Emerging dynamics of macrophage cellular senescence and immunosenescence in governing organismal aging and disease: Concepts and opportunities

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Abstract

An intricate relationship between impaired immune functions and the age-related accumulation of tissue senescent cells (SC) is rapidly emerging. The immune system is unique as it undergoes mutually inclusive and deleterious processes of immunosenescence and cellular senescence with advancing age. While factors inducing immunosenescence and cellular senescence may be shared, however, both these processes are fundamentally different which holistically influence the aging immune system. Immunosenescence is a well-characterized phenomenon, but our understanding and biological impact of cellular senescence in immune cells, especially in the innate immune cells such as macrophages, is only beginning to be understood. Tissue-resident macrophages are long-lived, and while functioning in tissue-specific and niche-specific microenvironments, senescence in macrophages can be directly influenced by senescent host cells which may impact organismal aging. In addition, evidence of age-associated immunometabolic changes as drivers of altered macrophage phenotype and functions such as inflamm-aging is also emerging. The present review describes the emerging impact of cellular senescence vis-à-vis immunosenescence in aging macrophages, its biological relevance with other senescent non-immune cells, and known immunometabolic regulators. Gaps in our present knowledge, as well as strategies aimed at understanding cellular senescence and its therapeutics in the context of macrophages, have been reviewed.

Keywords: Macrophages, Senescence, Immunosenescence, Immunometabolism, Aging

Abbreviations:
SC: senescent cells
SASP: senescence-associated secretory phenotype
SA-β-gal: senescence-associated β-galactosidase
mTOR: mammalian target of rapamycin
ROS: reactive oxygen species
NAD: nicotinamide adenine dinucleotide
1. Introduction

The last decade has seen rapid progress in our understanding of the molecular etiology and pathological effects of aging. There is increasing evidence that the stochastic phenomenon of aging is essentially a culmination of cellular and molecular damage over time that gradually manifests as the visible macrophenotype of aging (Bhatia-Dey et al., 2016; Chandrasekaran et al., 2017). Several different theories were formulated to explain the causes and mechanisms of aging, but gerontologists are now showing considerable interest in the cellular senescence centric interpretation of aging which is beginning to uncover an integrative and tangible understanding of the known effects of aging (Dodig et al., 2019; Borghesan et al., 2020). Aging itself is not a disease, but it rather predisposes elderly to several age-associated disorders including the characteristic hallmark diseases of 21st century, i.e., cancer and diabetes (Huo et al., 2018; DeSantis et al., 2019). In fact, a view is now emerging that the various age-related diseases should be studied in the purview of cellular senescence, and that targeting cellular senescence itself could be a novel approach in preventing frailty and the aggravation of a number of age-associated disorders (Borghesan et al., 2020; Report, 2021). This is supported by several studies which have demonstrated that accumulation of SC shorten healthspan and lifespan (Baker et al., 2016), and delaying the development of cellular senescence (Sharma et al., 2021) or lifelong removal of SC can enhance longevity (Baker et al., 2011), improve tissue regeneration (Krishnamurthy et al., 2006) and prevent the development of chronic age-associated disorders (Palmer et al., 2019; Lewis-McDougall et al., 2019; Cai et al., 2020; Ogrodnik et al., 2021). In fact, the targeted removal of SC using senolytics is an emerging area of research, and active preclinical search, as well as clinical assessment of putative senolytics is presently underway (Xu et al., 2018; Hickson et al., 2019; Wissler Gerdes et al., 2020).

The immune system is a complex, yet coordinated and highly plastic network of cells which has been traditionally recognized to protect the integrity of the organism from invading pathogens. However, it is now understood that the pattern recognition receptors of immune cells not only recognize the pathogen-associated molecular patterns, but also respond to cell-intrinsic damage-associated molecular patterns or DAMPs suggesting their broader and more significant physiological role (Mogensen, 2009). The innate and adaptive branches of the immune system communicate either directly or through a variety of signaling molecules, while the integrated neuroendocrine-immune system governs the interactions between immune cells and other regulatory systems. As observed with different physiological responses, the immune system also undergoes characteristic functional and phenotypical deterioration with age resulting in feeble response to pathogens and vaccination, and the development of chronic age-associated inflam-maging (Aw et al., 2007). These changes have been loosely described as ‘immunosenescence’, although whether it is the cause or a consequence of aging is still unclear. Aging in immune cells has traditionally been considered within the purview of immunosenescence; however, similar to other mammalian cells, immune cells are also liable to be affected by the different facets of cellular senescence which can have potentially deleterious impact on immune functions (Vicente et al., 2016). Although the triggers of immunosenescence and cellular senescence may overlap, but both these processes show fundamentally distinct characteristics, and yet have the potential to impair immune functions with age (Burton and Stolzing, 2018) (Table 1). However, unlike immunosenescence, our understanding of the role, and particularly the biological impact of cellular senescence in immune cells is only emerging. This is especially relevant for innate immune cells such as macrophages which form the basis of parenchymal homeostasis, integrity, and perform unique tissue-specific functions as resident immune cells. This review describes our present
understanding and relevance of cellular senescence in immune cells \textit{per se}, with an emphasis on macrophages. The lacunae in our present knowledge, as well as the potential for further research is discussed to understand how senescent macrophages contribute to the aging process.

**Table 1: Selected markers of immunosenescence and cellular senescence in macrophages**

| Immunosenescence                          | Cellular senescence |
|------------------------------------------|---------------------|
| Decreased activation response to LPS/IFN-γ | Cell cycle arrest; Persistent p53/p21 and/or p16 expression |
| Reduced TLR expression                   | DNA damage          |
| Impaired chemotaxis                       | Telomere attrition  |
| Decreased cytokine response              | Presence of SASP    |
| Impaired wound healing                    | Hypertrophy         |
| M1/M2 polarization                        | Reduced cell proliferation |
| Impaired antigen presentation             | Increased SA-β-gal activity |
| ROS burden*                              | ROS burden*         |
| Metabolic alterations*                    | Metabolic alterations* |

*Overlapping parameters

2. **Cellular senescence, immunity and aging: emerging concepts**

The fact that mitotic mammalian cells fail to divide after a certain finite number of cell divisions \textit{in vitro} was a seminal discovery in geroscience (Hayflick and Moorhead, 1961). The so-called ‘Hayflick limit’ seemingly provided a link between cell division and organismal aging, but this phenomenon remained controversial and was initially dismissed as a mere \textit{in vitro} artifact (Hayflick, 1998; de Magalhães, 2004). However, multiple studies have now confirmed that SC play a causal role in governing organismal aging and age-dependent disorders (Campisi and Robert, 2014; Childs et al., 2015; Katzir et al., 2021), and in fact, cellular senescence is established as one of the well-studied hallmarks of aging (López-Otín et al., 2013). Cellular senescence is characterized by cellular hypertrophy, permanent cell cycle arrest by the activation of cell cycle inhibitory pathways such as p53/p21\textsuperscript{WAF1} and/or p16\textsuperscript{Ink4a}/pRb, resistance to apoptosis by deregulation of Bax/Bcl-2 signaling, telomere attrition, development of a senescence associated secretory phenotype or SASP, and altered metabolic longevity pathways such as SIRT and mTOR (Carroll and Korolchuk, 2018; Hernandez-Segura et al., 2018; Maduro et al., 2021; Kumari and Jat, 2021). The induction of cellular senescence is intimately linked to external and intrinsic stressors. Age-dependent decline in cell stress response capacity results in impaired redox balance, augmentation of genotoxic stress and DNA damage that ultimately aids in cell cycle arrest and the development of cellular senescence (Robinson et al., 2018; Chen et al., 2020). Moreover, there is evidence that the characteristic telomere attrition, as observed in SC, could also be accelerated by...
oxidative damage (von Zglinicki, 2002; Ludlow et al., 2014). However, it is important to consider that SC are naturally occurring cells and their presence per se does not reflect on the pathological state of the organism, since SC are essential for processes such as wound healing and repair (Demaria et al., 2014). In younger organisms, SC are constantly replaced, but as we age, SC turnover decreases and tissue SC burden rapidly increases which contributes to age-related pathologies (Karin et al., 2019). More significantly, accumulation of SC result in the chronic presence of SASP which has pro-tumorigenic and pro-inflammatory paracrine effects on nearby healthier cells (Coppé et al., 2010). Thus, cellular senescence appears to be an antagonistic hallmark of aging and a candidate example of antagonistic pleiotropy theory of aging.

The exact causes of SC accumulation with age are yet to be completely understood. In this regard, the role of immune cells in maintaining SC cell homeostasis in tissues is being increasingly recognized (Prata et al., 2018). The SASP components of SC serve to chemotactically attract various immune cells such as macrophages, NK cells and cytotoxic T cells which subsequently recognize the upregulated immunogenic ligands on SC (such as MICA and ULBP2) resulting in their efficient removal (Coppé et al., 2010; Sagiv et al., 2016; Hall et al., 2016). That SC accumulate in aging organisms, could at least in part be ascribed to deficient immune functions/immunosurveillance associated with immunosenescence and/or cellular senescence (Figure 1). Indeed, a recent work showed that mice deficient in the cytotoxic potential of immune cells exhibited both higher SC tissue burden and chronic inflammation suggesting that weakened immunosurveillance could be a critical aspect regulating SC accumulation in vivo (Ovadya et al., 2018). Another study showed that impaired immunosurveillance in CD4+T cells results in the development of murine hepatocellular carcinomas, thus suggesting that senescence surveillance is important for tumor suppression in vivo (Kang et al., 2011). Similarly, it was reported that NK cells play a critical role in eliminating senescence activated stellate hepatic cells both in vitro and in vivo (Krizhanovsky et al., 2008). The role of macrophages in the removal of SC has long been recognized (Kay, 1975; Mevorach et al., 2010). Using a co-culture system, a recent study showed that macrophages can induce TNF-α mediated apoptosis in senescent fibroblasts followed by their phagocytic clearance (Ogata et al., 2021). Macrophages are also involved in the elimination of SC in the uterine stroma surrounding the embryo implantation site following parturition in mice (Egashira et al., 2017). In addition, the SASP of senescent hepatic cells not only attracts macrophages (Irvine et al., 2014), but also influences the M1/M2 polarization state of macrophages (Lujambio et al., 2013). It is also interesting to note that SC themselves can also develop features of immune evasion. It was demonstrated that SC can escape the immune system by upregulating the expression of non-classical MHC molecule HLA-E, which inhibits NK cells and CD8+ T cells mediated immune responses against SC, thereby aiding in inefficient removal and accumulation.
of SC (Pereira et al., 2019). This potentially suggests autonomous functioning of SC, much in line with tumorous cells and requires further exploration.

**Figure 1:** Immune cells mediated removal of SC is affected by cellular senescence and immunosenescence. In young organisms, SC secrete SASP factors which attract innate and adaptive immune cells resulting in the removal of SC. During aging, immunosenescence and cellular senescence collectively impair several immune functions which may together contribute to the inefficient clearance of SC and their gradual accumulation.

3. **Macrophage senescence: culmination of immunosenescence and cell senescence?**

The immune system reacts to exogenous and endogenous threats by initiating an inflammatory response. Macrophages are involved both during the onset of inflammation as well as its successful resolution. The recognition of a foreign particle prompts proinflammatory behavior in macrophages (also called classical activation or M1) characterized by increased production of inflammatory cytokines, respiratory burst and phagocytosis (Orecchioni et al., 2019). To resolve the latent inflammation and debris, macrophages switch to an anti-inflammatory M2 phenotype by initiating reconstruction of damaged tissue and by prompting synthesis of extracellular matrix and cell growth (Rőszer, 2015). As antigen presenting cells, macrophages are involved in the regulation and differentiation of adaptive immune responses, while scavenger receptors on macrophages recognize and remove oxidized proteins/lipoproteins expressing damaged or apoptotic cells (Krieger and Herz, 1994; Schliehe et al., 2011). These diverse functions of
macrophages are severely hampered with age and are often attributed to characteristic immunosenescence. For instance, there are reports suggesting that M1/M2 polarization of macrophages is affected during aging, although the nature of this skewness is debatable. Age-dependent M1 to M2 polarization of macrophages has been observed in aging skeletal muscles (Cui et al., 2019) and during macular degeneration (Zandi et al., 2015), while we previously observed an anti-inflammatory phenotype in aged animals (Sharma et al., 2014). On the other hand, an age-dependent shift towards M1 pro-inflammatory state was also observed in aged mice which was ascribed to the degeneration of enteric nervous system (Becker et al., 2018). These observed differences could be attributed to the heterogeneity and plasticity of macrophages, but regardless, any perturbations in the normal M1/M2 balance in macrophages may dysregulate the development of the host response, making elderly more susceptible to inflammatory and infectious diseases (Mahbub et al., 2012; Wynn et al., 2013). Studies have also shown that macrophage phagocytic ability (Wong et al., 2017), TLR expression (Renshaw et al., 2002) and LPS-induced cytokine production declines with age (Chelvarajan et al., 2006). Importantly, it was also identified that age-related decrease in phagocytic ability was not observed in bone marrow derived macrophages and monocytes suggesting that age-related changes in macrophage functions might not be intrinsic, but could be related to external environmental factors (Linehan et al., 2014). Aging also results in decreased antigen presenting capacity of macrophages which may compromise the swiftness and the potency of adaptive immune response. Studies have shown that old mice showed impaired stimulation as well as expression of MHC class II molecules compared to macrophages from young mice after stimulation with IFN-γ (Davila et al., 1990; Herrero et al., 2001). Aging has also been shown to reduce macrophage mediated tissue repair and regeneration both in mice and humans. It was observed that local injection of peritoneal macrophages isolated from young mice could accelerate wound healing in older animals (Danon et al., 1989). Age-related shifts in macrophage infiltration into wounds, alterations in chemokine content, and a concurrent decline in wound macrophage phagocytic function was also documented which may contribute to the delayed repair response during aging (Swift et al., 2001). A recent study showed that injured arteries in aging rats develop thicker neointimas, which were significantly correlated with a higher number of tissue macrophages and increased vascular IL-18 (Rodriguez-Menocal et al., 2014). Further, in vivo depletion of macrophages by clodronate liposomes ameliorated the vascular accumulation of IL-18 and development of thicker neointimas (Rodriguez-Menocal et al., 2014).

Unlike immunosenescence, conclusive evidence of cellular senescence in macrophages is only beginning to be elucidated. Prior studies in T cells have shown that immune cells do exhibit the features of cellular senescence both in vitro and in vivo, while such work on macrophages remained scanty (Vallejo et al., 1999; Effros, 2004; Chou and Effros, 2013). Nonetheless, in vitro studies have shown that on exposure to external stressors, macrophages can develop senescence like features including p21/p53 mediated cell cycle arrest (Nyunoya et al., 2003; Fong et al., 2007), altered morphology and the development of SA-β-gal activity (Kim et al., 2015). In a recent study, we observed that peritoneal macrophages isolated from old mice exhibit heterogenous and elongated morphology, enhanced expression of cell cycle inhibitors p53/p21 WAF1/p16Ink4a, and SA-β-gal activity as compared to macrophages isolated from younger animals (Kumar et al., 2020). In another study, age-related increase in the characteristic marker of senescence p16 Ink4a inhibited the normal phenotype of adipose tissue macrophages and contributed to type II diabetes risk (Fuentes et al., 2011). A negative correlation between Ki67 proliferation marker and cellular senescence marker SA-β-gal expression was observed in cultured macrophages suggesting that SA-β-gal activity labels macrophages with decreasing proliferation tendency (Holt and Grainger, 2012).
recent study observed that the majority of p16<sup>Ink4a</sup> and SA-β-gal positive cells in the tissues of aging mice were, in fact macrophages, indicating a significant role of senescent macrophages which previously had been attributed to non-immune SC only (Hall et al., 2016). However, in their follow up work, the same authors argued that p16<sup>Ink4a</sup> and SA-β-gal expression may be unrelated to senescence, and may only be a part of a reversible response of macrophages to their physiological stimuli (Hall et al., 2017). Altogether, these observations suggest an unexpected role of p16<sup>Ink4a</sup> expression in macrophages which warrants deeper characterization vis-à-vis other markers of cellular senescence. However, another hallmark indicator of cellular senescence, i.e., telomere attrition, has also been observed in macrophages both during aging and disease. Analyses of bone marrow-derived macrophages showed strong age-dependent shortening of telomere length in aged (19–24-mo-old) mice as compared to young (6-wk-old) mice (Sebastián et al., 2009). Murine peritoneal macrophages isolated from first-generation telomerase reverse transcriptase-deficient mice significantly upregulated SA-β-gal activity and the expression of genes regulating cell cycle arrest such as p16, p21, and the retinoblastoma protein (Gizard et al., 2011). Mean telomere length in peripheral blood leukocytes was significantly shorter in sickle cell disease patients which directly correlated with the disease genotype and inflammation markers (Colella et al., 2017). Some studies have suggested that a characteristic SASP is also prevalent in macrophages. A recent report has shown that LPS treated THP-1 macrophages develop features of cellular senescence and SASP as evident by expression of p53/p21/p16 cell cycle inhibitors and various SASP components (Wang et al., 2020). Using a diabetic mice model, it was observed that macrophages in diabetic kidneys showed archetypal features of cellular senescence as evident from increased mRNA expression pertaining to cell cycle arrest (p16/p21) and SASP (IL-6, TGF-β, PAI-1, MCP-1) suggesting that senescent macrophages are prevalent in vivo and could be a potential source of SASP in disease condition (Prattichizzo et al., 2018).

It is thus reasonable to envisage that both cellular senescence and immunosenescence can impact macrophage functions. However, either a conclusive correlation or a causative linkage between cellular senescence and immunosenescence in macrophages or other immune cells has not been established yet. Nonetheless, studies on telomere shortening have highlighted multiple negative effects on immune functions and disease which could be related to some aspects of immunosenescence. For example, telomerase knockout mice developed exaggerated lung inflammation and increased mortality upon respiratory staphylococcal infection (Kang et al., 2018). Authors observed that telomere dysfunction caused macrophage mitochondrial abnormality, oxidative stress, and hyperactivation of the NLRP3 inflammasome, suggesting that loss of telomeres is sufficient to drive mitochondria mediated redox imbalance, and thus contribute to macrophage functional incapacity which could be directly related to immunosenescence (Kang et al., 2018). Similarly, using telomerase knockout mice, it was observed that telomere loss is the cause for the enhanced oxidative stress, reduced Stat5a oxidation and phosphorylation and, ultimately, for the impaired GM-CSF-dependent macrophage proliferation (Sebastián et al., 2009). It was also demonstrated that monocytes (but not lymphocytes) of aged type II diabetic patients had significantly lower mean telomere length and a significant inverse relationship between oxidative DNA damage and telomere length was observed in diabetic group (Sampson et al., 2006). Another study showed a causative link between accumulating ROS and the development of senescent phenotype in macrophages (Singh et al., 2019). Together, it appears that some degree of causation between cellular senescence and immunosenescence does exist, but further deeper studies are required to completely understand this dichotomy.
4. Interactions between macrophages and senescent cells

Macrophages are present throughout the body, wherein they reside as specialized tissue resident cells and perform tissue-specific and niche-specific functions. As a result, different tissue resident macrophages may be functionally distinct, and respond differentially to effector functions (Minutti et al., 2017; Lacerda Mariano et al., 2020). Moreover, in terms of cellular senescence, recent studies have shown that different aging tissues experience varying degree of SC burden as the rate of cellular senescence is dependent on both the age of tissue, as well as the type of tissue (Karin et al., 2019; Tuttle et al., 2020; Sharma et al., 2021). Thus, it is conceivable that dynamic interactions between specialized long-lived macrophages and host tissues with varying degree of cellular senescence, may differentially influence their functional capacity and/or the rate of cellular senescence (Figure 2). Although these aspects are less explored, our previous ex vivo study showed that the secretory phenotype of senescent preadipocytes, but not proliferating preadipocytes, could induce a senescence-like phenotype in peritoneal macrophages isolated from young animals (but not from old animals) characterized by a strong SA-β-gal activity, cellular hypertrophy, increase in p16Ink4a expression and intracellular ROS production (Kumar et al., 2020). This implies that in a heterogenous and dynamic in vivo system, the highly senescence-prone adipose tissue (Minamino et al., 2009; Sharma et al., 2021) could directly influence cellular senescence in adipose tissue macrophages. A recent study showed that macrophages induced apoptosis in senescent fibroblasts and prevented their accumulation, however, chronic presence of fibroblast SASP factors downregulated macrophage mediated apoptosis and phagocytosis thereby potentially suggesting that senescent tissues can affect macrophage functions which can contribute to increased tissue SC burden (Ogata et al., 2021). Similarly, another report identified that lysophosphatidylcholines expression is elevated in the phospholipids of senescent dermal fibroblasts which interfere with the toll-like receptor 2 and 6/CD36 signaling and subsequent phagocytic capacity in macrophages (Narzt et al., 2021). Exposure of macrophages to the secretome of senescent hepatocytes induced an M2 phenotype, while an M1 phenotype was observed in macrophages exposed to secretory factors of non-senescent hepatocytes (Sen et al., 2021). Similarly, exposure of macrophages to the conditioned media of senescent thyrocytes induced an anti-inflammatory M2 phenotype in macrophages showing high CD206 and low MHC II markers, and upregulated CCL17 secretion which ultimately promoted tumor progression (Mazzoni et al., 2019). It is thus evident that senescent cells can directly influence the functional and phenotypical aspects of aging in macrophages which can accelerate SC burden in tissues.
Figure 2: Bidirectional relationship between resident macrophages and tissues. Dynamic interactions between senescent host tissues and long-lived tissue resident macrophages may impact senescence and aging. SASP of host tissues can modulate macrophage functions and decrease its potency to remove damaged cells. Senescent macrophages can directly augment SC accumulation owing to impaired immunosurveillance.

5. Immunometabolism and macrophage senescence

Understanding the metabolic regulation of immune functions, especially during aging, is an emerging area of research (Lee et al., 2021). To precisely perform their diverse functions, immune cells like macrophages need to quickly shift between various activation states which are subject to tight regulation by metabolic pathways and nutrient availability. For instance, upon activation, M1 like macrophages rapidly convert arginine to nitric oxide through inducible NO synthase activity, while M2 macrophages convert arginine to ornithine for effector functions (Galván-Peña and O'Neill, 2014). Several studies suggest that nutrient signaling longevity pathways, such as mTOR, SIRT and Insulin/IGF-1 are the key modulators and therapeutic targets of health and lifespan (Mazucanti et al., 2015). Aged macrophages generally show heightened inflammatory metabolic pathways, despite reduced phagocytic ability, antigen presentation and mitochondrial dysfunction (van Beek et al., 2019; Lee et al., 2021). Using mass spectrometry-based proteomic approach, age-dependent alterations in inflammatory signaling, mitochondrial function, and cellular metabolic pathways were observed in aged microglia cells, while a component of the mTORC2 complex was identified as a novel upstream regulator of several biological functions (Flowers et al., 2017). Rapamycin-induced inhibition of mTOR signaling accelerated diabetic encephalopathy in rats through the augmentation of macrophage autophagy (Wang et al., 2018). Knockdown of SIRT1 in the mouse macrophage RAW264.7 cell line and in intraperitoneal macrophages broadly activated the JNK and IKK inflammatory pathways, and
increased LPS-stimulated TNFα secretion (Yoshizaki et al., 2010). On the other hand, overexpression of SIRT1 during bone marrow-derived macrophage differentiation increased their proliferative capacity and self-renewal (Imperatore et al., 2017). Sirtuin family of enzymes depend on the coenzyme NAD for their activation and function. NAD levels decline with age in different tissue types, including macrophages, which have been shown to increase inflammation and impair oxidative metabolism (Minhas et al., 2019). It is plausible that declining NAD levels in aged macrophages could also compromise SIRT activity that may negatively impact inflammatory homeostasis during aging (He et al., 2020). Interestingly, senescent macrophages have recently been implicated in the immunometabolism of NAD (Covarrubias et al., 2020). It was observed that NAD consuming CD38+ M1 macrophages with upregulated markers of senescence (p16/p21) and inflammation accumulated in adipose tissue which were primarily responsible for decreased NAD availability in tissues during aging, suggesting a causative relationship between CD38+ macrophages, NAD consumption, Sirtuins and aging (Covarrubias et al., 2020). In addition, M2 phenotype macrophages were identified as novel sources of IGF1, and ablation of IGF1 receptors from myeloid cells reduced phagocytosis, increased macrophages in adipose tissue, elevated adiposity, lowered energy expenditure, and led to insulin resistance in mice fed a high-fat diet indicating that IGF1 signaling shapes the macrophage-activation phenotype (Spadaro et al., 2017). A recent study showed the regulatory role of IGF1 signaling in cellular senescence and macrophage polarization wherein mice deficient for IGF1 receptors showed attenuated premature senescence in pneumocytes, reduced M2 macrophage polarization and lung fibrosis (Chung et al., 2020). Together, it is evident that understanding the metabolic control of cellular senescence mediated by nutrient sensing pathways is essential for completely comprehending the etiology and effects of macrophage senescence.

6. Unanswered questions and niche research areas

There are several lacunae in our present understanding of the causes and biological effects of cellular senescence in macrophages. This is of particular importance since tissue resident macrophages, similar to memory T cells, are considered to be long lived (Bain et al., 2016), and are thus liable to accumulate cellular and biochemical insults over time leading to genotoxic stress and the development of cellular senescence (Sedelnikova et al., 2004). Some of the key aspects to be investigated are:

- The extent and depth of in vivo cellular senescence in macrophages is unclear. There is a need to characterize macrophages for age-dependent as well as tissue-dependent changes pertaining to the development of cellular senescence using the established markers of immune cell senescence (Zhou et al., 2021). Given the high heterogeneity amongst macrophages, using multi-omics approach at single cell resolution would be a prudent approach.

- Macrophage secretory phenotype changes with age, but whether it is related to the SASP of non-immune cells, or if senescent macrophage secretome contributes to inflamm-aging remains to be fully understood. In this context, whether senescent macrophages play any direct role in driving age-related inflammatory disorders such as type II diabetes or osteoarthritis is also unknown.

- More information is required to understand the interrelationship between tissue resident macrophages and SC. How SC or their secretome influence immunosenescence and/or
cellular senescence in macrophages \textit{in vivo}, and whether senescent macrophages can also impact the development of senescence in non-immune cells remains to be explored.

- Immunometabolism of senescent macrophages is also poorly studied. Whether the longevity pathways play a similar role in macrophage senescence, and if these pathways could be therapeutic targets regulating macrophage functions during aging and disease remains to be elucidated.

- Much information about the mechanisms and ligands of macrophage mediated immunosurveillance is not known. It needs to be determined whether senescent macrophages preserve their capacity of immunosurveillance which can directly impact tissue SC accumulation.

- Finally, nutraceuticals such as probiotics and polyphenols, as well as micronutrients are known to affect immunosenescence (Sharma et al., 2013; Wu et al., 2019), but whether the underlying mechanisms are relatable to cellular senescence is little understood (Sharma and Padwad, 2020) (Figure 3). In this regard, our previous work showed that green tea catechin EGCG can strongly attenuate the development of senescence-like features such as expression of p53/p21^{WAF1}/p16^{Ink4a}, and SA-β-gal activity in macrophages exposed to conditioned media of senescent preadipocytes (Kumar et al., 2020). This suggests that nutraceuticals can impact senescence induction in macrophages, although further elaborative studies are warranted.

![Figure 3: Multifaceted effects of nutraceuticals in modulating different aspects of cellular senescence and immunosenescence.](image-url)
7. Concluding remarks

Aging, cellular senescence and immunosenescence appear to be intimately linked (Sharma and Padwad, 2020). As most diverse and plastic immune cells, delineating the dichotomy of cellular senescence and immunosenescence in macrophages is necessary for comprehending the relationship between aging and immune functions. Evidence that cellular senescence is an active force driving the aging of macrophage and which may contribute to age-related pathologies is already emerging. In addition, senescent non-immune cells can also impair macrophage immune functions through paracrine effects and augment SC burden suggesting an intricate relationship between host tissues and macrophages senescence which needs further characterization. In short, it is imperative that future studies consider macrophages in the purview of cellular senescence which may ultimately provide an integrative and deeper understanding of organismal aging and the pathology of age-dependent diseases.

Funding sources

This work was supported by grant from the Department of Science & Technology, Government of India under the INSPIRE Faculty scheme (grant no. IFA17-LSPA79).

Declarations of interest: none

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