Deletion of the deISGylating enzyme USP18 enhances tumour cell antigenicity and radiosensitivity

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BACKGROUND: Interferon (IFN) signalling pathways, a key element of the innate immune response, contribute to resistance to conventional chemotherapy, radiotherapy, and immunotherapy, and are often deregulated in cancer. The deubiquitylating enzyme USP18 is a major negative regulator of the IFN signalling cascade and is the predominant human protease that cleaves ISG15, a ubiquitin-like protein tightly regulated in the context of innate immunity, from its modified substrate proteins in vivo.

METHODS: In this study, using advanced proteomic techniques, we have significantly expanded the USP18-dependent ISGylome and proteome in a chronic myeloid leukaemia (CML)-derived cell line. USP18-dependent effects were explored further in CML and colorectal carcinoma cellular models.

RESULTS: Novel ISGylation targets were characterised that modulate the sensing of innate ligands, antigen presentation and secretion of cytokines. Consequently, CML USP18-deficient cells are more antigenic, driving increased activation of cytotoxic T lymphocytes (CTLs) and are more susceptible to irradiation.

CONCLUSIONS: Our results provide strong evidence for USP18 in regulating antigenicity and radiosensitivity, highlighting its potential as a cancer target.

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identified only in vertebrates; moreover, human and murine ISG15 protein sequences only share 64% and 74% homology and similarity, respectively.17

Usp18−/− mice present a strong accumulation of ISG15 conjugates in tissues as well as a prolonged increase in STAT1-dependent activation.18 These mice are hypersensitive to IFN and Poly I:C (a synthetic double-stranded RNA) stimulation and are more resistant to intracerebral infection. In some mice genetic backgrounds, Usp18 deficiency has been linked to brain abnormalities.15,16,19

USP18 functions that are independent of ISG15 seem to be crucial in mice since the phenotype observed in Usp18−/− animals, including brain inflammation and deregulated STAT1 signalling, cannot be rescued by lsg15 deletion.20 In humans, however, free ISG15 stabilises USP18 and acts as a negative regulator of the IFN pathway.21,22 ISG15-null patients present with inflammation due to deregulation of the type I interferon response and, interestingly, are susceptible to mycobacterial, but not viral, diseases.21,22

It has also been reported that silencing USP18 increases surface expression of peptide-loaded MHC-II,24 supporting its role as a negative regulator of immunogenicity. USP18 has been linked to cancer development, being overexpressed in lung, colon, pancreas and breast cancer, but also associated with autoimmune diseases and neuroinflammation.15,23,26 Notably, USP18 has also been described to have a role as a resistance factor to traditional chemotherapy agents Bortezomib and Mafosfamide in tumour cells,28,29 suggesting that its inhibition may serve as a boost for chemotherapy agents Bortezomib and Mafosfamide in tumour such therapies.

Protein ISGylation profiles in human cells have been previously reported, but with limited depth and predominantly based on overexpression systems.30–35 To obtain deep coverage and modified residue information in a more physiologically relevant context, we decided to combine advanced proteomic techniques with the immunoaffinity purification of GlyGly tryptic peptides in wild-type (WT) and USP18−/− knockout (KO) HAP1 cells, in order to analyse the first USP18-dependent ISGylome of human cancer cells.

METHODS

Cell lines and T cells

Chronic myeloid leukaemia (CML)-derived HAP1 wild-type (WT; Horizon #C631) cells and HAP1 USP18−/− (KO; Horizon #HZGHC000492c011) were a kind gift from the laboratory of Sebastian Nijman and were cultured in IMDM media (Gibco #12440-53) supplemented with 10% FBS (Gibco #10500-64) at 37 °C in a humidified 5% CO2 atmosphere. HAP1 USP18 KO cells were generated by editing parental HAP1 cells with CRISPR/Cas9 to contain a 4 bp deletion in the coding exon 3 of the USP18 gene. Melan-A specific CTL lines were isolated from healthy blood donors as described.36 Colorectal carcinoma-derived HCT116 wild-type (ATCC-CCL-247) and HCT116 USP18 KO cells were cultured in DMEM media (Gibco #10313) supplemented with 10% FBS (Gibco #10500-64) at 37 °C in a humidified 5% CO2 atmosphere. HCT116 KO cells were generated using the commercially available USP18 CRISPR/Cas9 KO (Santa Cruz #sc-402259) and USP18 homology-directed DNA repair HDR (Santa Cruz #sc-402259-HDR) plasmids, following the manufacturer’s recommendations. PBMC were isolated from healthy blood purchased through the UK National Blood Service (agreement T293). Briefly, dendritic cells pulsed with Melan A peptides (ELA26-35 ELAGIGILTV, Sigma) were incubated with autologous PBMCs; after 12 days melan-A specific CD8 T cells were sorted with HLA-A2-ELA class I tetramers and expanded.

Stimulation of cells

Cells were treated with 1000 U/mL of human interferon α 2 (α2b) from PBL Assay Science (Cat. No. 11105-1) for the indicated times. DNA plasmids and generation of stable cell lines

The Flag-HA-USP18 WT construct was purchased from Addgene (#22572) and the Flag-HA-USP18C64R/C65R construct was generated using the QuikChange Site-Directed Mutagenesis Kit (NEB, E0554S) and the primers OL87: TGGACAGACCGcgccCTTAACTCTTGT and OL88: ATGTGTGTAACAAACC. Primers were designed using the NEBaseChanger online tool: http://nebasechanger.neb.com/ and the successful clone confirmed by sequencing (Eurofins).

Transfected HAP1 and HCT116 cells were selected, pooled, and subsequently cultured in growth media supplemented with puromycin (Gibco) at 1 µg/mL.

siRNA reagents

ISG15 siRNA on-TARGET plus SMART plus (Dharmacon #L-004235-03-0005) and ADAR siRNA on-TARGET plus SMART plus (Dharmacon #L-008630-00-0005).

Cell transfection

USP18 cDNA, CRISPR/Cas9, and HDR plasmids were transfected with Lipofectamine LTX and Plus (Invitrogen #15338-100) following the manufacturer’s instructions. The indicated siRNA sequences were transfected with Lipofectamine RNAmax (Invitrogen #13778-150) following the manufacturer’s instructions.

Antibodies

See separate Table S1.

Western blotting

Cells were washed with ice-cold PBS and lysed with NP-40 lysis buffer (50 mM Tris pH 7.4, 0.5% (v/v) NP-40, 150 mM NaCl and 20 mM MgCl2) buffer containing protease and phosphatase inhibitors. For Western blotting, 25 µg of protein was then fractionated on Tris–glycine SDS-PAGE gradient (4–15% acrylamide) gels (BioRad; #3450123), transferred onto PVDF membranes (Millipore; IFPL00010), and detected with the indicated antibodies using a LI-COR detection system.

USP18 ABP

ISG15-PA was synthesised in the Kessler lab using the ISG15 C-termin (79-156) construct from Addgene (#110760) and DUB ABP assays were performed as previously described.37 Briefly, HAP1 cells were lysed in glass beads lysis buffer (GIBL: 50 mM Tris, pH 7.5, 5 mM MgCl2, 0.5 mM EDTA, and 250 mM Sucrose), the extracts were subsequently labelled with ISG15-PA for 45 min at 37 °C, and immunoblotted with specific anti-USP18 antibodies.

Cell growth, morphology and proliferation assays

10000 cells were seeded in 6-well or 12-well plates, and cells were imaged in an IncuCyte Zoom Imager (Essen Bioscience) for live-cell imaging during the indicated times, at the indicated intervals, in the phase contrast channel. Cell growth was determined by measuring the percent confluence over time. End point cell viability was measured using the Resazurin assay (Alamar Blue; Thermo Fisher #DAL1025), following the manufacturer’s instructions.

Immunofluorescence microscopy

HAP1 control or HAP1 USP18 KO cells were seeded in fluorescence compatible 96-well plates (Corning), treated with IFN for the indicated times, subsequently washed with PBS, fixed using 4% paraformaldehyde (sc-281692, ChemCruz) for 10 min, permeabilised using 100% methanol for 10 min at −20 °C, and blocked with 5% BSA in PBS for one hour at room temperature. The primary antibodies were diluted in PBST (PBS + 0.05% Tween20) to the following concentrations: USP18 (1:100), ISG15 (1:100) and pSTAT1 (1:400), added and incubated overnight at 4 °C. Cells were then washed with PBST and subsequently incubated with a secondary antibody mix consisting of Goat anti-rabbit conjugated alexa-647 (1:10,000) and 4',6-diamidino-2-phenylindole (Dapi) (2.5 µg/mL),
for 1 h at room temperature. Plates were imaged using the Opera Phenix imaging platform (Perkin Elmer).

For dsRNA analysis, the same workflow was used with the following modification: The cells were fixed and permeabilised using Cytofix/Cytoperm buffers (BD biosciences), according to the manufacturer’s instructions. dsRNA antibody was used at a 1:100 dilution and the secondary donkey anti-mouse at a 1:500 dilution.

Identification of ISGylated proteins using GlyGly peptidomics and matching proteomics

HAP1 lysates were used for GlyGly immunoprecipitation using the PTMScan Ubiquitin Remnant Motif Kit (Cell Signaling), according to the manufacturer’s protocol. Briefly, 20 mg of extracts were solubilised and denatured in 10 mL lysis buffer (20 mM HEPES, pH 8.0, 9 M urea, 1 mM sodium orthovanadate, 2.5 mM sodium
pyrophosphate, 1 mM β-glycerophosphate), reduced using dithiothreitol (4.5 mM final) for 20 min at 55 °C. This was followed by alkylation using iodoacetamide (100 mM final) for 15 min at room temperature in the dark. Samples were subsequently diluted fourfold in 20 mM HEPES, pH 8.0 (~2 M urea final), followed by digestion with trypsin-TPCK (Worthington, LS003744, 10 mg/mL final) overnight at 37 °C. Samples were then acidified using trifluoroacetic acid (1% final) and desalted using C-18 Sep-Pak (Waters) cartridges according to the manufacturer’s protocol. At this point, 20 μg of digested protein were separated for matching proteome control analysis. Peptides were lyophilized and re-suspended in 1.4 mL immunoprecipitation IAP buffer (PTMscan), and the remaining insoluble material cleared by centrifugation and anti-GlyGly antibody beads added followed by rotation and kept 4 °C for 2 h. Beads were subsequently washed twice using 1 mL IAP buffer, followed by three water washes. Immunoprecipitated material was eluted twice in 55 and 50 μL 0.15% trifluoroacetic acid in water. Peptide sample was desalted and concentrated using C-18 Sep-Pak (Waters) cartridges according to the manufacturer’s protocol. Purified GlyGly-modified peptide eluates and matching proteome material were dried by vacuum centrifugation and re-suspended in buffer A (98 % MilliQ-H2O, 2% CH3CN and 0.1% TFA).

ISG15 interactome immunoprecipitation
HAP1 cells were lysed with Co-IP lysis buffer (20 mM Hpes pH 8.0, 150 mM NaCl, 0.2% NP-40, 10 % glycerol, 5 mM NEM, phosphatase and protease inhibitor cocktails; 25 ×106 cells per condition) and subjected to immunoprecipitation using 5 μg of ISG15 antibody (Boston Biochem #A-380) plus 25 μL of protein G Sepharose slurry (Invitrogen; #15920-10) for 16 h at 4 °C. Beads were washed 4 times with Co-IP lysis buffer and immunocomplexes were eluted with 2X Laemmli. 10% of the eluates was used for immunoblotting with the indicated antibody. The remaining eluate was prepared for MS analysis as previously described. Briefly: immunoprecipitated sample eluates were diluted to 175 μL with ultra-pure water and reduced with 5 μL of DTT (200 mM in 0.1 M Tris, pH 7.8) for 30 min at 37 °C. Samples were then alkylated with 20 μL of iodoacetamide (100 mM in 0.1 M Tris, pH 7.8) for 15 min at room temperature (protected from light), followed by protein precipitation using a double methanol/chloroform extraction method. Protein samples were treated with 600 μL of methanol, 150 μL of chloroform, and 450 μL of water, followed by vigorous vortexing. Samples were centrifuged at 17,000 × g for 3 min, and the resultant upper aqueous phase was removed. Proteins were pelleted following the addition of 450 μL of methanol and centrifugation at 17,000 × g for 6 min. The supernatant was removed, and the extraction process was repeated. Following the second extraction process, precipitated proteins were re-suspended in 50 μL of 6 M urea and diluted to <1 M urea with 250 μL of 20 mM HEPES (pH 8.0) buffer. Protein digestion was carried out by adding trypsin (from a 1 mg/ml stock in 1 mM HCl) to a ratio 1:100, rocking at 12 rpm and room temperature overnight. Following digestion, samples were acidified to 1% trifluoroacetic acid and desalted on C18 solid-phase extraction cartridges (SEP-PAK plus, Waters), dried, and re-suspended in buffer A.

Liquid chromatography-tandem mass spectrometry (LC-MS/MS) analysis
LC-MS/MS analysis was performed using a Dionex Ultimate 3000 nano-ultra high-pressure reverse-phase chromatography coupled online to a Q Exactive HF (GlyGly), Fusion Lumos (ISG15 interactome) or a Q Exactive (Matching proteome) mass spectrometer (Thermo Scientific) as described previously. In brief, samples were separated on an EASY-Spray PepMap RSLC C18 column (500 mm × 75 μm, 2 μm particle size, Thermo Scientific) over a 60 min (120 min in the case of the matching proteome) gradient of 2–35% acetonitrile in 5% dimethyl sulfoxide (DMSO), 0.1% formic acid at 250 nL/min. MS1 scans were acquired at a resolution of 60,000 at 200 m/z and the top 12 most abundant precursor ions were selected for high collision dissociation (HCD) fragmentation.

Data analysis
From raw MS files, searches against the UniProtKB human sequence database (92,954 entries) and label-free quantitation were performed using MaxQuant Software (v1.5.5.1). Search parameters include carbamidomethyl (C) as a fixed modification, oxidation (M) and deamidation (NQ as variable modification), maximum 2 missed cleavages (3 for the GlyGly peptidome analysis), matching between runs, and LFQ quantitation was performed using unique peptides. Label-free interaction data analysis was performed using Perseus (v1.6.0.2), and volcano and scatter plots were generated using a t-test with permutation FDR = 0.01 for multiple-test correction and s = 0.1 as cut-off parameters.

Other graphs were generated using GraphPad PRISM 8 and Excel and for the statistical analysis, we applied two-way ANOVA tests, including multiple comparison testing via the Dunnett method available through the GraphPad Prism software. P value style is GraphPad: NS, *P = 0.01234, *P = 0.0332, **P = 0.0021, ***P = 0.0002, ****P < 0.0001.

Gene ontology and pathway enrichment analysis were performed using STRINGdb (https://string-db.org/cgi/input.pl?sessionId=9c1fKQ0VOpdjY&input_page_show_search=on) and g: PROFILER (https://biit.cs.ut.ee/gprofiler/gost).

Flow cytometry data were analysed with Flowjo 10, upon gating on live singlets.

FACS/T cell cell activation pulsing with Melan-A peptides
50,000 HAP1 WT and USP18−/− cells (+/− IFNo-2a, 1000 U/mL; 24 h) were plated in 96-well plates (flat bottom, Costar) and pulsed with the indicated amounts of Melan-A peptides (ELAGILTV, Sigma) for 2 h at 37 °C. Cells were washed three times with RPMI media (Sigma) prior to addition of melan-A-specific T cells (healthy donor-derived) at a ratio 2 tumour cells: 1 immune cell for 18 hours. T cell activation (FACS) and cytokine release in the supernatant (ELISA) were analysed subsequently.

For FACS analysis, triplicate wells were pooled, cells were stained for 20 min at room temperature with a near infrared live/dead marker (Biolegend) and then with a cocktail of titrated
antibodies, 30 min on ice. Samples were acquired on a BD Symphony flow cytometer.

ELISA
Enhanced ELISA plates (Corning) were coated with diluted anti-IFNγ antibodies (2 μg/mL) in coating buffer (100 mM NaHCO3) and left overnight at 4 °C. Plates were washed 6 times with PBS/0.01 % Tween and blocked with PBS/10% FCS for 2 h at room temperature. Diluted standards and samples in PBS/10% FCS were added to the plate and incubated for 3 h at 37 °C. Plates were washed 6 times with PBS/0.01% Tween prior addition of biotinylated anti-cytokine detecting mAb (2 μg/mL in PBS/10%
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**Fig. 2** USP18 is the major cellular de-ISGylase. **a** Workflow for analysis of GlyGly-modified peptides. **b** Comparative table of the C-terminal motifs of the different ubiquitin-like proteins. ISG15, Ubiquitin, and NEDD8 leave a unique GlyGly motif (in green) after trypsin digestion of modified proteins. **c** Ubiquitin-conjugated and **d** NEDD8-conjugated proteins do not increase after stimulation with type I interferon (IFN) in USP18 KO cells. **e** USP15 modification is accumulated in the same conditions as in (c). **f** Comparative volcano plot of the GlyGly-modified peptides in HAP1 WT and USP18 KO cells treated with IFN for 48 h showing a strong and significant USP18-dependent upregulation of GlyGly-modified peptides in the KO cells treated with IFN for 48 h. In red are shown the upregulated peptides in the KO cells, in green the upregulated peptides in WT cells, and in blue the modified peptides from an ISG-cancer signature (the statistical cut-off values used for all the proteomic analyses performed in this study are FDR: 0.01 and s0: 0.1). Pathway enrichment analysis showed the upregulated peptides in the USP18 KO cells come from proteins involved in the indicated biological processes. 

**RESULTS**

USP18 catalytic activity is essential for cell survival in the presence of type I IFN

We decided to study the effects of type I interferon α 2 (IFNα2; IFN hereafter) on human HAP1 cells, a chronic myeloid leukaemia (CML)-derived cell line, in the presence or absence of the USP18 gene. Under normal culture conditions, both cell lines proliferate at the same rate (Fig. 1b). However, while the parental cells are completely insensitive to treatment with 1000 U/ml of IFN (Fig. 1c), the USP18-deficient cells stop growing after 24 h (Fig. 1c and Fig. S1A) and start to die as visualised by microscope phase-contrast imaging Fig. S1B. The same results were observed using the colorectal cancer cell line HCT116, suggesting an essential role for USP18 under an innate immune stimulation context, in different cancer cell types (Fig. S2A and S2B). Interestingly, the sensitivity to IFN correlated with the expression of USP18 in three different KO clones, expressing different amounts of the protein (Fig. S2C, top panel). Supporting this phenotype, a molecular indicator of programmed cell death, the cleavage of the protein PARP, is only observed in the KO cells treated with IFN (Fig. 1d). Consistent with previous literature, deletion of USP18 also led to the accumulation of ISG15 conjugates in our cellular model, shown as a strong smear after 48 h of treatment with IFN (Fig. 1e) and an increase in ISG15 levels and STAT1 phosphorylation in immunofluorescence analysis after 24 hours of IFN stimulation (Fig. S1C). Enhancement of STAT1 phosphorylation in tyrosine 701 after USP18 deletion or depletion was also detected by immunoblotting in both, HCT116 (Fig. S2C), and HAP1 (Fig. S2D) cells. Strikingly, the accumulation of ISGylated proteins is barely noticeable in HAP1 WT cells after IFN treatment (Fig. 1e).

USP18 functions as a negative regulator of the IFN pathway have been linked not only to its ability to remove ISG15 from proteins, but also to inhibitory protein-protein interactions with IFNAR2 and STAT2. In order to evaluate the importance of USP18 catalytic activity in our model, we re-expressed the wild-type (WT) protein and a catalytically inactive mutant, where we mutated the catalytic cysteine and a second adjacent cysteine (C64R/C65R), in HAP1 USP18 KO cells. Stable transfectants expressing the WT protein rescued KO cells from the toxic effects of IFN (Fig. 1f and Fig. S1D) and prevented the accumulation of ISGylated proteins (Fig. 1g). Unexpectedly, and in contrast with what has been published in murine systems, re-expression of USP18 C64R/C65R did not rescue the KO cells from IFN toxicity (Fig. 1f and Fig. S1D) or from the accumulation of ISGylated proteins (Fig. 1g). As a control to evaluate the activity of the two proteins, we performed deISGylating activity assays using ISG15 activity-based probes that bind irreversibly to catalytically active USP18 and allow the visualisation of the active enzyme as an increase in the molecular weight ~15 kDa (USP18 plus ISG15 probe) by immunoblotting. We observed that only wild-type USP18 is able to bind the activity-based probe, whereas the C64R/C65R mutant is completely inactive (Fig. 1g, middle panel). As a control for inhibition of the IFN-receptor by direct binding of WT or C64R/C65R USP18, we performed immunoblots against the ISGs HERCS, IFIT3, and ISG15. Both, WT and C64R/C65R USP18 are able to inhibit the expression of these three ISGs, although the inhibitory effect seems to be smaller for the catalytically inactive mutant (Fig. S1E), perhaps pointing at a slightly impaired ability to bind IFNAR2 or STAT2 of this mutant due to structural changes induced by the mutations.

USP18-dependent GlyGly peptidom proteome reveals tumour cell ISGlyome

Digestion with trypsin of proteins modified either with ubiquitin, ISG15 or NEDD8 leaves a unique GlyGly motif on the modified lysine (Fig. 2a, b). Enriching for GlyGly-modified peptides followed by mass spectrometry (MS) analysis provides access to information on not only the prevalence/amount of the modification but also on its exact position in the protein sequence. A recent report exploited this to analyse cellular ISGlyomes in mice, although using a different genetic cellular background strategy to the one used here and in the context of infection. In this study, we employed a similar approach to conduct the first comprehensive analysis of the USP18-dependent ISGlyome in human cancer cells. Even though GlyGly peptidomics is not specific for ISG15, we hypothesised that KO of USP18 (the main deISGylating enzyme in vivo) in cells should lead to accumulation of ISGylated substrates upon type I IFN stimulation that would be suitable for identification using GlyGly immunoaffinity purification of the generated tryptic peptides. To validate our hypothesis, we first wanted to verify, by immunoblotting, that ubiquitination and NEDDylation were not affected across our four experimental conditions (HAP1 WT and USP18 KO in the presence/absence of IFN). As shown in Fig. 2c, d, ubiquitination, and NEDDylation did not seem to be affected. If something, we saw a small reduction of ubiquitylated proteins in the USP18 KO cells treated with IFN. In contrast, IFN treatment of the KO cells led to a massive accumulation of conjugated ISG15 material (Fig. 2d). Based on these observations, we would expect to detect mainly the accumulated ISGylated proteins when applying the GlyGly peptidomics in these particular experimental conditions. 2341 GlyGly-modified peptides were identified in this mass spectrometry-based experiment, corresponding to 2172 protein groups. As the volcano plot in Fig. S3A shows, there are no
significant changes in the comparative GlyGly peptidomes of HAP1 cells treated with and without IFN, consistent with the results from the immunoblots. However, and again in line with the gel results, following treatment of USP18 KO cells with IFN and comparison of their GlyGly peptidome to that of the WT cells (also treated with IFN), we observed a massive USP18-dependent, increase in GlyGly peptides, corresponding to 476 proteins, in the KO cells (Fig. 2f and Data Table S1). After gene ontology/pathway enrichment analysis, we could reinforce the role of USP18 as a major regulator of the innate immune response with significant
enrichment of ISG15-modified peptides for dsRNA binding and closely related proteins such as ADAR, DDX58, DDX60, DHX58, OAS1/2, EIF2K2A (PKR), IFIH1, DDX3X, DHX9, STAT1, and others that are part of innate immunity pathways. In addition, we observed a strong upregulation of ISGylated (immuno-)proteasomal subunits PSMA7, PSMB9, PSMB10, and PSME2 as well as TAP1, all of which are part of the antigen presentation pathway. Proteins involved in translation and glucose metabolism processes were also represented with numerous ISGylated protein members (Fig. 2f). Interestingly, we also found that proteins from a recently published signature of interferon-stimulated genes (ISGs) expressed constitutively in a subset of cancer cell lines were also strongly upregulated and represented in our ISGylome (Fig. 2f). The global proteome of the same samples was analysed in parallel (Figs. S3B, 3C, and 3D). USP18 is expressed in very low amounts in cancer ISG signature (Fig. S3D). In agreement with the ISGylome and proteome (in IFN-treated cells) data sets, we could observe that in our case ISGylation is not leading to a decrease in protein levels (Fig. 2g; in green) and an accumulation of modified proteins, mainly ISGs (Fig. 2g; in blue). The latter effect may putatively not be due to ISGylation leading to a stabilisation of proteins but might instead reflect the roles of USP18 as an inhibitor of late-stage IFN signalling. However, the potential role of ISG15 as stabiliser of a certain set of ISGs cannot be excluded at this point.45

USP18 controls ISGylation of the dsRNA modifier ADAR (p150) and the innate ligand sensor PKR and regulates their activity

To verify these results, we performed immunoblotting with specific antibodies against a group of selected ISGylated hits such as ADAR, PKR, HERCS, and IFIT3, and we could observe additional bands for these proteins (consistent with, at least, an increase in the MW of ~15 kDa) only in the KO cells treated with IFN (Fig. 3a, d). ISGylated HERCS material was also detected in the colorectal cancer cell line (Fig. S2c). To further validate ISGylation as the observed modification, we silenced the ISG15 gene using specific siRNA sequences in the USP18-deficient cells and we treated them with IFN. As predicted, the amount of modified ADAR (p150) was significantly reduced after partial silencing of ISG15 (Fig. 3b). As the ultimate validation technique for our GlyGly-based ISGylome, we decided to perform a classic interactome analysis after immunofluorescence purification (IAP-MS) of endogenous ISG15 proteins in the same experimental conditions. After optimisation of the immunoprecipitation (IP) conditions, we could efficiently purify ISG15 and its conjugated substrates (Fig. 3c). Again, we did not observe an increase of ubiquitylated proteins in the inputs, and we did not immunoprecipitate ubiquitin or ubiquitylated proteins together with ISG15 (Fig. 3c, lower panel). In the same IP eluates, we observed an enrichment of modified HERCS, ADAR, PKR and IFIT3, all ISGylated in the GlyGly data, in the USP18 KO cells treated with IFN (Fig. 3d). Mass spectrometry analysis of the eluates showed enrichment of very similar proteins to the matching GlyGly peptide in the USP18 KO cells treated with IFN (Fig. 3e). From a total of 312 ISGylated proteins found to interact with ISG15 in a USP18-dependent manner (Data Table S3), 110 proteins overlapped when comparing the two techniques (Fig. S4A), primarily the ones showing a stronger enrichment in the volcano plot in Fig. 3e (highlighted and labelled in green) and in the comparative scatter (ISG15 interactome versus GlyGly peptidome) plot in Fig. 3f, upper right area. These complementary experiments validated our approach for analysis of the site-specific ISGylome using GlyGly peptidomics.

The double-stranded RNA-specific adenosine deaminase ADAR (adenosine deaminase acting on RNA) is an essential gene for tumour cells that express ISGs.5,46 Even more importantly, ADAR has been shown to be involved in resistance to ICB therapy and irradiation in preclinical models.5 The molecular basis of these observations is to the excessive accumulation of dsRNA induced by the absence of ADAR and the subsequent activation of the sensors for this innate ligand, mainly PKR and the axis DDX58/IFIH1, leading to cell death and tumour inflammation.5,46 As we found ADAR to be highly modified in the KO cells treated with IFN, we decided to check if in our cellular model ADAR is also essential upon activation of the IFN pathway. To do so, we silenced ADAR with specific siRNA sequences in HAP1 WT cells and we subsequently treated them with IFN. In a similar fashion to USP18 depletion, ADAR genetic inactivation also sensitises HAP1 cells to IFN (Fig. 4a). In addition to its effects on cell growth, we evaluated the activation of the dsRNA sensor PKR, measured by phosphorylation at threonine 446, in WT and USP18 KO cells treated with IFN for 0, 6, 24 and 48 h. Both, ADAR and PKR are ISGylated upon IFN treatment but only in the USP18 KO cells (Fig. 4b). ISGylation of PKR fits with its markedly higher phosphorylation in the KO cells compared with a relatively modest activation in the WT cells because ISGylation of PKR (although on different residues) has been previously reported to activate PKR.44 We identified 7 lysine residues in the sequence of ADAR that are potentially ISGylated (K433, K637, K763, K781, K798, K895 and K996; Fig. 4c). When highlighting these residues in the structure of the deaminase domain of ADAR2 bound to RNA, three of them seem to directly contribute to the binding of the enzyme to RNA (K637, K781, and K996; Fig. 4d). Finally, we measured the levels of the innate ligand dsRNA in our cells using specific antibodies against dsRNA by immunofluorescence as an indication of ADAR activity. ADAR enzymes are responsible for binding to double-stranded RNA (dsRNA) and converting adenosine
Fig. 4  **USP18-dependent ISGylation of ADAR leads to enzymatic activity inhibition and dsRNA accumulation.** ADAR knockdown using siRNA renders HAP1 WT sensitive to growth inhibition by type I IFN as shown by the growth curves in (a). b Immunoblots of a time-course experiment showing an increase in modified ADAR and PKR and activation of PKR (phosphorylated PKR in Thr 446) USP18-deficient cells treated with IFN for the indicated times. c A plot of the ADAR GlyGly-modified lysine residues and their ion intensities measured by MS. d dsRNA bound ADAR2 deaminase domain structure (PDB code 5ED2). Lysine residues corresponding to the ADAR1 GlyGly sites are represented with sticks. The Zn$^{2+}$ cation is represented as a grey sphere (top panel). Table showing the ADAR1 potential GlyGly sites indicating their domain location and their contribution to RNA binding. A score of 1 indicates the highest contribution and 3 the lowest and away from the RNA binding pocket. DD = deaminase domain. dsRBD = dsRNA binding domain (lower panel). e, f Analysis of the dsRNA levels after IFN treatment in WT and KO cells by immunofluorescence using specific antibodies ($n = 6$ per condition).
(A) to inosine (I) by deamination, thus reducing the intracellular levels of dsRNA. The results shown in Fig. 4e, f illustrate the accumulation of dsRNA in a time-dependent manner only in the USP18 KO cells treated with IFN for 0, 24 and 48 h. This accumulation matches well with the ISGylation levels of ADAR p150 seen in Fig. 4b and points at USP18 and ISG15 as key regulators of ADAR deaminase activity.

USP18 regulates MHC class I antigen presentation, PD-L1 expression and stimulation of a T cell functional response. In the USP18-dependent ISGylome and proteome of cancer cells treated with type I IFN, we observed ISGylated components of the antigen presentation pathway (Figs. 2f, g, and 3e). To test whether USP18 modulates antigen presentation and T cell stimulation, we...
USP18 regulates major histocompatibility complex class I (MHC-I) antigen presentation, PD-L1 expression and stimulation of a T cell response. Wild-type or USP18-deficient HAP1 cells were pre-treated with or without IFN-a, then pulsed with Melan-A26–35 peptide at the indicated concentrations and co-cultured with Melan-A-specific T cells. a Upregulation of activation markers on T cells was assessed by flow cytometry. Flow plots show representative examples of CD25 and CD137 expression on live CD8+ cells (T cells) following co-culture with the indicated peptide-pulsed HAP1 cells and the proportion of T cells expressing both CD25 and CD137 is plotted below. b T cell release of IFNγ into HAP1-T cell co-culture supernatants was measured by ELISA. Bar graph shows mean values from all experimental repeats, and for the statistical analysis, we applied two-way ANOVA tests, including multiple comparison testing via the Dunnett method available through the GraphPad Prism software. P value style is GraphPad: NS, P > 0.05; *P < 0.05; **P < 0.01; ***P < 0.001; ****P < 0.0001. Expression of c HLA-A2 and d PD-L1 on HAP1 cells following co-culture with T cells was measured by flow cytometry. Histogram plots of the data are shown above, and the plots below show the percentage of cells (c) expressing high levels of HLA-A2 (defined as greater than those expressed on non-peptide-pulsed cells not pre-treated with type 1 IFN and (d) expressing PD-L1. e PD-L1 expression on T cells following co-culture with the indicated HAP1 cells. Representative histogram plots are shown above, and the geometric mean fluorescence intensity (gMFI) of PD-L1 expression on T cells from each co-culture condition is plotted below. The results shown in this figure are representative of data from 1 of 3 experiments and the FACS shows information of three replicates pooled into one. Event counts on histogram plots have been normalised to unit area to enable comparison between samples.

Pulsed the HLA-A2 positive HAP1 cell line (WT and USP18 KO; with and without pre-treatment with IFN for 24 hours) with different concentrations of the HLA-A2-restricted Melan-A26–35 peptide and then co-cultured them with Melan-A specific CTLs for 18 h.37,47

Peptide-pulsed WT or USP18 KO HAP1 cells (+/− type I IFN pre-treatment) stimulated phenotypic activation of T cells (as assessed by upregulation of the activation markers CD25 and CD137) with equal efficiency (Fig. 5a). However, both with and without IFN pre-treatment, the USP18 KO cells triggered a more robust functional response by the T cells, as evidenced by greater T cell production of IFNγ following peptide recognition (Fig. 5b). Since the strength of the activating stimulus received by T cells determines the nature of their response, with phenotypic activation and production of chemokines such as MIP1β being more readily triggered than production of cytokines such as IFNγ,46 this suggests that USP18-deficient tumour cells exhibit superior antigenicity. Non-peptide-pulsed HAP1 USP18 KO cells expressed a similar level of HLA-A2 to the parental cells and HLA-A2 was equivalently upregulated on both following exposure to type 1 IFN (Fig. 5c; top left histogram plot). HLA-A2 was further upregulated on both WT and KO cells after pulsing with the Melan-A26–35 peptide and co-culture with T cells, likely in response to IFNγ secreted by the T cells (Fig. 5c; top right and lower histogram plots). However, a proportion of the peptide-pulsed cells exhibited downregulation of HLA-A2 to levels below those expressed on non-peptide-pulsed cells following co-culture with T cells, potentially because of HLA-A2 internalisation and/or degradation (Fig. 5c; top right and lower histogram plots). Notably, high levels of HLA-A2 expression were retained on a greater proportion of the USP18 KO cells than their WT counterparts (Fig. 5c, bottom graph), putatively contributing to their greater antigenicity.

Strikingly, we observed a strong upregulation of PD-L1 on both tumour cells (Fig. 5d) and T cells when USP18 was absent (Fig. 5e). This effect was antigen dependent, although the KO cells pre-treated with type I IFN presented a high basal expression of PD-L1 in the absence of peptide. PD-L1 protein expression in tumours has been described as a potential predictive biomarker for sensitivity to immune checkpoint blockade (ICB) with PD1/PD-L1 inhibitors.50 Also, we observed an upregulation of PD-L1 levels on the T cells when co-cultured with USP18 KO cancer cells, regardless of treatment. In contrast, very little upregulation of PD-L1 was noted on T cells stimulated with WT cells (+/− IFNγ), after addition of peptide (Fig. 5e). PD-L1 is expressed on activated T cells and is required for T cell conditioning and dendritic cell maturation.50 However, its expression on T cells has been mostly linked to inhibition of their responses.51 Other studies indicates that expression of PD-L1 on CD8+ and CD4+ T cells correlates with patient response to ICB, suggesting PD-L1 expression in CD8+ T cells as a prognostic marker in melanoma.9,52

USP18 deficiency sensitises cancer cells to radiation therapy (RT) Efficacy of local radiotherapy (RT) has been linked to the ability of the irradiated cells to repair lethal DNA damage but also to the induction of T cell-dependent cell killing. This immune attack has been linked to the induction of the type I IFN response after tumour RT10 and also seems to be ADAR-dependent.4 As shown above, USP18-decient HAP1 cells displayed greater cellular death following activation of IFN signalling by exposure to type I IFN in tissue culture. Thus, we hypothesised that loss of USP18 in cancer cell could also augment their sensitivity to ionising radiation. Indeed, USP18 KO cells exhibited a significantly lower clonogenic survival rate after radiation in tissue culture compared with WT (representative images in Fig. 6a and summarised in Fig. 6b). In addition, the average colony diameter of USP18 KO cells was smaller than that of WT (Fig. 6c) indicating that USP18 deficiency also results in greater growth delay after exposure to ionising radiation. Consistently, following a single dose of 5 Gy irradiation, the cell number ratio of WT over USP18 KO cells increased over time (Fig. 6d).
Deletion of the deISGylating enzyme USP18 enhances tumour cell radiosensitivity (IR). Many of these proteins are ISGs, and a number of them are part of a signature of genes that are upregulated in a subset of tumour cells, independent of immune infiltration. ISG-positive cancer cells are sensitive to ADAR loss, another ISG with deaminase activity, controlling the levels of dsRNA, an innate ligand. We identified USP18-dependent ISG15ylation of seven lysine residues in ADAR (K433, K637, K763, K781, K798, K895 and K996),
localised within the dsRNA binding (dsRBD) and deaminase domains (DD), which may potentially affect both enzymatic activity and substrate binding affinity. ADAR has been recently described as a major factor for resistance to immunotherapy and radiotherapy. 4 In our model, ADAR gene silencing using siRNA sequences sensitises cancer cells to IFN stimulation in a similar fashion as USP18 does. We could detect ISGylated ADAR by immunoblotting in USP18-deficient cells after 24 hours treatment with IFN, and this time point matched the time when the growth of the cells started to be inhibited by this cytokine. The described effects of ADAR loss in overcoming immunotherapy and radiotherapy have been linked to its ability to reduce the levels of innate ligand dsRNA and subsequent inhibition of the dsRNA sensors PKR, and MDAS/RIG-I, involved in growth inhibition and tumour inflammation, respectively. 4,5 ISGylation of PKR has been reported to activate its activity, and we could also see ISGylation and activation of PKR in the USP18 KO cells treated with IFN. Our data, therefore, suggested that USP18-dependent ISGylation of ADAR was inhibiting its activities, inducing the accumulation of dsRNA, resulting in further activation of PKR. Immunofluorescence analysis using specific dsRNA antibodies showed a significant and time-dependent accumulation of dsRNA in the USP18 KO cells after IFN treatment, indicating that USP18-dependent ISGylation under these conditions could inhibit ADAR activity.

In addition to ADAR, PKR, RIG-I and MDAS, we found other proteins involved in antigen presentation and resistance to immunotherapy, such as TAP1, GBP1, STAT1, IFIT1, PSMB10, PSMB9, GBP2, MAGE and PARP14, also regulated by USP18-dependent ISGylation. Many of these genes are part of processing MHC class I antigens, suggesting that the USP18 KO cells could possibly be more antigenic. Indeed, our results confirmed that USP18 KO cells had an enhanced ability to stimulate IFNγ secretion by Melan-A antigen-specific T cells, which was associated with elevated HLA-A2 expression levels. Interestingly, PD-L1 levels, a prognostic factor for ICB therapy when using PD1 inhibitors, were also higher in USP18 KO as compared to WT cells, and PD-L1 was also upregulated on the T cells following interaction with these cells. This is noteworthy as other studies have linked the expression of PD-L1 on CD8+ T cells, and CD4+ T cells with patient response to ICB, suggesting PD-L1 expression in CD8+ T cells as a prognostic marker in melanoma. As the innate immune response is activated by radiotherapy and ADAR recently described as an important factor in resistance to RT, 4,5 we decided to perform a second round of functional experiments consisting of irradiating our cancer cells and evaluating viability and clonogenicity after RT. The results clearly suggest that USP18 deficiency sensitises tumour cells to RT in a dose-dependent manner, most likely by interfering with end-stage IFN signalling.

Directing the immune system against tumours, also known as cancer immunotherapy, is supplementing or replacing more conventional forms of therapy as the first line of treatment for many tumour types due to striking curative effects in a subset of the treated patients. However, resistance mechanisms prevent this therapy from being 100% effective. The innate immune response has been described as one of the main resistance mechanisms against resistance to ICB and the discovery of factors involved in reactivation of the innate response, such as ADAR, has been a priority for the cancer research community. So far, inhibition of ADAR has been unfruitful. Our data suggest that USP18 inhibition represents a novel strategy to block ADAR functions, to activate the immunoproteasome/antigen presentation machinery and innate ligand sensors. As a consequence, this may help to overcome resistance to immunotherapy and RT, potentially turning ‘cold’ tumours into ‘hot’ tumours. 

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AUTHOR CONTRIBUTIONS
A.P.F., E.A., N.J., T.R.H., K.H., R.M., P.B., V.C. and B.M.K. directed this study. Most experiments were devised by A.P.F. and B.M.K. and carried out by A.P.F. M.S. helped, supervised, and assisted with the T cell experiments. T.P. helped, supervised and assisted with the T Cell experiments. J.C. helped, supervised and assisted with the radiation therapy experiments. G.V. performed immunoblot and helped with the published ISGyme comparison. H.G. carried out cell viability experiments. C.S.O. performed immunofluorescence experiments. A.D. generated the catalytically inactive USP18 mutant. H.C.S. assisted with the ISG-PA activity-based probe synthesis. H.J.P., B.P. and A.C. helped to establish the ISG15 immunoprecipitation method. L.D.S. and P.S. helped with ADAR structural analyses. C.G.L. provided support with the T cell experiments. A.P.F. wrote the paper with the help of B.M.K., P.B. and T.P. All authors commented on the text.

ADDITIONAL INFORMATION
Ethics approval and consent to participate HAP1 cells (WT and KO) are a kind gift from the laboratory of Sebastian Nijman and the HCT116 cell line was purchased from ATCC (ATCC-CCL-247).

Consent to publish Not applicable.

Data availability The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository with the data set identifier PXD018299.

Competing interests The authors declare no competing interests.

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