observed higher levels of phospho-Ser2-His-CTD, using a specific phospho-RNAPII (Ser2) antibody, in both the KO clones as compared to the WT cells (Supplementary Fig. 6A). To verify whether Caspase-8 manifests these inhibitory effects within cells as well, we next over-expressed increasing amounts of Flag-Casp8 in HeLa KO cells and observed a Flag-Casp8 amount-dependent decrease in Ser2 phosphorylation of RNAPII (henceforth phospho-RNAPII) and Thr186 phosphorylation of CDK9 (henceforth pCDK9) (Fig. 5D, bar graph). On the other hand, down-regulating CASP8 in HeLa WT cells showed a siCasp8 concentration-dependent increase in phospho-RNAPII and pCDK9 (Fig. 5E, bar graph). Additionally, the cytoplasm and nuclear fractions of WT and KO HeLa (Fig. 5F) and SiHa (Supplementary Fig. 6B) cells also showed increased levels of pCDK9 and phospho-RNAPII in the nuclei of the KO cells as compared to the WT cells.

In summary, these experiments have confirmed Caspase-8’s role in negatively regulating the Thr186 phosphorylation of CDK9 and its kinase activity on the Ser2 phosphorylation of RNAPII within cells.

Due to phospho-RNAPII’s role in transcriptional regulation, we next investigated whether Caspase-8 expression altered overall transcription in the cells. For this, we performed an EU assay, which relies upon incorporating the uridine analogue—5-ethynyl uridine (EU) into the nascently synthesized RNA [45]. Interestingly, we observed a significant increase in the EU uptake in the KO vs. WT cells, suggesting an alteration in overall transcription in the absence of Caspase-8 expression (Fig. 5G). To further confirm Caspase-8’s role in altering overall transcription, we repeated the EU assay by expressing increasing amounts of Flag-Casp8 or siCasp8 in the KO and WT cells, respectively. We observed a significant Flag-Casp8 amount-dependent reduction of EU uptake, as indicated by the decrease in fluorescent signal, as compared to Empty Flag-Vector (EV, negative control) (Supplementary Fig. 6C). In contrast, we observed a significant siCasp8 concentration-dependent increase in fluorescent signal, as compared to siCtrl transfected WT cells (Supplementary Fig. 6D).

In summary, the EU experiments suggested that the loss of Caspase-8 expression altered the overall transcription in HeLa and SiHa cells, possibly by increasing the phosphorylation of CDK9 at Thr186 and CDK9’s targeted phosphorylation of SPT5 and RNAPII.

**Caspase-8 inhibits the phosphorylation of CDK9 in primary cervical cancer samples**

Our work so far has demonstrated a direct correlation between Caspase-8 expression and pCDK9 levels in cervical
A. GST and GST-Casp8 with + CDK9 kinase.

B. GST and GST-Casp8 with + CDK9 kinase + His CTD.

C. Input and IP: SPT5 with + CDK9 kinase.

D. HeLa KO with EV 0.5, 1, 1.5, 2, 3 µg.

E. siCasp8 with 5, 10, 20, 30, 40 nM.

F. WT and KO with pCDK9 (42 KDa), CDK9 (55 KDa), CDK9 (42 KDa), Cyclin T1 (81 KDa), phospho-RNAPII (250 KDa), RNAPII (250 KDa), BRD4 (152 KDa), NELF-A (57 KDa), Casp8 (55/53 KDa), β-Tubulin (55 KDa), and Histone H3 (15 KDa).

G. WT and KO with normalized EU intensity fold change, ****.
The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser2…

Due to the expression revealed a significant correlation with pCDK9 levels in cervical cancer patients. Low expression (≤ 6) was associated with increased or decreased pCDK9 levels and, by extension, phospho-RNAPII levels, respectively. These results are consistent with our cellular observations and strongly suggest that low Caspase-8 expression in cervical cancer patients might stimulate enhanced cell-migration and cell-invasion.

**Caspase-8 negatively regulates the transcription of cell-migration- or cell-invasion-associated genes**

Having demonstrated the altered regulation of Caspase-8 on RNAPII-mediated global transcription, our next question was whether this could lead to alterations in the transcription of genes associated with cell-migration and cell-invasion. This might explain the enhancements in these cellular behaviors in the Caspase-8 KO cells (Fig. 2F, G). To determine whether the differential transcription of genes between the two data sets, of which 10, like SLI9, were similarly over-expressed in the KO cells (Fig. 7E), and others were classified as “low” and > 6 as “high” expression. The median pCDK9 index for all patients was 2.0% (range of 0%—50%). The Weighted Score (WS) of Caspase-8 expression in primary cells derived from a cervical tumor. Immunoblotting revealed enhanced pCDK9 and phospho-RNAPII in the siCasp8-treated cells compared to siCtrl transfected cells (Fig. 6D, bar graph). In addition, the knock-down of Caspase-8 expression in cervical cancer patient-derived cells also led to a significant increase in their invasiveness compared to non-transfected (NT) or siCtrl transfected cells (Fig. 6E).

In summary, these data validated that, in cervical cancer patients, low or high Caspase-8 expression significantly correlated with increased or decreased pCDK9 levels and, by extension, phospho-RNAPII levels, respectively. These results are consistent with our cellular observations and strongly suggest that low Caspase-8 expression in cervical cancer patients might stimulate enhanced cell-migration and cell-invasion.

**Caspase-8 negatively regulates the transcription of cell-migration- or cell-invasion-associated genes**

Having demonstrated the altered regulation of Caspase-8 on RNAPII-mediated global transcription, our next question was whether this could lead to alterations in the transcription of genes associated with cell-migration and cell-invasion. This might explain the enhancements in these cellular behaviors in the Caspase-8 KO cells (Fig. 2F, G). To determine whether the differential expression of genes between the two data sets, of which 10, like SLC9A3R1, were similarly over-expressed in the KO cells (Fig. 7E), and compared to non-synchronized (NS) and S/G2-phase synchronized (synch.) KO HeLa cells (Fig. 7A). After normalization, ~300 transcripts were obtained using the cut-offs of Log10 (p value) of ≤ 1.30 and Log2 FC of ≥ ± 0.5, whose expressions were altered either in the WT [non-synchronized (NS) and S/G2-phase synchronized (synch.)] or KO [non-synchronized (NS) and S/G2-phase synchronized (synch.)] cells. The DAVID GÖTERM_BP analysis of the ~300 transcripts revealed several cell-migration associated functions (Fig. 7B, highlighted in black). To determine whether the differential mRNA expressions of these genes, due to the presence/absence of Caspase-8 expression, also translated to their protein expression, we also performed proteomics analysis of non-synchronized (NS) and S/G2-phase synchronized (synch.) KO HeLa cells (Fig. 7C). Once again, using the cut-offs of Log10 (p-value) of ≤ 1.30 and Log2 FC [non-synchronized (NS) and S/G2-phase synchronized (synch.)] of ≥ ± 0.5, we identified 60 proteins whose DAVID GÖTERM_BP analysis revealed several cell-migration associated functions (Fig. 7D, highlighted in black). Comparison of the 60 proteins and ~300 genes registered 14 common genes between the two data sets, of which 10, like STOM, SLC9A3R1 (NHERF1), were similarly over-expressed in the WT and 4, like TGM2 and EPB2, in the KO cells (Fig. 7E), whose DAVID GÖTERM_BP analysis again revealed
A high expression low expression

Casp8

pCDK9 (%)

p = 0.050

C low Casp8 high Casp8

pCDK9 (%)

p = 0.004

B

pCDK9 (%)

Casp8 weighted score

p = 0.004

D

E

Mean number of invading cells

NT siCtrl siCasp8

phospho-RNAPII (250 KDa)
Casp8 (55/53 KDa)
phospho-RNAPII (250 KDa)
RNAPII (250 KDa)
pCDK9 (42 KDa)
CDK9 (42 KDa)
GAPDH (35 KDa)

pCDK9 rel. phosphorylation

NT siCtrl siCasp8

20 µm
The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser2...

### Table 1

|               | CASP8 low | CASP8 high | p value | pCDK9 low | pCDK9 high | p value |
|---------------|-----------|------------|---------|-----------|------------|---------|
| CASP8         | n (%)     | n (%)      |         | n (%)     | n (%)      |         |
| Low (WS≤6)    | 42        | 19 (27.7)  | 0.018   | 23        | 33.3       |         |
| High (WS>6)   | 27        | 20 (30.0)  |         | 7         | 17.9       |         |
| pCDK9         | n (%)     | n (%)      |         | n (%)     | n (%)      |         |
| Low (≤med)    | 39        | 19 (27.7)  | 0.018   | 20        | 30.0       |         |
| High (>med)   | 30        | 27 (39.1)  |         | 7         | 10.3       |         |

WS Weighted Score, med median

Examples of cervix cancer biopsies with high and low immunohistochemical detection of Caspase-8 and pCDK9 (Thr186). Original magnification ×100, scale bar: 100 μm.

A correlation between Caspase-8 expression and pCDK9 in cervical cancer patients. A Examples of cervix cancer biopsies with high and low immunohistochemical detection of Caspase-8 and pCDK9 (Thr186). Original magnification ×100, scale bar: 100 μm. B The association of the immunohistochemical detection of pCDK9 (percentage of positive cells) and the Caspase-8 weighted score (intensity of staining×percent of positive tumor cells) in pretreatment biopsies of 69 patients. C The association of the immunohistochemical detection of pCDK9 with high (score 8–12) and low (score 0–6) Caspase-8 expression. The tick line is the median value, the solid box is the interquartile range, and the whiskers are the 5th and 95th percentiles. D Immunoblot of the lysate of a cervical cancer primary patient cell line, in which Caspase-8 expression was either knocked down using siCasp8 or transfected with siCtrl. The immunoblot was checked for the levels of Caspase-8, phospho-RNAPII, RNAPII, pCDK9, CDK9, and GAPDH. The levels of pCDK9 and phospho-RNAPII in siCtrl and siCasp8 transfected cells, normalized to their respective CDK9 and RNAPII levels, have been shown graphically. E 3D cell-invasion assay of patient-derived cervical cancer primary cells transfected with either siCasp8 or siCtrl or was non-transfected (NT). Invaded cells were stained with DAPI, and their quantification was graphically represented. The bottom panel shows the fluorescent images of DAPI stained invaded cells of each type [n=3; mean ± SD; p value (paired t test, two-tailed); **** = < 0.0001]

To confirm our findings, we checked the protein expressions of two of these genes—STOM and TGM2 in HeLa WT and KO cells due to their reported roles in promoting metastasis in different cancer entities [65, 66]. We observed that TGM2 expression was up-regulated, while Stomatin was down-regulated in the KO cells (Fig. 7G), precisely matching our analysis (Fig. 7E). Finally, the knock-down of TGM2 (siTGM2) in HeLa and SiHa WT and KO cells significantly reduced their 2D migration, as compared to their respective siCtrl transfected counterparts. This suggested that the enhanced expression of TGM2 is, at least in part, responsible for the enhanced migration of cervical cancer cells, lacking Caspase-8 expression (Supplementary Figs. 7A, B).

This finding is crucial as we could demonstrate that when the HeLa WT and KO cells were treated with increasing concentrations of the CDK9 inhibitor BAY1251152 [38] and determined their proliferation over 120 h (Fig. 8A), HeLa KO cells were significantly less sensitive to BAY1251152 than their WT counterpart (IC50 WT vs. KO: 84.15 vs. 219.3 nM). BAY1251152 also induced significantly lower levels of apoptosis in KO cells, as measured by Caspase-3/7 activities (Fig. 8B) and Annexin V/7AAD staining (Fig. 8C). This suggests that the absence of Caspase-8 expression de-sensitizes cervical cancer cells to CDK9 inhibitors.

To rule out the possibility that the loss of sensitivity of the KO cells to the CDK9 inhibition is not due to the loss of the apoptotic function of Caspase-8, we rescued the KO cells with either Flag-Casp8-WT or its catalytically inactive mutant (Flag-Casp8-C360A) [69], and subjected them
AC DB
0 0,5 1 1,5 2 2,5 3 3,5
extracellular matrix organization
angiogenesis
type I interferon signaling pathway
immune response
response to drug
positive regulation of signal
response to hypoxia
cell migration
cAMP-mediated signaling
positive regulation of inflammatory
processes
histone H3 acetylation
angiogenesis
actin filament depolymerization
response to drug
cell migration
Casp8 (55/53 KDa)
CPS1
STOM
C9A3R1
GSTM1
PC
VAMP8
SNCG
ALS3BP
THY1
LCO2A1

DAVID GOTERM_BP
WT vs KO Transcriptome

DAVID GOTERM_BP
WT vs KO Proteome

Common genes between WT vs KO UT proteomics and transcriptomics

Common genes between proteome and transcriptome

HeLa
TGM2 (78 KDa)
Stomatin (31 KDa)
Casp8 (55/53 KDa)
GAPDH (37 KDa)
The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser29...

The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser29...

We next wanted to investigate the long-term efficacy of BAY1251152 on the 3D-growth of cervical cancer cell lines, which mimic the physiological conditions more accurately [70, 71]. Long-term treatment (up to 144 h) of the 3D-spheroids, derived from HeLa WT and KO cells, with increasing concentrations of BAY1251152, showed that the lack of Caspase-8 expression significantly attenuated the response of HeLa KO cells to CDK9 inhibition. Accordingly, BAY1251152 reduced the growth/volumes of the 3D-spheroids derived from HeLa KO cells significantly less than those derived from WT cells (down to 25% for the WT spheroids as compared to only 60% for the KO spheroids

using 200 nM BAY1251152) (Fig. 8D, E). Additionally, we performed a 3D colony-formation assay on HeLa WT and KO cells treated with BAY1251152 for up to 10 days. The calculated IC50 values of BAY1251152 further confirmed the loss of sensitivity of the Hela KO-derived colonies as compared to their WT-derived counterparts (IC50: 11.30 vs. 21.07 nM, respectively) (Fig. 8F). These results are consistent with our previous findings, which indicated that the loss of Caspase-8 causes a cellular over-activity of CDK9 and highlights the critical role of Caspase-8 in improving the response and sensitivity of cervical cancer to CDK9 inhibition.

**Fig. 7** Identification of differentially expressed genes in the presence/absence of Caspase-8. A Comparison of differentially expressed transcripts in HeLa WT and KO cells, as detected through HumanHT-12 v3 microarrays. > 48,000 transcripts were detected when not adjusted for FDR q value. The red dashed lines represent a cut-off of − Log10 (p value) of ≤ 1.30 and Log2 FC [non-synchronized (NS) and S/G2-phase synchronized (synch.)] of ≥ 0.5 (over-expressed in the KO cells) and ≥ − 0.5 (over-expressed in the WT cells) (x-axis). Normalization of the raw data by adjusting for FDR q value of ≤ 0.05 resulted in 302 transcripts, with an over-expression of 78 transcripts in the KO and 224 in the WT cells. (B) The top 15 functions regulated by these 302 transcripts were determined using the DAVID GOTERM_BP analysis. Highlighted in black are the processes associated with cell-migration. C Comparison of proteins differentially expressed in Tandem Mass Tag (TMT) labeled HeLa WT and KO cells detected through MS analysis. > 5000 proteins were initially detected, of which 60 proteins (33 in WT and 27 in KO cells) passed the cut-offs of − Log10 (p value) of ≤ 1.30 and Log2 FC [non-synchronized (NS) and S/G2-phase synchronized (synch.)] of ≥ ± 0.5, as represented by the red dashed lines. D The top 15 functions regulated by these 60 proteins, as determined using the DAVID GOTERM_BP analysis. Highlighted in black are the processes associated with cell-migration. E Of the 302 genes from transcriptome analysis and 60 protein-encoding genes from the proteome analysis, 14 were shared between the two data sets. Representation of the expression profiles of the 14 common genes in the transcriptome and proteome data sets. F The top 10 functions are regulated by the 14 common genes, as determined using the DAVID GOTERM_BP analysis. Highlighted in black are the processes associated with cell-migration. G Immunoblots of HeLa WT and KO cells representing the protein levels of TGM2, Stomatin, Caspase-8, and GAPDH

**Combinations of Cisplatin with BAY1251152 synergistically inhibit the 2D cell-proliferation and 3D clonogenic-growth of cervical cancer cells**

The gold standard of cervical cancer treatment includes Carboplatin or Cisplatin in combination with radiotherapy. However, cervical cancers frequently develop resistance to platinum-based chemotherapeutics [72, 73], severely undermining their treatment. Thus, we wondered whether targeting CDK9 would improve the response to Cisplatin-based chemotherapy in Cervical cancer cells.

For this purpose, HeLa WT and KO cells were treated with increasing concentrations of BAY1251152 or Cisplatin as single agents or in combination, and their 2D cell-proliferation was assessed over 120 h. Expectedly, the sensitive Caspase-8 WT cells reacted to the BAY1251152/Cisplatin combination by demonstrating significantly enhanced growth inhibition compared to the single treatments at all indicated concentrations. Furthermore, Combination Indices (CI) revealed that BAY1251152 synergistically enhanced the growth inhibitory effect of Cisplatin at all combinations tested (Fig. 9A). Remarkably, even the resistant Caspase-8 KO cells responded to BAY1251152/Cisplatin combination by also demonstrating significantly reduced survival, as compared to the single treatments, at all indicated concentrations. CI values once again revealed that BAY1251152 synergistically enhanced the growth inhibitory effect of Cisplatin at all combinations tested (Fig. 9A).

We also similarly treated HeLa WT and KO cells with BAY1251152 or Cisplatin as single agents or in combination and assessed their 3D clonogenic-growth over 10 days. As before, the BAY1251152/Cisplatin combination significantly decreased the clonogenic-growth of HeLa KO cells, but only the lowest combination of Cisplatin with BAY1251152 did so synergistically (0.75 µM + 10 nM; CI 0.80), as the higher combinations were either additive (1.0 µM + 20 nM; CI 1.03) or antagonistic (1.5 µM + 30 nM; CI 1.33) (Fig. 9B, C). Here again, the HeLa WT cells demonstrated a significant and synergistic decrease in clonogenic-growth at all
A  HeLa WT vs. KO

Normalized cell viability (%) vs. Log [BAY1251152]

- WT 120 h: LogIC₅₀=1.925 nM, IC₅₀=84.15 nM
- KO 120 h: LogIC₅₀=2.341 nM, IC₅₀=219.3 nM

B  BAY1251152

Caspase 3/7 activity (RLU x 10⁴ fold change)

- WT 10 days: LogIC₅₀=1.053 nM, IC₅₀=11.30 nM
- KO 10 days: LogIC₅₀=1.324 nM, IC₅₀=21.07 nM

C  HeLa WT vs. KO

Annexin positive (%) vs. BAY1251152

- WT: DMSO 75 nM, 100 nM, 200 nM
- KO: DMSO 75 nM, 100 nM, 200 nM

D  HeLa WT vs. KO

Normalized % of spheroid volume vs. BAY1251152

- WT: NT DMSO 75 nM, 100 nM, 200 nM
- KO: NT DMSO 75 nM, 100 nM, 200 nM

E  HeLa WT vs. KO

Annexin positive (%) vs. BAY1251152

- WT 200 nM BAY
- KO 200 nM BAY

F  HeLa WT vs. KO

Normalized cell viability (%) vs. Log [BAY1251152]

- WT 10 days: LogIC₅₀=1.053 nM, IC₅₀=11.30 nM
- KO 10 days: LogIC₅₀=1.324 nM, IC₅₀=21.07 nM
The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser2...

In summary, the results of our 2D- and 3D-cellular experiments demonstrated that the response of cervical cancer to Cisplatin-based standard chemotherapy might significantly improve by concurrently blocking CDK9 activity using small-molecule inhibitors. This renders CDK9 a promising molecular target for treating cervical cancer patients with low Caspase-8 expression.

**Discussion**

Concurrent chemoradiation (CCRT) with Cisplatin or Carboplatin alone or combined with Paclitaxel, Gemcitabine, and Vincristine is the standard treatment approach for late-stage cervical cancer [74]. However, cervical cancer frequently develops chemotherapeutic resistance, which along with metastases, lowers the 5-year survival rate to just 17% [3, 5]. Therefore, investigating the molecular mechanisms underlying cervical cancer metastasis, the root causes of treatment resistance, and discovering new prognostic and predictive biomarkers may help to develop novel therapeutic strategies for this lethal malignancy. In addition to its traditional function of initiating cell death, Caspase-8 is also implicated in several non-apoptotic processes in various cancers, including cell-cycle control, proliferation, migration, invasion, and angiogenesis [10, 24–31, 74]. Furthermore, by modulating the enzymatic activities of CDK9 and RNAPII, our current investigation added a novel and crucial role for Caspase-8 in mediating transcription elongation.

First, our meta-analysis revealed that Caspase-8 expression levels correlate with tumor development and clinical outcome (Fig. 1). We discovered that low Caspase-8 expression is associated with poor OS and PFS in cervical cancer patients. Furthermore, the loss of Caspase-8 expression promotes migration and invasion in patient-derived cervical cancer cells (Fig. 6E), suggesting a correlation between Caspase-8 expression and elevated cervical tumor malignancy and aggression. These results suggested that cervical cancer patients' Caspase-8 protein expression profile could be used as a predictive biomarker.

Additionally, concurrently with the patient data, transcriptome and proteomic profiling of Caspase-8-depleted cervical cancer cell lines (HeLa and SiHa) identified numerous genes with altered expression, including STOM and TGM2, which may boost the propensity for metastasis and invasion. In cells missing Caspase-8, TGM2 levels increased, whereas stomatin levels declined (Fig. 7G). This finding was intriguing because stomatin has been shown to prevent metastasis in NSCLC cells, and its downregulation during EMT stimulates metastasis and is associated with poor prognosis [64]. Contrarily, the up-regulation of TGM2 expression/activity has been linked to a greater incidence of metastasis in various cancer entities [75–77]. In light of these findings, we could show that the knock-down of TGM2 expression/activity has been linked to a greater incidence of metastasis in various cancer entities [75–77]. In light of these findings, we could show that the knock-down of TGM2 expression/activity has been linked to a greater incidence of metastasis in various cancer entities [75–77]. In light of these findings, we could show that the knock-down of TGM2 expression/activity has been linked to a greater incidence of metastasis in various cancer entities [75–77].
Fractionation experiments confirmed that the physical association between caspase-8 and CDK9 takes place within the nucleus (Fig. 4J), even though Caspase-8 has previously been reported to be a cytoplasmic protein [10, 18] and even though it also exhibits a nuclear localization sequence (NLS) in its p10 domain [78]. This finding is in line with previous studies reporting the accumulation of Caspase-8 in the nucleus of cervical cancer cells with high-risk human papillomaviruses (HPVs) expressing E6 and E7 proteins such as HeLa cells [75–77]. The nuclear accumulation of Caspase-8...
The non-apoptotic function of Caspase-8 in negatively regulating the CDK9-mediated Ser2...

...is due to its stimulated recruitment by the E6 oncoprotein, such that Caspase-8 exerts beneficial functions for the viral life cycle [77]. Here, the oncoprotein E6-Caspase-8 association appears to have little impact on promoting Caspase-8’s apoptotic process, most likely caused partly by Caspase-8’s changed cellular location [77].

CDK9 is primarily activated by forming a heterodimeric complex involving Cyclin T1 to form P-TEFb, although minor cyclins like Cyclins T2a and T2b can also perform this function [78]. While around 80% of CDK9 remains as P-TEFb, sequestered in the nucleus either as an inactive complex 7SK snRNP or an active complex with BRD4 [38, 79–83], the remaining ~20% of inactive CDK9 remains within the cytoplasm, bound to two chaperones—HSP70 and the HSP90/CDC37 complex, and undergoes cytoplasmic to nuclear shuttling [38, 62, 84, 85]. The phosphorylation of the CDK9 Thr186 residue, either through autophosphorylation or by CDK7 [86], is critical for its association with Cyclin T1 [38, 87] and its dissociation from the HSP90/CDC37 complex [88]. Based on this, our data suggest that Caspase-8 allosterically inhibits the activity of CDK9, as measured by its autophosphorylation. This would occur by multiple mechanisms like modifying the conformation of CDK9 to block the Thr186 site or the ATP binding site, the hinge region, or by competing directly with CDK7 to prevent it from phosphorylating CDK9. Interestingly, we have been able to confirm here that, under in vitro conditions, Caspase-8 binds to CDK9 directly (Fig. 4C). Furthermore, as our ex vivo experiments have demonstrated a nuclear interaction between Caspase-8 and CDK9, taken together, these experiments have convincingly established that these two proteins interact, within the nucleus, without the presence of any bridging molecules like BRD4 or members of the 7SK snRNP complex like—HEXIM1/2, LARP7 and MePCE [38].

In addition, the loss of Caspase-8 caused what we can describe as a hyperactive state of CDK9, resulting in the increased phosphorylation of key CDK9 targets. Thus, we showed that Caspase-8 depletion increased CDK9-mediated phosphorylation of crucial substrates like RNAPII and SPT5 in vitro and in cells. This increase in RNAPII activity consequently caused an alteration in cellular transcription, as demonstrated by the increase in EU and the alteration of the expression of several genes (Fig. 7E, G), notably those involved in metastasis and invasion. Similarly, our investigation of a cohort of primary cervical cancer patient-derived tissues also confirmed a significant correlation between high Caspase-8 expression and low CDK9 Thr186 phosphorylation, and low RNAPII Ser2 phosphorylation (Fig. 6A–D).

This novel mechanism of Caspase-8-dependent inhibition of CDK9 is a singular finding that differs entirely from what has been reported in the literature about caspase-mediated regulation of protein kinases [18, 20, 89]. More specifically, many kinases, particularly cell-cycle-associated ones, have been described to undergo caspase-dependent cleavage activation [89]. Accordingly, an increased CDK2 kinase activity has long been closely linked to apoptosis. Furthermore, this activation has been shown to be triggered by a caspase-dependent cleavage cascade, directly by triggering the cleavage of Cyclin A, creating a proteasome-resistant and kinase-active form of Cyclin A/CDK2 involved in the apoptotic response to ionizing radiation [90], and indirectly by inducing the cleavage of the CDK inhibitors, p21Cip1/Waf1, and p27Kip1 [91], or WEE1 inactivation [92], which promotes inhibitory phosphorylations of CDK2 during apoptosis, and consequently leading to increased activity of CDK2 [91].

One of the significant findings of our study can be summarized by the fact that in cervical cancer cell lines, as well as in patient-derived cells, loss of Caspase-8 induces an over-activation of CDK9, which consequently alters the cellular transcriptional landscape (transcriptional re-programming), by increasing RNAPII activity. This represents, most likely, one of the potential resistance mechanisms of advanced cervical cancer to standard chemotherapy, similar to what has been described in medulloblastoma resistance to conventional therapy [93].

This significant finding has given us a solid rationale for targeting low Caspase-8 expressing cells with increased CDK9 kinase activity, using small molecule CDK9 inhibitors, either alone (Fig. 8) or in combination with Cisplatin (Fig. 9). We could demonstrate that combining Cisplatin with the small-molecule CDK9 inhibitor—BAY1251152, could overcome the chemoresistance and restore the sensitivity of HeLa KO cells toward Cisplatin, synergistically and significantly, again under both 2D- and 3D-conditions (Fig. 9). Not surprisingly, the HeLa WT cells, with low CDK9 activity, were synergistically targeted by a combination of BAY1251152 and Cisplatin, under both growth conditions. This finding is important because several third-generation CDK9 inhibitors like BAY1251152, AZD-4573, and
JSH-150 are already undergoing pre- or clinical trials [38], and newer strategies are being applied for the identification of novel inhibitors against transcription-associated CDKs like CDK9 [94]. Moreover, there are excellent precedents for combining standard chemotherapeutics with small-molecule inhibitors of CDKs like those against CDK 4/6 [95, 96].

Additionally, our observation that in cervical cancer patients with low TMB, low Caspase-8 expression elicits a significantly better prognosis than in patients with high Caspase-8 expression (Fig. 1B) could be explained by the possibility that during the early stages of cervical cancer development, localized cervical cancer cells are not overtly dependent on the altered expressions of metastasis inducing genes. Therefore, the presence of Caspase-8 in these cells is not a serious impediment to the development of this cancer. Only during invasive stages the expression of Caspase-8 is down-regulated to aid in the process. Indeed, it has been reported that TMB in cervical cancer significantly correlates with its stages [97]. The presence of Caspase-8 in the earlier stages probably also assists the chemotherapeutic treatment of this cancer, eliciting a better prognosis [16].

In conclusion, we believe that the data presented by this study support further pre-clinical and clinical evaluations of CDK9 inhibitors in advanced and resistant cervical cancer. However, a more rigorous assessment of Caspase-8 expression and the corresponding stratification of patients will be of utmost importance in determining the clinical success of the CDK9-based targeted strategies in cervical and other cancer entities in the future.

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1007/s00018-022-04598-3.

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Author contributions The execution and analyses of most of the experiments were performed by RM, MR, and AK. FR and IK participated in the analysis of IHC of cervical cancer patient-derived materials. Transcriptome and TCGA analyses were performed by SPL. Proteome analysis was performed by BH and TO. RM, MR, MS, and KS participated in the discussion of the data. KS and SB supervised and guided the entire project. RM, MS, and KS wrote the manuscript.

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Data availability All materials used in this work will be provided upon request. Illumina microarrays were deposited in the Gene Expression Omnibus (GEO) under the accession number GSE217360. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRoteomics IDEntifications Database (PRIDE) partner repository with the dataset identifier PXD038102.

Declarations

Conflict of interest The authors have no relevant financial or non-financial interests to disclose. T.O. received research funding from Gilead and Merck KGaA (unrelated to this work). T.O. is a consultant for Roche and Merck KGaA (both unrelated to this work).

Ethical approval This study was approved by the Institutional Review Boards of the UCT and the Ethical Committee at the University Hospital Frankfurt (Project Nr.: UCT-13-2021).

Consent to participate Cervical cancer tissue/tumor samples used in this study were provided by the University Cancer Center Frankfurt (UCT). In addition, written informed consent was obtained from all patients.

Consent for publishing All co-authors have read and approved the final version of the manuscript and its submission.

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