In serum veritas—in serum sanitas? Cell non-autonomous aging compromises differentiation and survival of mesenchymal stromal cells via the oxidative stress pathway

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Even tissues capable of complete regeneration, such as bone, show an age-related reduction in their healing capacity. Here, we hypothesized that this decline is primarily due to cell non-autonomous (extrinsic) aging mediated by the systemic environment. We demonstrate that culture of mesenchymal stromal cells (MSCs) in serum from aged Sprague–Dawley rats negatively affects their survival and differentiation ability. Proteome analysis and further cellular investigations strongly suggest that serum from aged animals not only changes expression of proteins related to mitochondria, unfolded protein binding or involved in stress responses, it also significantly enhances intracellular reactive oxygen species production and leads to the accumulation of oxidatively damaged proteins. Conversely, reduction of oxidative stress levels in vitro markedly improved MSC function. These results were validated in an in vivo model of compromised bone healing, which demonstrated significant increase regeneration in aged animals following oral antioxidant administration. These observations indicate the high impact of extrinsic aging on cellular functions and the process of endogenous (bone) regeneration. Thus, addressing the cell environment by, for example, systemic antioxidant treatment is a promising approach to enhance tissue regeneration and to regain cellular function especially in elderly patients.

**Keywords:** aging; adult stem cells; cell–microenvironment interactions; tissue regeneration; oxidative stress; cellular stress response

**Abbreviations:** MSCs, mesenchymal stromal cells; ROS, reactive oxygen species; PD, population doublings; p21WAF1/CIP1, cyclin-dependent kinase inhibitor 1; p16INK4A, cyclin-dependent kinase inhibitor 2A; OM, osteogenic media; AM, adipogenic media; AR, Alizarin Red; OR, Oil Red O; MS, mass spectrometry; HSP, heat shock protein; micro-CT, microcomputed tomography; NAC, N-acetyl-L-cysteine; MTF, maximum torque at failure; TS, torsional stiffness; ETF, energy to failure; BV/TV, bone volume fraction; Tb.Th, trabecular thickness; Tb.N, trabecular number; Tb.Sp, trabecular separation; ANOVA, analysis of variance

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Extrinsic aging affects MSC survival and function

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age-related influence of the systemic environment on cellular function. Conboy et al. performed parabiotic pairing experiments between young and aged mice. Results of these experiments suggest that satellite cells and liver progenitor cells of aged mice can be rejuvenated in vivo by exposure to a young systemic milieu. Recently, it was also shown that such heterochronic parabiosis reverses age-related cardiac hypertrophy. Thus, we hypothesize that extrinsic (cell non-autonomous) aging has a higher impact on the function of MSCs than intrinsic aging.

To explore potential mechanisms and consequences by which an age-altered systemic environment affects young and aged MSC functions, we studied concurrently cellular and molecular changes in response to serum derived from young and aged Sprague–Dawley rats. Our results show that the systemic environment modulates age-dependent MSC survival and differentiation. Our protein expression and cell assay data identified increased intracellular (oxidative) stress as a potential cause for the altered MSC function. Conversely, antioxidant treatment markedly improved age-altered MSC function and in vivo bone regeneration. In summary, we propose that the systemic environment crucially contributes to the age-related decline in bone regeneration by increasing intracellular ROS levels, hence compromising viability and function of mesenchymal (progenitor) cells.

Results

Age-altered systemic environment reduces proliferation, increases cell cycle inhibitor expression and apoptosis of MSCs. Since our previous results indicate a gradual decline in MSC number and function with aging, we used MSCs and serum from 3 weeks (yMSCs; ySerum) to 12 months (aMSCs, aSerum) old male Sprague–Dawley rats for in vitro investigations. To determine the influence of ySerum and aSerum on the growth dynamics of yMSCs and aMSCs, we assessed the number of population doublings (PD) in short-term proliferation assays (Figure 1a). Both yMSCs and aMSCs grown in aSerum displayed significantly reduced proliferation rates compared with the corresponding cultures in ySerum (yMSCs: PD_{aSera} = 1.68, PD_{ySera} = 2.16, \( P = 0.005 \); aMSCs: PD_{aSera} = 1.36, PD_{ySera} = 1.93, \( P = 0.016 \)). To examine the underlying molecular causes for the reduced cell proliferation in aSerum, the expression of cell cycle inhibitors in yMSCs and aMSCs was analyzed by western blot technique (Figures 1b–d). This analysis revealed a significantly enhanced cyclin-dependent kinase inhibitor 1 (p21^{WAF1/CIP1}) and cyclin-dependent kinase inhibitor 2A (p16^{INK4A}) expression in yMSCs (3.1-fold and 2.6-fold, respectively) cultured in aSerum compared with their counterparts grown in ySerum.

Figure 1 Effects of serum from young (ySerum) and aged (aSerum) Sprague–Dawley rats on proliferation and survival of MSCs. (a) Proliferation rates were determined by using CyQuant assay and obtained values were used to calculate the number of PD. Growth of MSCs from young (yMSCs) and aged (aMSCs) animals cultured in aSerum was significantly lower than in parallel cultures with ySerum. (b) and (c) Compared with ySerum, MSC culture in aSerum caused an upregulation of the cell cycle inhibitors p21^{WAF1/CIP1} and p16^{INK4A}. Graphs illustrate quantified signal intensities relative to GAPDH and (d) shows the corresponding representative pictures of the western blot analysis. (e) Apoptosis rates of MSCs cultured in ySerum and aSerum, respectively, were determined by caspase-Glo 3/7 Assay. Graph shows the obtained fluorescence value normalized to total DNA content. (For (a–d), \( n = 6 \), \(^* P < 0.05\), Mann–Whitney U test; (e), \( n = 6 \), \(^* P < 0.05\), \(^\# P < 0.05\)—positive control (Etoposide) versus all other treatment groups; ANOVA with Bonferroni correction)
To test whether the increased cell cycle inhibitor expression correlates with higher apoptosis rates, we determined the caspase-3/7 activity as surrogate marker for apoptotic cell death, and used 50 μM etoposide as a positive control (Figure 1e). Caspase-3/7 activity in yMSCs and aMSCs cultured in aSerum were 39% (P = 0.010) and 35% (P = 0.006) higher, respectively, than in the parallel cultures with ySerum. Notably, no significant difference in proliferation rate, cell cycle inhibitor expression and apoptosis rate were observed between aMSCs and yMSCs cultured in the same serum.

Age-altered systemic environment affects MSCs osteogenic and adipogenic differentiation capacity. To further investigate the impact of an age-altered systemic environment on the progenitor phenotype of yMSC and aMSCs, we compared their osteogenic and adipogenic differentiation capacity. Osteogenic differentiation was quantified by matrix mineralization using Alizarin Red (AR) and normalized to the number of viable cells (Figures 2a and b). MSCs cultivated in aSerum showed a significantly lower capability to differentiate into the osteogenic lineage (yMSCs: 3.0 ± 0.3; aMSCs: 2.0 ± 0.1) than the parallel cultures with ySerum (yMSCs: 6.0 ± 0.4, P < 0.001; aMSCs: 3.6 ± 0.2, P < 0.001). In addition, we also found significant differences in matrix mineralization between aMSCs and yMSCs grown in the same serum (ySerum: P < 0.001; aSerum: P = 0.006).

To examine the effect of an age-altered systemic environment on adipogenic differentiation, we used Oil Red O (OR) staining to quantify lipid vacuoles and normalized obtained values to the number of viable cells (Figures 2c and d). A trend of increased adipogenic differentiation was observed for yMSCs cultured in aSerum (1.34 ± 0.20) compared with their counterparts in ySerum (1.11 ± 0.07, P = 0.112), but did not reach statistical significance. However, aMSCs cultured in aSerum (1.71 ± 0.14) displayed a significantly higher adipocytes/viable cell ratio compared with their corresponding cultures in ySerum (1.14 ± 0.08, P < 0.001) as well as to all other treatment groups. Together these results indicate that age-related changes in the systemic environment reduce MSC proliferation and survival, compromise their osteogenic differentiation ability, and promote their differentiation into the adipogenic lineage.

Age-altered systemic environment alters expression of proteins related to mitochondria, unfolded protein binding and stress response. In order to identify underlying molecular changes responsible for the observed functional alteration in an age-altered systemic environment, we profiled the protein expression of yMSCs and aMSCs cultured in aSerum and ySerum, respectively. Two-dimensional gel electrophoresis detected a total of 147 protein spots to be modulated by the age-altered systemic environment in MSCs. Using mass spectrometry (MS), 105 of these spots were identified as 89 different proteins and 16 isoforms (Supplementary Table 1). Among the detected proteins, 49 were significantly and reproducibly upregulated and 34 downregulated in yMSCs grown in aSerum compared...
with corresponding cultures in ySerum (Figure 3a). In aMSCs grown in aSerum, 53 proteins were increased and 33 proteins decreased in their expression compared with the parallel cultures with ySerum. Taken together, the expression of 80 proteins were serum-dependently modulated in both MSC groups \((n_{\text{upregulated}} = 48, n_{\text{downregulated}} = 32)\).

Subsequently, we performed functional annotation clustering of the differentially expressed proteins according to cellular component, biological process and pathway (Figure 3b). Assessment of cellular component by gene ontology analysis indicated that a large number of proteins were part of an organelle (GO:0005739, 'organelle part', \(n = 40, \ P < 0.001\)), including mitochondria, endoplasmic reticulum and the cytoskeleton (Supplementary Table 2). Clustering according to biological processes revealed that differentially regulated proteins are also related to a 'cellular metabolic process' (GO:0044237, \(n = 19, \ P = 0.017\); Supplementary Table 3). Analysis in respect to the pathways revealed that seven of these proteins are part of the glycolysis/gluconeogenesis pathway \((P < 0.001)\).

In accordance to the altered expression of proteins of the endoplasmic reticulum (GO:0005783, \(n = 8, \ P = 0.009\)), the cluster 'cellular metabolic process' also contains proteins involved in unfolded protein binding (GO:0051082, \(n = 11, \ P < 0.001\)). The majority of these proteins are molecular chaperones \((n = 10, \ P < 0.001)\). Interestingly, chaperones like the heat shock protein (HSP) 90 kDa beta member 1 (HSP90b1), HSP70 and HSP40, which are known to be upregulated in response to stress, are more highly expressed in MSCs cultured in aSerum. In line with these result, we found that the majority of proteins clustered into the biological process 'response to stress' \((13 \ out \ of \ 18)\) were upregulated in response to aSerum. These results led us to assume that the age-altered systemic environment may directly or indirectly provoke enhanced intracellular stress, which would be detrimental to MSC function and survival.

Age-altered systemic environment enhances intracellular oxidative stress and thereby indirectly compromises MSC differentiation. Exemplarily, differentially expressed proteins involved in 'unfolded protein binding' were validated by western blot of whole MSC lysates to confirm the proteome data (Figure 4). A significant increase in the expression of HSP90 (yMSCs: 1.3-fold \(P = 0.003\); aMSCs: 1.4-fold \(P = 0.002\)), HSP70 (yMSCs: 1.3-fold \(P = 0.010\); aMSCs: 1.6-fold \(P = 0.030\)), HSP40 (yMSCs: 1.7-fold \(P = 0.012\); aMSCs: 1.8-fold \(P = 0.030\)) and PDI (yMSCs: 1.7-fold \(P = 0.030\); aMSCs: 1.7-fold \(P = 0.014\)) was detected in MSCs cultured in aSerum. In accordance with our proteome analysis, expression of the mitochondrial chaperone HSP60 was significantly decreased in yMSCs (0.6-fold \(P = 0.009\)) and aMSCs (0.8-fold \(P = 0.010\)) grown in aSerum.

To test whether this enhanced HSP/chaperone expression is correlated with higher levels of intracellular oxidative stress, we incubated yMSCs and aMSCs with the fluorescent ROS indicator CM-H2DCFDA and normalized obtained values to DNA content. MSCs pre-treated with the intracellular ROS generator pyocyanin (500 \(\mu\)M) were used as positive controls (Figures 5a and b). Cultivation of MSCs in aSerum lead to an

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**Figure 3** Serum-dependent proteomic changes in yMSCs and aMSCs. Two-dimensional gel electrophoresis and MS were used to analyze the protein expression pattern of MSCs cultured in ySerum and aSerum, respectively. (a) Graph depicts the number differentially expressed proteins of MSCs cultured in aSerum compared with corresponding cultures in ySerum. (b) Pie charts show the results of functional annotation clustering according to (left) cellular component and (right) biological process of the serum-dependently expressed proteins. Major clusters are indicated by bold font. (At least three independent experiments with two technical replicates were carried out for all treatment groups)
Figure 4  MSC culture in aSerum modulates the expression of HSP and molecular chaperones. (a) Representative western blots for HSP90, HSP70, HSP60, HSP40, protein disulfide isomerize (PDI), and the corresponding γ-Adaptin control blot. (b) Quantification of western blot analysis depicting the expression level of the indicated protein normalized to γ-Adaptin and relative to corresponding MSCs in ySerum. (For (b), n = 6, *P < 0.05; Mann–Whitney U-test)

Figure 5  Intracellular ROS concentrations and the amount of oxidatively modified proteins are increased in MSC cultures with aSerum. Both, cultures of (a) yMSCs and (b) aMSCs with aSerum displayed higher ROS production than corresponding cultures in ySerum. Supplementation of the antioxidant NAC attenuated the aSerum-induced ROS generation in both MSC populations. Intracellular ROS level were determined using CM-H2DCFDA and normalized to total DNA content. Cell cultures pre-treated with pyocyanin (500 μM) and a combination of pyocyanin and NAC (2 mM) served as controls. (c) Cultivation of MSCs in aSerum results in significantly higher levels of oxidatively modified proteins. ROS-related damages to proteins were analyzed by using OxyBlot Protein Oxidation Detection Kit. Graph shows quantified signal intensities normalized to loading control (γ-Adaptin). (d) To investigate a potential relationship between oxidative stress and altered MSCs differentiation ability, yMSCs were cultured in OM with pyocyanin (5 μM) and/or NAC (10 μM). Graph depicts quantitative analysis of the amount of AR staining normalized to cell number (osteogenic differentiation index) for each treatment group. (For (a–c), n = 6, (d) n = 3; *P < 0.05, **P < 0.01—pyocyanin group versus all other treatment groups; (a, b and d) ANOVA with Bonferroni correction; (e) Mann–Whitney U-test)
increased accumulation of ROS compared with the corresponding cells grown in ySerum, indicating enhanced intracellular oxidative stress. Notably, supplementation of the potent antioxidant N-acetyl-L-cysteine (NAC) attenuated the pyocyanin-induced as well as the aSerum-induced ROS generation. Enhanced ROS production has been previously shown to induce global macromolecular damages in cellular constituents like cytosolic proteins, DNA and lipids. To elucidate if the aSerum-induced ROS levels are sufficient to cause protein damages, the levels of protein oxidation (carbonyl groups) were determined as a surrogate marker (Figure 5c). In line with the enhanced ROS levels in cells cultured in aSerum, also the levels of carbonyls in yMSCs and aMSCs were 1.7-fold ($P = 0.013$) and 2-fold ($P = 0.038$) higher, respectively. These results strongly suggest that growth of MSCs in aSerum promotes oxidative stress, and further strengthen our assumption that ROS might be the principal cause for the functional alteration in response to an age-altered systemic environment. To validate this hypothesis, we cultivated yMSCs in osteogenic media (OM) supplemented with pyocyanin (5 μM) or a combination of pyocyanin and NAC (2 mM; Figure 5d). Supplementation of pyocyanin significantly reduced matrix mineralization (1.9 ± 0.3) compared with corresponding osteogenic controls without supplements (5.5 ± 0.4, $P < 0.001$). Similar to the results from the ROS measurement, the additional supplementation of NAC could at least partially compensate the pyocyanin effect (3.5 ± 0.4, $P < 0.007$).

**Oral antioxidant treatment enhances bone regeneration in aged animals.** Our *in vitro* analysis suggested that the age-altered systemic environment compromises MSC function via the induction of intracellular (oxidative) stress, which could be (at least partially) reversed by treatment with the antioxidant NAC. Next, we investigated whether systemic antioxidant administration is capable to improve bone healing in aged Sprague–Dawley rats. In parallel, we monitored NAC therapy in middle-aged animals with suboptimal fixation stiffness, which also delays bone regeneration despite the presence of a biologically competent environment.

Six weeks after surgery, qualitative radiologic analyses were performed using microcomputed tomography (micro-CT; Figures 6a–f). This analysis revealed that NAC treatment of middle-aged and aged animals results in an increased bone fraction volume (BV/TV) compared with the corresponding untreated controls ($P = 0.038$ and $P = 0.032$, respectively). On the microstructural level, NAC treatment led to a significantly increased trabecular number (Tb.N; $P = 0.014$ and $P = 0.009$, respectively) and reduced trabecular separation (Tb.Sp; $P = 0.010$ and $P = 0.009$, respectively) compared with the untreated controls. No difference in trabecular thickness (Tb.Th) was observed between the regenerated tissue of NAC-treated and untreated animals ($P = 0.254$ and $P = 0.520$, respectively).

To investigate whether the improved mineralization or tissue microstructure of regenerated bone after NAC treatment is also accompanied by an enhanced biomechanical tissue quality, qualitative biomechanical testing was performed (Figures 6g–i). Similar to the microCT data, biomechanical testing showed that NAC treatment significantly increases maximum torque at failure ($MTF = 36 ± 9$ (% contralateral)), torsional stiffness ($TS = 27 ± 8$ (% contralateral)) and energy to failure ($ETF = 68 ± 19$ (% contralateral)) of the regenerated bone of aged animals compared with the untreated group ($MTF = 12 ± 3%$ $P = 0.023$; $TS = 7 ± 3%$ $P = 0.025$; and $ETF = 16 ± 6%$ $P = 0.013$). In addition, also in the middle-aged animal group with suboptimal fixator stiffness, NAC treatment led to superior healing ($MTF = 60 ± 10%$; $TS = 50 ± 7%$; and $ETF = 54 ± 10$) compared with the untreated controls ($MTF = 25 ± 4%$ $P = 0.014$; $TS = 18 ± 4%$ $P = 0.010$; and $ETF = 21 ± 4%$ $P = 0.014$).

**Discussion**

There is evidence that tissue regeneration is impaired in aged individuals, although details of the underlying mechanisms remain unclear. Moreover, conventional therapies seems to be less effective in aged than in young patients, resulting in a clear demand for new regenerative approaches to overcome the pathophysiological challenges in the elderly. Thus, the aspect of aging represents one of the major issues in the field of regenerative medicine. Apart from an incomplete understanding of the exact mechanisms by which certain adult progenitor cells, like MSCs, contribute to regeneration, there is a paucity of data about how growing old influences their regenerative potential. In this study, we focused our analyses on bone regeneration as this is one of the few tissues lacking scar formation and allowing complete restoration of initial structure and function under optimal biomechanical and biochemical conditions. More importantly, previous investigations showed an age-related decline in bone regeneration, which can be traced back to a decline in both the number and function of bone marrow MSC. However, underlying mechanisms for age-associated decrease in MSC regeneration potential are still elusive. In addition to intrinsic alterations, extrinsic systemical factors seem to have an important role in stem cell aging.

In this study, we showed that MSC proliferation and differentiation *in vitro* is highly affected by the systemic environment provided by the surrounding serum. MSCs cultured in aSerum exhibited a decreased proliferation and osteogenic differentiation ability and an increased differentiation into adipocytes compared with cells in ySera. These observations were independent from the intrinsic age of the analyzed MSCs. Our results are in concordance with another *in vitro* study demonstrating a decreased osteoblast differentiation of human MSCs from women in aSera compared with ySera.

Altered proliferation and differentiation of MSCs could be because of (a) systemic factors in ySera, which directly promote these processes or (b) systemic factors in aSera, which indirectly compromise these processes via increased cellular stress. Enhanced MSC apoptosis and the higher expression of cell cycle inhibitors in response to aSera led us to assume the latter. These observations are in line with the results from other studies showing an excessive *in vivo* and *in vitro* expression of cell cycle inhibitors in satellite cells of aged muscle fibers. Correspondingly, our proteome analysis revealed altered expression of proteins associated with the mitochondria, protein (re)folding and the general...
Mitochondria are the major source of endogenous ROS and mitochondrial dysfunction leads to elevated oxidative stress levels, which compromise the overall ability of cells to produce and maintain properly folded proteins. Thus, enhanced oxidative stress results in accumulation of misfolded or damaged proteins, which induces the unfolded protein response and enhances the production of molecular chaperones. Under normal conditions, molecular chaperones such as HSP90, HSP70, HSP60 and HSP40 have numerous roles in cellular processes, including normal protein folding during translation, refolding of accumulated proteins and the regulation of protein degradation. In response to various environmental stresses, HSPs are upregulated to refold damaged molecules and/or to prevent their irreversible aggregation with other proteins. Accordingly, HSPs are abundantly expressed (1–2% of the total protein) in unstressed cells but their levels increase to 4–6% upon proteotoxic stresses. In this study, we observed elevated expressions of HSP90, HSP70 and HSP40 in response to an age-altered systemic environment, which coincide with enhanced intracellular ROS levels and increased amounts of oxidatively modified proteins. HSP90 and HSP70 are highly conserved proteins and represent two different classes of molecular chaperone. HSP70 prevents irreversible aggregation of unfolded proteins and catalyzes their refolding in an energy- and HSP40-dependent reaction. HSP90 also prevents the aggregation of intermediately folded proteins, but lacks the ability to refold denatured proteins. In addition, HSP90 and HSP70 exhibit anti-apoptotic functions and promote cell survival via inhibiting caspase activation or subsequent cleavage of their substrates. However, if HSP response fails to restore correct protein conformation or to degrade these proteins via the proteasomal system, it triggers cell cycle arrest, apoptosis or senescence.
we found that the expression of HSP60 is reduced in MSCs cultured in aSerum. A similar finding was reported by another study, showing elevated HSP90, HSP60, HSP40 levels but reduced expressions of HSP60 in MSCs upon temperature stress. HSP60 primarily localizes to the mitochondrial matrix and is involved in proper folding of newly translated mitochondrial proteins. The role of HSP60 in the process of cell death is controversial, since pro-survival and pro-apoptotic functions have been reported. For example, it prevents stress-induced apoptosis of epithelial cells via direct inhibition of caspase 3 and is upregulated in several types of cancer. On the other hand, HSP60 promotes proteolytic maturation of caspase 3 and enhances apoptosis in HeLa and Jurkat cells. Thus, the function of HSP60 in the process of cell death seems to depend on the apoptotic context and the reduced expression in response to aSer might indicate an altered mitochondrial structure and function. Collectively, our findings of altered HSP expressions, intracellular ROS levels and the amount of oxidatively modified proteins in MSCs provide further evidence for increased intracellular oxidative stress in response to an age-altered systemic environment, which in turn compromises MSC proliferation and differentiation.

The causal relationship between oxidative stress and altered MSCs differentiation ability is further supported by the finding that acute exposure to pyocyanin, known inducer of ROS, significantly reduced matrix mineralization, which could be at least partially reversed by enhanced oxidative stress protection (NAC treatment). Accordingly, other studies also reported that enhanced oxidative stress levels inhibited osteogenic differentiation and triggered the adipogenic differentiation of adipose tissue-derived stem cells and skeletal muscle precursors. Thus, our in vitro results led us to conclude that age-related changes of the systemic environment have a higher impact on MSCs than intrinsic aging. Irrespective of the cell donor age, aSerum compromises MSC function and survival via enhanced oxidative stress. This mechanism might also affect other mesenchymal cell types required for bone regeneration and may result not only in delayed healing, but also in a limited quality of the regenerated tissue.

Verifying this hypothesis, we subsequently investigated to which extend a systemic antioxidant administration enhance the in vivo bone regeneration in impaired healing situations. We showed that oral treatment with the potent antioxidant NAC is sufficient to improve the mechanical properties and the microstructure of the regenerated bone tissue in aged animals. However, previous studies showed that the long-term bone-healing outcome is not only dependent on the animal age but also significantly influenced by local mechanical conditions. A certain degree of movement within the fracture gap is required for uneventful healing, but high magnitudes of mechanical strain (e.g., as a consequence of a suboptimal fixator stiffness) prolong the initial inflammation phase and delay healing, potentially also mediated via locally enhanced oxidative stress levels. Therefore, we used middle-aged rats with semi-ridge fixator system and could show that treatment with NAC is also a sufficient approach to restore appropriate tissue regeneration in a mechanically challenging environment. On the basis of these results, systemic NAC administration seems to be a potent option to prevent poor healing outcomes under both age- and mechanically compromising conditions.

In summary, we propose that the systemic environment crucially contributes to the age-related decline in endogenous (bone) regeneration by increasing intracellular oxidative stress levels, hence compromising viability and function of mesenchymal (progenitor) cells. From the clinical perspective, our study suggests that systemic antioxidant treatment is a promising opportunity to restore the tissue regeneration potential, especially in elderly patients. From the research perspective, our study prompts for the identification of age-related changes in the serum, but the nature of the causative factor(s), for example, whether they are proteins, peptides or small molecules, is still elusive. We currently aim to validate our findings in the human system and to improve our MALDI-MS proteomic methods to elucidate biochemical alteration in serum from young and aged human donors. Further studies have to be performed to validate our findings in the human system and to identify the serum factor(s) causative for the observed cellular and molecular alterations.

Materials and Methods

Ethics statement. All experiments involving the use of animals were in compliance with the policies and principles established by the Animal Welfare Act, the NIH Guide for Care and Use of Laboratory Animals and the National Animal Welfare (TierSchG) and were approved by State Office of Health and Social Affairs Berlin (permit number: T0434/08, T0232/07 and G0091/10).

MSC and serum isolation. Bone marrow and serum were isolated from 3 weeks to 12 months old male Sprague–Dawley rats (Harlan Winkelmann, Eysinga, Germany, www.harlan.com). The serum from 18 animals per age group was combined in one serum pool and each pool was considered as one biological replicate. For all in vitro assays using rat serum, a minimum of six biological replicates in at least three technically independent experiments were performed on separate days.

MSCs were isolated from the bone marrow of the Sprague–Dawley rats and selected by plastic adherence. For cultivation of MSCs, Dulbecco’s modified Eagle’s medium (DMEM; Life Technologies, Darmstadt, Germany) supplemented with 10% fetal calf serum (Biochrom, Berlin, Germany) and 100 U/ml penicillin – 100 μg/ml streptomycin was used (expansion medium). Culture medium was substituted twice a week and cells were harvested after reaching 70–80% confluence. Cells were subcultured by trypsinization and seeded at 3300 cells/cm² on uncoated culture dishes. Cell number and viability of trypsinized MSCs were determined using the cell counter CASY TT (Roche, Mannheim, Germany, www.roche-applied-science.com).

Functional assays. All assays were carried out with primary MSCs in passages 2–4. In order to investigate the age-dependent influence of serum on primary MSCs, functional assays were carried out as described below with 10% serum from either young or aged rats.

Proliferation. For short-term proliferation assays, 2000MSCs/cm² were seeded onto 96-well plates (96-MTP). Cell number was measured 1 and 4 days after seeding using CyQuant assay (Promega, Mannheim, Germany, www.promega.com) according to the manufacturer’s instructions.

Differentiation. Osteogenic differentiation of confluent MSCs was induced by using OM (DMEM, 10% rat serum, 50 μM ascorbic acid, 10 mM β-glycerophosphate and 0.1 μM dexamethasone). The matrix mineralization was visualized with AR staining. Quantification was achieved by measuring the absorbance of AR (optical density AR) that was normalized to number of viable cells determined by AlamarBlue (Life Technologies). If indicated, pyocyanin (5 μM) and NAC (10 μM) were added to the culture medium. Adipogenic differentiation was induced by using adipogenic medium (AM; DMEM, 10% rat serum, 2 μM insulin,
100 μM indomethacin, 500 μM isobutylmethylxanthine and 1 μM dexamethasone) and quantified after OR staining, which was normalized to cell number. **Capase-3/7 activity assay.** For short-term proliferation assays, 2000 MSCs/cm² were seeded onto 96-well plates (96-MTP). Capase-3/7 activity was measured after 4 days of culture using Apo-ONE Homogeneous Caspase-3/7 (Promega) according to the manufacturer’s instructions.

**Western blot.** The Novex system was used according to the Invitrogen NuPAGE protocol. Primary antibodies were mouse (2-μg rat CDK1acin (Promega) at 1:1000, Abcam, Cambridge, UK, www.abcam.com) and mouse (2-μg GAPDH) at 1:7000, Abcam). As secondary antibody goat (2-μg IgG) peroxidase was utilized. HSP90/chaperone expression was determined using HSP90/chaperone Antibody Sampler Kit according to the supplier’s protocol (New England Biolabs, Frankfurt am Main, Germany, www.neb-online.de). Amount of total oxidized proteins was determined using the OxylBlot Protein Oxidation Detection Kit (Merck Millipore, Darmstadt, Germany, www.merckmillipore.de) according to the manufacturer’s instructions. Bound secondary antibodies were detected using the chemiluminescent substrate ECL (GE Healthcare Piscataway, NJ, USA, http://www.gelifesciences.com). Band intensities were quantified by NIH ImageJ software package (http://rsb.info.nih.gov/ij/).

**2D gel electrophoresis and MS.** MSCs were cultivated for 5 days in expansion medium containing serum from aged and young rats, respectively. Subsequent to washing cells with phosphate-buffered saline (PBS), they were harvested with PBS (containing 5 mM EDTA and a phosphatase/protease inhibitor cocktail) and by cautious scraping on ice. Cells were washed again with PBS and the remaining buffer was quantitatively removed before shock freezing of cells in liquid nitrogen. Seventy micrograms of protein lysates were separated by 2DE and harvested with PBS (containing 5 mM EDTA and a phosphatase/protease inhibitor cocktail) and by cautious scraping on ice. Cells were washed again with PBS and the remaining buffer was quantitatively removed before shock freezing of cells in liquid nitrogen. Seventy micrograms of protein lysates were separated by 2DE and harvested with PBS (containing 5 mM EDTA and a phosphatase/protease inhibitor cocktail) and by cautious scraping on ice. Cells were washed again with PBS and the remaining buffer was quantitatively removed before shock freezing of cells in liquid nitrogen.

Protein spot of interest were excised from gels and subjected to in-gel tryptic digestion. Tryptic fragments were analyzed by nanoflow high-performance liquid chromatography (nanoHPLC; Proxeon Easy-nLC, Odense, Denmark)/electrospray ionization—MS/MS on a LCQ Deca XP ion trap instrument (Thermo Finnigan, Waltham, MA, USA) and MALDI TOF/TOF on AutoFlex III (Bruker Daltonik, Bremen, Germany). Mass spectra were analyzed using our in-house MASCOT software package license version 2.2 (Matrix Science, London, UK) automatically searching the SwissProt database for rate (SwissProt 51.8). MS/MS ion search was performed with this set of parameters: (i) taxonomy: Ratus norvegicus, (ii) proteolytic enzyme: trypsin, (iii) maximum of accepted missed cleavages: 1, (iv) mass value: monoisotopic, (v) peptide mass tolerance: 0.8 Da, (vi) fragment mass tolerance: 0.8 Da and (vii) variable modifications: oxidation of methionine and acrylamide adducts (propionamide) on cysteine. Only proteins with individual ions scores >31 or extensive homology (>0.05), with at least two independent peptides identified were considered. To conduct functional categorizing, all differentially expressed proteins were submitted to the Database for Annotation, Visualization and Integrated Discovery (DAVID). Proteome analysis was done in at least two technically independent experiments with three biologically independent replicates for each MSC/serum combination.

**Reactive oxygen species (ROS) measurement.** Concentrations of ROS were determined using CM-H2DCFDA (Life Technologies) as described previously. Cells were cultivated in expansion medium containing either aSera or ySera. Subsequently, cells were incubated with medium supplemented with 10 μM CM-H2DCFDA for 20 min at 37 °C. The resulting fluorescent signal was measured using Infinite 200Pro plate reader (Tecan, Mainz, Germany, www.tecan.com). Cells pre-treated with 500 μM pyocyanin and/or 2 mM NAC for 20 min were used as positive or negative control, respectively. The fluorescent intensities were normalized to the cell number determined by CyQuant.

**Animal surgical procedure and NAC treatment.** Operations and postoperative care were performed according to a previously published protocol and it used a standardized biomechanically validated external fixation device. The experimental model has been previously described and is briefly summarized here. Sixteen aged (52 weeks) and 16 middle-aged (12 weeks) Sprague–Dawley rats were divided into subgroups of eight animals depending on the administration of NAC (NAC−/NAC+). Animals were not restricted in weight bearing. Based on the mean drinking volume of 25 ± 3 ml/day (95% CI) of adult Sprague–Dawley rats, as evaluated in about 100 animals of our previous studies, a high dosage of NAC was administered via freely available drinking water (1.4 g/kg of body weight/day; ACC 600, Hexal AG, Holzkirchen, Germany) in the NAC+ subgroups from surgery until killing. All other animals (NAC−) received plain water for drinking. Daily, well-being and drinking amount were controlled.

**Surgical procedure.** Using an anterolateral approach, the left femur was osteotomized at the mid-shaft, distracted to a gap of 1.5 mm and externally fixated using a previously described fixation system. The distance between fixator and bone (offset) was set to 7.5 mm in the rigid configuration (aged animals) and 15 mm in the semi-rigid (middle-aged rats) configuration. Two-weekly, in general anesthesia, a radiographical examination was performed regarding the occurrence of implant failures and bony consolidation. One aged (untreated group) and two middle-aged (untreated group) animal were excluded from further analysis as a result of infection or implant failure, respectively. Total follow-up of the study was 6 weeks. After animal killing, both femurs were harvested.

**3D in vitro micro-CT.** Directly after harvesting the osteotomized and contralateral femur, the K-wires were carefully removed and the specimens were placed in cooled (4 °C) PBS solution in acrylic custom-made batch scanning tubes. Subsequently, bones were scanned on a Viva40 micro-CT (Scanco Medical AG, Brüttisellen, Switzerland) and evaluated as described previously, except voxel size was selected to be isotropic and fixed at 10.5 μm and the fixed global threshold was 445.5 mg HA/cm³. Using the binarized images, the following morphometric key parameters for bone microarchitecture were determined: (i) bone volume fraction (BV/TV (%)—ratio of segmented bone to total volume), (ii) Tb.Th (mm), (iii) Tb.N (1/mm)—average number of trabeculae per unit length, (iv) Tb.Sp (mm)—mean distance between trabeculae. All analyses were performed on the digitally extracted callus tissue using 3D distance techniques (Scanco Software, Scanco Medical AG).

**Biomechanical testing.** Immediately after scanning, at least 4 h after killing, both femurs were embedded with methyl methacrylate (Technovit 3040, Heraeus Kulzer, Hanau, Germany) in custom-made casting containers leaving the fracture zone between the inner pin holes and an equivalent zone at the contralateral bones for torsional testing (ElectroForce TestBench LM1, Bouse, Framingham, MA, USA) under an axial pre-load of 0.3 N, an adjustment time of 0.1 N/s, and at a torsion speed of 0.5/°s until failure. The M1F, the TS and the ETF of the operated bone was reported as percentage of intact contralateral bones.

**Statistical analysis.** The SPSS 18.0 software package (SPSS Inc., Chicago, IL, USA) was used for statistical evaluation. If not stated otherwise, results from at least six independent experiments were analyzed for statistical significance using the Mann–Whitney U-test (two groups). Multiple pairwise comparisons were performed by one-way analysis of variance (ANOVA, repeated measures) and P-values were adjusted using Bonferroni’s P-value adjustment multiple comparison procedure. Unless otherwise specified, results are presented as mean ± S.E.M. All tests were analyzed two-sided and P<0.05 was regarded as significant.

**Conflict of Interest**

The authors declare no conflict of interest.

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**Author contributions**

SG, MT, GK, GND and PS designed research; SG, MT, OK, KS-B, AE, DJ and PS performed research; SG, MT, KS-B, DJ, AO, JK, AD, GND and PS analyzed data, SG, MT, AE, DJ and GK contributed reagents/materials/analysis tools and SG wrote paper.
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