MicroRNA-34a: a new player in arterial inflammaging

Ileana Badi, Angela Raucci

Experimental Cardio-Oncology and Cardiovascular Aging Unit, Centro Cardiologico Monzino-IRCCS, Milan, Italy

Correspondence: Angela Raucci
E-mail: araucci@ccfm.it
Received: March 26, 2015
Published online: April 13, 2015

Arterial inflammaging highly contributes to cardiovascular morbidity and mortality. As vascular cells age they become senescent and sustain a chronic low grade sterile inflammation by acquiring a senescence-associated secretory phenotype (SASP). The molecular mechanisms leading to the phenotypic changes affecting endothelial cells (ECs) and vascular smooth muscle cells (VSMCs) are also relevant for the pathogenesis of vascular diseases, such as atherosclerosis and hypertension. Therefore, unravelling the etiology of vascular inflammaging becomes of crucial importance. MicroRNAs (miRNAs) are small non-coding negative post-transcriptional regulator that are emerging as promising drug targets. MicroRNA-34a (miR-34a) had been implicated in tissues aging and endothelial and endothelial progenitor cells senescence. Our recent work showed that this miRNA is upregulated in aged mouse aortas as well as in senescent VSMCs. Conversely, its target SIRT1 is downregulated in the same specimens. We also found that miR-34a can inhibit VSMCs proliferation and induce VSMCs senescence, the latter by the direct regulation of SIRT1. Notably, for the first time, we demonstrated that miR-34a is also able to modulate the SASP by inducing the transcriptional expression of a subset of pro-inflammatory factors in a SIRT1-independent manner. These data support a model in which the age-dependent upregulation of miR-34a, by affecting senescence and inflammation of vascular cells, could play a causal role to arterial dysfunctions. Hence, further studies are necessary to unravel miR-34a-dependent mechanisms leading to arterial inflammaging in order to develop an effective strategy for age-related cardiovascular complications.

Keywords: miR-34a; Inflammaging; SIRT1; SASP; Vascular Aging

pathogenesis of vascular diseases such as atherosclerosis and hypertension. Therefore, elucidating the cellular and molecular processes underlying vascular inflammation and related dysfunctions is of high scientific interest.

MicroRNAs (miRNAs) are small non-coding RNA molecules that usually negatively regulate the post-transcriptional expression levels of multiple target genes and they are emerging as a promising class of drug targets. MicroRNA-34a (miRNA-34a) was firstly described as a tumor suppressor miRNA, direct target of p53 that, when up-regulated, can induce apoptosis, cell cycle arrest and senescence in several cancers. Recently miRNA-34a expression has been shown to increase with age in different tissues and organs of humans and mice. Notably, miR-34a levels increase with aging in human peripheral blood and murine plasma and peripheral blood mononuclear cells (PBMCs); furthermore, Schipper and colleagues found augmented miR-34a levels in PBMCs of patients with the inflamming-associate Alzheimer’s disease compared to normal elderly controls. Concerning vascular cells, Ito and collaborators demonstrated that miR-34a can induce endothelial cells (ECs) senescence at least in part through the modulation of the longevity-associated SIRT1 gene and Zhao and co-workers observed increased senescence and decreased SIRT1 protein levels upon miR-34a overexpression in endothelial progenitor cells. However, the miR-34a role in whole aorta and specifically VSMCs aging and associated inflammation has been unexplored so far.

We, therefore, undertook a study aimed to understand whether miR-34a could affect VSMCs senescence and their SASP. Firstly, we demonstrated that miR-34a expression levels were increased in aged aortas of old mice compared to young animals while the contractile smooth muscle cells marker (SM22a) appeared downregulated. Interestingly, among all the assessed miR-34a targets (SIRT1, Bcl2 and Axl) only SIRT1 protein levels were significantly downregulated during aging. We also observed that SIRT1 was expressed in both ECs and VSMCs in young aortas, while it was barely detectable in both cell types in the aged vessels. In accordance to the in vivo data, miR-34a expression increased whereas SIRT1 protein levels decreased during replicative senescence of cultured VSMCs. Moreover, overexpression of miR-34a in proliferative VSMCs caused at first, G0/G1 cell-cycle arrest along with p21 upregulation, followed by SIRT1 downregulation and senescence. While miR-34a-induced inhibition of VSMCs proliferation was SIRT1-independent, induction of cell senescence resulted, at least in part, directly regulated by this sirtuin. These results suggest that miR-34a could be an important player in vascular aging and its regulation of senescence as well as SIRT1 expression appear cell type-independent. However, since a considerable number of genes involved in cellular growth and senescence are direct or indirect targets of miR-34a, it is unlikely that this miRNA can affect vascular aging only through SIRT1. Further studies are needed to shed more light on this aspect.

Further, we demonstrated that miR-34a is not only a senescence-associated miRNA, but it may influence inflamming also by modulating other processes than senescence, such as inflammation, and in particular VSMCs SASP. Indeed, for the first time, we found that miR-34a could promote the expression of a subset of SASP molecules, specifically, the pro-inflammatory factors, IL1β, IL6, IL8, BMP2, and MCP1, and the soluble adhesion molecule ICAM1. In contrast, no effect was observed on growth factors and their regulators IGFBP4/6, GRO/α, and VEGF, and osteoprotegerin RNA levels. Accordingly, expression and secretion of pro-inflammatory SASP molecules have been found increased in VSMCs from aged animals as well as in replicative senescent VSMCs where the miR-34a is up-regulated. Furthermore, a very recent work from Fan and colleagues supports our findings leading to a pro-inflammatory role for miR-34a in vascular cells; indeed, they showed that this miRNA is involved in the flow-dependent regulation of endothelial inflammation by affecting VCAM1 and ICAM1 expression in ECs and consequently monocyte adhesion.

The precise regulatory mechanisms responsible for SASP acquisition is currently unknown. In our in vitro experiments, the miR-34a-mediated transcriptional activation of pro-inflammatory SASP factors was not prevented by SIRT1 ectopic overexpression. Interestingly, very recently, Hayakawa et al. demonstrated that in x-radiation-induced senescent human fibroblasts, depletion of SIRT1 enhanced the transcriptional levels of SASP components (IL6 and IL8) through epigenetic gene regulation. The opposite was not observed in SIRT1 overexpression conditions. Yet, although, the SIRT1 activator resveratrol has been shown to reduce the secretion of IL1β, MCP1 and IL6 by VSMCs, recently, Bollmann and colleagues demonstrated that resveratrol-dependent inhibition of cytokines expression occurs through the reduction of mRNA stability by the KH-type splicing regulatory protein (KSRP) activity and, is indeed, SIRT1-independent. Moreover, other factors rather than SIRT1 have been implicating in SASP regulation. Csiszar and co-workers, for instance, showed that the secretion of IL1β, MCP1, TNFα and IL6 in aged VSMCs correlates with increased NF-κB activation. Fan and colleagues showed that miR-34a affects ECs inflammation namely, VCAM1 and ICAM1 expression, through enhanced acetylation of the RelA/p65 subunit of NF-κB and thereby
activation of NF-κB signaling \[^{21}\]. They also observed that miR-34a-induced expression of these cell adhesion molecules is partially prevented by SIRT1 \[^{21}\]. Since SIRT1 can regulate the acetylation of RelA/p65 subunit and thus decrease NF-κB activation \[^{25}\], we speculate that in ECs miR-34a could influence NF-κB signaling at least partially via SIRT1. Hence, unrevealing the exact role of SIRT1 in SASP and whether miR-34a induces SASP factors expression in VSMCs by activating NF-κB or other pathways calls for further investigations.

Taken together, all these experimental evidences indicate that miR-34a could be an important player in arterial inflammaging, since an age-dependent increase of its levels in vascular cells could enhance senescence and inflammation and thus cause arterial dysfunctions such as atherosclerosis. Accordingly, miR-34a expression has found augmented in human atherosclerotic lesions and in the aortas of an animal model of atherosclerosis \[^{26, 27}\].

Nonetheless, more studies are needed to deepen our knowledge on miR-34a involvement in arterial inflammaging, as this could be of extreme importance in order to develop an effective strategy beneficial for old people.

**Conflicting interests**

The authors declare that they have no conflicting interests.

**Acknowledgements**

We thank our funding agencies: Fondazione Monzino and Centro Cardiologico Monzino-IRCCS (RC2012-2015). I.B. is supported by Fondazione Umberto Veronesi (Post-doctoral Fellowship - 2015) and A.R. by Young Investigator Grant, Ministry of Health (GR-2010-2312693).

**References**


