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MicroRNAs
From Basic Mechanisms to Clinical Application in Cardiovascular Medicine

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MicroRNAs (miRs) are small noncoding RNAs (≈23 nucleotides) that regulate gene expression at a post-transcriptional level by degradation or translational inhibition of target mRNAs. Initially discovered as regulators of development in plants, worms, and fruitflies, miRs are emerging as pivotal modulators of cardiovascular biology and disease. Thus, the cell types, namely endothelial cells, smooth muscle cells, and macrophages. A review by Schober et al consequently surveys the differential roles of distinct miRs during the pathogenesis of atherosclerosis, which encompass downregulation of miR-145 controlling smooth muscle cell differentiation, delivery of miR-126 in endothelial cell-derived microparticles to signal the need for endothelial repair, or an upregulation of miR-155 relevant in proinflammatory macrophage polarization. The identification of this miR triad sheds light on the current concepts of atherogenesis and establishes novel treatment options. An important contribution of miRs to the regulation or alteration of lipid metabolism and glucose homeostasis may determine the predisposition to cardiometabolic disease and...
atherosclerosis. For instance, miR-33 controls cellular cholesterol export and fatty acid degradation, which are stimulated by its host genes, whereas miR-122 can limit cholesterol synthesis and lipoprotein secretion in the liver. Underlying mechanisms are detailed by Fernandez–Hernando et al.\textsuperscript{7} in a review, which also discusses the modulation of miRs as a strategy to treat metabolic diseases.

Membrane-derived vesicles and lipoproteins can serve to carry and transport miRs between cells with distinct signatures altered by cardiovascular pathologies. A review by Boon and Vickers\textsuperscript{8} is focused on the complexities and multitude of open questions remaining about the roles of miRs intercellular communication and their applicability to cardiovascular disease. A review by Fish et al.\textsuperscript{9} highlights how miRs regulate multiple aspects and functions of the vascular endothelial growth factor signaling pathway in vasculogenesis and angiogenesis, in particular providing insights into the role of miRs and downstream effectors in modulating vascular endothelial growth factor output during development. This knowledge may be relevant to therapeutic targeting of pathological blood vessel growth and identifying clinical applications for miR manipulation. Myocardial infarction as a common and severe manifestation of advanced atherosclerosis is characterized by altered gene expression and dysregulation of underlying signaling pathways, which may involve an induction or repression of miRs affecting cell-specific downstream effects on cardiac function. A review by Fiedler and Thum\textsuperscript{10} summarizes the current knowledge about the mechanistic importance of several miRs and novel miR-based therapeutic approaches to counteract maladaptive remodeling after myocardial infarction. Postgenomic technologies enable an interrogation of complex pathophysiological perturbations of gene regulatory networks by introducing a systems context to biomarker discovery. As cardiovascular diseases develop over decades, a multi-biomarker panel including miRs may be required to detect and monitor different stages of disease, example, vulnerable atherosclerotic plaques. A review by Mayr et al.\textsuperscript{11} discusses strategies for biomarker discovery using postgenomic technologies with a particular focus on circulating miRs. This may eventually serve and reveal distinctive cardiovascular phenotypes and identify biomarker signatures that complement the traditional scores in clinical risk prediction.

A key regulatory event in miR pathways is Ago loading, when Ago proteins undergo conformational opening to bind double-stranded small RNA duplexes forming a pre-RISC, and subsequently mediate unwinding to finally leave a single strand stably incorporated in a mature RISC. The N domain of Ago2 has been required for active wedging and unwinding of the duplex during RISC assembly but not for precedent duplex loading or subsequent target cleavage.\textsuperscript{12} Moreover, the evolutionarily conserved PAZ domain is dispensable for Ago loading of slicing-competent RISC but in the absence of slicer activity or slicer-substrate, duplex miRs is required for ejecting the passenger strand and forming functional RISC complexes.\textsuperscript{13} Although it was thought that, depending on the thermodynamic stability of the strands in a precursor miR, cells preferentially select the less stable one (guide strand) and degrade the other (passenger strand), expression profiling analyses indicate that both strands can accumulate as miR pairs in some tissues but are subject to strand selection in others.\textsuperscript{1} Furthermore, target prediction and validation assays demonstrated that both strands of a miR pair can target equal numbers of genes to suppress their expression. Because the number of context- and tissue-specific targets may therefore vary, likely being higher and more versatile than expected, it will be important to determine the mechanisms involved in tissue-dependent miR biogenesis, expression, and degradation, as well as those contributing to context-dependent strand and target selection. It may be envisioned that a specific regulation of the N and PAZ domains of Ago within the RISC may give rise to such selectivity, depending on the duplex pair and the target cell. Given the abundant involvement of miRs in posttranscriptional regulation that has been revealed in various models, the elucidation of the tissue-specific strand selectivity seems to be of particular relevance for future studies in the cardiovascular system, both to unravel the basic mechanisms and to improve the options for therapeutic applicability in clinical cardiovascular medicine.

References


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