

# Chronic obstructive pulmonary disease: MicroRNAs and exosomes as new diagnostic and therapeutic biomarkers

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Chronic obstructive pulmonary disease (COPD) is known as a progressive lung disease and the fourth leading cause of death worldwide. Despite valuable efforts, there is still no accurate diagnostic and prognostic tool for COPD. Hence, it seems that finding new biomarkers could contribute to provide better therapeutic platforms for COPD patients. Among various biomarkers, microRNAs (miRNAs) have emerged as new biomarkers for the prognosis and diagnosis of patients with COPD. It has been shown that deregulation of miRNAs targeting a variety of cellular and molecular pathways such as Notch, Wnt, hypoxia-inducible factor-1 $\alpha$ , transforming growth factor, Kras, and Smad could be involved in COPD pathogenesis. Multiple lines of evidence have indicated that extracellular vesicles such as exosomes could carry a variety of cargos (i.e., mRNAs, miRNAs, and proteins) which transfer various cellular and molecular signals to recipient cells. Here, we summarized various miRNAs which could be applied as diagnostic and prognostic biomarkers in the treatment of patients with COPD. Moreover, we highlighted the role of extracellular vesicles containing miRNAs as diagnostic and prognostic biomarkers in COPD patients.

**Key words:** Biomarker, chronic obstructive pulmonary disease, diagnosis, exosome, microRNA, therapy

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## INTRODUCTION

Chronic obstructive pulmonary disease (COPD) is a disease due to chronic inflammation of the airways. This disease is accompanied by the presence of chronic obstruction.<sup>[1]</sup> Multiple lines of evidence have indicated that a wide range of factors (i.e., genetical and environmental) are involved in the initiation and progression of COPD. Smoking is known as one of the main factors which could be associated with chronic inflammation of the airways.<sup>[1]</sup> Inflammation is an important factor which has critical role in the initiation and progression of COPD.<sup>[2]</sup> It has been shown that various immune cells and a variety of cellular and molecular pathways are involved

in inflammation and play critical roles in COPD pathogenesis.<sup>[3,4]</sup> Among many cellular and molecular targets, microRNAs (miRNAs) have emerged to be involved in COPD pathogenesis.<sup>[5,6]</sup> Deregulation of miRNAs is associated with the initiation and progression of several diseases such as stroke, cardiovascular diseases, diabetes, cancer, and inflammatory diseases.<sup>[7-14]</sup> Many miRNAs including miR-223, miR-1274a, miR-101, and miR-144 exert their effects via inhibition/activation of a sequence of cellular and molecular pathways (i.e., Smad, transforming growth factor  $\beta$  [TGF- $\beta$ ], Kras, Notch, and Wnt) involved in COPD.<sup>[5,6]</sup> Hence, it seems that these molecules could be used as new diagnostic and therapeutic biomarkers for patients with COPD. Alterations of miRNA expression could occur due to various events such as mutations and

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inhibition/activation of various targets.<sup>[15-17]</sup> Exosomes are extracellular vesicles which could change the expression of miRNAs in recipient cells. These nanocarriers could transfer various signals via cargos such as messenger RNAs (mRNAs), miRNAs, and proteins to recipient cells.<sup>[18,19]</sup> It has been shown that several exosomal miRNAs such as miR-100, miR-21, and miR-181a could be used as diagnostic and prognostic biomarkers in COPD.<sup>[19-21]</sup> In the current review, we focus on various miRNAs that could be utilized as diagnostic and prognostic biomarkers for COPD patients. Moreover, we highlighted the role of exosomes containing various cargos, especially miRNAs as diagnostic and therapeutic biomarkers for patients with COPD.

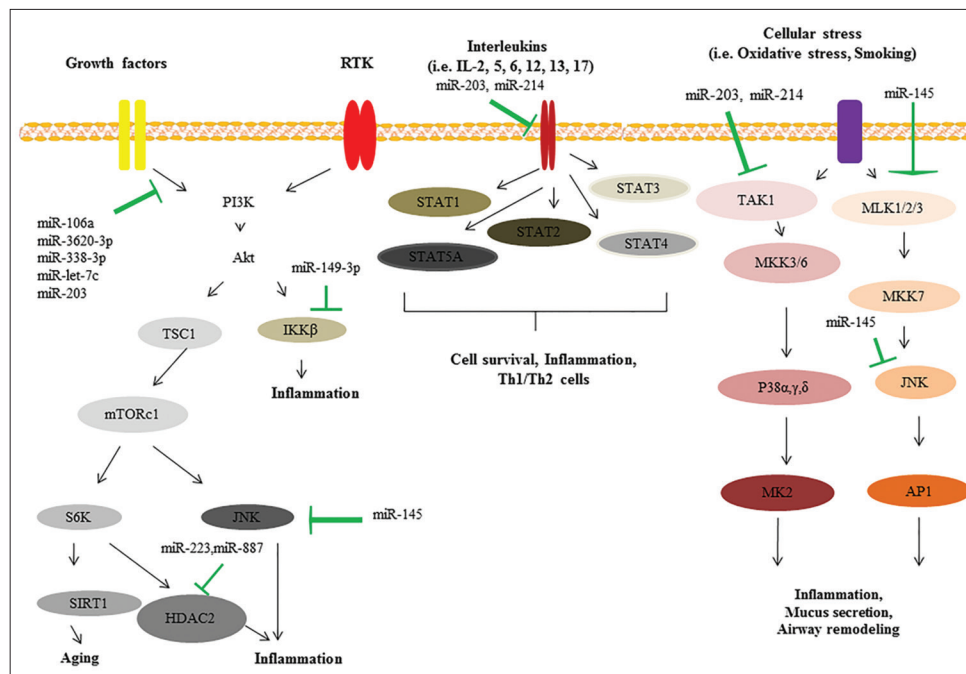
## MICRO-RNA AS DIAGNOSTIC AND THERAPEUTIC BIOMARKERS IN CHRONIC OBSTRUCTIVE PULMONARY DISEASE

COPD is a chronic heterogeneous disease of the lungs which could be characterized by persistent and excessive inflammation, alveolar lesions, accelerated decrease in lung function, and airflow limitation.<sup>[22]</sup> Many factors such as inflammation and cell structure changes could be involved in the initiation and progression of COPD. miRNAs have also been suggested to be involved in the COPD pathogenesis [Figure 1]. These molecules are known as epigenetic regulators which exert their effects via targeting a variety of cellular and molecular pathways.<sup>[19,23-29]</sup> It has been shown that deregulation of miRNAs could be associated with the initiation and progression of several diseases such as cardiovascular diseases, stroke, diabetes, cancer, and COPD.<sup>[5,30-35]</sup> Multiple lines of evidence have revealed

that up/down-regulation of various miRNAs (i.e., miR-21, miR-145, miR-181, miR-1, miR-144, and miR-101) could be related to COPD pathogenesis.<sup>[5,6]</sup> It has been showed that a variety of external factors (i.e., smoking and oxidative stress) and internal factors (i.e., deregulation of various growth factor ligands, interleukins [ILs], and receptor tyrosine kinase) could lead to deregulation of many cellular and molecular pathways and miRNA expression.<sup>[36-38]</sup> These events are associated with small and large alterations in molecular and cellular levels and could contribute to the progression of COPD.<sup>[36]</sup>

MiRNAs are involved in various signaling pathways such as TGF- $\beta$  signaling pathway. Several studies have confirmed that TGF- $\beta$  pathways play critical roles in COPD pathogenesis.<sup>[39,40]</sup> Baraldo *et al.* confirmed that downregulation of TGF- $\beta$  type II receptor (TGF- $\beta$ 2) is associated with the progression of COPD.<sup>[41]</sup> Their results indicated that the expression profile of TGF- $\beta$ 1 in bronchial glands was similar in the two groups of cases while there was a decreased in expression of TGF- $\beta$ 2 in smokers with COPD than in smokers with normal lung function. Expression of TGF- $\beta$ 2 was associated with the values of Reid's index which is known as a measure of gland size. These findings suggested that TGF- $\beta$  signaling pathway has key roles in COPD pathogenesis.<sup>[41]</sup>

Ezzie *et al.* assessed gene expression profiles of many miRNAs and mRNAs in 26 patients with COPD.<sup>[5]</sup> They showed that miR-146a, miR-15b, miR-223, and miR-1274a were upregulated in COPD samples. Moreover, they showed that a variety of genes (e.g. BMP5 and BMP6,



**Figure 1:** Various cellular and molecular pathways involved in chronic obstructive pulmonary disease

TGF- $\beta$ 1 and TGF- $\beta$ 2, and SMAD7) involved in TGF- $\beta$  signaling pathway were downregulated in patients with COPD. Bioinformatics analysis indicated that various miRNAs could target these genes in COPD patients. For example, upregulation of miR-15b in a bronchial epithelial cell line could be associated with downregulation of SMAD7, SMURF2, and downstream decorin protein. These results proposed that many miRNAs are involved in TGF- $\beta$  signaling pathway via targeting various genes and thus could contribute to the initiation and progression of COPD.<sup>[5]</sup>

O'Leary *et al.* indicated that miR-145 targets SMAD3 which is known as one of the important downstream signaling molecules in the TGF- $\beta$  pathway and plays critical roles in the initiation and progression of COPD.<sup>[42]</sup> They showed that TGF- $\beta$  could induce the expression of SMAD3 via increasing miR-145 in COPD patients. It has been shown that of miR-145 could be regulated by the MAP kinases, MEK-1/2, and p38 MAPK pathways. Upregulation of miR-145 in patients with COPD could inhibit the release of IL-6 and CXCL8. These findings suggest that miR-145 could regulate pro-inflammatory cytokine release from airway smooth muscle cells in COPD via targeting SMAD3.<sup>[42]</sup>

Kusko *et al.* assessed several genes and pathways involved in idiopathic pulmonary fibrosis (IPF) and COPD.<sup>[43]</sup> They showed that various members of the signaling pathways including hypoxia-inducible factor-1 $\alpha$ , MDM2, and NFKBIB are involved in COPD and IPF pathogenesis. Moreover, they revealed that alternative splicing of p53/hypoxia pathway-related molecules NUMB and PDGFA occurred more frequently in IPF or COPD compared with the control group. RNA-seq analysis indicated that among many miRNAs, miR-96 acts as a major regulatory hub in the p53/hypoxia gene-expression network. The regulation of miR-96 *in vitro* recapitulates disease-related gene-expression network.<sup>[43]</sup> These findings suggest that a variety of signaling pathways that regulate miRNAs network are involved in COPD pathogenesis and miRNAs could be utilized as potential candidates for the diagnosis and prognosis of COPD.

Accumulating evidence indicates that smoking is one of the major risk factors for COPD which could be associated with the progression of COPD in various stages.<sup>[44]</sup> Numerous studies have confirmed that smoking can down/up-regulate many miRNAs and could thus affect COPD progression.

Du *et al.* assessed the expression levels of miR-181c in 34 patients with COPD (smoking cases) compared with healthy controls.<sup>[45]</sup> Their results indicated that miR-181c could be significantly downregulated in patients with COPD versus healthy controls who had never smoked. They also

showed that upregulation of miR-181c could be associated with several effects such as reduction of the inflammatory response, neutrophil infiltration, reactive oxygen species generation, and inflammatory cytokines production. Downregulation of miR-181c could be associated with opposite effects. Moreover, they revealed that miR-181c exerts its effects via targeting CCN1. Downregulation of miR-181c could lead to an increase in CCN1 expression in the lung tissues of COPD patients compared with healthy controls. These findings suggested that miR-181c could be used as a therapeutic target for the treatment of patients with COPD.<sup>[45]</sup>

Multiple lines of evidence have indicated that DNA damage pathways are main players for aging disorders such as COPD.<sup>[46]</sup> This damage could be due to many factors such as oxidative stress and activates ataxia telangiectasia-mutated (ATM) kinase which has critical roles in the DNA damage response.<sup>[47]</sup> It has been shown that the increasing of DNA in the lung biopsies from smokers and COPD patients may accelerate lung aging and pathogenesis of COPD.<sup>[48,49]</sup>

Paschalaki *et al.* indicated that deregulation of miR-126 is associated with activation of ATM kinase.<sup>[50]</sup> They showed that the levels of miR-126 were downregulated in blood outgrowth endothelial cells from smokers and COPD patients compared with nonsmoker subjects. These results suggested that downregulation of miR-126 via targeting ATM could promote tissue aging and dysfunction in smoker and COPD subjects. Hence, this miRNA could be utilized as a novel therapeutic target in patients with COPD.<sup>[50]</sup>

Besides the role as therapeutic targets, miRNAs could be employed as prognostic and diagnostic biomarkers in various diseases such as COPD.<sup>[6]</sup> COPD is a multifactorial disease, and various efforts to find effective diagnostic and therapeutic platforms for treatment of patients with COPD have been made. It has been shown that the levels of various cytokines (i.e., IL-6, IL-8, IL-10, IL-17, IL-12p70, and IL-1 $\beta$ ), cysteinyl-leukotrienes (LTs), LTB4, prostaglandin E(4), hydrogen peroxide (H(2)O(2)), and 8-isoprostane could be associated with COPD pathogenesis.<sup>[51]</sup> Despite many efforts, effective diagnostic biomarkers are still rare. Previous studies confirmed that miRNAs play critical roles in COPD pathogenesis. Hence, it seems that these molecules may be utilized as new prognostic and diagnostic biomarkers for monitoring patients with COPD.<sup>[6]</sup>

Soeda *et al.* investigated the expression levels of various circulating miRNAs in the plasma of 40 COPD patients.<sup>[52]</sup> They indicated that deregulation of many miRNAs could be associated with progression of COPD. Their results revealed that a variety of miRNAs including miR-29b, miR-483-5p,

miR-152, miR-629, miR-26b, miR-101, miR-106b, miR-532-5p, and miR-133b were significantly downregulated in the plasma from COPD patients compared with control group. Moreover, they showed that there was negative association between the levels of miR-106b and duration of disease since the diagnosis in COPD ex-smokers and duration of smoking in COPD current smokers. These findings suggested that downregulation of miR-106b could reflect persistent and systemic alteration even after the discontinuation of smoking in patients with COPD. Hence, miRNAs may be utilized as diagnostic biomarkers for COPD patients.<sup>[52]</sup>

Kara *et al.* assessed the expression levels of several miRNAs including miR-16, miR-17, miR-29c, miR-92, miR-125, miR-126, miR-146, miR-155, miR-181, and miR-122 using real-time polymerase chain reaction in 60 patients with COPD.<sup>[6]</sup> Their results indicated that miR-29c and miR-126 were upregulated in COPD patients Stage III compared with healthy controls. These findings propose that these miRNAs may be utilized as diagnostic biomarkers for patients with COPD.<sup>[6]</sup> Table 1 lists various miRNAs involved in COPD pathogenesis which could be used as diagnostic and therapeutic biomarkers for the treatment of patients with COPD.

MiR-218-5p is another important miRNA which has a main role in COPD pathogenesis and it seems that it could be used

as an effective candidate for the diagnosis and monitoring of patients with COPD.<sup>[53]</sup> Conickx *et al.* indicated that various miRNAs such as miR-218-5p were deregulated in COPD patients.<sup>[53]</sup> They showed that the expression of miR-218-5p could be decreased in smokers without airflow limitation and in COPD patients compared with never-smokers. These results suggest that miR-218-5p has a protective role in cigarette smoke (CS)-induced inflammation and COPD patients and could be used as diagnostic and therapeutic biomarker for the detection and monitoring of patients with COPD.<sup>[53]</sup>

## EXOSOME AND CHRONIC OBSTRUCTIVE PULMONARY DISEASE

Many cells could change their behavior via receiving a variety of cellular and molecular signals. Extracellular vesicles (i.e., exosomes, apoptotic bodies, and microvesicles) are able to carry several signals to recipient cells.<sup>[12,92]</sup> Exosomes are known as one of the important nanovehicles which have critical roles in various physiological events. These vesicles could carry many cargos such as DNAs, mRNAs, miRNAs, and proteins.<sup>[18,19]</sup> Accumulating evidence has indicated that these extracellular vesicles could be employed as diagnostic and therapeutic biomarkers in the treatment of various diseases such as cancer and COPD [Table 2].<sup>[19-21]</sup>

**Table 1: Several microRNAs involved in chronic obstructive pulmonary disease pathogenesis**

miRNA	Expression in COPD	Detection method	Target gene	Model ( <i>in vivo</i> /human)	Sample (n)	Reference
miR-146a	Downregulation	qRT-PCR	COX-2	Human	14	[54]
miR-1	Downregulation		IGF-1, IGF-1R	Human	31	[55]
let-7c	Downregulation		TNFR-II	Human	32	[56]
miR-125b	Downregulation			Human	32	[56]
miR-30e-3p	Downregulation			Human	12	[56]
miR-101	Upregulation		CFTR	<i>In vivo</i> , human	16	[57]
miR-34a	Upregulation		HIF-1 $\alpha$	<i>In vivo</i> , human	55	[58]
miR-34a	Upregulation		Sirtuin-1	Human, cell line	23	[59]
miR-199a-5p	Upregulation		HIF-1 $\alpha$	<i>In vivo</i> , human	55	[58]
miR-328	Downregulation			Human	10	[60]
miR-21	Downregulation			Human	10	[60]
miR-106b	Downregulation		TGF- $\beta$ R	Human	40	[52]
miR-29b	Downregulation			Human	40	[52]
miR-483-5p	Downregulation			Human	40	[52]
miR-152	Downregulation			Human	40	[52]
miR-629	Downregulation			Human	40	[52]
miR-26b	Downregulation			Human	40	[52]
miR-532-5p	Downregulation			Human	40	[52]
miR-133b	Downregulation			Human	40	[52]
miR-106b	Downregulation			Human	40	[52]
miR-499	Upregulation			Human	103	[61]
miR-206	Upregulation			Human	103	[61]
miR-133	Upregulation			Human	103	[61]
miR-223	Upregulation			Human	26	[5]

*Contd...*

Table 1: Contd....

miRNA	Expression in COPD	Detection method	Target gene	Model ( <i>in vivo</i> /human)	Sample (n)	Reference
miR-223	Upregulation		HDAC2	<i>In vivo</i>	-	[62]
miR-1274	Upregulation			Human	26	[5]
miR-923	Downregulation			Human	26	[5]
miR-937	Downregulation			Human	26	[5]
miR-125b-1	Downregulation			Human	26	[5]
miR-101	Upregulation		CFTR	<i>In vivo</i> , human	16	[57]
miR-144	Upregulation		CFTR	<i>In vivo</i> , human	16	[57]
miR-452	Downregulation		MMP 12	<i>In vivo</i>	-	[63]
miR-20a	Downregulation			Human	20	[64,65]
miR-28-3p	Downregulation			Human	20	[64,65]
miR-34c-5p	Downregulation			Human	20	[64,65]
miR-100	Downregulation			Human	20	[64,65]
miR-7	Upregulation			Human	20	[64,65]
miR-145-5p	Downregulation		IFI30	Human	25	[66]
miR-338-3p	Downregulation		LTK, TNFR	Human	25	[66]
miR-3620-3p	Upregulation		TNFR	Human	25	[66]
miR-132-212 cluster	Upregulation		$\alpha_1$ -antitrypsin	Human	10	[67]
miR-342-5p	Upregulation			Human	10	[67]
miR-422a	Upregulation			Human	10	[67]
miR-423-5p	Upregulation			Human	10	[67]
miR-425	Upregulation			Human	10	[67]
miR-486-3p	Upregulation			Human	10	[67]
miR-98	Downregulation			Human	10	[67]
miR-193a-5p	Downregulation			Human	10	[67]
miR-324-5p	Downregulation			Human	10	[67]
miR-342-3p	Downregulation			Human	10	[67]
miR-342-5p	Upregulation			Human	10	[67]
miR-365	Downregulation			Human	10	[67]
miR-133	Downregulation			Human	17	[68]
miR-206	Downregulation			Human	17	[68]
miR-1 <sup>4</sup>	Downregulation			Human	17	[68]
miR-21	Upregulation			Human, <i>in vivo</i>	49	[69]
miR-181a	Downregulation			Human, <i>in vivo</i>	49	[69]
miR-7	Upregulation		Epac1	Cell line	-	[70]
miR-34c	Downregulation		SERPINE1	Human, cell line	29	[71]
miR-let7c	Downregulation		TNFR2	-	-	[72]
miR-183	Upregulation		BKCa $\beta$ 1	Human, cell line	45	[73]
miR-200b	Upregulation			Human, cell line	45	[73]
miR-200c	Upregulation			Human, cell line	45	[73]
miR-15b	Upregulation		SMAD7	Human	26	[5]
miR-218	Downregulation		CCR6	Human, <i>in vivo</i>	12	[74]
miR-145	Upregulation		SMAD3	Cell line	-	[42]
miR-203	Upregulation		TAK1, PI3KCA	Human, <i>in vivo</i>	17	[75]
Let-7g	Downregulation		KRAS	Human, <i>in vivo</i>	554	[76]
miRNA-181a-2-3p	Downregulation			Human	58	[77]
miR-181d-5p	Upregulation			Human	58	[77]
miR-501-3p	Upregulation			Human	58	[77]
miR-769-5p	Upregulation			Human	58	[77]
miR-191-5p	Upregulation			Human	58	[77]
miR-125a/b	Upregulation		MAVS	Human	10	[78]
miR-125a	Downregulation			Human	-	[79]
miR-130a	Downregulation			Human	-	[79]
miR-301a	Downregulation			Human	-	[79]
miR-424-5p	Downregulation			Human	-	[79]

Contd...

Table 1: Contd....

miRNA	Expression in COPD	Detection method	Target gene	Model ( <i>in vivo</i> /human)	Sample (n)	Reference
miR-887	Downregulation		HDAC6	Cell line	-	[80]
miR-126	Upregulation			Human	60	[6]
miR-29c	Upregulation			Human	60	[6]
miR-210	Downregulation			Human	50	[81]
miR-218-5p	Downregulation			Human	30	[82]
miR-130a	Upregulation		ATG 16L	Human, cell line	43	[83]
miR-146b-5p	Upregulation			Human	18	[84]
miR-141-3p	Upregulation			Human	18	[84]
miR-186-5p	Upregulation			Human	18	[84]
miR-4532	Upregulation			Human	18	[84]
miR-4635	Upregulation			Human	18	[84]
miR-2681-3p	Upregulation			Human	18	[84]
miR-4503	Upregulation			Human	18	[84]
miR-196b-5p	Downregulation			Human	18	[84]
miR-149-3p	Downregulation		TLR-4/NF-κB	Human, cell line	9	[85]
miR-214	Upregulation		CCNL2	Human, cell line	18	[86]
miR-3177-3p	Downregulation			Human	141	[87]
miR-106b-5p	Downregulation			Human	141	[87]
miR-125a-5p	Upregulation			Human	141	[87]
miR-183-5p	Downregulation			Human	141	[87]
miR-218-5p	Downregulation		CLK3, CYP1B1, LIF and DUSP5	Human, <i>in vivo</i>	12	[53]
miR-21	Upregulation			Human	49	[88]
miR-518	Upregulation			Human	24	[89]
miR-1233	Upregulation			Human	24	[89]
miR-106a	Upregulation			Human	24	[89]
miR-20b	Upregulation			Human	24	[89]
miR-1291	Upregulation			Human	24	[89]
miR-377	Upregulation			Human	24	[89]
miR-124	Upregulation			Human	24	[89]
miR-216b	Upregulation			Human	24	[89]
miR-646	Upregulation			Human	24	[89]
miR-548o	Upregulation			Human	24	[89]
miR-631	Upregulation			Human	24	[89]
miR-518b	Upregulation			Human	24	[89]
miR-216a	Upregulation			Human	24	[89]
miR-182	Upregulation			Human	24	[89]
miR-188-5p	Upregulation			Human	24	[89]
miR-492	Upregulation			Human	24	[89]
miR-662	Upregulation			Human	24	[89]
miR-200a	Upregulation			Human	24	[89]
miR-23b	Downregulation			Human	24	[89]
miR-888	Downregulation			Human	24	[89]
miR-641	Downregulation			Human	24	[89]
miR-184	Downregulation			Human	24	[89]
miR-95	Downregulation			Human	24	[89]
miR-224	Downregulation			Human	24	[89]
miR-369-5p	Downregulation			Human	24	[89]
miR-140-5p	Downregulation			Human	24	[89]
miR-613	Downregulation			Human	24	[89]
miR-531c	Downregulation			Human	24	[89]
miR-155-5p	Upregulation			Human	10	[90]
miR-20a-5p	Downregulation			Human	10	[90]
miR-126-3p	Downregulation			Human	10	[90]

Contd...

**Table 1: Contd....**

miRNA	Expression in COPD	Detection method	Target gene	Model ( <i>in vivo</i> /human)	Sample (n)	Reference
Let-7f-5p	Downregulation			Human	10	[90]
miR-1909	Upregulation			Human	15	[91]
miR-29a	Upregulation			Human	15	[91]
miR-632	Upregulation			Human	15	[91]
miR-720	Upregulation			Human	15	[91]
miR-605	Upregulation			Human	15	[91]
miR-99a	Upregulation			Human	15	[91]
miR-132	Upregulation			Human	15	[91]
miR-940	Upregulation			Human	15	[91]
miR-497	Upregulation			Human	15	[91]
miR-29c	Upregulation			Human	15	[91]
miR-424	Upregulation			Human	15	[91]
miR-222	Upregulation			Human	15	[91]
miR-22	Upregulation			Human	15	[91]
miR-142-5p	Upregulation			Human	15	[91]
miR-150	Upregulation			Human	15	[91]
miR-186	Downregulation			Human	15	[91]
miR-185	Downregulation			Human	15	[91]
miR-19b	Downregulation			Human	15	[91]
miR-584	Downregulation			Human	15	[91]
miR-652	Downregulation			Human	15	[91]
miR-191	Downregulation			Human	15	[91]
miR-151-3p	Downregulation			Human	15	[91]
Let-7i	Downregulation			Human	15	[91]
Let-7b	Downregulation			Human	15	[91]
Let-7d	Downregulation			Human	15	[91]
miR-339-3p	Downregulation			Human	15	[91]
miR-25	Downregulation			Human	15	[91]
miR-24	Downregulation			Human	15	[91]
miR-18b	Downregulation			Human	15	[91]
miR-30e	Downregulation			Human	15	[91]
miR-331-3p	Downregulation			Human	15	[91]
miR-320a	Downregulation			Human	15	[91]
miR-140-3p	Downregulation			Human	15	[91]
miR-1974	Downregulation			Human	15	[91]
miR-484	Downregulation			Human	15	[91]
miR-374b	Downregulation			Human	15	[91]
miR-486-3p	Downregulation			Human	15	[91]

COPD = Chronic obstructive pulmonary disease; COX-2 = Cyclooxygenase; IGF-1 = Insulin-like growth factor 1; IGF-1R = Insulin-like growth factor 1 receptor; TNFR II = Tumor necrosis factor receptor 2; CFTR = Cystic fibrosis transmembrane conductance regulator; HIF-1  $\alpha$  = Hypoxia-inducible factor 1-alpha; TGF- $\beta$ R = Transforming growth factor beta receptor; HDAC2 = Histone deacetylase 2; MMP12 = Matrix metalloproteinase 12; IFI30 = Gamma-interferon-inducible protein 30; LTK = Leukocyte tyrosine kinase; EPAS1 = Endothelial PAS domain protein 1; SERPIN1 = Serine protease inhibitor superfamily 1; BKCa $\beta$ 1 = Ca<sup>2+</sup>-activated K<sup>+</sup> channels  $\beta$ 1 subunit; CCR6 = CC chemokine receptor 6; TAK1 = Transforming growth factor beta-activated kinase 1; PI3KCA = Phosphatidylinositol 3-kinase; MAVS = Mitochondrial antiviral-signaling protein; ATG16L = Autophagy-related protein 16-1; TLR-4 = Toll-like receptor 4; NF- $\kappa$ B = Nuclear factor-kappaB; CCNL2 = Cyclin L2; CLK3 = CDC-like kinase 3; CYP1B1 = Cytochrome P450 Family 1 Subfamily B Member 1; DUSP5 = Dual specificity protein phosphatase 5; miRNA = MicroRNA

Several studies have indicated that extracellular vesicles such as exosomes are able to carry various cargos. Among several cargos, miRNAs are important and can transfer various cellular and molecular signals to recipient cells.<sup>[97]</sup> Hence, exosomal miRNAs could be used as new tools for diagnosis and treatment of various diseases such as COPD.

Fujita *et al.* investigated the effect of exosomal miRNAs in the suppression of autophagy in COPD patients.<sup>[98]</sup> It has been shown that CS exposure could lead to emphysema, increasing of myofibroblast, and airway remodeling,

which all contribute to COPD progression. They showed that CS could induce upregulation of exosomal miR-210 in bronchial epithelial cells. Exosomal miR-210 could induce myofibroblast differentiation in lung fibroblasts (LFs). Exosomal miR-210 could directly control autophagy processes via affecting ATG7. The exosomal miR-210 expression is associated with downregulation of ATG7 in LFs. They reported decreased autophagy in LFs from COPD patients, and also, the silencing of ATG7 in LFs could lead to myofibroblast differentiation. These results indicated that CS exposure induces the modification of exosome

**Table 2: Several cargos which carried by different microparticles such as exosomes**

Cargo	Expression in COPD	Detection method	Model	Sample (n)	Reference
Protein					
CD31	Upregulation	Flow cytometry	Human	92	[93]
CD62E/CD31	Upregulation	Flow cytometry	Human	49	[94]
CD144	Upregulation	Flow cytometry	Human	27	[95]
CD66	Upregulation	Flow cytometry	Human	18	[96]
CD235ab	Upregulation	Flow cytometry	Human	18	[96]
miRNA					
miR-1	Upregulation	qRT-PCR	Human	103	[61]
miR-499	Upregulation	qRT-PCR	Human	103	[61]
miR-133	Upregulation	qRT-PCR	Human	103	[61]
miR-206	Upregulation	qRT-PCR	Human	103	[61]
miR-21/ miR-181	Upregulation	qRT-PCR	Human	49	[69]
miR-20a	Downregulation	qRT-PCR	Human	20	[64]
miR-28-3p	Downregulation	qRT-PCR	Human	20	[64]
miR-34c-5p	Downregulation	qRT-PCR	Human	20	[64]
miR-100	Downregulation	qRT-PCR	Human	20	[64]
let-7c	Downregulation	qRT-PCR	Human	32	[56]
miR-125b	Downregulation	qRT-PCR	Human	32	[56]
let-7a	Downregulation	qRT-PCR	Human	10	[60]
miR-21	Downregulation	qRT-PCR	Human	10	[60]
miR-328	Downregulation	qRT-PCR	Human	10	[60]

miRNA = MicroRNA; qRT-PCR = Quantitative real-time polymerase chain reaction; COPD = Chronic obstructive pulmonary disease

components and exosomal miR-210 acts as a paracrine autophagy regulator of myofibroblast differentiation. This molecule could be employed as a therapeutic biomarker for patients with COPD.<sup>[97]</sup>

Skeletal muscle weakness is an important systemic complication of COPD which could affect exercise capacity and mortality. Burke *et al.* assessed the role of exosomal miRNAs in COPD patients who had skeletal muscle weakness.<sup>[98]</sup> They isolated exosomal miRNAs from serum and bronchoalveolar lavage fluid (BALF) of four patients with COPD. Their results indicated that one exosomal miRNA was upregulated in serum of COPD patients and four exosomal miRNAs were downregulated in BALF of patients with COPD. *In silico* analysis indicated that these miRNAs could exert their effects via targeting many genes including S6K involved in the mTORC1 signaling pathway which serves as a key regulator of skeletal muscle wasting. These results indicated that exosomal miRNAs play critical roles in skeletal muscle wasting in patients with COPD and could be utilized as diagnostic and prognostic biomarkers for the detection, treatment, and monitoring of patients with this disease.<sup>[98]</sup>

Exosomal proteins (i.e., CD31, C-reactive protein [CRP], soluble tumor necrosis factor receptor 1 [sTNFR1], CD114, and CD66) are other types of biomarkers which could be employed for diagnosis and monitoring of COPD patients [Table 2].<sup>[99]</sup> Recently, Tan *et al.* assessed the expression levels of exosomes in plasma of patients

with acute exacerbations of COPD (AECOPD) ( $n = 20$ ), stable COPD (sCOPD;  $n = 20$ ), and nonsmoking healthy group ( $n = 20$ ).<sup>[99]</sup> Their results revealed that plasma levels of exosomes in AECOPD and sCOPD were significantly increased compared with healthy controls. Moreover, they showed that expression levels of exosomes were associated with plasma levels of CRP, sTNFR1, and IL-6. These findings suggested that exosomes could be anticipated in various pathogenic events involved in AECOPD and sCOPD. Hence, it seems that exosomes and their cargo could be utilized as diagnostic and prognostic biomarkers for the treatment of patients with COPD.<sup>[99]</sup>

It has been shown that persistence of inflammation is one of the important characteristics of COPD.<sup>[100]</sup> Exosomes could be involved in these events via targeting various cellular and molecular pathways involved in inflammation.<sup>[101]</sup> Multiple lines of evidence have revealed that several factors such as smoking, acceleration of epithelial cell senescence, airway epithelial cell injury, destruction of pulmonary capillary vasculature, and airway remodeling are associated with COPD pathogenesis.<sup>[102]</sup> Among various factors, airway epithelial cell injury is known as an important player in COPD pathogenesis. Several studies have indicated that injured lung epithelial cells could be an important source for inflammatory mediators such as granulocyte-macrophage-colony-stimulating factor, IL-1 $\beta$ , tumor necrosis factor- $\alpha$ , TGF- $\beta$  and CXCL-8.<sup>[102]</sup> These mediators could exert their autocrine and paracrine effects on several cells. For example, TGF- $\beta$  could stimulate



remodeling of airway cells via modulation and induction of myofibroblast differentiation which is known as one of the main causes of fibrosis development during airway remodeling.<sup>[102]</sup> It has been revealed that the expression levels of TGF- $\beta$  in the small airway epithelium of patients with COPD could be associated with the severity of airway obstruction.<sup>[103]</sup> Li *et al.* indicated that paracrine activity of various mediators is mediated by various exosomes released from human macrophages.<sup>[104]</sup> Hence, it seems that chronic exposure to CS could lead to epithelial cell death and lung tissue loss via targeting various exosomes containing mediators.<sup>[104]</sup>

Another study indicated that prolonged exposure to CS could induce the release of CCN1-enriched exosomes from lung epithelial cells. It has been indicated that CCN1 has critical roles in tissue remodeling and repair process as an extracellular matrix protein.<sup>[105]</sup> This protein could improve release of CXCL-8 from cells via targeting a variety of cellular and molecular signaling pathways such as Wnt signaling pathway.<sup>[106]</sup> These findings suggested that exosomes containing CCN1 could be associated with the paracrine stimulation of CXCL-8 in the lung mesenchyme or parenchyma and the subsequent recruitment of inflammatory cells.<sup>[105,107,108]</sup> These physiological events could lead to lung tissue fibrosis.<sup>[105,107,108]</sup> Letsiou *et al.* revealed that CS could increase the numbers of circulating lung epithelial cell-derived exosomes.<sup>[109]</sup> It has also been shown that the degree of lung endothelial injury in patients with COPD could be used as a diagnostic biomarker.<sup>[109]</sup> These studies suggested that various exosomes could be employed as diagnostic and therapeutic biomarkers in patients with COPD.<sup>[109,110]</sup>

## CONCLUSION

COPD is known as a multifactorial disease, in which many factors such as smoking play critical roles. These factors could exert their effect via targeting vital pathways involved in inflammation. Among many molecules and pathways involved in COPD pathogenesis, miRNAs and exosomes have emerged as important players. It has been indicated that modulation of miRNAs via targeting various cellular and molecular pathways involved in COPD could contribute to the initiation and progression of COPD. The recognition of new markers that related with prognosis, diagnosis, therapy, and response to therapy may help improve and monitor of disease progression in COPD-afflicted patients. Among of various markers, miRNAs and exosomes have been emerged as new tools for using as diagnostic and therapeutic biomarkers in the treatment of COPD.

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## Conflicts of interest

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