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Metformin attenuates hyperoxia-induced lung injury in neonatal rats by reducing the inflammatory response

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1Department of Pediatrics, Division of Neonatology, Leiden University Medical Center, Leiden, The Netherlands; 2Department of Pediatrics, Los Angeles Biomedical Research Institute at Harbor-University of California, Los Angeles Medical Center, Torrance, California; 3Department of Pharmacology, Utrecht Institute for Pharmaceutical Sciences, Utrecht University, Utrecht, The Netherlands; and 4Department of Pulmonology, Leiden University Medical Center, Leiden, The Netherlands

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Chen X, Walther FJ, Sengers RM, Laghmani EH, Salam A, Folkerts G, Pera T, Wagenaar GT. Metformin attenuates hyperoxia-induced lung injury in neonatal rats by reducing the inflammatory response. Am J Physiol Lung Cell Mol Physiol 2015; 309: L262–L270. First published June 5, 2015; doi:10.1152/ajplung.00389.2014. —Because therapeutic options are lacking for bronchopulmonary dysplasia (BPD), there is an urgent medical need to discover novel targets/drugs to treat this neonatal chronic lung disease. Metformin, a drug commonly used to lower blood glucose in type 2 diabetes patients, may be a novel therapeutic option for BPD by reducing pulmonary inflammation and fibrosis and improving vascularization. We investigated the therapeutic potential of daily treatment with 25 and 100 mg/kg metformin, injected subcutaneously in neonatal Wistar rats with severe experimental BPD, induced by continuous exposure to 100% oxygen for 10 days. Parameters investigated included survival, lung and heart histopathology, pulmonary fibrin and collagen deposition, vascular leakage, right ventricular hypertrophy, and differential mRNA expression in the lungs of key genes involved in BPD pathogenesis, including inflammation, coagulation, and alveolar development. After daily metformin treatment rat pups with experimental BPD had reduced mortality, alveolar septum thickness, lung inflammation, and fibrosis, demonstrated by a reduced influx of macrophages and neutrophils and hyperoxia-induced collagen III and fibrin deposition (25 mg/kg), as well as improved vascularization (100 mg/kg) compared with control treatment. However, metformin did not ameliorate alveolar enlargement, small arteriole wall thickening, vascular alveolar leakage, and right ventricular hypertrophy. In conclusion metformin prolongs survival and attenuates pulmonary injury by reducing pulmonary inflammation, coagulation, and fibrosis but does not affect alveolar development or prevent pulmonary arterial hypertension and right ventricular hypertrophy in neonatal rats with severe hyperoxia-induced experimental BPD.

infants and frequently results in neonatal chronic lung disease [bronchopulmonary dysplasia (BPD)]. BPD leads to permanently enlarged alveoli, caused by lung damage, an arrest in alveolar and vascular development, and a subsequent reduction of the alveolar surface and lung function (5, 19). BPD is seriously complicated in the perinatal period by inflammation and oxidative stress and at later stages associated with pulmonary arterial hypertension (PAH)-induced right ventricular hypertrophy (RVH) and lung fibrosis (1, 5, 8, 34). Similar to premature infants at risk for developing BPD, neonatal rats are born in the saccular stage of lung development. After exposure to hyperoxia, neonatal rats develop chronic lung inflammation, followed by persistent alveolar simplification, fibrosis, PAH, and RVH (8, 9).

Because inflammation plays an important role in the pathogenesis of BPD, anti-inflammatory agents may have therapeutic potential by reducing inflammation-induced tissue damage in the lung as demonstrated in multiple studies of experimental BPD (10, 46). Metformin, an effective therapeutic option for type 2 diabetes by lowering blood glucose levels (39), has potent anti-inflammatory properties as well, both in vitro (3, 14, 26) and in vivo (28, 38). In addition metformin has protective effects on vascular function by improving capillary blood flow and limb perfusion (35, 43), and disease, including atherosclerosis, vascular remodeling, revascularization after ischemia, and apoptosis (16, 35). These pleiotropic effects of metformin are probably mediated via activation of AMP-activated protein kinase (AMPK; 20, 29). AMPK agonists have anti-inflammatory properties in vitro, as shown in multiple cell types exposed to lipopolysaccharides (LPS), including macrophages (17, 18, 45) and airway epithelial cells (26), and in vivo, as demonstrated in obese mice with allergic eosinophil inflammation (7) and in mice with inflammatory bowel disease (4).

The role of metformin in chronic lung disease, including BPD is unknown. Because inflammation is an important contributor to the pathogenesis of chronic lung disease, treatment with the potent anti-inflammatory AMPK agonist metformin may result in the identification of a novel therapy for BPD and chronic obstructive pulmonary disease. To advance our knowl-
edge on AMPK activation in neonatal cardiopulmonary disease in vivo, we studied the effects of metformin in neonatal rats with experimental BPD, induced by prolonged exposure to hyperoxia, by investigating inflammation, coagulation, alveolarization, and PAH in the lung (9).

MATERIALS AND METHODS

Animals. The research protocol was approved by the Institutional Animal Care and Use Committee of the Leiden University Medical Center. For each experiment, newborn wild type Wistar rat pups from two to three litters were pooled and equally distributed over two experimental groups: an oxygen group (n = 12) and two room air (RA)-exposed control groups (n = 6 each). For the intervention experiments, newborn rat pups were distributed over two groups, i.e., an oxygen-NaCl (n = 6) and an oxygen-metformin group (n = 6), and two RA-exposed control groups (n = 6 each) were daily injected subcutaneously either with 100 μl 0.9% NaCl or metformin (25 or 100 mg/kg; D150959; Sigma-Aldrich, St. Louis, MO). All pups were fed by Wistar foster dams. Foster dams were rotated daily between the oxygen-exposed pups and two groups of RA-exposed pups to avoid oxygen toxicity: 24 hr in 100% oxygen and 48 h in RA. Oxygen concentration, body weight, evidence of disease, and mortality were monitored daily. Pups were continuously exposed to 100% oxygen for 10 days. Lung and heart tissue was collected on day 10. Separate groups were daily injected with 10 –250 mg/kg metformin or NaCl. Because metformin reduced fibrin deposition in a broad concentration range, we used fibrin deposition in the lung as a readout. We found that metformin reduced fibrin deposition in a broad concentration range.

RESULTS

Fibrin detection assay. Quantitative fibrin deposition was determined in lung tissue homogenates by Western blotting (41, 42; n = 12). Lung tissue homogenates for quantitative fibrin deposition by Western blotting were pretreated as described previously (41). Tissue samples, dissolved in reducing sample buffer (10 mM Tris pH 7.5, 2% SDS, 5% glycerol, 5% β-mercaptoethanol, and 0.4 mg/ml of bromophenol blue) were subjected to SDS-PAGE (7.5% gel; 5% stacking gel) and blotted onto PVDF membrane (Immobilon-FL: Millipore, Bedford, MA). The 56-kDa fibrin β-chains were detected with monoclonal 59D8 (Oklahoma Medical Research Foundation, Oklahoma City, OK; diluted 1:1,000), infrared labeled goat-anti-mouse secondary antibody (IRDye 800CW; Thermo Fisher Scientific, Waltham, MA). The 56-kDa fibrin β-chains were detected using an infrared detection system (Odyssey infrared imaging system, Licor Biosciences). Fibrin deposition was quantified using rat fibrin as a reference.

Bronchoalveolar lavages and protein assay. Total protein content in lung lavages was determined as an indicator of vascular leakage using a standard protein assay (DC protein assay; Bio-Rad, Veendael, The Netherlands), according to the manufacturer’s instructions, using bovine serum albumin (fraction V; Roche Diagnostics, Almere, The Netherlands) as previously described (9; n = 10).

Real-time RT-PCR. Total RNA isolation from lung tissue homogenates (RNA-Bee, Tel-Test; Bio-Connect, Huisven, The Netherlands), first-strand cDNA synthesis (SuperScript Choice System; Life Technologies, Breda, The Netherlands), and real-time quantitative PCR, using β-actin as a housekeeping gene reference were performed on a Light Cycler 480 (Roche, Almere, The Netherlands) of the Leiden Genome Technology Center (Leiden, The Netherlands) as described previously (9; n = 10). Primers are listed in Table 1.

Statistical analysis. Values are expressed as means ± SE. Differences between groups were analyzed by one-way ANOVA for independent samples, followed by Tukey’s multiple comparison test, using the GraphPad Prism version 6 software package (San Diego, CA). Differences at P < 0.05 were considered statistically significant.

RESULTS

Dose finding for metformin in experimental BPD. To determine the optimal dosing of metformin, we performed a pilot experiment in which hyperoxia-exposed rat pups were treated daily with 10–250 mg/kg metformin or NaCl. Because metformin has anti-inflammatory and -coagulant properties (3, 29), we used fibrin deposition in the lung as a readout. We found that metformin reduced fibrin deposition in a broad concentra-
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**Table 1. Sequences of oligonucleotides for forward and reverse primers for real-time RT-PCR**

<table>
<thead>
<tr>
<th>Gene Product</th>
<th>Forward Primer</th>
<th>Reverse Primer</th>
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<tbody>
<tr>
<td>AMPK</td>
<td>5'-TGAGCAAGCTGAGTTTAAA-3'</td>
<td>5'-TTTCCAGTAATTGAGATCA-3'</td>
</tr>
<tr>
<td>CIN1</td>
<td>5'-GGAGGCAACTTTGAGATCA-3'</td>
<td>5'-GGGAGGACCTTTGAGATCA-3'</td>
</tr>
<tr>
<td>FGF4</td>
<td>5'-GTGGACGAGGCAAGCTTTT-3'</td>
<td>5'-GCCAGAAGCATGCACTTTT-3'</td>
</tr>
<tr>
<td>MCP-1</td>
<td>5'-AGTCCAGGAATGGAGATCA-3'</td>
<td>5'-TTCCAGGAGGACTTTGAG-3'</td>
</tr>
<tr>
<td>PAI-1</td>
<td>5'-GGTCCAGATGCACTTTT-3'</td>
<td>5'-GCTGCTGCTGCTGCAAGAAG-3'</td>
</tr>
<tr>
<td>TF</td>
<td>5'-GGGCGAGCAGATGCACTTTT-3'</td>
<td>5'-GGTCCAGAAGCATGGAATTT-3'</td>
</tr>
<tr>
<td>β-Actin</td>
<td>5'-TGGTGGAGGCAAGCTTTT-3'</td>
<td>5'-AGTGGGACTGCAAGGCAATCA-3'</td>
</tr>
</tbody>
</table>

AMPK, AMP-activated kinase; CIN1, chemokine-induced neutrophil monocyte chemoattractant-1; FGF4, fibroblast growth factor receptor type 4; MCP-1, monocyte chemoattractant protein-1; PAI-1, plasminogen activator inhibitor-1; TF, tissue factor.

...tion range of 10–250 mg·kg⁻¹·day⁻¹ with 25 mg·kg⁻¹·day⁻¹ of metformin as the most optimal dose to reduce fibrin deposition in the lung in hyperoxia-induced BPD (Fig. 1; n = 6). Because 100 mg·kg⁻¹·day⁻¹ of metformin, administered intraperitoneally, is effective in attenuating experimental pulmonary hypertension in adult rats (2), we used 25 and 100 mg·kg⁻¹·day⁻¹ of metformin in our experiments.

**Effects of metformin on growth and survival.** Body weight on day 10 was comparable in NaCl treated rat pups kept in RA (19.5 g; Fig. 2A) and in 100% oxygen (13.5 g). Exposure to hyperoxia resulted in a 40% survival on day 10 in NaCl-treated rat pups (Fig. 2B). Treatment of experimental BPD with 25 mg·kg⁻¹·day⁻¹ of metformin increased survival to 61% (P < 0.05, compared with hyperoxia-exposed controls), whereas 100 mg·kg⁻¹·day⁻¹ had no beneficial effects on hyperoxia-induced mortality. All RA-exposed pups showed no morbidity or mortality during the experimental period of 10 days.

**Effects of metformin on lung airway development and inflammation.** Oxygen exposure for 10 days resulted in lung edema, a heterogeneous distribution of enlarged air spaces with a decreased number of alveolar crests (2.4-fold; P < 0.001; Figs. 3B and 4A), surrounded by septa with increased thickness (1.7-fold; P < 0.001; Figs. 3B and 4D), reduced pulmonary vessel density (2.4-fold; P < 0.001; Figs. 3B and 4B), and increased pulmonary arterial wall thickness (2.5-fold; P < 0.001; Figs. 3F and 4C). Exposure to hyperoxia also induced an inflammatory response, characterized by an influx of macrophages (10.3-fold; P < 0.001; Figs. 3J and 4E) and neutrophils (13.1-fold; P < 0.01; Figs. 3N and 4F). Administration of metformin to RA-exposed controls did not have an effect on the parameters investigated (not shown). Compared with oxygen-exposed controls, administration of 25 mg·kg⁻¹·day⁻¹ of metformin during oxygen exposure reduced alveolar septal thickness (1.3-fold; P < 0.05; Figs. 3C and 4D) and the influx of macrophages (1.8-fold; P < 0.01; Figs. 3K and 4E) and of neutrophils (2.1-fold; P < 0.05; Figs. 3O and 4F). Administration of 100 mg·kg⁻¹·day⁻¹ of metformin during oxygen exposure increased the number of blood vessels compared with hyperoxia-exposed controls (1.4-fold; P < 0.001; Fig. 4B). However, metformin did not affect hyperoxia-induced inhibition of alveolarization and increase of arterial medial wall thickness.

**Effects of metformin on pulmonary deposition of collagen, elastin, and fibrin, and vascular leakage.** In rat pups kept in normoxia, collagen III was only present at high levels in the perivasculature of large and small blood vessels (Fig. 5A). Expression was low or absent in alveolar septa. In lungs of pups exposed to hyperoxia for 10 days, collagen III deposition increased 9.4-fold (P < 0.001; Fig. 5D) and was present in the perivasculature of blood vessels and in thick alveolar septa (Fig. 5B). Treatment with 25 mg·kg⁻¹·day⁻¹ of metformin for 10 days reduced collagen III expression by 29% (P < 0.05; Fig. 5J) and was present in the wall of blood vessels (Fig. 5E). Elastin expression decreased 1.5-fold under hyperoxia (P < 0.001; Fig. 5J) and was predominantly present on the septal tips and in the wall of blood vessels (Fig. 5E). Elastin expression decreased 1.5-fold under hyperoxia (P < 0.001; Fig. 5J) and was predominantly present on the alveolar walls rather than on septal tips and in the wall of small blood vessels (Fig. 5F). Treatment with 25 mg·kg⁻¹·day⁻¹ of metformin decreased elastin expression further: 2.2-fold (P < 0.001, compared with RA controls) and 1.5-fold (P < 0.01, compared with oxygen-exposed controls) in blood vessels and alveolar walls (Fig. 5G and J).

Pulmonary fibrin deposition was studied in homogenates as a readout for lung damage (Fig. 5K). Fibrin deposition was at reference levels during normal neonatal pulmonary development on day 10 (<10 ng fibrin/mg tissue) and increased 44-fold (P < 0.01) in lungs of pups exposed to 100% oxygen for 10 days. Administration of 25 mg·kg⁻¹·day⁻¹ of metformin reduced hyperoxia-induced fibrin deposition by 73% (p < 0.05), whereas administration of 100 mg·kg⁻¹·day⁻¹ of metformin showed a tendency towards lower levels compared with hyperoxia-exposed controls. Total protein concentration in bronchoalveolar lavage fluid was determined to establish the effect of pulmonary edema by capillary-alveolar leakage (Fig. 5L). Protein concentration on postnatal day 10 showed a
tendency to increase after hyperoxia but was not affected by metformin. Administration of 100 mg·kg⁻¹·day⁻¹ of metformin did not have beneficial effects on hyperoxia-induced pulmonary deposition of collagen, elastin, and fibrin, and vascular leakage compared with hyperoxia-exposed controls (Fig. 5, D and H–L).

Effects of metformin treatment on mRNA expression in lung tissue. Administration of metformin for 10 days during normal neonatal development in RA increased AMPK mRNA expression 1.6-fold (p < 0.05; not shown) but did not affect expression of the proinflammatory factors monocyte chemoattractant protein (MCP)-1 (Fig. 6A) and the chemokine-induced neutrophilic chemoattractant-1 (CINC1; Fig. 6B), the procoagulant factor tissue factor (TF; Fig. 6C), antifibrinolytic protein plasminogen activator inhibitor-1 (PAI-1; Fig. 6D), and fibroblast growth factor receptor type 4 (FGFR4; Fig. 6E). Ten days of oxygen exposure resulted in an increase in mRNA expression of MCP-1 (17-fold; P < 0.001), CINC-1 (10-fold; P < 0.001),
RVH compared with hyperoxia-exposed controls. (Fig. 7; TF (4.2-fold; \( P < 0.001 \)), and PAI-1 (62-fold; \( P < 0.001 \)), and FGFR4 mRNA expression was reduced in lungs of oxygen-exposed pups (10-fold; \( P < 0.001 \)), whereas AMPK expression was not different from RA controls (Fig. 6F). Treatment of oxygen-exposed pups with metformin for 10 days did not affect mRNA expression compared with NaCl-treated oxygen-exposed pups.

**Effects of metformin on RVH.** RVH was investigated using HE-stained heart sections. Exposure to hyperoxia for 10 days resulted in a 1.4-fold increase in the RV/LV free wall thickness ratio in Wistar control pups compared with RA controls (Fig. 7; \( P < 0.01 \)). Metformin had no beneficial effects on RVH compared with hyperoxia-exposed controls.

**DISCUSSION**

Treatment of rat pups with hyperoxia-induced lung injury, an in vivo model for experimental BPD (41), with metformin prolongs survival; reduces lung injury by attenuating lung inflammation, coagulation, septal thickness, and collagen III expression; and stimulates vascularization. Metformin had no beneficial effects on alveolarization, capillary alveolar leakage, arterial medial wall thickness (PAH), and RVH, and no adverse effects on normal lung and heart development. These data demonstrate that metformin may be a suitable candidate to reduce lung inflammation, coagulation, and fibrosis in preterm infants with severe BPD.

Metformin is a potent antidiabetic drug that is commonly used in type 2 diabetic patients to lower glucose levels in the circulation. Compared with other treatment modalities for type 2 diabetes, patients treated with metformin were protected against mortality in cardiac disease and had less cancer, suggesting that metformin has various biological functions other than its blood glucose-lowering effect in diabetic patients (12, 13, 33). Many of the pleiotropic effects of metformin are related to reduced inflammation, cancer, and cardiovascular disease and improved vascular function. Evidence is accumulating that these beneficial effects of metformin on inflammation and the cardiovasculature are mediated via activation of AMPK-dependent signaling (29, 35) and subsequent inhibition of mammalian target of rapamycin (mTOR) and NF-\( \kappa \)B signaling (27, 30, 32). Inflammation is an important player in the pathogenesis of BPD, because it may contribute to severe tissue damage and fibrosis, and because treatment with anti-inflammatory agents provides protection against hyperoxia-induced neonatal lung disease or experimental BPD (10, 46). Metformin protected against hyperoxia-induced neonatal lung injury in rat pups by reducing mortality, fibrosis, coagulation, and inflammation, as demonstrated by a reduced pulmonary influx of inflammatory cells, including macrophages and neutrophils. This anti-inflammatory effect of metformin in neonatal rats with experimental BPD is supported by observations in vivo in adult mice, in which AMPK agonist treatment, including metformin, suppressed ovalbumin-induced allergic eosinophilic lung inflammation (7, 29) and inflammatory bowel disease (4). In addition, the anti-inflammatory properties of metformin and other AMPK agonists were observed in vitro, in proinflammatory monocytes and macrophages, vascular smooth muscle cells, umbilical vein endothelial cells, and primary bronchial epithelial cells (3, 14, 17, 20, 26, 45). BPD is a multifactorial disease in which lung immaturity, aberrant alveolar development, inflammation, coagulation, vascular leakage, and pulmonary hypertension contribute to injury,
Metformin had a small, but significant beneficial effect on disease severity, and mortality (1, 5, 19). Mortality associated with experimental BPD may also be related to outcomes. Metformin does not influence, including PAH and RVH. Indeed, we demonstrated in a previous study that treatment of experimental BPD with the endothelin receptor antagonist ambrisentan had no beneficial effects on alveolarization, vascularization, inflammation, and coagulation but attenuated PAH, RVH, and fibrosis, and also improved survival (42).

The anti-inflammatory effect of metformin demonstrated by reduced pulmonary fibrin deposition in neonatal rats with experimental BPD, may, at least in part, be explained by the anti-inflammatory properties of metformin. Inflammation and coagulation are two closely related processes. Activated inflammatory cells, including macrophages, may activate coagulation directly by upregulating endogenous TF expression or triggering the release of TF-bearing microparticles (6), resulting in alveolar thrombin generation and fibrin deposition (15, 25). In addition, proinflammatory cytokines, including tumor necrosis factor (TNF)-α and interleukin (IL)-1β, may have procoagulant effects. In addition, metformin decreases the activity of the potent fibrinolytic inhibitor PAI-1, thereby promoting fibrin degradation and reducing fibrin deposition (24). Moreover, neutrophils may worsen pulmonary fibrin deposition by increased degradation of the important anticoagulant factor-activated protein C by neutrophil elastase (11). These anti-inflammatory and -coagulant effects were not confirmed by the expression of the mRNAs of the inflammatory markers MCP-1 and CINC1, the procoagulant factor TF and the regulator of fibrinolysis PAI-1, suggesting the involvement of other chemokines and cytokines, other mechanisms involved in inflammation, including cell adhesion, secretion, and migration or regulation of gene expression at a posttranscriptional level. Although neonatal exposure to hyperoxia does not affect AMPK transcription in the lung, protein activity may be reduced by proinflammatory mediators, including TNFα and IL-6, and activated by the anti-inflammatory mediator IL-10 (30). Stimulation of NF-κB signaling by oxidative stress plays an important role in inflammatory responses, including hyperoxia-induced neonatal lung injury. The anti-inflammatory and -coagulant effect of metformin may be explained by activation of AMPK and its downstream targets, including sirtuin 1 (SIRT1), peroxisome proliferator-activated receptor gamma coactivator-1α (PGC-1α), p30, and forkhead box O (FOXO) factors, thereby inhibiting NF-κB signaling (30). Metformin reduced hyperoxia-induced fibrosis in rats with severe experimental BPD by attenuating extravascular collagen III deposition. This antifibrotic effect of metformin-induced AMPK activation is supported by the beneficial effects of metformin in mice with bleomycin-induced lung fibrosis (29) and an aggravated pulmonary fibrotic response towards bleomycin in heterozygous AMPKε1-deficient mice (29).

Metformin had a small, but significant beneficial effect on hyperoxia-induced reduced vascularization, which may be explained by a proangiogenic effect observed previously in vitro.
in pulmonary arterial endothelial cells from sheep with pulmonary hypertension after AMPK activation (36) and in vivo in mice in which metformin stimulated angiogenesis and recovery after experimental stroke (40) and revascularization after hindlimb ischemia via an endothelial nitric oxide synthase (eNOS)-dependent pathway (35). This observation is supported by the beneficial effects of stimulation of the NO-eNOS-cGMP pathway with inhaled NO, apelin, and sildenafil on experimental lung and heart disease in animal models of experimental BPD by us and others (8, 9, 21, 37). Because metformin had a beneficial effect on reduced vascularization and proangiogenic factors have beneficial effects on aberrant alveolar development in (experimental) BPD (5, 9, 19, 21), we expected to find reduced alveolar enlargement as well after metformin treatment. We speculate that the proangiogenic effect of metformin is probably too small to trigger alveolarization of the simplified lung.

The absence of a beneficial effect of metformin on PAH-induced RVH in neonatal rats with hyperoxia-induced lung disease is in contrast to adult rats in which metformin protected against monocrotalin or hypoxia-induced PAH (2) and in vitro studies in which metformin inhibited endothelin-1- or hypoxia-induced proliferation of pulmonary arterial vascular smooth muscle cells (23, 44). This discrepancy in response of metformin towards PAH in neonates and adults may be explained by differences in age, development of disease in different animal models, or concentration of metformin used in the experimental rat models of PAH.

Daily treatment of neonatal rats with 25 mg/kg of metformin was the most optimal dose to attenuate hyperoxia-induced neonatal lung injury and is a similar dose used for prolonged oral daily treatment of diabetic type 2 patients (2–3 g/day in adolescents and adults). In adult rats and mice, in which metformin was administered orally via gavage or gastric tube, or intraperitoneally, the dose was higher and ranged from 100 to 350 mg·kg⁻¹·day⁻¹ to treat ovalbumin- and fungal-associated allergenic protease-induced asthma (29), bleomycin-induced fibrosis (29), allergic eosinophilic inflammation in obese mice (7), hypoxia- or monocrotalin-induced pulmonary hypertension (2), reduced vascularization in hindlimb ischemia (35), and pancreatic cancer (27). The efficacy of the relatively low dose of metformin, administered subcutaneously, in neonatal rats with BPD compared with adult rats and mice may be
explained by age, route of drug administration, species, or
development of disease in an immature lung. A relatively low
effective dose of metformin in neonatal rats will help to
prevent or reduce potential adverse effects including abdomi-
nal discomfort and diarrhea and potential bronchial edema
development due to inhibition of the epithelial Na+ channel
ENaC (26).

Because therapeutic options are lacking for BPD, there is an
urgent medical need to discover novel targets/drugs to treat this
neonatal chronic lung disease. If the absence of adverse effects
of treatment with metformin in rats can be confirmed in
newborn infants, extrapolation of the beneficial effects of
metformin and other AMPK activators in rat pups with exper-
imental BPD to premature infants with respiratory failure may
provide in a new treatment option to attenuate lung inflamma-
tion and fibrosis, which are major reasons for mortality or
mobidity in preterm infants with severe BPD.

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DISCLOSURES
No conflicts of interest, financial or otherwise are declared by the author(s).

AUTHOR CONTRIBUTIONS
Author contributions: X.C., R.M.S., E.H.L., A.S., and G.T.W. performed
experiments; X.C., R.M.S., E.H.L., A.S., and G.T.W. analyzed data; X.C.,
R.M.S., E.H.L., A.S., and G.T.W. interpreted results of experiments; X.C. and
approved final version of manuscript; F.J.W., G.F., T.P., and G.T.W.
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REFERENCES
1. Abman SH. Role of endothelin receptor antagonists in the treatment of
2. Agard C, Rolli-Derkinderen M, Dumas-de-La-Roque E, Rio M, Sagan
C, Savineau JP, Loirand G, Pacaud P. Protective role of the antiadhesive
drug metformin against chronic experimental pulmonary hypertension. Br J
Metformin, an antiabetic agent, suppresses the production of tumor
necrosis factor and tissue factor by inhibiting early growth response
factor-1 expression in human monocytes in vitro. J Pharmacol Exp Ther
4. Bai A, Yong M, Ma AG, Ma Y, Weiss CR, Guan Q, Bernstein CN,
Peng Z. Novel anti-inflammatory action of 5-aminoimidazole-4-carbox-
amide ribonucleoside with protective effect in dextran sulfate sodium-
induced acute and chronic colitis. J Pharmacol Exp Ther 333: 717–725,
2010.
5. Baralde E, Filippone M. Chronic lung disease after premature birth. N
6. Bastarache JA, Fremont RD, Kropski JA, Bossert FR, Ware J.B.
Procoagulant alveolar microparticles in the lungs of patients with acute
respiratory distress syndrome. Am J Physiol Lung Cell Mol Physiol 297:
7. Calixto MC, Lintomen L, André DM, Leiría LO, Ferreira D, Lellis-
Santos C, Anhê GF, Bordin S, Landgraf RG, Antunes E. Metformin
attenuates the exacerbation of the allergic eosinophilic inflammation in
8. de Visser VP, Walther FJ, Laghaeni EH, Boersma H, van der Laar
A, Wagenaar GT. Sildenafil alleviates pulmonary inflammation and
fibronectin deposition, mortality and right ventricular hypertrophy in neonatal
9. de Visser VP, Walther FJ, Laghaeni EH, van der Laar A, Wage-
naar GT. Apeilin attenuates hypoxic lung and heart injury in neonatal
10. de Visser VP, Walther FJ, Laghaeni EH, Steendijk P, Middeldorp M,
vander Laar A, Wagenaar GT. Phosphodiesterase-4 inhibition attenu-
ates persistent heart and lung injury by neonatal hypoxia in rats. Am J
dation in vitro by neutrophil elastase. Biol Chem Hoppe Seyler 372:
12. Elrich DT, Majumdar SR, McAlister FA, Tsuusty RT, Johnson JA.
Improved clinical outcomes associated with metformin in patients with
13. Evans JM, Donnelly LA, Emslie-Smith AM, Alessi DR, Morris AD.
Metformin and reduced risk of cancer in diabetic patients. BMJ 330:
induced nuclear factor kappaB activation via AMP-activated protein
kinase activation in vascular endothelial cells. Hypertension 47: 1183–
1188, 2006.
15. Iddel S, James KK, Levin EG, Schwartz BS, Manchanda N, Maunder
RJ, Martin TR, McIardy J, Fair DS. Local abnormalities in coagulation
and fibrinolytic pathways predispose to alveolar fibrin deposition in the
16. Isoda K, Young JI, Zilrik A, MacFarlane LA, Tsuboi N, Berdes G,
Schönbeck U, Libby P. Metformin inhibits proinflammatory responses
and nuclear factor-kappaB in human vascular wall cells. Arterioscler
Kim JB. Berberine suppresses proinflammatory responses through AMFk
activation in macrophages. Am J Physiol Endocrinol Metab 296: E955–
E964, 2009.
18. Jhun BS, Jin Q, Oh YT, Kim SS, Kong Y, Cho YH, Ha J, Baik HH,
Kang I. 5-Aminimidazole-4-carboxamide riboside suppresses lipopoly-
saccharide-induced TNF-alpha production through inhibition of phospa-
thidininositol 3-kinase/Akt activation in RAW 264.7 murine macrophages.
20. Kim SA, Choi HC. Metformin inhibits inflammatory response via
AMPK-PTEN pathway in vascular smooth muscle cells. Biochem Biophys
21. Kunig AM, Balasubramaniam V, Markham NE, Morgan D, Mont-
gomery G, Grover TR, Abman SH. Recombinant human VEGF treat-
ment enhances alveolarization after hypoxic lung injury in neonatal
22. Koppel R, Han RN, Cox D, Tanswell A, Robinovich M. Alpha
1-antitrypsin protects neonatal rats from pulmonary vascular and paren-
PN, Veasey SC, Ihida-Stansbury K, Jones PL, Goncharova EA. Bidirectional
relation between beta3-adrenergic receptor agonist and beta3-adrenergic
receptor antagonist in vascular smooth muscle cells. J Pharmacol Exp
24. Landin K, Tengborn L, Smith U. Effects of metformin and metoprolol
CR on hormones and fibrinolytic variables during a hyperinsulinemic,
26. Myerburg MM, King JD Jr, Oyster NM, Fitch AC, Magill A, Baty CJ,
Watkins SC, Kolls JK, Pilewski JM, Hallows KR. AMPK agonists
ameliorate sodium and fluid transport and inflammation in cystic fibrosis
27. Nair V, Sreevalsan S, Bashra R, Abdelrahim M, Abudayyeh A, Ro-
drigues Hoffman A, Safe S. Mechanism of metformin-dependent inhibi-
tion of mammalian target of rapamycin (mTOR) and Rap activity in

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