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Title:

Cellular senescence inhibits renal regeneration after injury in mice, and senolytic treatment promotes repair

One Sentence summary:

Senescent renal epithelia promote fibrosis and functional loss after injury whereas depletion improves regeneration in aged or irradiated kidneys.

Authors

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Abstract

The ability of the kidney to regenerate successfully after injury is lost with advancing age, chronic kidney disease and after irradiation. The factors responsible for this reduced regenerative capacity remain incompletely understood, with increasing interest in a potential role for cellular senescence in determining outcomes after injury. Here, we demonstrated correlations between senescent cell load and functional loss in human aging and chronic kidney diseases including radiation nephropathy. We dissected the causative role of senescence in the augmented fibrosis occurring after injury in aged and irradiated murine kidneys. In vitro studies on human proximal tubular epithelial cells, and in vivo mouse studies demonstrated that senescent renal epithelial cells produced multiple components of the senescence associated secretory phenotype including transforming growth factor beta-1, induced fibrosis and inhibited tubular proliferative capacity after injury. Treatment of aged and irradiated mice with the B-cell lymphoma (Bcl) 2/w/xL inhibitor ABT-263 reduced senescent cell numbers and restored a regenerative phenotype in the kidneys with increased tubular proliferation, improved function and reduced fibrosis after subsequent ischaemia-reperfusion injury. Senescent cells are key determinants of renal regenerative capacity in mice and represent emerging treatment targets to protect aging and vulnerable kidneys in man.

Introduction

Hundreds of millions of people worldwide live with chronic kidney disease (CKD), a condition characterised by progressive kidney fibrosis, renal functional decline and premature cardiovascular death (1). CKD prevalence is increasing, and aging adults (>65 years old) are known to be at particular risk (2). Aging adults and patients with CKD are also more susceptible to acute kidney injury (AKI), a life-threatening complication which itself can accelerate progression of underlying renal fibrosis and loss of function (3). Therefore, incompletely understood factors in the aging and chronically injured kidney promote fibrosis and inhibit renal regeneration. Understanding these processes and modifying them to protect vulnerable aging and injured kidneys represents an unmet research and clinical need.

Senescent cells (SCs) are permanently growth arrested and phenotypically altered cells which accumulate in vivo with aging, organ injury and after genotoxic insults such as ionizing radiation. Senescence is associated with worsened damage and reduced regenerative capacity following tissue injury (4-6). Recent evidence implicated SC accumulation as contributing to aging and age-related diseases in fast-aging and naturally aged mice, with depletion strategies extending healthy lifespan (7, 8). SCs have also been reported to be present in increased numbers in human and rodent kidneys with aging, CKD and after transplantation (9-12). Whether SCs actively contribute to the increased fibrosis and reduced regenerative capacity seen in aged and injured kidneys subject to further injury is a topic of major clinical and research interest(13).

SCs are attractive therapeutic targets as they remain metabolically active with constitutive nuclear factor kappa-light-chain-enhancer of activated B cells (NF- κ B) activation driving

expression of the senescence-associated secretory phenotype (SASP) with release of cytokines (Interleukin (IL)-6, Tumour necrosis factor (TNF- α), chemokines (chemokine (C-C motif) ligand (CCL)2, chemokine (C-X-C motif) ligand CXCL8) and pro-fibrotic transforming growth factor beta (TGF)- β (14). ABT-263, a potent B-cell lymphoma (Bcl)2/w/xL inhibitor, effectively depletes SCs when administered orally to irradiated or normally aged mice (15). Depletion reduces irradiation-induced premature haematopoietic aging and rejuvenates aged tissue stem cells (15). The efficacy of ABT-263 treatment on kidney regeneration and fibrosis after injury in aged animals and animals with prematurely senescent kidneys (via total body irradiation, TBI) remains untested.

In this study, we tested the hypothesis that senescent renal epithelia are key determinants of fibrosis and regenerative capacity after renal injury and can be therapeutically targeted. Senescent proximal tubular epithelial cells (PTECs) were identified in naturally aged mice, were also induced by radiation exposure in vitro and in vivo and were depleted using ABT-263. By doing so, we demonstrate the importance of epithelial senescence in kidney fibrosis, damage response, function and repair after injury, as well as the impact of their depletion on successful regeneration.

Results

Markers of renal senescence are increased in patients with age and chronic kidney disease. Using two separate human datasets of renal transcriptomic data available on the *Nephroseq* repository (16, 17), we determined expression of markers of permanent cell cycle arrest in healthy and diseased kidneys using *CDKN1A* (P21CIP1) and *CDKN2A* (P16INK4A) expression (Fig 1A). Both *CDKN1A* and *CDKN2A* were increased in diseased kidneys vs healthy controls (Fig 1B; *CDKN1A*: $p=5.52 \times 10^{-9}$, *CDKN2A*: $p=1.2 \times 10^{-8}$; based on median centred Log2 expression) and rose progressively with age, with *CDKN1A* expressed more highly (Fig 1C; $p=0.004$; *CDKN2A* $p=0.048$). Of note, only *CDKN1A* was significantly associated with falling kidney function in chronic kidney disease (measured by estimated renal glomerular filtration rate; eGFR) (Fig 1D; $p=0.0015$); validating *CDKN1A* as a marker of both age- and injury-related renal senescence. Multivariate regression demonstrated that age and eGFR were independently associated with changes in *CDKN1A* whereas the univariable models associations of age with *CDKN2A* was lost in the multivariable model (table S1). The transcriptome analysis of kidney biopsies in human CKD patients and healthy donors also identified upregulation in multiple transcripts of members of the SASP, fibrosis and BCL family members (Fig. S1)

Immunohistochemistry (IHC) staining of a local Edinburgh dataset performed in our United Kingdom Accreditation Service (UKAS)-certified histology lab demonstrated that nuclear P21CIP1 staining was induced in diseased kidneys versus healthy controls (Fig 1E; $p<0.05$). Immunohistochemistry for P16INK4A and P21CIP1 demonstrated minimal staining in the young healthy kidney but increased P21CIP1 expression was detected in disease [early glomerulonephritis, radiation nephropathy and in advanced renal fibrosis (Fig 1F)]. In each

case P21CIP1 staining appeared confined to tubular epithelia, in contrast to P16INK4A which was expressed at much lower concentrations in epithelial and interstitial cells (Fig 1G-H).

ABT-263 specifically targets senescent human renal proximal tubular epithelial cells (RPTECs). Human RPTECs exposed in vitro to 10 Gray (Gy) γ -radiation developed reduced total cell number ($p < 0.05$) and increased cell size ($p < 0.05$) compared to non-irradiated controls (Fig 2A-C), consistent with induction of senescence. Quantitative polymerase chain reaction (qPCR) analysis at d7 post-irradiation revealed significantly decreased proliferation as measured by *MKI67* (KI67) expression ($p < 0.005$), loss of *LMNB1* (Lamin B1; $p < 0.05$), increased cyclin dependent kinase inhibitors *CDKN2A* ($p < 0.05$) and *CDKN1A* (P21CIP1; $p < 0.005$) and increases in SASP markers *IL1B* ($p < 0.05$), *TNF* ($p < 0.05$) and *TGFBI* ($p < 0.0005$), *MMP1* (matrix metalloprotease 1) and neutrophil chemotactic factor, *CXCL8* (Fig 2D). Because ABT-263 targets BCL-2 family members, *BCL2* (BCL-2), *BCL2L1* (BCL-xl) and *BCL2L2* (BCL-w) were measured and all demonstrated increased mRNA expression in irradiated/senescent RPTECs vs control (Fig 2d; *BCL2*; $p < 0.01$, *BCL2L*; $p < 0.01$ and *BCL2L2*; $p < 0.01$). Irradiated RPTECs also increased senescence-associated (SA)- β -Galactosidase expression (Fig 2E, $p < 0.001$). Collectively these findings indicated that 10Gy irradiation induces senescence and its associated secretory phenotype in human RPTECs.

We then tested the specificity of ABT-263 to induce cell death in human senescent RPTECs. ABT-263 selectively killed senescent cells in a dose-dependent manner, with higher concentrations of ABT-263 (1 μ M and 5 μ M), reducing viable SCs to 2% and 0% (relative to vehicle-treated SCs) (Fig 2F). There was no toxicity to non-irradiated cells at even the highest concentrations of ABT263 (Fig 2F), which was unexpected, as previous studies had

found some toxicity at higher concentrations in non-senescent human renal epithelial cells(15).

Aged mice have higher expression of SASP factors and worsened fibrotic responses after IRI. Baseline expression senescence/fibrosis in kidneys of old and young mice were measured (Fig 3A). qPCR analysis revealed upregulation of senescence-associated *Cdkn1a* (P21CIP1; $p<0.05$), *Cdkn2a* ($p<0.005$), as well as SASP genes (*Il6*; $p<0.05$, *Tnf*; $p<0.01$, *Tgfb1*; $p<0.05$) in old compared with young kidneys (Fig 3B). Fibrosis, as measured by Collagen I immunofluorescence (IF) was higher in old kidneys ($p<0.01$; Fig 3C) with an equivalent rise in total Collagen seen on picrosirius red (PSR) staining ($p<0.05$; Fig 3D). γ H2A.X staining showed evidence of increased DNA damage and DNA damage response (DDR) in the outer medulla and cortex of old kidneys compared to the young kidneys (Fig 3E). Lamin B1 antibody IF was performed, confirming that Lamin B1 was lost in kidney tubular nuclei with age (Fig 3F). To assess the impact of age on renal regeneration after injury the outcome of unilateral ischemia reperfusion injury (IRI) was tested in old and young mice (Fig 3G), demonstrating that old mice develop greater fibrosis after identical durations of injury ($p<0.005$; Fig 3H), with IRI resulting in an increase in expression of senescence-associated *Cdkn1a* by qPCR ($p<0.01$; Fig 3I).

Clearance of SC in aged mice protects the kidney after AKI. To test whether pre-existing SCs present at increased numbers in old animals promote fibrosis and inhibit regeneration after subsequent AKI, old female mice were treated with ABT-263 to deplete SCs before unilateral IRI surgery (Fig 4A). 5 weeks after IRI kidneys were harvested and analysed. Senolytic pre-treatment significantly reduced the numbers of senescent cells present in the uninjured injury ($p<0.05$; Fig S2A) and reduced post-IRI fibrosis ($p<0.01$; Fig S2B-C)

Key markers of senescence, SASP release and *Bcl2* family transcripts were measured by qPCR, compared to old naïve kidneys (with naïve kidney expression shown by the red line, Fig. 4B and S2D). These demonstrated induction of multiple senescence-associated transcripts, with *Cdkn1a* expression reduced by ABT-263 treatment (Fig 4B) and *Bcl2l2* (BCL-W; $p < 0.05$) also showing reduction. Post-ischaemic kidney weight (vs the contralateral kidney from the same mouse, Fig 4C) was increased in senolytic-treated compared to control treated animals, although this remained below healthy weight (shown by red line). Given that IRI leads to tubular loss (19), this implicated senescent cells in impeding tubular recovery and survival after injury, with senolytic treatment potentially restoring regenerative capacity by increasing proliferative capacity in remaining renal tubules (20). To dissect these processes, further experiments were undertaken.

Kidney Injury Molecule 1 (KIM-1/*Havcr1*) is upregulated acutely in damaged proximal tubules in humans (21) and chronic expression in CKD predicts likelihood of progression to end stage-renal disease (ESRD) in man (22). We quantified KIM-1 protein 5 weeks post-IRI as a marker of ongoing tubular injury and progressive fibrosis in diseased kidneys. KIM-1 was reduced in ABT-263-treated IRI vs vehicle-treated IRI kidneys ($p < 0.01$; Fig 4D), illustrating that senescent tubular cells drive chronic KIM-1 release in diseased kidneys, and their depletion in vivo was safe and opposed ongoing injury. Haematoxylin and eosin (H&E) staining revealed improved renal architecture in ABT-263 vs vehicle-treated mice (Fig S2E).

Kidney sections were then stained with *Lotus tetragonolobus* lectin (LTL), a marker of differentiated renal tubules (23) and the proliferation marker Ki67 (Fig 4E-F). IRI induces loss of tubules (19). Proliferating KI67+ LTL+ tubules were counted (Fig 4F; white arrows in

Fig 4E). ABT-263 pre-treated old IRI kidneys exhibited significantly increased tubular proliferation compared to control ($p < 0.01$; Fig 4F). Thus, senescence inhibits epithelial proliferation in response to injury. As TGF- β 1 is a recognised driver of renal fibrosis post-IRI (24-26) and is produced by SC (27, 28), this protein was measured in the kidney. Senolytic treatment reduced TGF- β 1 in diseased kidneys ($p < 0.05$; Fig 4G), consistent with a role of senescent epithelia in controlling renal TGF- β 1 through direct production \pm recruitment of TGF- β 1 producing cells. TGF- β 1 is thought to drive renal fibrosis in part through mediating macrophage infiltration and switching to a pro-fibrotic CD206 $^{+}$ phenotype (26, 29). ABT-263 treatment reduced the overall number of F480 $^{+}$ macrophages ($p < 0.05$; Fig S2G-H) and percentage of F480 $^{+}$ /CD206 $^{+}$ macrophages in the post-IRI diseased kidney ($p < 0.01$; Fig S2I). Therefore, senescence affects macrophage phenotype in post-IRI kidney disease, increasing pro-fibrotic CD206 $^{+}$ macrophages. Whereas SA- β -galactosidase was not reduced (Fig S2J), a marked reduction in γ H2A.X foci in the cortex ($p < 0.05$; Fig S2K) and increase in Lamin B1-positive nuclei ($p < 0.01$; Fig S2L) were both noted with ABT-263 treatment .

To establish whether the improvement in multiple markers of senescence and fibrosis translated into enhanced renal functional recovery after IRI, further experiments used a bilateral IRI injury to generate a model of functional kidney injury and impairment (Figure 5A). Aged mice treated with ABT-263 prior to bilateral IRI had significantly improved renal function measured by serum Cystatin C ($p < 0.01$ at both d7 and 14; Fig 5B) and reduced numbers of p21 CIP1 $^{+}$ senescent cells prior to injury ($p < 0.05$; Fig 5C). This functional protection was accompanied by reduced renal fibrosis by both PSR staining ($p < 0.01$; Fig 5D-E) and Collagen I analysis by immunofluorescence ($p < 0.01$; Fig 5F-G). Hence, senescent cell depletion improves the subsequent structural and functional regenerative capacity of the naturally aged kidney.

Total body irradiation induces cellular senescence and fibrosis in the murine kidney in vivo. To distinguish the specific effects of senescence from those of the general aging phenotype, γ -irradiation was used to induce senescence in the young murine kidney (15). Young C57BL/6 mice underwent 7Gy total body irradiation (TBI) with tissue harvest 8 and 16 weeks later (Fig 6A), with imaging of whole kidneys to determine patterns of DNA damage by γ H2A.X staining (Fig 6B-C). qPCR analysis of kidneys showed that the senescence marker *Cdkn1a* (P21CIP1) increased over time after TBI reaching significance by 16 weeks after TBI ($p < 0.05$; Fig 6D) with a similar pattern seen for *Cdkn2a*. Analysis of markers of the SASP, fibrosis and *Bcl2* family (measured by qPCR; Fig 6E) and the senescence marker SA- β -Gal (Fig 6F) all had higher means at week 16 post irradiation, but with a high biological variation.

IF staining demonstrated increased P16INK4A protein expression by 16 weeks post-TBI ($p < 0.01$; Fig 6G). In line with previous observations in the lung after TBI (15), senescence in the kidneys rose modestly by 8 weeks post-TBI, reaching significance by 16 weeks, a finding compatible with ongoing SASP activation driving progressive late senescence distinct from the initial genotoxic insult. Consistent with these findings, TBI drove initiation of the DDR in the diseased kidney, with maximal increases seen in the cortex, mirroring findings in aged kidneys (Fig 6H). Collagen I increased at 16 weeks post-TBI ($p < 0.01$; Fig 6I) with a similar pattern seen on PSR staining (Fig 6J).

To test the sensitivity of irradiation induced SCs to senolytic therapy, mice were treated with ABT-263 before kidney harvest at 20 and 25 weeks post-TBI (Fig S3A). *Cdkn1a* (P21CIP1) mRNA was significantly decreased with senolytic treatment as measured by qPCR ($p < 0.05$;

Fig S3B). Whilst no change in *Cdkn2a* transcript levels was detected P16INK4A protein abundance also fell post-ABT-263-treatment (Fig S3C-D). *Lmnbl* was found to increase in the post-TBI kidney in ABT-263 treated mice (Fig S3E) at 20 weeks post-TBI. ABT-263 reduced evidence of the DDR in the irradiated kidney, with a reduction in positive γ H2A.X positive nuclei in the cortex of treated kidneys ($p < 0.05$; Fig S3F). Fibrosis as measured by PSR staining for collagen did not differ between groups, indicating that ABT-263 had no detectable impact on established fibrosis (Fig S3G).

Irradiation worsens fibrotic responses to subsequent renal injury, which is prevented by ABT-263 pre-treatment. To dissect the effect of senescence on subsequent injury responses from generic aging effects, we compared post-IRI kidney disease outcomes in young irradiated mice with increased numbers of senescent cells (+/- ABT-263, Fig 7A).

PSR staining for collagen showed that previous TBI increased fibrosis in post-IRI kidney disease, which was prevented with senolytic treatment (Fig S4A-B). qPCR analysis of the profibrotic factors *Colla1*, *Col3a1*, *Fnl* (Fibronectin 1) and *Acta2* (α -SMA) demonstrated that post-IRI fibrosis/matrix deposition was exacerbated by prior TBI, with ABT263 pretreatment providing protection (Fig S4C-F). IRI induced renal *Cdkn1a* (P21CIP1) and *Cdkn2a* expression in age-matched control mice (in the young range ~30 weeks old), showing that senescence is induced in young kidneys after IRI, in agreement with published reports (6, 30). However, these were higher in pre-irradiated mice ($p < 0.05$) unless pre-treated by ABT-263 ($p < 0.01$) (Figs 7B and S4G). A similar pattern was found in P16INK4A protein abundance (IF; Fig S4H).

SA- β -Galactosidase quantification in diseased kidneys post-IRI confirmed that senescence was further increased by previous TBI and was reversed with ABT-263 (Fig S4I). qPCR showed that SASP markers (*Il6*, *Tnf*, and *Tgfb1*) also increased in post-IRI kidney disease, were higher with irradiation, and reduced with senolytic treatment (Fig S4J-L). *Bcl2* was also reduced by ABT-263 treatment in irradiated animals (Fig S4M). There were significantly more Lamin B1 positive tubular nuclei in the kidneys of irradiated mice pre-treated with the senolytic before IRI ($p < 0.01$; Fig S4N). γ H2A.X quantification post-IRI confirmed that DNA damage and DDR were further increased by previous TBI and were reversed with ABT-263, with maximal changes seen in the renal cortex (Fig S4O-P).

Post-IRI kidney disease decreases kidney mass due to destruction of tubules without appropriate repair. By weighing the post-IRI and contralateral kidneys we demonstrated that IRI reduces kidney weight, with prior irradiation causing greater weight loss. ABT-263 pretreatment led to significantly less weight loss reflecting improved tubular regenerative capacity ($p < 0.01$; Fig 7C; dashed line represents healthy weight ratio). We also examined ongoing renal damage by analysing chronic KIM-1 expression in the diseased kidney post-IRI. KIM-1 was elevated after irradiation and lowered by ABT-263 administration ($p < 0.01$; Fig 7D).

LTL⁺ tubules were co-stained with Ki67 to measure proliferation (Fig 7E). ABT-263 treatment increased numbers of healthy LTL⁺ tubules present post-IRI (Fig 7F). Many LTL⁺ cells had Ki67⁺/DAPI⁺ nuclei in post-IRI kidney disease representing proliferating interstitial cells (Fig 7E, red arrows). The number of Ki67⁺ nuclei associated with LTL⁺ tubules were counted (Fig 7E, yellow arrows). ABT-263 administration significantly increased Ki67⁺ tubules ($p < 0.05$; Fig 7G), indicating that senescence impairs tubular

regeneration after injury, with senolytic treatment restoring regenerative capacity to previously irradiated kidneys. Dual staining with Ki67 and p21CIP1 demonstrated that these two markers were mutually exclusive, confirming that Ki67 did not reflect the amount of senescence (Fig S4Q).

TGF- β 1 protein, as measured in injured kidney lysates, was highest after irradiation and was reduced with senolytic treatment ($p < 0.05$; Fig 7H), in agreement with reductions observed in transcript expression (Fig S4L) and in aged mice (Fig 4G), indicating that SCs are a major determinant of renal TGF- β 1. Due to the effect of TGF- β 1 on macrophages post-IRI, we next investigated macrophage numbers and phenotype. F4/80 IHC and *Cd68* qPCR analysis (Fig S5A-C) confirmed an increase in macrophage influx after IRI post-TBI, compared to non-irradiated kidneys. Senolytic treatment reduced CD206+ pro-fibrotic macrophages in the IRI kidney of TBI mice ($p < 0.05$; Fig S5D).

Last, we undertook bilateral IRI studies to establish whether senescent cell depletion restored functional regeneration in the young, irradiated kidney (Fig 8A). These studies demonstrated that irradiated mice treated with ABT-263 prior to IRI had improved renal function measured by serum Cystatin C ($p < 0.01$ both for d7 and d35; Fig 8B) and reduced numbers of p21CIP1+ senescent cells after injury ($p < 0.01$; Fig 8C). This functional protection was accompanied by reduced renal fibrosis by both PSR staining (Fig 8D-E) and Collagen I analysis by IF (Fig 8F-G). Dual staining of P16INK4A with Collagen I and p21CIP1 with Collagen I demonstrated that markers of both arms of the senescence induction pathway correlated with collagen deposition in the immediate surrounding tissue of the kidney (Fig 8H-I).

Of note, age-matched young non-irradiated mice were included in all the groups above as additional controls. In all cases, ABT-263 therapy had no impact on these mice compared to vehicle treatment in any parameter studied (Figs S6A-G), so for simplicity they have otherwise been omitted from the presented data. This finding illustrates that in the absence of increased numbers of pre-injury senescent cells ABT-263 exerts no other protective effect on non-senescent cells such as epithelia, mesenchyme or leukocytes.

The presence of senescent cells impaired structural and functional recovery after injury and promoted fibrosis in both naturally aged and young irradiated mice. Furthermore, pre-injury depletion of senescent cells improved renal structural and functional regenerative capacity.

Discussion

Aged patients are known to be at elevated risk of acute kidney injury and subsequent progressive chronic kidney disease (3). Several studies have shown correlations between increased senescence within the kidney and progression of CKD, or with adverse outcomes after transplantation (9, 31-35). Whether senescent cells themselves directly influence impaired repair and drive fibrosis after subsequent injury remains a key unanswered question. Determining the safety and efficacy of senescent cell depletion prior to renal injury would clarify the contribution of senescent cells to repair and fibrosis and would be of translational importance for aged patients at risk of AKI and CKD.

Whereas aged patients are most susceptible to AKI and CKD, murine models of AKI and CKD are typically undertaken in young healthy animals. In this study, we tested the impact of senescence on renal regenerative capacity by using both naturally aged and young irradiated mice, demonstrating that cellular senescence increased fibrosis and inhibited regeneration following AKI. Pre-treatment with the senolytic ABT-263 reduced senescent cell number, augmented epithelial proliferation and reduced deposition of fibrosis in the aftermath of renal injury in both aged and young irradiated mice, similar to healthy young mice. These findings are consistent with and extend previous interventions with senolytics on the uninjured kidney (7, 18). Based on these data and supporting evidence from the literature that senescent epithelial cell growth arrest suppresses regenerative capacity (36, 37) we propose that senescent cells in the kidney represent a key determinant of renal regenerative capacity in vivo. These findings mirror those recently reported in other organ systems including the lung and liver (38, 39). SASP factor release has been shown to activate myofibroblasts to secrete matrix with TGF- β 1, a prime candidate as a known fibroblast activator and mitogen (40). In

this study, TGF- β 1 decreased after IRI (RNA and protein) with senolytic pretreatment, implicating senescence as a major regulator of TGF- β 1 expression and release.

These data support the hypothesis that chronically senescent cells are deleterious to longer term renal homeostasis and function. In the models of aging and injury studied, senescent cell depletion pre-injury improved subsequent renal regenerative capacity and reduced fibrosis. Further studies would be indicated including assessment of long-term pharmacological treatment of senescence on kidney structure and function, and trials of clinically licenced agents capable of suppressing SASP factor release (41). Given the reported importance of senescence in acute wound repair it will be important to assess the optimal timing of senescent depletion, as the safety and efficacy of depletion in the aftermath of acute injuries such as renal transplantation and post-surgical AKI remains unknown.

ABT-263 has been found to be selectively toxic to senescent cells via its actions on Bcl2/w/xL (15), but prolonged dosing has been complicated by thrombocytopaenia occurring as an on-target effect of Bcl-2 inhibition (42). Importantly, the selectivity of ABT-263 for senescent human epithelia in vitro could facilitate its use in carefully selected patients, where the limited duration of dosing required for senolysis minimises bleeding risk. In the work presented here ABT-263 was safe and efficacious in senescent cell depletion and preservation of renal regenerative capacity with no gastrointestinal distress, toxicity or bleeding problems detected. On the basis of the results reported here, approval has been granted for a pre-clinical study to quantify the efficacy and safety of ABT-263 administration to ex-vivo human kidneys that have been rejected for transplantation in man. Should efficacy and safety in human kidneys be similar to the results in experimental animals, further studies should

investigate treating kidneys prior to transplantation, and high-risk patients with CKD prior to surgical procedures with high risks of AKI.

There are several limitations to this study which should be addressed in future research. Whereas two distinct modes of senescent cell generation (irradiation and natural aging) were studied, this does not encompass the full range of senescence-generating stimuli present in human aging and kidney disease. Our study focused on the role of senescent cells present prior to a further renal injury and did not address the therapeutic efficacy of post-injury senescent cell depletion. Finally, our study used ABT-263 as our senolytic agent for all in vivo work, and the efficacy of alternative senolytic drugs with different modes of action remain to be determined in both murine models of disease and in human patients. The findings presented here support further studies using alternative senolytic drugs, timings of administration and modes of senescence induction to establish the breadth of potential benefits for senescent cell depletion.

This work demonstrates cellular senescence to be a key determinant of renal regenerative capacity and a mediator of post-injury fibrosis and function. Furthermore, our data show that senolytic treatment may be efficacious as an intervention in populations with pre-existing SCs including the elderly, those with CKD and previously irradiated patients. With several agents, including ABT-263, entering clinical trials, there is the potential for senolysis to fill an unmet need for those with elevated senescence who are heightened risk of progressive fibrotic renal disease.

Materials and Methods

Study Design

This study addressed the hypothesis that senescent cells present in the aged and young, prematurely senescent kidney negatively impact the ability of the kidney to regenerate from injury, and that their therapeutic depletion would therefore be protective. A series of controlled laboratory experiments were performed to determine the degree of fibrosis present in the kidney after injury with and without treatment with ABT-263 *in vivo*. All studies were conducted in accordance with ARRIVE guidelines (Animal Research: Reporting of In Vivo Experiments) (43). Based on previous work a two-sided two sample test was used to generate group sizes for all studies. All studies were performed at the University of Edinburgh, UK, with data collection performed at pre-specified endpoints unless limited by animal distress as identified by facility staff and/or veterinary surgeons. No available data was excluded from the analysis. Primary and secondary endpoints were pre-specified. Each mouse represented one experimental unit. In all experiments, there was randomised allocation to groups. Drug administration and surgical timing was rigorously rotated across all groups. Tissue was collected by two staff members blinded to the treatment groups, using a coding system for each animal. Semi-blinded staining and image analysis was undertaken with unblinding only being performed after all data acquisition was complete.

Human biopsies

After appropriate ethical approval from Tissue Governance at the Royal Infirmary of Edinburgh, sections were obtained from renal biopsy tissue from n=10 renal biopsies, selected to encompass a range of renal function spanning from normal to stage IV chronic kidney disease. Slides were cut and immunohistochemistry performed for P16INK4A and

P21CIP1. Staining was performed in the UKAS (United Kingdom Accreditation Service) certified laboratory of Edinburgh Royal Infirmary stained on Leica Bond III autostainers using Bond Refine detection kits (Peroxidase-DAB detection). Antibodies to P16INK4A were produced by Cintech, supplied by Roche/Ventana and used at 1:1 dilution. Antibodies to P21CIP1 were produced by Leica (code NCL-L-WAF1(P21)) and used at 1:20 dilution. Both underwent high pH antigen retrieval for 20 minutes. Transcriptomic datasets were obtained from published and publicly available resources from 95 renal transcriptomes from eligible studies in Nephroseq (University of Michigan, Ann Arbor, MI) (16, 17).

Cell culture

Human renal proximal tubular epithelial cells (hRPTEC; ATCC) were maintained in Dulbecco's Modified Eagle Medium (DMEM)-F-12 + Glutamax-1 supplemented with hTERT immortalised RPTEC growth kit (ATCC) and 50mg/ mL Geneticin (Gibco-life Technologies). Cells were grown at 37°C in 5% CO₂. hRPTEC cells were plated at 2 x 10⁵ cells per well of a 6-well plastic culture dish. After overnight culture, these cells were exposed to 10 Gy radiation. Cells were incubated for 7 days with media change every 3-4 days. For senolytic treatment with ABT-263 100,000 PTECs were seeded onto a 12-well plate. The following day, they were exposed to 10Gy Gamma radiation. For ABT-263 treatment, on day 5, 0.1, 1 and 5uM ABT-263 was added to the culture (replenished every 48 hours) until 12 days post-irradiation when the cells were washed with PBS. Cells were analysed on day 14.

Random fields of view within the plate (at x20 magnification) were captured using light microscopy on an Evos microscope. Scaling was performed using a graticule. Lines were drawn around 5 random cells per well, with area calculated using ImageJ (NIH, USA) (44).

For each well, multiple random fields of view were captured for analysis and the results averaged prior to cell lysis in trizol for RNA extraction.

Senescence associated (SA)- β -Galactosidase staining

Staining was performed as per manufacturer's instructions (Cell Signalling Technology, London, UK). Briefly, samples were fixed using reagents provided before staining (pH 6), in low CO₂ incubators. RPTECs were imaged after 3 hours of incubation. For kidney, frozen sections were air dried before staining for 12 hours. Nuclear fast red was used to counterstain slides, before dehydration and mounting. To analyse RPTEC staining, total cells per field were counted and then SA- β -Galactosidase⁺ cells (blue) were counted, and % SA- β -Galactosidase⁺ calculated. For kidney sections, fields of view at x10 magnification were captured. % positivity (areas of blue) was calculated over multiple random fields of view using ImageJ imaging software.

Experimental mice

All mice were on either a C57BL/6 or a mixed genetic background. All experiments comprised male mice, unless otherwise stated. Ages ranged from 6-8 weeks for young and 18-24 months for old. Mice were bred and maintained at the University of Edinburgh or purchased from Harlan, UK. Paraffin embedded samples of kidney tissue from P16INK4A WT and P16INK4A KO mice were sourced from our collaborators Dr Krimpenfort at the Netherlands Cancer Institute and Professor Melk and Dr Schmitt at Hannover Medical School. All animal work was compliant with IACUC guidelines, conducted in accordance with the UK Government Animals (Scientific Procedures) Act 1986, in conventional barrier unit facilities with conventional bedding, 12:12h light:dark cycle, ambient temperature control and access to food and water ad libitum and was approved by the University of

Edinburgh Ethical Review Committee.

ABT-263 senolytic treatment

ABT-263 (Chemgood, VA, USA) was prepared in 10% Ethanol, 30% polyethylene glycol (PEG) 400 and 60% Phosal 50PG. ABT-263 was administered by gavage (50 mg/kg/day) daily for 2 cycles, each of 7 days.

Mice that had undergone TBI were treated with ABT-263 or vehicle at 15 and 18 weeks post-TBI, or at an equivalent age for age matched controls. At week 20, mice underwent unilateral ischaemia-reperfusion injury surgery (IRI) or were culled for tissue analysis. A subset of mice that had undergone TBI were treated with ABT-263 or vehicle at 15, 18 and 35 weeks post-TBI. ABT-263 was administered by gavage (50 mg/kg/day) daily for 3 cycles, each of 7 days. At week 37, mice underwent bilateral IRI for kidney function measurement.

Ischaemia Reperfusion Injury (IRI)

IRI surgery was performed as previously described (45). Anesthesia was induced with 2% iso-fluorane. Buprenorphine analgesia was administered subcutaneously. A mid-line laparotomy was performed and the left renal pedicle identified and clipped using an atraumatic clamp for 15 min. During the ischemic period, body temperature was maintained using a heating blanket with homeostatic control (Harvard Apparatus) via a rectal temperature probe. After clamp removal, peritoneal closure with 5/0 suture and skin closure with clips was performed prior to subcutaneous administration of 1mL 0.9% saline. The right kidney was uninjured in some surgeries (unilateral IRI) and is referred to as the contralateral kidney. In later experiments, both kidneys were clamped (bilateral IRI). Animals were maintained for 2-5 weeks post-IRI before tissue harvest. Kidneys were weighed at time of sacrifice. Tissue was fixed in methcarn (Methanol 60%, Chloroform 30%, Glacial Acetic

Acid 10%) or snap frozen in liquid nitrogen/dry ice. In later experiments, tissue was also fixed in 4% paraformaldehyde, as this was found to be optimal for p21CIP1 antibody staining allowing us to carry out IHC for this extra marker, which was not possible in earlier experiments.

Murine irradiation

Young mice were exposed to a sub-lethal dose (7 Gy) of total body irradiation (TBI) using a closed caesium-137 gamma source. Age-matched control mice were not irradiated but housed in the same facility and conditions. A subset of irradiated mice were culled 8, 16, 20 and 25 weeks after exposure for tissue analysis of radiation-induced renal senescence.

Immunohistochemistry (IHC)

Picrosirius red: For picrosirius red (PSR) staining, sections were deparaffinised and rehydrated before treatment in haematoxylin for 8 minutes. After washing, these were stained in picrosirius red for 1 hour, before acidified water washing, dehydration and mounting.

H&E: sections were deparaffinised and rehydrated before staining with haematoxylin and eosin, before acidified water washing, dehydration and mounting.

DAB Staining: Tissue was fixed in methacarn overnight, changed to 70% ethanol and processed. Briefly, paraffin embedded tissue sections (5 µM) were deparaffinised and rehydrated. Antigen unmasking was undertaken (see table S2). Endogenous peroxidase was quenched with 3% H₂O₂ (Sigma Aldrich) for 15 minutes, before avidin/biotin (Vector) and protein blocking (DAKO) for 10 minutes each. Slides were then incubated with antibody overnight at 4°C. Species-appropriate biotinylated secondary antibodies (1:200) were added and staining developed using the Vector ABC kit and 3,3'-diaminobenzidine (DAB).

To investigate the distribution of DNA damage/senescence in young, aged and irradiated kidneys, pre- and post-IRI, γ H2A.X IHC was carried out. Images of whole kidneys were collected as .CZI files on the Zeiss AxioScan. This allowed visualisation of full kidney sections using QuPath(46). Sections were divided into Inner Medulla, Outer Medulla and Cortex, and areas within these analysed for % γ H2A.X positivity.

Immunofluorescence (IF)

For IF staining, paraffin embedded tissue sections were deparaffinized and rehydrated. Antigen unmasking was undertaken (see table S2). Slides were incubated with protein block for 30 minutes before addition of the primary antibodies overnight at 4°C. Secondary antibodies are specified in table S2 (1:200, 1:500 or 1:1000 dilution). The slides were mounted with Vectashield with DAPI (Vector).

Kidney sections were visualized and captured using the Zeiss Axioskop 2mot+ or on an Axioscan slidescanner. % area of positivity was calculated in ImageJ or in QuPath (46). In light of concerns about the specificity of antibodies to P16INK4A, staining performed in normally aged and post-IRI kidneys from P16INK4A $-/-$ and WT mice confirmed that AHP1488 detected no p19ARF protein in aged or injured kidneys (Fig S7). To measure correlation between SCs and collagen staining, dual IF was carried out for the P21CIP1 or P16INK4A proteins (red fluorescence) and COL I (green fluorescence) on all relevant kidney sections. Images of these kidneys were collected as CZI files on the AxioScan. This allowed visualisation of full kidney sections in the QuPath programme. % positivity for p21CIP1 and P16INK4A proteins (red) and Col I (green) was assessed from 3-7 random fields per kidney (ImageJ macro). These values were collated and analysed together to assess any linear correlation. A scatter plot was generated, a trendline added and, R^2 and P values calculated in

Microsoft Excel. For Lamin B1 calculations, total tubular (LTL+) DAPI+ and Lamin B1+ nuclei were counted per field and % Lamin B1+ nuclei per tubule calculated. For *Lotus tetragonolobus* lectin (LTL)/Ki67 dual staining, total LTL+ tubules and LTL+ tubules containing Ki67+ nuclei were counted and % Ki67+ tubules calculated.

RNA extraction and qPCR

RNA was extracted from human PTECS or murine kidneys using TRIZOL (Invitrogen, Carlsbad, CA) according to the manufacturer's instructions and reverse transcribed to cDNA (Qiagen Quantitect Reverse Transcription Kit). qPCR was carried out using VWR Perfecta qPCR mastermix with TAQman[®] gene expression assays on a Roche Lightcycler 480 using standard protocol.

For human RPTEC cells, the TAQman[®] gene expression assays used were glyceraldehyde 3-phosphate dehydrogenase (Hs02758991_g1 GAPDH), hypoxanthine-guanine phosphoribosyltransferase (Hs02800695 HPRT1), peptidylprolyl isomerase A (Hs04194521 PPIA), MKI67 (Hs04260396_g1, KI67), Laminb1 (Hs01059210_m1 LMNB1), *Cdkn2a* (Hs00923894_m1 CDKN2A), P21CIP1 (Hs00355782_m1 CDKN1A), Interleukin (Il)-1b (Hs01555410_m1 IL1B), Il6 (Hs00174131_m1 IL6), tumour necrosis factor (Hs00174128_m1 TNF), transforming growth factor (Hs00998133_m1 TGFB1), Cxcl8 (Hs00174103_m1 CXCL8), Matrix metalloproteinase 1 (Hs00899658 MMP1), Bcl (B cell lymphoma)-2 (Hs04986394_s1 BCL2), Bcl-xl (Hs00236329_m1 BCL2L1), Bcl-w (Hs01573809_g1 BCL2L2), prob FOXO4 (Hs00172973 FOXO4). mRNA expression was normalized for GAPDH, HPRT1 and PPIA (average) expression and presented as fold increases over day 7 non-irradiated cells analysed in parallel.

For murine kidneys, the following TAQman[®] gene expression assays were used: House keeping genes: hypoxanthine-guanine phosphoribosyltransferase (Mm03024075_m1 Hprt), Gapdh (Mm9999915_g Gapdh), Peptidylprolyl Isomerase A (Mm0234230 PPia) Phosphoglycerate kinase 1 (Mm00435617_m1 Pgk1); Senescence and SASP markers: *Cdkn2a* (Mm00494449_m1 Cdkn2a), P21CIP1 (Mm04205640_g1 Cdkn1a), Il-6 (Mm00446190_m1 Il6), Tnf (TNF; Mm00443258_m1 Tnf), transforming growth factor (Tgf; Mm01178820_m1 Tgfb1), Lamin B1 (Mm00521949 Lmnb1) BCL-2 family members: Bcl-2 (Mm0477631 Bcl2), Bcl-x1 (Mm00437783 Bcl2l1), Bcl-w (Mm0043054 Bcl2l2); Macrophage marker: CD68 (Mm03047343_m1 Cd68), Fibrosis markers: collagen (col)1a1 (Mm00801666_g1 Colla1), Col3a1 (Mm01254476_m1 Col3a1), fibronectin (Mm01256744_m1 Fn1), α -smooth muscle actin (Mm00725412_s1 Acta2). mRNA expression was normalized for HPRT or average HK expression in the case of old IRI tissue and presented as fold increases over naïve or age-matched control analysed in parallel.

ELISAs

TGF- β 1 ELISA: Protein was extracted from whole kidneys using complete RIPA buffer containing protease inhibitors (Santa Cruz). Briefly, kidney tissue was weighed and then placed in 300 μ l RIPA buffer with a 5 mm steel ball in a 2 ml eppendorf tube. The tissue was disrupted by high-speed shaking using a tissue lyser (Qiagen). Samples were placed on ice for 10 minutes before centrifugation for 20 minutes at 10,000g at 4°C prior to supernatant collection. Latent TGF- β 1 was activated using 1N HCL for 10 minutes, neutralization with 1.2N NaOH/0.5M HEPES, then analysed on a TGF- β 1 ELISA (R&D; DuoSet ELISA kit) according to the manufacturer's instructions.

Cystatin C ELISA for Kidney Function: Tail vein blood was recovered from mice that had undergone bilateral IRI 7, 14 or 35 days previously in an equal volume of 4% Citrate buffer before centrifugation at 5000g for 5 minutes at 4°C. 2ul of the clear phase was transferred into 250ul PBS +/- with 0.5% BSA. The mouse cystatin C ELISA (R&D; DuoSet ELISA kit) was carried out according to the manufacturer's instructions.

Statistical Analysis

Data were analysed with Microsoft Excel, Prism software (GraphPad) or R version 4. Multivariable linear regression was performed on R using the finalfit package (1.0.2). All values are expressed as mean values with error bars representing standard deviations. For all in vivo studies n numbers reflect biological replicates. Depending on the results of normality testing groups were compared using unpaired Student's *t*-test, Mann-Whitney or one/two-way ANOVA (Tukey post-hoc test) were used for analysis. P-values < 0.05 denote statistical significance, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.005$, **** $p < 0.001$.

List of supplementary materials

Figure S1. Expression of selected cyclin dependent kinase inhibitors, SASP factors, fibrotic and Bcl-2 family transcripts detected on microarray analysis in CKD patients and healthy controls, along with adjusted q-values and fold-change (adapted from Nakagawa et al(17))

Figure S2. Pre treatment with ABT-263 in aged murine kidneys results in a reduction of fibrosis, DNA damage and macrophages, and favours a reduction of SC/SASP/BCL2 family members and a preservation of LTL+ tubules after injury.

Figure S3. ABT-263 treatment reduces expression of *Cdkn1a* and *Cdkn2a* transcripts, reduces cortical γ H2A.X expression and increases Lamin B1 in the murine kidney.

Figure S4. Irradiation-induced epithelial senescence promotes fibrosis and senescence in murine kidneys with ABT-263 treatment protective.

Figure S5: Treatment with ABT-263 alters qPCR and protein markers of macrophage infiltration within the murine kidney.

Figure S6: The efficacy of pre-treatment with ABT-263 is lost in young mice with low quantities of pre-IRI senescent cells.

Figure S7: BioRad Antibody AHP1488 anti-*Cdkn2a* detects P16INK4A but not p19ARF protein expression in aged and fibrotic murine kidneys.

Table S1. Multivariate analysis of CDKN1A and CDKN2A and their effect on estimated glomerular filtration rate (egfr) in humans.

Table S2. Primary and secondary antibodies used for IHC and IF.

Data file S1. Raw data for all figures.

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Author contributions:

KJM carried out the experimental design, carried out in vivo experiments, collected the data, performed the analysis, wrote and revised the paper. EDO assisted with data collection and analysis for various figures but in particular Figure 1. They also reviewed the paper. DH assisted with collection of data for Figure 2. DB assisted with collection of the data for Figure 2. MD assisted with collection of the data of IF of p21CIP1 for figures. SAN assisted with

collection of the data for various qPCRs. PJK provided reagents and reviewed the paper. AM provided reagents and reviewed the paper. RS provided reagents and reviewed the paper. SF-G provided reagents and reviewed the paper.

SJF provided supervision and reviewed the paper. JH conceived and designed the analysis, provided supervision and reviewed the paper. DAF conceived and designed the analysis, provided supervision, performed the analysis, assisted with writing and revision of the paper.

Competing interests.

All authors declare no competing interests.

Data and materials availability.

All data associated with this study are present in the paper or supplementary materials.

Figure Captions:

Figure 1. Markers of cellular senescence increase at transcript and protein abundance in human renal disease. Datasets published on Nephroseq were examined for human *CDKN1A* and *CDKN2A* transcript expression, and in-house human biopsy samples were stained for P16INKA and P21CIP1 protein expression (A). Analysis of two published transcriptomic datasets show that both *CDKN1A* and *CDKN2A* rise in kidneys with chronic kidney disease. Fig 1B: n=48 patients with CKD, n=5 normal controls (17) (B). In a separate cohort, *CDKN1A* and *CDKN2A* both increase in the kidney with increasing age, n=42, (16) (C). *CDKN1A* alone correlated with advancing renal impairment, n=42, (16) (D). Staining performed in a UKAS-certified histology facility demonstrated P16INK4A and P21CIP1 protein expression in patients with CKD (n=10). Sample images are shown from normal

kidneys, early CKD, young patients with radiation nephropathy and from older patients with CKD (F; scale bar = 10 μ m). Expression patterns differed between P21CIP1 and P16INK4A with P21CIP1 expressed predominantly in tubular epithelial nuclei, here shown in a patient with advanced renal fibrosis (G). ns=non-significant, * P <0.05 by unpaired t-test.

Figure 2. Senescent human tubular epithelial cells demonstrate selective sensitivity to ABT-263 in vitro. Senescence was induced in human proximal tubular epithelial cells via 10Gy γ -irradiation (A). Irradiation resulted in reduced cell number per field (B) and increased average cell size (C). qPCR analysis of markers of senescence, SASP factors and BCL-2 family members including *MKI67*, *LMNB1*, *CDKN2A*, *CDKN1A*, *TNF*, *IL1B*, *TGFBI*, *BCL2*, *BCL2L1* and *BCL2L2* in irradiated cells (D). SA- β -galactosidase (SA- β -Gal) staining in irradiated cells (E, scale bar represents 100 μ M). RPTECs (control and irradiated) were treated with ABT-263 at various concentrations (0.1, 1 and 5 μ Mol), and cells per field counted and analysed compared to vehicle controls (F). Representative micrographs are shown in lower panel (40 μ M scale bar). * P <0.05, ** P <0.01. *** P <0.005, **** P <0.001. n=4/group. Unpaired t -tests used for experimental comparisons in all groups except F : 2-way ANOVA.

Figure 3. Aged murine kidneys display more senescence, DDR and fibrosis than young kidneys and are more susceptible to post-IRI fibrosis. Experimental schema of sample collections from young and old mice (A). Whole kidney qPCR on samples from young (6-10 weeks) and old mice (>18 months) measuring senescence and SASP markers (B). Markers of tissue fibrosis and senescence associated DNA damage were quantified using IF for Collagen I (C), PSR (D) and γ H2A.X IHC (E). Representative micrographs are shown in right panels (scale bar represents 50 μ M). The % of Lamin B1⁺ (red) nuclei (DAPI; blue) in kidney

tubules (LTL: green) of young vs old kidneys (**F**, n=3-4 group). Representative micrographs are shown in right panels (scale bar represents 20 μ M). Old vs young IRI kidneys analysed for collagen deposition (PSR) (**G-H**, comparisons made using one-way ANOVA test with Tukey post-test used). Representative micrographs are shown in right panel (scale bar represents 100 μ M). Quantitative PCR of *Cdkn1a* in IRI kidneys from old and young mice compared to naïve kidneys illustrates marked upregulation in both IRI groups (**I**). All groups are n=4-8 with unpaired *t*-tests used for experimental comparisons unless otherwise stated. * P <0.05, ** P <0.01. *** P <0.005.

Figure 4. ABT-263 treatment reduces fibrosis and increases the regenerative capacity of aged murine kidneys post-IRI. Diagram detailing ABT-263 treatment regime and unilateral IRI surgery in old female mice (**A**). qPCR analysis of *Cdkn1a* expression in vehicle and ABT-263-treated mice (**B**; red line=expression in old age-matched control tissue). The weight ratios of IRI/CLK kidneys were also calculated in ABT-263-treated and vehicle-treated mice (**C**; dashed line represents left/right (l/r) kidney ratio in healthy animal). Kidney injury molecule 1 (Kim1/Havcr1) IF analysis of post-IRI kidneys +/- ABT-263 (**D**; representative micrographs are shown; scale bar represents 50 μ M). Kidney sections were co-stained for proximal tubule marker LTL (green) and the proliferation marker Ki67 (red). Representative micrograph (**E**) shows the Ki67+ nuclei in LTL+ tubules (white arrows; scale bar represents 50 μ M). The % LTL+ tubules positive for Ki67 was calculated in both groups (**F**) TGF- β protein abundance was measured in whole IRI kidney lysates by ELISA (**G**, n=4-5/group).

Figure 5. ABT-263 treatment improves kidney function in the bilateral IRI model of renal injury. Diagram detailing ABT-263 treatment regime and bilateral IRI surgery in old

female mice (A). Serum Cystatin C is shown at days 7 and 14 post IRI in ABT263 treated mice (B; n=7-8/group, Mann-Whitney test). ABT-263 treatment effect on p21CIP1⁺ senescent cells in baseline kidneys (C, Mann-Whitney test). PSR staining after IRI to evaluate fibrosis in ABT-263 treated animals (D, Mann-Whitney test, n=4-7/group, E; representative micrographs are shown; 50 μ M scale bar). Collagen I staining by IF after IRI in ABT-263 treated animals compared to controls (F, p Mann-Whitney test, n=4-7/group, G; representative micrographs are shown; 50 μ M scale bar). All n=4-6 per group unless stated, unpaired *t*-tests used for all comparisons unless stated. **P*<0.05, ***P*<0.01.

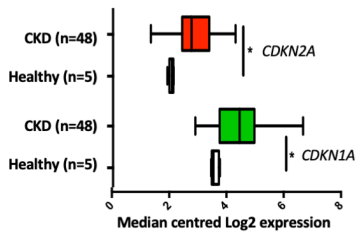
Figure 6. Total body irradiation (TBI) drives senescence and fibrosis in young murine kidneys. Diagram detailing irradiation of mice and analysis of kidneys 8 and 16 weeks post-irradiation (A). IHC and whole kidney imaging was used to identify areas of γ H2A.X induction (B, C). qPCR analysis of whole kidneys for *Cdkn1a* (P21CIP1) and *Cdkn2a* after irradiation (D, n=3/6/7 for age matched control (AMC) vs 8 weeks vs 16 weeks post-irradiation, comparisons by one-way ANOVA test with Tukey post-test). qPCR whole kidney for various SASP and BCL-2 family members was carried out (E). SA-B-Gal staining and analysis of kidney sections after irradiation (F). P16INK4A IF on irradiated kidneys at 8 and 16 weeks post injury (G; representative micrographs are shown; 100 μ M scale bar). γ H2A.X staining of the irradiated kidneys to evaluate DNA damage (H; representative micrographs; scale bar represents 50 μ M). Collagen I IF of irradiated kidneys at 8 and 16 weeks after irradiation (I; representative micrographs are shown; scale bar represents 100 μ M). A similar pattern of PSR staining was seen in naïve and post-irradiated kidneys (J). For panels E-F, n=6-8, comparisons by one-way ANOVA test with Tukey post-test. Unpaired *t*-tests used for all comparisons unless otherwise stated. **P*<0.05, ***P*<0.01. ****P*<0.005.

Figure 7. Irradiation-induced epithelial senescence impairs post injury renal regeneration and promotes fibrosis in murine kidneys, with ABT-263 pre-treatment providing protection. Diagram detailing ABT-263 treatment regime and unilateral IRI surgery in irradiated mice (A). Whole kidney qPCR for *Cdkn1a* in pre-irradiated kidneys + IRI, compared to IRI alone (B), ABT-263 treatment effects on post-ischaemic kidney weight (C; dashed line represents l/r kidney ratio in healthy animal) and tubular injury marker KIM1 by IHC (D). Dual IF staining was performed for Ki67 and the tubular marker LTL (E, yellow arrow=Ki67⁺LTL⁺; red arrow=Ki67⁺LTL⁻). Quantification of LTL⁺ tubules (F), tubular proliferative capacity (G) and renal TGFβ1 protein (H).

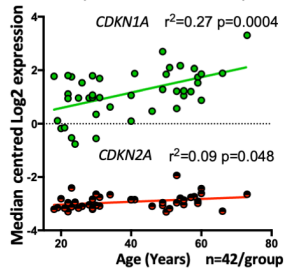
Figure 8. Depletion of irradiation-induced senescent cells improves kidney function after subsequent bilateral IRI. Diagram detailing ABT-263 treatment regime and bilateral IRI surgery in irradiated mice (A). Serum Cystatin C concentrations at days 7 and 35 post IRI in ABT263 treated mice (B). ABT-263 treatment effects on p21CIP1⁺ senescent cells in pre-IRI and post-IRI kidneys (C). PSR staining after IRI shows effects on fibrosis in ABT-263 treated animals (D-E; representative micrographs are shown; 100 μM scale bar, D). Collagen I staining by IF after IRI shows effects on fibrosis in ABT-263 treated animals (F-G; representative micrographs are shown; 50 μM scale bar, F). Dual IF staining (P16INK4A red and collagen I; green) was carried out in irradiated IRI kidneys and age matched controls, and regression analysis undertaken to determine correlation between areas of positivity of both proteins (H). Further dual IF staining (P21CIP1; red and collagen; green) was carried out in irradiated IRI kidneys and age-matched controls, and regression analysis undertaken to determine correlation between areas of positivity of both proteins (I). Key: IRI= ischemia reperfusion injury, Irrad= total body irradiation. n=4-8 per group. Unless otherwise stated comparisons made by unpaired *t*-test. **P*<0.05, ***P*<0.01. ****P*<0.005, *****P*<0.001.



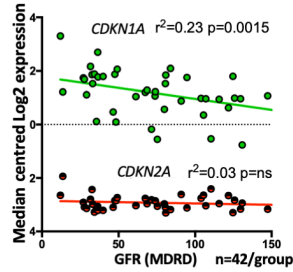
B *CDKN1A/2A* are elevated in CKD (data from Nakagawa et al. 2015)



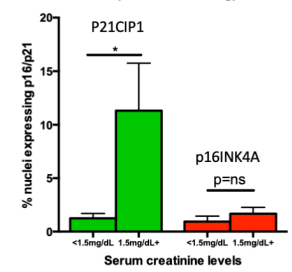
C Renal *CDKN1A/2A* rises with age (data from Ju et al. 2015)



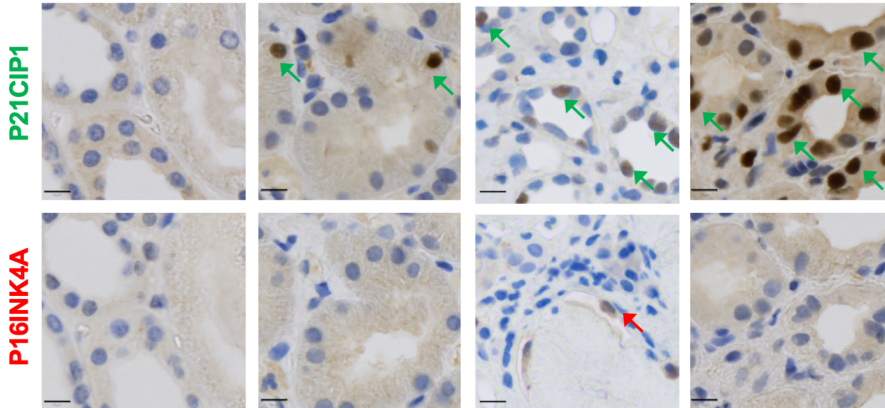
D *CDKN1A* rises with falling GFR (data from Ju et al. 2015)



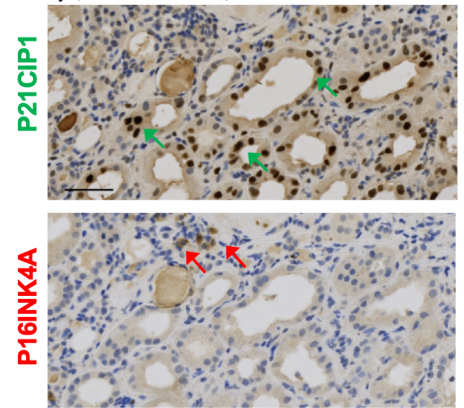
E P21CIP1 rises in human CKD (in house staining)

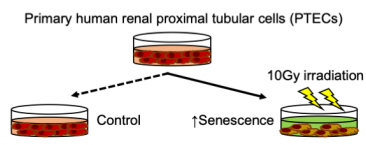
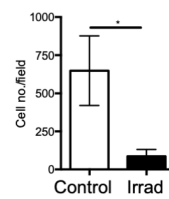
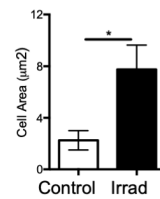


F 19yo normal kidney | 19yo GN, low fibrosis | 29yo previous irradiation | 63yo, advanced fibrosis

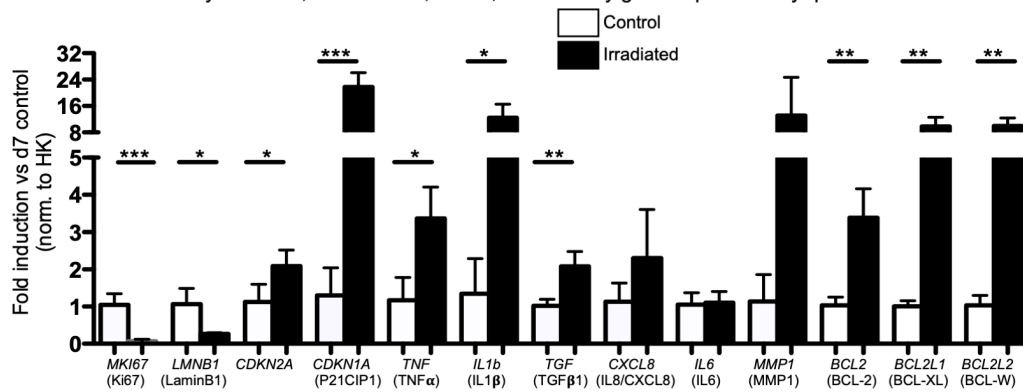
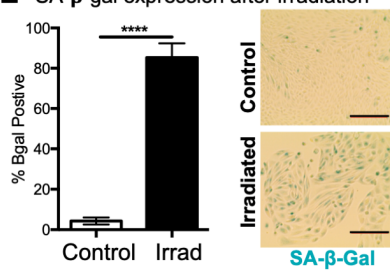
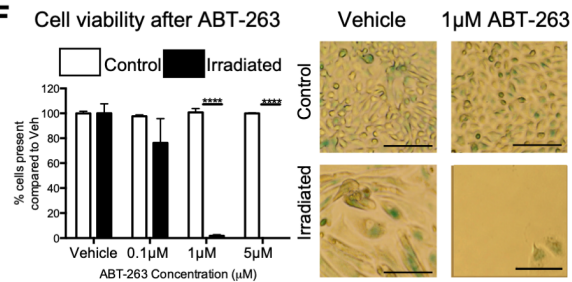


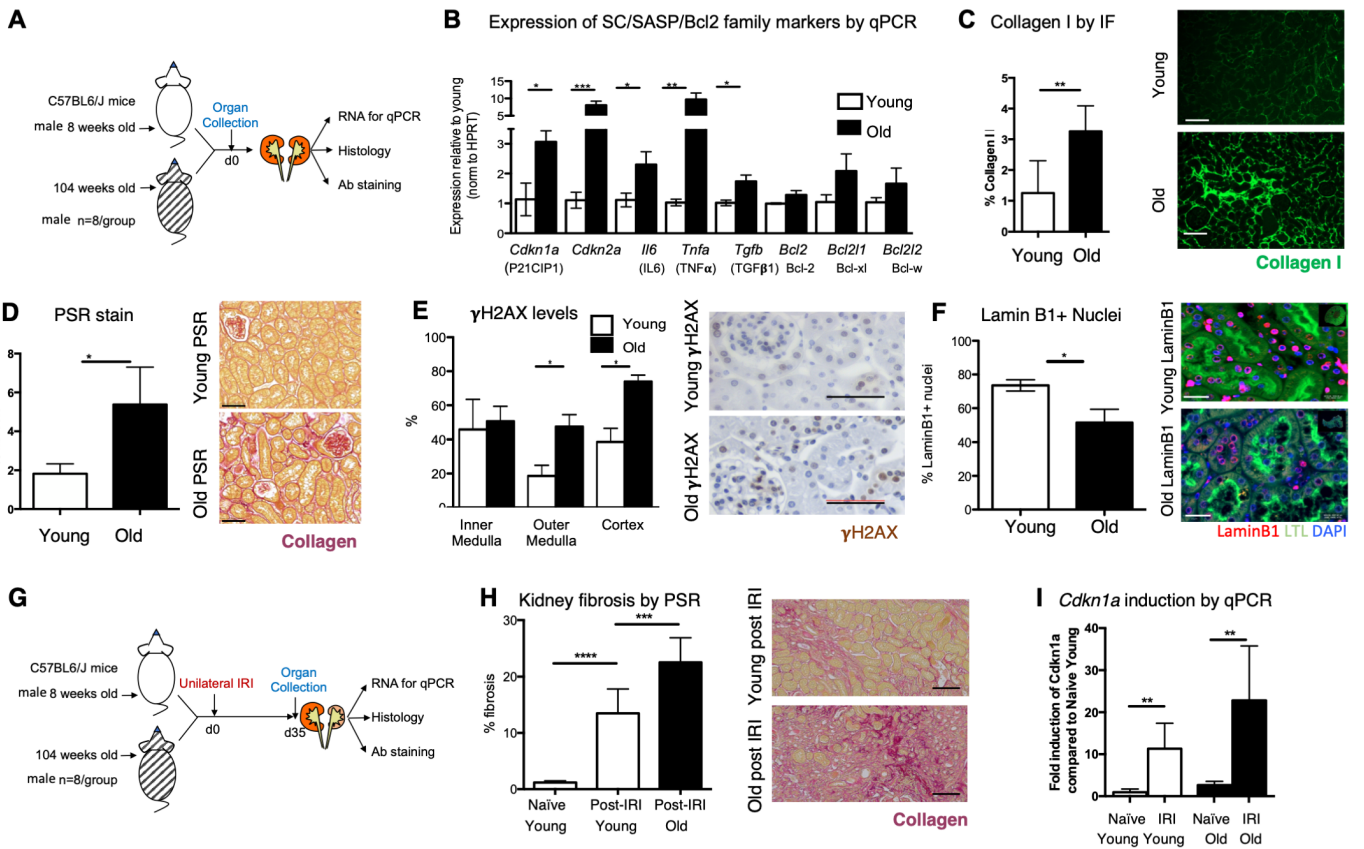
G 63yo, advanced fibrosis, matched P21CIP1+P16INK4A

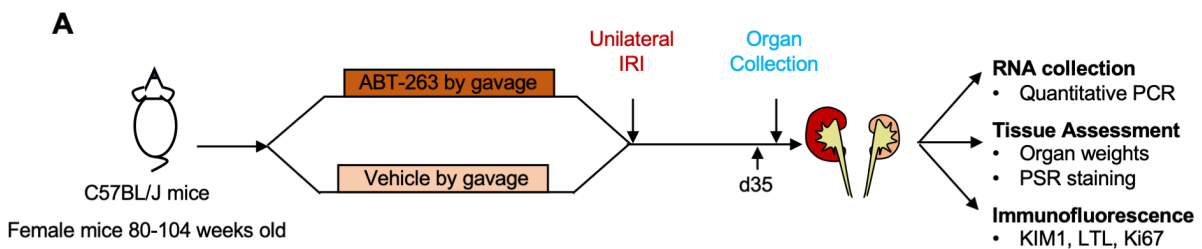


A**B** Cell number/field**C** Mean cell size**D**

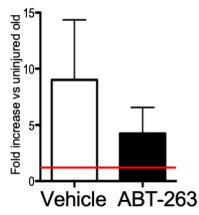
Cell cycle arrest, senescence, SASP, BCL2 family gene expression by qPCR

**E** SA-β-gal expression after irradiation**F** Cell viability after ABT-263

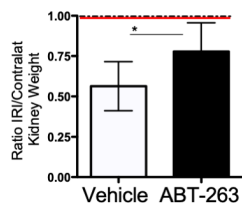




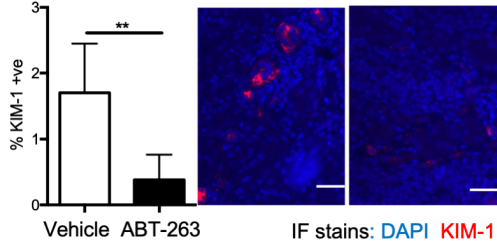
B *Cdkn1a* by PCR



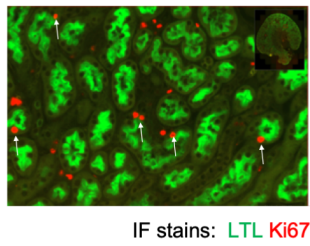
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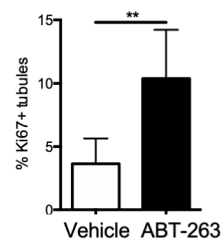
D KIM-1 levels



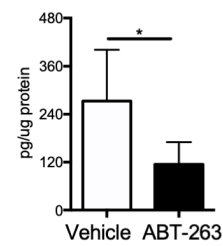
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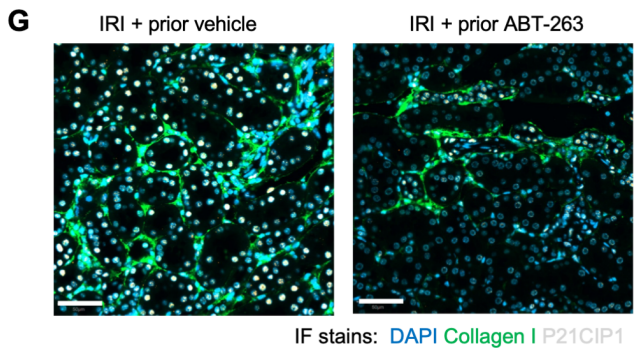
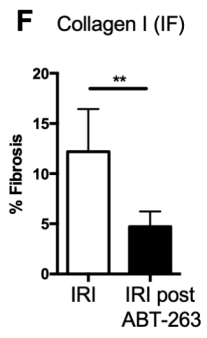
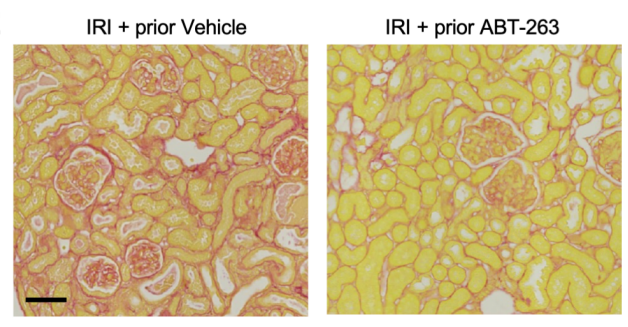
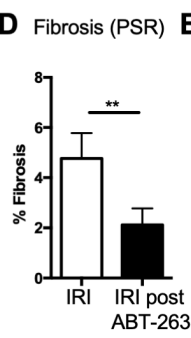
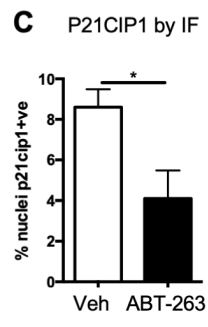
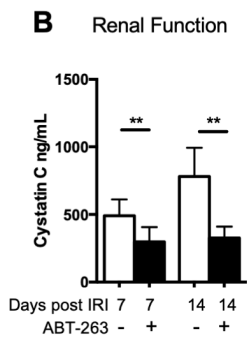
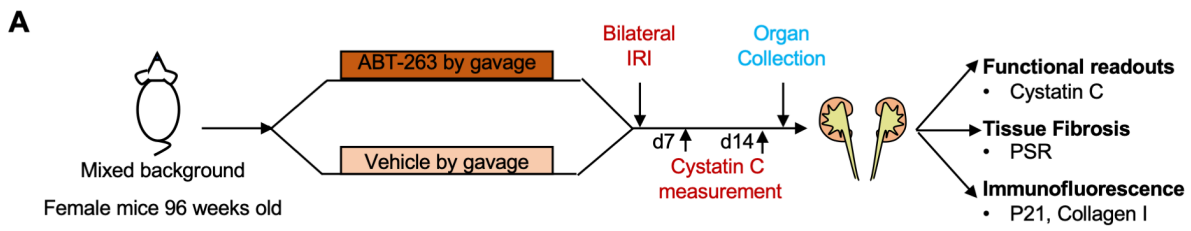


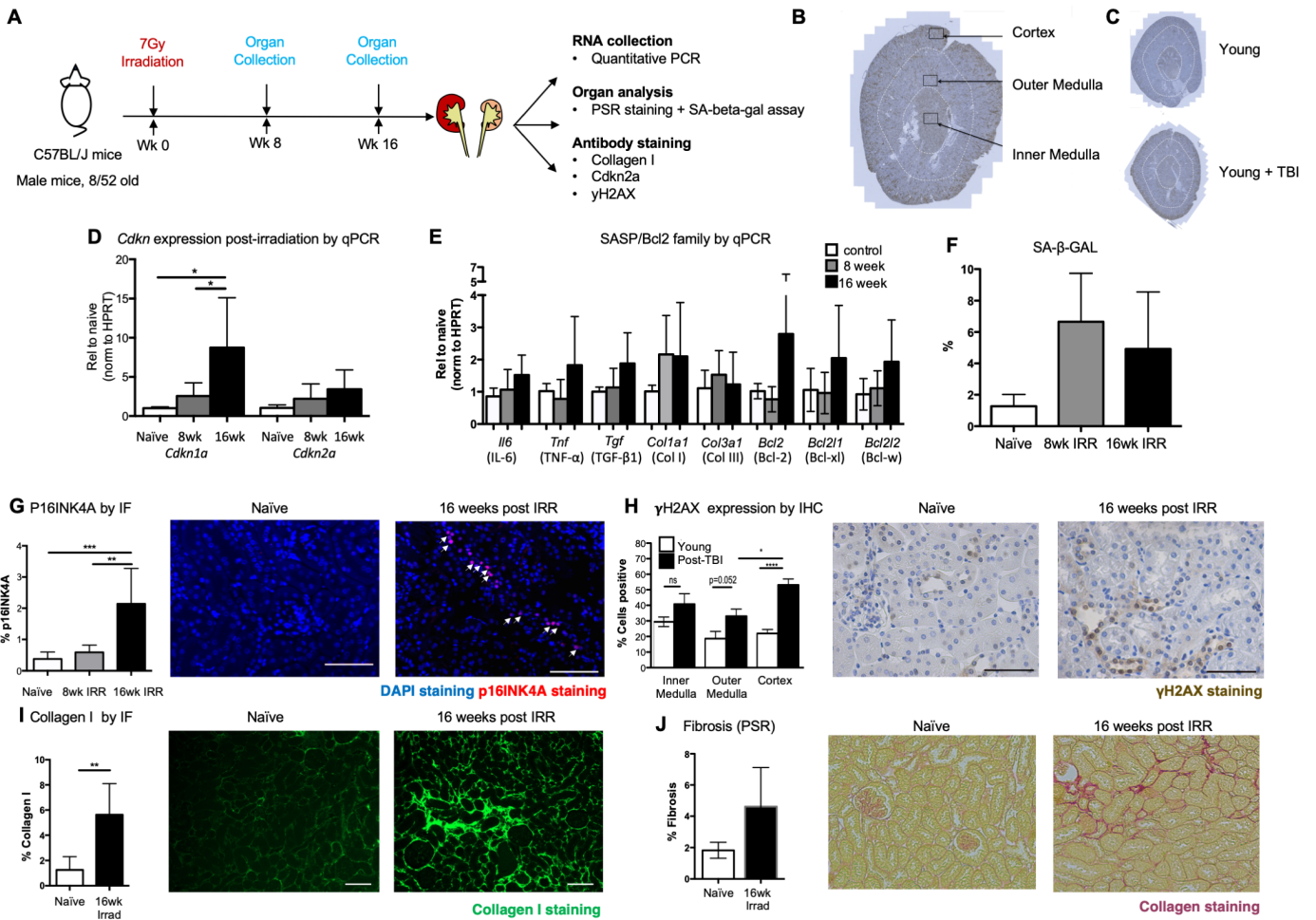
F Proliferation

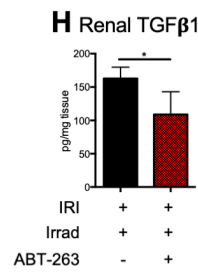
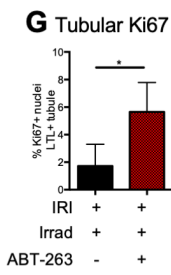
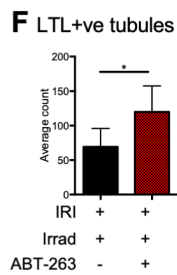
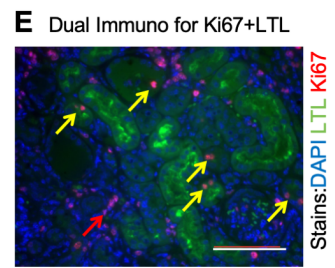
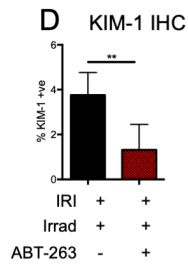
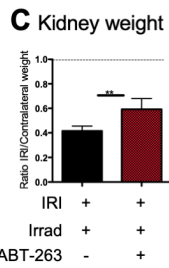
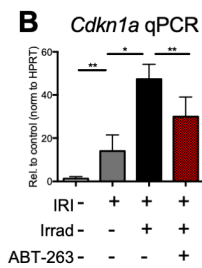
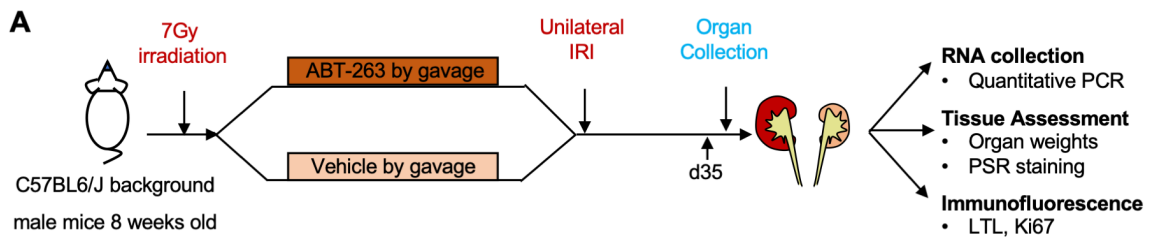


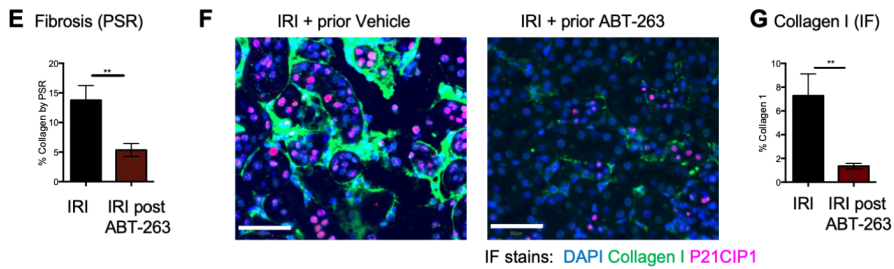
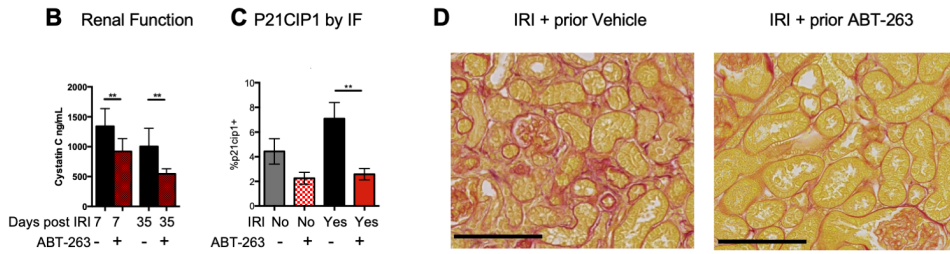
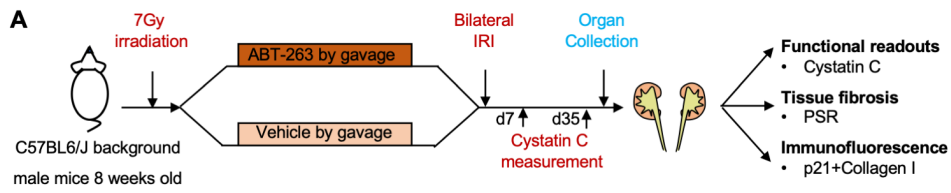
G Tissue TGFβ1



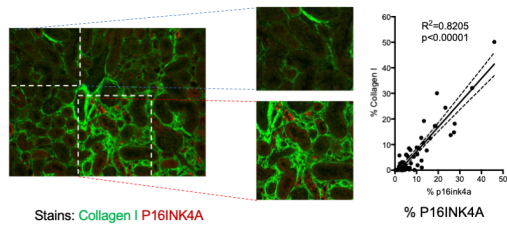




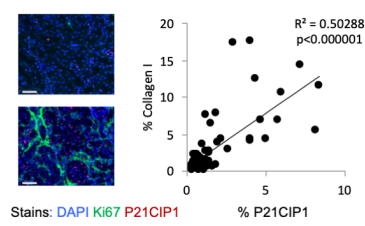




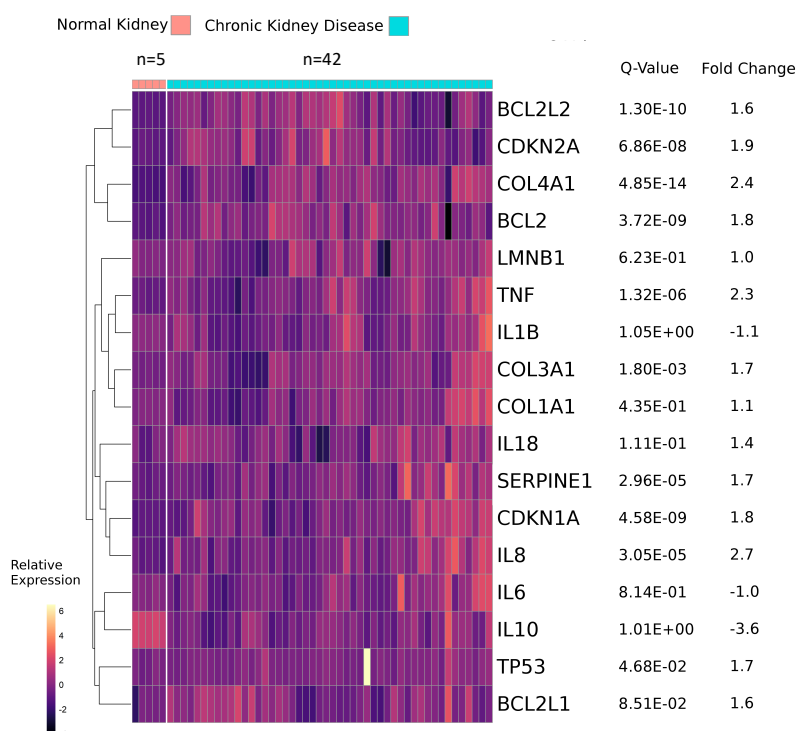
H Dual IF correlation between P16INK4A and Coll I proteins



I Dual IF correlation between P21CIP1 and Coll I proteins



Selected Senescence, SASP, Fibrosis and Bcl-2 family transcript levels (from human CKD and healthy controls)



From Nakagawa et al Plos ONE 2013

Figure S1. Expression of selected cyclin dependent kinase inhibitors, SASP factors, fibrotic and Bcl-2 family transcripts detected on microarray analysis in CKD patients and healthy controls, along with adjusted q-values and fold-change (adapted from Nakagawa *et al*(17))

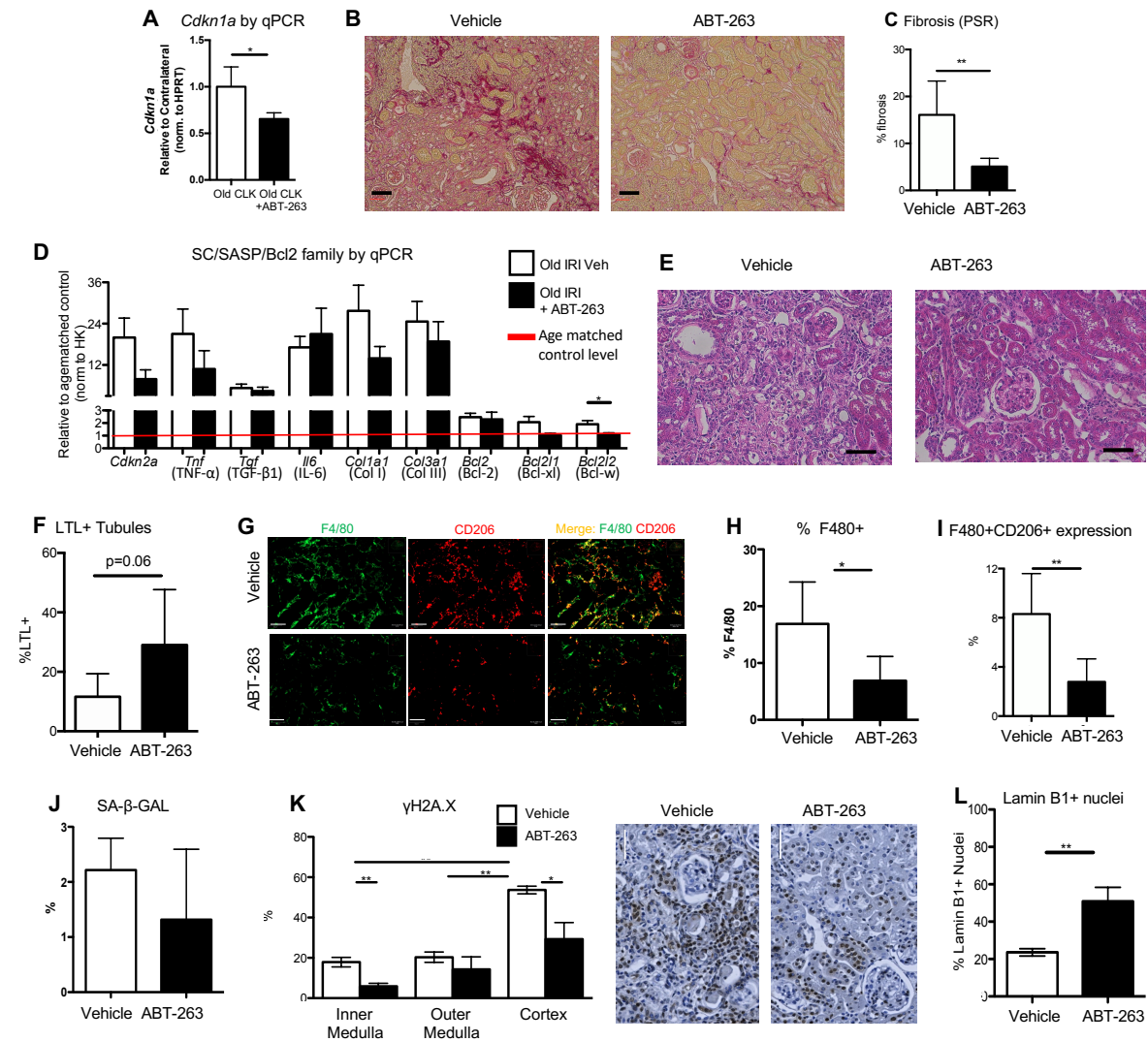


Figure S2. Pretreatment with ABT-263 in old murine kidneys results in a reduction of fibrosis, DNA damage and macrophages, and favours a reduction of SC/SASP/BCL2 family members and a preservation of LTL+ tubules after injury. ABT-263 treatment reduces *Cdkn1a* expression in the uninjured kidney (A; Mann-Whitney test). Post-IRI PSR IHC (B; representative micrographs are shown; 50 μ M scale bar), with analysis showing significant reduction in fibrosis in ABT-263 treated animals (C). Whole kidney qPCR on samples from old IRI vs old IRI + ABT-263 measuring senescence, SASP markers, fibrosis and BCL2 family members (D; red line = expression in old age-matched control tissue) showed that *BCL2L2* (BCL-w) was reduced with ABT-263 treatment (D). H&E staining revealed healthier renal architecture in ABT-263 vs vehicle-treated mice (E; scale bar represents 50 μ M). Staining for

the marker of tubular differentiation (LTL) showed trends toward increase in ABT-263 treated animals (F) F4/80 and CD206 double IF was carried out (G; representative micrographs are shown; 50 μ M scale bar). A reduction in F4/80⁺ macrophages was measured by IF (H). % CD206⁺ macrophages present in vehicle- vs ABT-263-treated IRI kidneys calculated (I). SA-B-Gal staining was carried out between old IRI vs old IRI + ABT-263 (J). γ H2A.X staining of the kidneys shows evidence of a reduction of DNA damage with ABT-263 treatment after IRI in old mice, most marked in the cortex (K). Pre-treatment with ABT-263 in old kidneys increased the proportion of Lamin B1⁺ tubular nuclei (L). Unless otherwise stated comparisons made by unpaired t-test. * P <0.05, ** P <0.01. *** P <0.005.

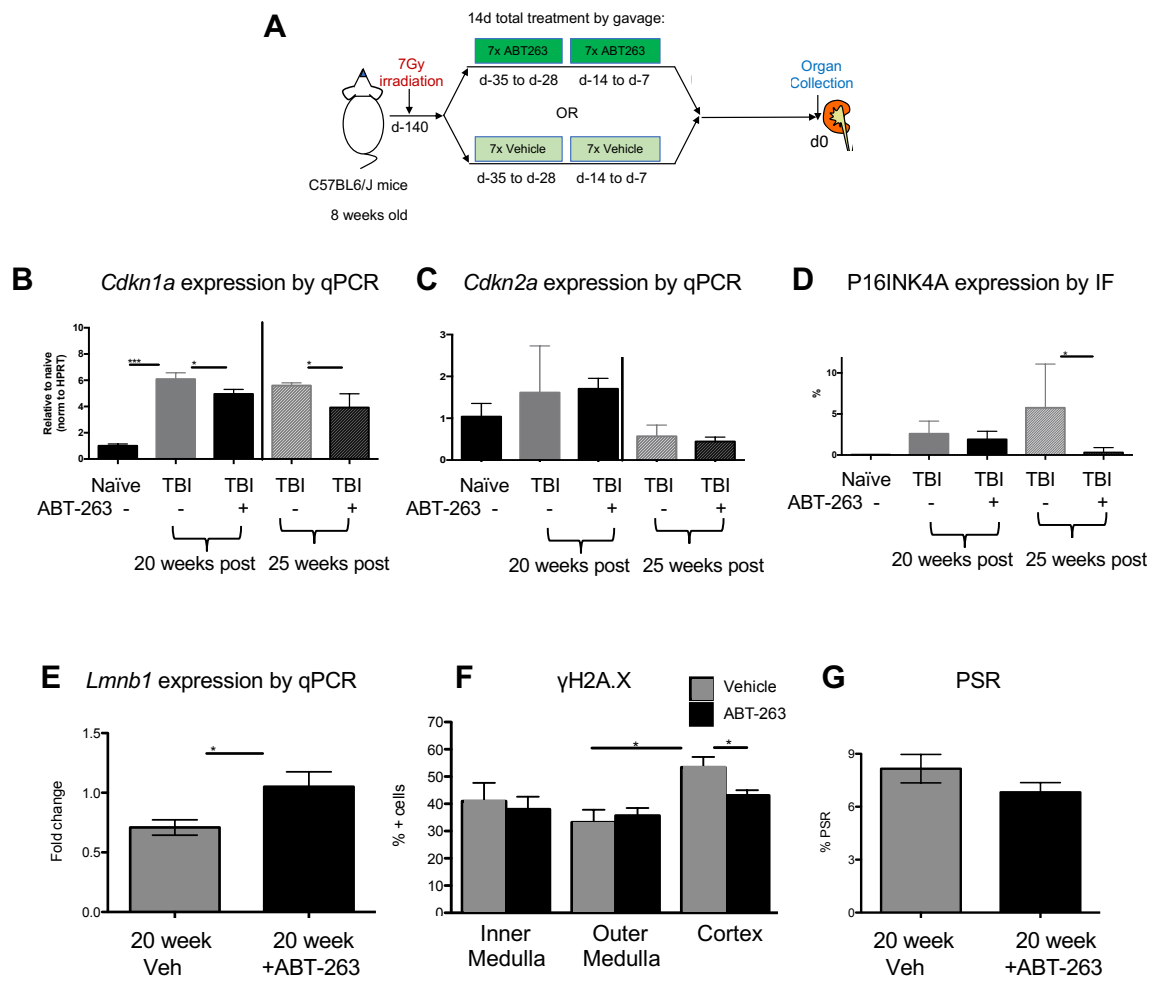


Figure S3. ABT-263 treatment reduces expression of *Cdkn1a* and *Cdkn2a* transcripts, reduces cortical γ H2A.X expression and increases Lamin B1 in the murine kidney.

TBI at 7Gy was carried out on young mice at day 0, treatment with ABT-263 was at weeks 15 and 18 for 7 days, kidneys were recovered at 20 and 25 weeks (A). qPCR was carried out for *Cdkn1a* (P21CIP1; B) and *Cdkn2a* (C) and IHC for P16INK4A (D). At 20 weeks post-TBI, qPCR was also carried out for *Lmn1* (E), as was IHC for γ H2A.X staining (F) and PSR staining for collagen (G). Unless otherwise stated comparisons made by unpaired *t*-test.

* $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$.

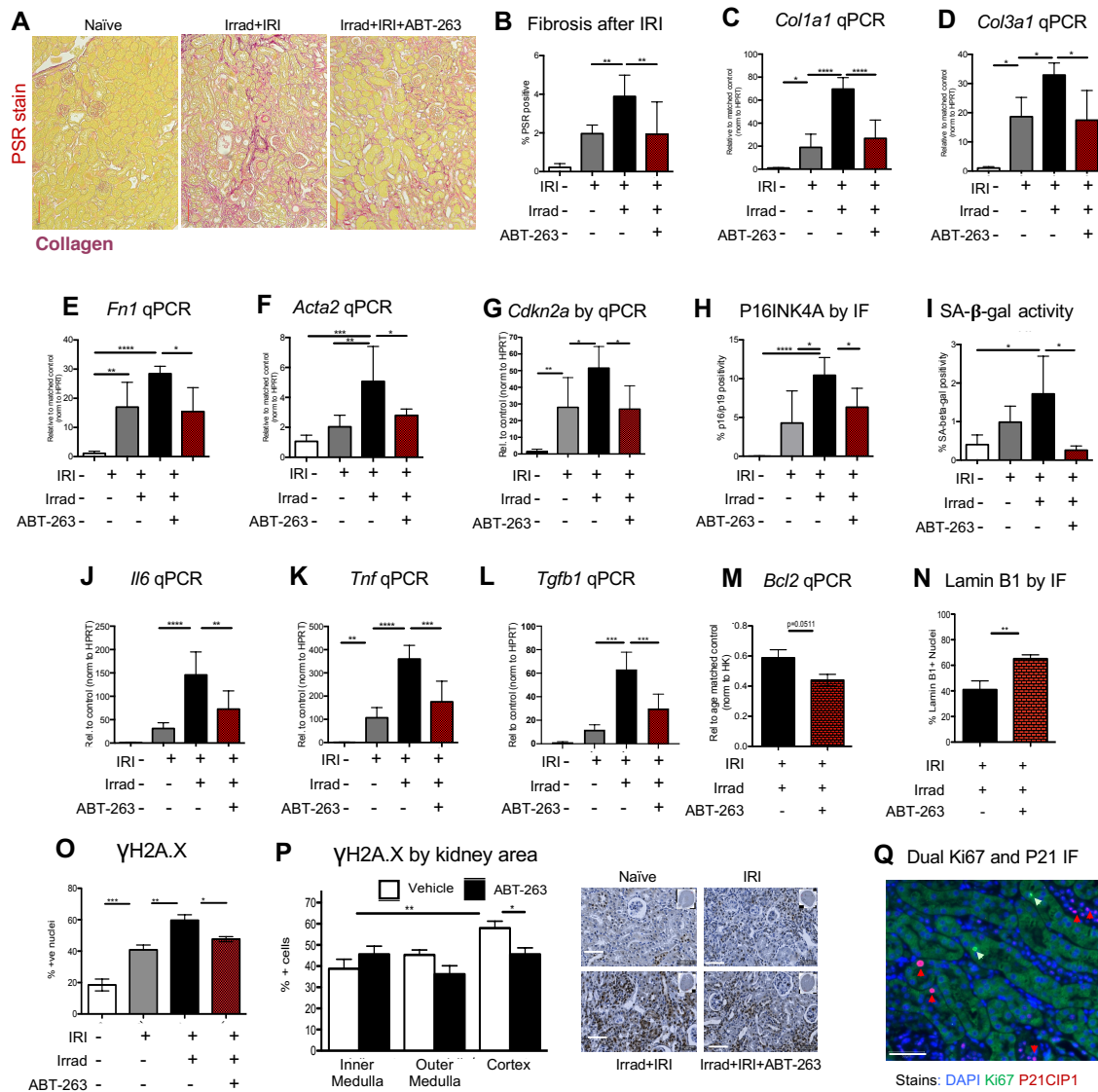


Figure S4. Irradiation-induced epithelial senescence promotes fibrosis and senescence in murine kidneys with ABT-263 treatment protective.

PSR staining showed significant reduction in Collagen in ABT-263 pre-treated mice (A; representative images, 100 μm scale bar, B; comparisons by one-way ANOVA test with Tukey post-test). Fibrosis markers *Colla1*, *Col3a1*, *Fn1* and *Acta2* mRNA showed maximal rises in IRI + irradiation mice, with ABT263 pre-treatment protective (C-F). Senescence associated *Cdkn2a* was measured at transcript level by qPCR (G), and P16INK4A protein quantified by immunofluorescent staining (H; comparisons by one-way ANOVA test with Tukey post-test). SA-B-Gal staining (I) and SASP markers *Il6*, *Tnf* and *Tgfb1* (J-L) were all augmented in pre-irradiated kidneys + IRI, compared to IRI alone, with ABT-263 pre-treatment normalising the

post-injury expression. qPCR analysis of *BCL2* revealed a decrease between vehicle-treated vs ABT-263-treated pre-irradiated IRI kidneys (**M**). Lamin B1⁺ tubular nuclei were counted comparing vehicle-treated versus ABT-263-treated pre-irradiated IRI kidneys (**N**). γ H2A.X staining (**O**) indicated that DNA damage was increased in pre-irradiated kidneys + IRI, compared to IRI alone, with ABT-263 pre-treatment normalising the post-injury expression. Changes in γ H2A.X staining were maximal in the cortex, with significant reduction after ABT-263. Representative micrographs are shown (**P**; scale bar represents 50 μ M). Dual IF staining for Ki67 and P21CIP1 demonstrated that these markers are mutually exclusive (**Q**, scale bar represents 50 μ M). Unless otherwise stated comparisons made by unpaired *t*-test. * P <0.05, ** P <0.01. *** P <0.005.

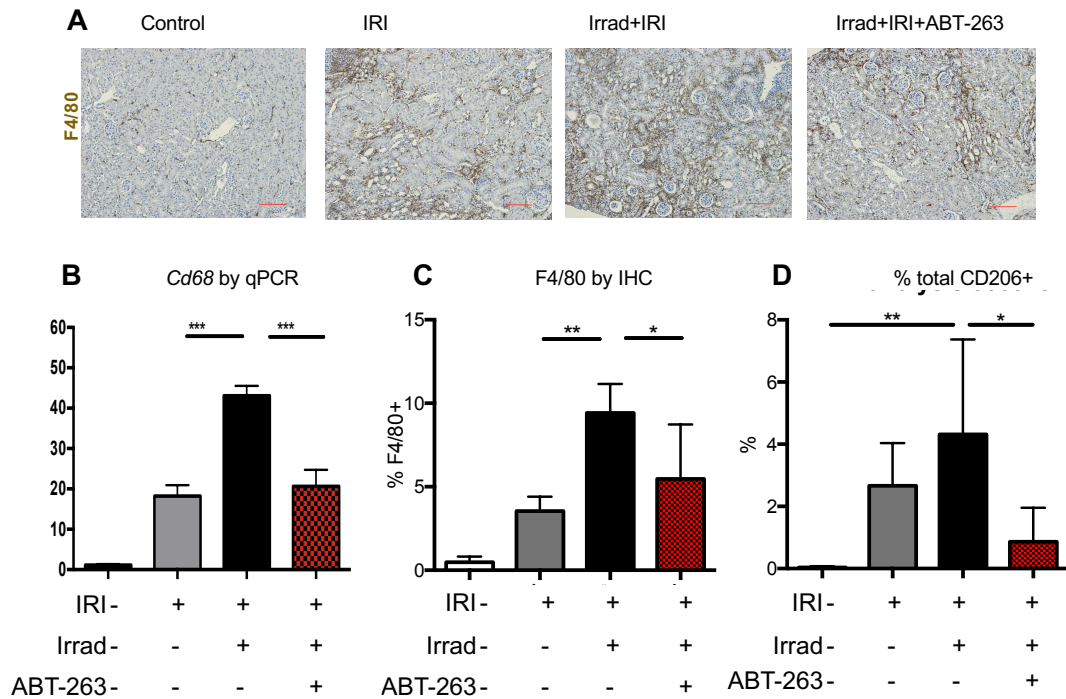


Figure S5. Treatment with ABT-263 alters qPCR and protein markers of macrophage infiltration within the mouse kidney. *Cd68* qPCR and F4/80 IHC analysis and representative micrographs (A-C; 100 μ M scale bar). Dual F4/80 and CD206 (MMR) IF analysis showing the % CD206⁺ macrophages present 5 weeks post-IRI (D). Unless otherwise stated comparisons made by unpaired t-test. * P <0.05, ** P <0.01. *** P <0.005.

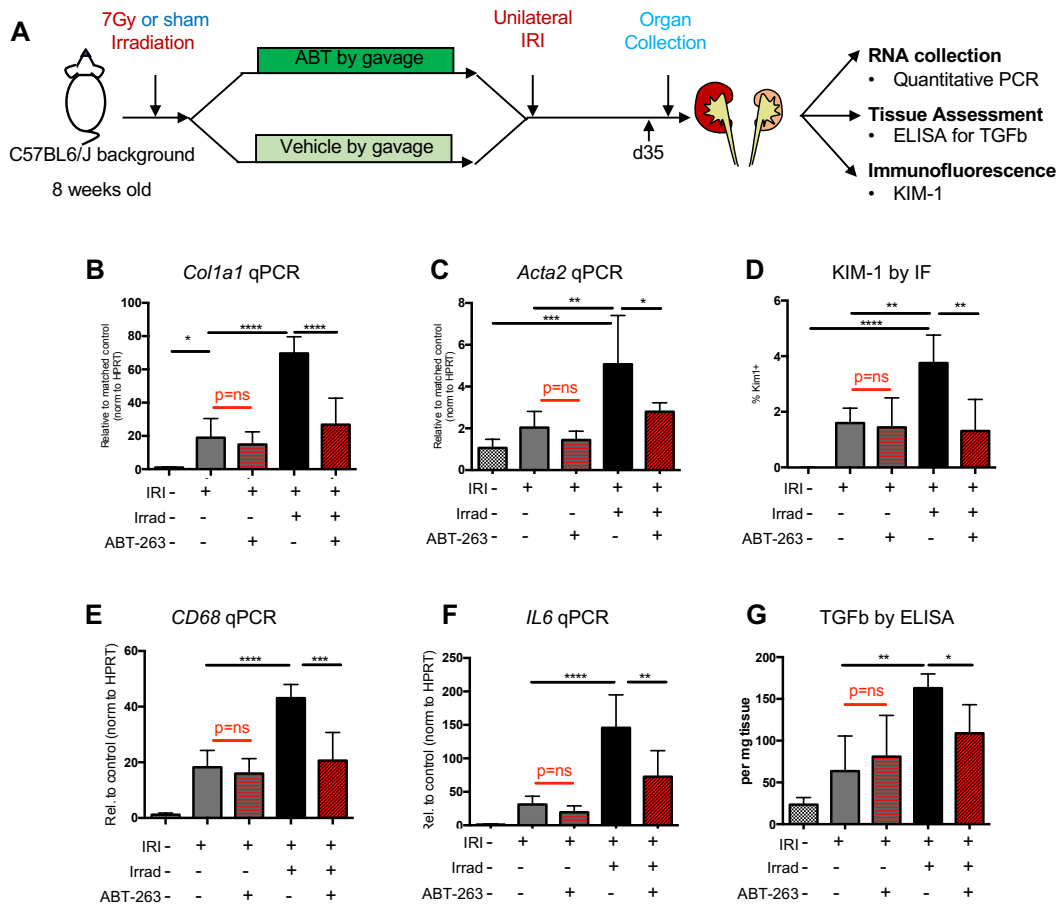


Figure S6. The efficacy of pre-treatment with ABT-263 is lost in young mice with low quantities of pre-IRI senescent cells. Young male control mice underwent renal IRI with and without ABT263 and without prior irradiation (A). Whilst ABT-263 improved many markers of regeneration in irradiated mice when given prior to IRI, in healthy young mice no protective effect was seen in any assay performed. Plots are shown with equivalent outcomes for *Colla1* (B) and *Acta2* by qPCR (C). There was no alteration in KIM-1 by IF (D), in macrophage gene expression by qPCR (*Cd68*, E), or the key SASP gene *Il6* by qPCR (F) or TGF-beta on whole kidney ELISA (G). ns=non-significant. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$, **** $P < 0.001$

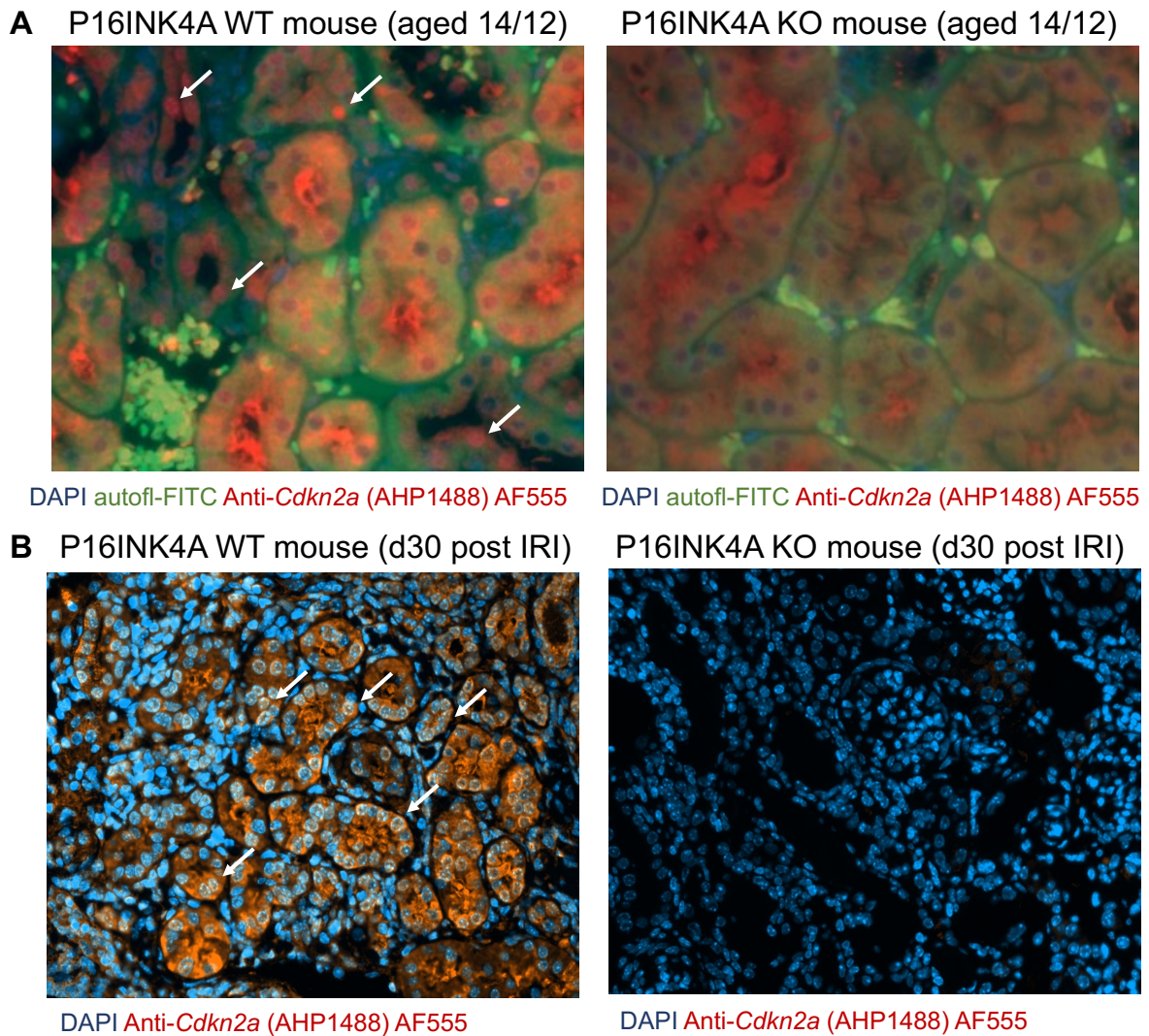


Figure S7. BioRad Antibody AHP1488 anti-*Cdkn2a* detects P16INK4A but not p19ARF protein expression in aged and fibrotic kidneys. Kidney samples from naturally aged (14 month) mice or young mice at d30 post renal IRI (with or without a targeted deletion within an exon specific to P16INK4A protein) were stained with AHP1488 (an antibody generated from a P16INK4A protein fragment). Multiple sites of nuclear positivity were seen in aged and fibrotic kidneys from WT mice (**A** and **B**, left panels). In an equivalently aged kidney, or a kidney at the same timepoint after injury with targeted deletion of P16INK4A and preserved p19ARF, no staining was seen (**A** and **B**, right panels). White arrows mark P16INK4A⁺ nuclei.

Table S1. Multivariate analysis of CDKN1A and CDKN2A and their effect on estimated glomerular filtration rate (egfr) in humans.

CDKN1A

		unit	value	Coefficient (univariable)	Coefficient (multivariable)
age	[18.0,73.0]	Mean (sd)	1.2 (0.9)	0.03 (0.01 to 0.04, p<0.001)	0.02 (0.00 to 0.04, p=0.018)
sex	Female	Mean (sd)	1.1 (1.0)	-	-
	Male	Mean (sd)	1.2 (0.7)	0.01 (-0.56 to 0.58, p=0.978)	-0.15 (-0.63 to 0.34, p=0.547)
egfr	[12.0,299.0]	Mean (sd)	1.2 (0.9)	-0.01 (-0.01 to -0.00, p=0.001)	-0.01 (-0.01 to -0.00, p=0.049)
Adjusted R-squared = 0.29					

CDKN2A

		unit	value	Coefficient (univariable)	Coefficient (multivariable)
age	[18.0,73.0]	Mean (sd)	-2.9 (0.3)	0.01 (0.00 to 0.01, p=0.048)	0.00 (-0.00 to 0.01, p=0.119)
sex	Female	Mean (sd)	-2.9 (0.3)	-	-
	Male	Mean (sd)	-2.9 (0.2)	-0.03 (-0.21 to 0.14, p=0.714)	-0.05 (-0.23 to 0.13, p=0.571)
egfr	[12.0,299.0]	Mean (sd)	-2.9 (0.3)	-0.00 (-0.00 to 0.00, p=0.241)	-0.00 (-0.00 to 0.00, p=0.706)
Adjusted R-squared = 0.033					

Table S2. Primary and secondary antibodies used for IHC and IF

Antibody (anti-)	Section Type	Dilution	Company	DAB/IF	Antigen Retrieval	Secondary Antibody
Coll I	MFPE	1:200	Southern Biotech	IF	Heat: TE	Anti-goat-AF488
F4/80	MFPE	1:50	AbCam	DAB, IF	Proteinase K (RT)	Biot-anti-rat, anti-rat AF488
γ H2A.X	MFPE	1:100	AbCam	DAB	None	Biot-anti-rabbit
Kim1	MFPE	1:50	R&D	IF	Heat: TE	Anti-goat AF555
Ki67	MFPE, PFAPE	1:100	Cell Sig Tech/AbCam	IF	Heat: TE or CB	Anti-rabbit or sheep-AF555
LTL-FITC	MFPE	1:150	Vector	IF	Heat: TE/CB	None
MMR	MFPE	1:100	R&D	IF	Proteinase K (RT)	Anti-goat-AF555
P16INK4A	MFPE	1:100	BioRad	IF	Heat: TE or CB	Anti-rabbit AF555
P21CIP1	PFAPE	1:100	Abcam	IF	Heat: CB	Anti-rabbit AF555
Lamin B1	MFPE	1:100	AbCam	IF	Heat: CB	Anti-rabbit AF555

CB: Citrate buffer; DAB: Diaminobenzidine; IF: Immunofluorescence; IHC:

Immunohistochemistry; MFPE: methacarn fixed paraffin embedded; PFAPE: 4%PFA fixed paraffin embedded; RT: Room Temperature. TE: Tris EDTA.

Data file S1: Raw data for all figures.