The impact of vitamin D on fetal and neonatal lung maturation. A systematic review

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1Hans Christian Andersen Children’s Hospital, Odense University Hospital, Odense, Denmark; 2Clinical Institute, Faculty of Health Sciences, University of Southern Denmark, Odense, Denmark; 3Institute of Molecular Medicine, Department of Cardiovascular and Renal Research, Faculty of Health Sciences, University of Southern Denmark, Odense, Denmark

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Lykke Sorensen GL, Beck-Nielsen SS, Christesen HT. The impact of vitamin D on fetal and neonatal lung maturation. A systematic review. Am J Physiol Lung Cell Mol Physiol 308: L587–L602, 2015. First published January 16, 2015; doi:10.1152/ajplung.00117.2014.—Respiratory distress syndrome (RDS) and bronchopulmonary dysplasia (BPD) are major complications to preterm birth. Hypovitaminosis D is prevalent in pregnancy. We systematically reviewed the evidence of the impact of vitamin D on lung development, surfactant synthesis, RDS, and BPD searching PubMed, Embase, and Cochrane databases with the terms vitamin D AND (surfactant OR lung maturation OR lung development OR respiratory distress syndrome OR fetal lung OR prematurity OR bronchopulmonary dysplasia). Three human studies, ten animal studies, two laboratory studies, and one combined animal and laboratory study were included. Human evidence was sparse, allowing no conclusions. BPD was not associated with vitamin D receptor polymorphism in a fully adjusted analysis. Animal and laboratory studies showed substantial positive effects of vitamin D on the alveolar type II cell, fibroblast proliferation, surfactant synthesis, and alveolarization. These data support the hypothesis of hypovitaminosis D as a frequent, modifiable risk factor of RDS and BPD, which should be tested in randomized controlled trials on pregnant women, those with threatening preterm delivery, or in the preterm neonates. Future experimental and human studies should aim to identify optimal time windows, vitamin D doses, and cut-off levels for 25-hydroxyvitamin D in interventions against RDS, BPD, and later adverse respiratory outcomes.

vitamin D; fetus; lung; surfactant; preterm; neonate
at the end of this phase during the 24th week and increases as the pulmonary parenchyma grow during the first half of the saccular phase with completion by week 34. In the alveolar phase, more alveoli are formed, providing an increase in the gas-exchange surface area, beginning week 38 and continuing postpartum, especially during the first 1–3 yr of life, with completion when the child is ~8 yr old (14).

Contrary to humans, the expression of surfactant in rats starts in the saccular phase on the 19th gestational day and is completed on the 21st-22nd day (at term) (86). The alveolar phase in rats does not begin until postpartum at postnatal days 4–5 and slowly progresses throughout life (7, 66, 72) (Fig. 1). Accordingly, rats are born at a lung developmental stage equivalent to that of a preterm human neonate (72), making rats highly suitable as an animal model for human lung diseases of the preterm.

Pulmonary surfactant is a lipoprotein complex located on the surface of the lung alveoli. Surfactant reduces surface tension and participates in host defense and the control of inflammation in the lung (75). Surfactant is synthesized and secreted by the ATII pneumocytes. In addition, the ATII cells are responsible for the regulation of alveolar ion transport. The cells are characterized by specific organelles, the osmiophilic lamellar bodies (OLBs), which represent the intracellular storage site of lung surfactant. The ATII cells are in direct contact with the ATI cells above the basement membrane and fibroblasts and other interstitial cells beneath the membrane. Fibroblasts are recognized as influencing surfactant biosynthesis through paracrine actions. Before parturition, fibroblasts secrete a polypeptide that stimulates the rate-limiting enzyme for surfactant synthesis by the ATII cells (34, 60, 70). Perinatal lung maturation is a highly regulated process, and alveolar epithelial-mesenchymal interactions play a critical role (90, 96).

Surfactant consists of 90% phospholipids and 10% proteins. The major phospholipid component dipalmitoylphosphatidylcholin reduces the surface tension in the alveoli and maintains alveolar expansion at the end of expiration. The proteins are mainly the four apoproteins, surfactant protein A, B, C, and D (SP-A, SP-B, SP-C, and SP-D). The hydrophilic SP-A and SP-D apoproteins are a defense against infections and inflammaion (1, 10). Furthermore, SP-A regulates the secretion of surfactant by binding phospholipids in a calcium-dependent manner and by helping the formation of tubular myelin at the air-liquid interphase (1). SP-A regulates the homeostasis of surfactant phospholipids through a negative feedback loop and enhancement of the recycling of phospholipids (48). SP-D is involved in activation of alveolar macrophages and in protection against pathogens but is also present in nonpulmonary tissues, such as the gastrointestinal tract and genital organs. This suggests additional actions of the protein located to other mucosal surfaces (1, 93). SP-B and SP-C are hydrophobic, and especially SP-B has a pivotal role in the absorption and distribution of surfactant (8). SP-B is required for the synthesis of lamellar bodies, the aggregation of phospholipids into the OLB, and for the reduction of surface tension at the air-liquid interphase (1, 57, 58). SP-A and SP-B (and, to a lesser extent, SP-C) promote the formation of the surfactant monolayer, which is the functionally active form of surfactant and possesses the surface-active properties (1). Whereas mutations in the SP-C-gene have been associated with neonatal RDS and interstitial lung disease, the absence of SP-B has led to respiratory failure and death shortly after birth (10, 57, 58). Concerning SP-D, animal studies show that SP-D does not participate in the normal development of the lung but may have a role in the development of emphysema and fibrosis later in life (10).

In humans, premature birth before the 34th gestational week is associated with RDS, a disease characterized by structural lung immaturity and inadequate surfactant production. RDS is a major cause of death in extremely premature neonates, where birth before the 24th gestational week is at the limit of survival despite modern intensive care. In humans, premature infants born before 32 wk of gestation have low concentrations of SP-A and SP-B and barely detectable levels of SP-C (8, 10). After birth, the concentration of SP-A increases rapidly, whereas SP-B and particularly SP-C increase more slowly (8).

Antenatal corticosteroids protect to some degree against RDS when given to mothers with potential preterm labor from 24–34 wk of gestation (86). The corticosteroids stimulate fetal lung maturation, including ATII cell differentiation and pul-
Vitamin D, also known as cholecalciferol, is a fat-soluble secosteroid and is either synthesized from 7-dehydrocholesterol in the skin after ultraviolet B (sunlight) exposure or obtained from food of animal origin. Vitamin D2 or ergocalciferol is obtained only from food of plant origin. Both forms of vitamin D may also be obtained by vitamin D supplementation. Vitamin D is converted in the liver to 25(OH)D, and the further classical pathway involves renal conversion of 25(OH)D to the biologically active form 1,25(OH)2D, also known as calcitriol (38, 45). A more recently described pathway involves nonrenal uptake of 25(OH)D, either free or bound to vitamin D-binding protein (VDBP, also known as Gc-globulin), followed by intracellular 1α-hydroxylation to 1,25(OH)2D in almost all tissues (38, 45, 80). Metabolites in the vitamin D pathway are transported in the circulation predominantly bound to VDBP and albumin, with <1% in the free, bioavailable form (17, 38). VDBP regulates availability in tissues, allowing only the small free fraction of vitamin D metabolites to passively enter cells through diffusion across cell membranes (17).

Moreover, VDBP has immunomodulatory functions primarily related to macrophage activation and neutrophil chemotaxis, and variations in the gene are associated with airway diseases, such as asthma and chronic obstructive pulmonary disease (17). The human VDBP gene is highly polymorphic, and the ability to bind vitamin D and respond to vitamin D supplementation varies according to genotype. Largely because of two frequent VDBP polymorphisms, black Americans had lower VDBP concentrations compared with whites but similar estimated bioavailable 25(OH)D despite lower total serum 25(OH)D concentrations (77). As a consequence, the widely accepted serum total 25(OH)D cut-off values of 25, 50, and 75 nmol/l for vitamin D deficiency, insufficiency, and optimal status, respectively (3, 45, 46, 83, 98), should be used with caution across racial genetic differences. Moreover, use of older routine 25(OH)D assays may differ 20% below or above the modern liquid chromatography-tandem mass spectrometry golden-standard method (26). Given these limitations, the vitamin D supplementation dose is a superior measure compared with achieved 25(OH)D concentrations in evaluation of effects in randomized controlled trials (RCTs) on vitamin D, especially across races (12).

In white Europeans, genome-wide association studies have shown that variants near genes involved in cholesterol synthesis (the 7-dehydrocholesterol reductase gene DHCR7/NADSYN1), 25-hydroxylation (CYP2R1), vitamin D transport (the VDBP gene GC), and degradation (the 24-hydroxylase gene CYP24A1) affect the S-25(OH)D concentration (101).

The actions of 1,25(OH)2D3 are mediated through binding to the nuclear VDR, a member of the superfamily of steroid receptors (45, 68, 69). VDR is a ligand-dependent transcription factor that alters the expression of select target genes in response to 1,25(OH)2D3 (38, 76). The VDR-ligand complex forms a heterodimer with the retinoid-X receptor (RXR) and binds vitamin D-responsive elements (VDREs) located predominantly in the promoter regions of target genes (53, 76). Likewise, the vitamin A-derivate retinoic acid (RA) binds to the retinoic acid receptor (RAR). The 1,25(OH)2D3-VDR complex and RA-RAR complex can form heterodimers and thereby transduce the hormonal signal into altered gene transcriptional events. In combination with transcription factors and coregulatory proteins, the 1,25(OH)2D3-VDR complex promotes or suppresses the transcription of a wide range of genes (76). VDR knockout in mice results in early-onset emphysema and decline in lung function, with lymphoid aggregate formation secondary to inflammatory changes (94). More subtle mutations or polymorphisms in the VDR gene, which may affect gene expression, RNA transcription efficacy, or protein structure, have been associated with a variety of diseases, such as multiple sclerosis, type I diabetes mellitus, osteoporosis, and colon, breast, and prostate cancer (49, 74, 97). The impact of VDR knockout on the fetal and early postnatal lung has, to our knowledge, not been studied.

Early lung development studies on the effect of DNA variations in the vitamin D synthesis, transport, degradation, and action from polymorphisms to gene knockout are generally still missing. Taken together, data on common vitamin D-related polymorphisms should optimally be included in studies reporting on vitamin D doses or serum 25(OH)D levels.

During pregnancy, the maternal requirements of vitamin D are increased because of several modifications of the vitamin D metabolism. Maternal plasma levels of 25(OH)D do not change significantly during normal pregnancy if the intake of vitamin D and sun exposure remain unchanged. In contrast, circulating levels of active 1,25(OH)2D3 increase several times over from early pregnancy and remain high during the entire pregnancy (32, 45, 73). The fetus has no endogenous production of 25(OH)D and is fully dependent on the transfer from the mother. Whereas 25(OH)D passes the placenta, 1,25(OH)2D3 is restricted to the maternal circulation and is believed to be produced de novo in the placenta and in the fetus itself (23, 32, 33, 45, 104). During pregnancy and in early childhood, the interpretation of serum 25(OH)D levels may be complicated by the presence of the inactive or less biologically active C3-epimer of 25(OH)D (6, 106). It is suggested that the C3-epimer is generated within the fetal-placental unit and not transferred across the placenta as efficiently as 25(OH)D3 (6). Hypovitaminosis D is frequent in pregnant women and in neonates (3, 22, 39, 45, 46, 84, 98), and, because the transplacental transfer of 25(OH)D mainly occurs during the last trimester, preterm infants are particularly at risk of vitamin D deficiency (65).

Taking the knowledge of a potential role of vitamin D in lung development and maturation into account, we raised the hypothesis that hypovitaminosis D could be a frequent yet largely unrecognized and modifiable risk factor of RDS and BPD in premature neonates. We therefore aimed to systematically review the evidence on the impact of vitamin D on lung development and maturation, including synthesis of surfactant, and the impact on the development of RDS and BPD.
Methods

Our systematic review was based on the guidelines of Transparent Reporting of Systematic Reviews and Meta-analysis (PRISMA statement 2009) (52).

Search strategy. For the systematic review, the databases PubMed and Embase were searched using the term vitamin D AND (surfactant OR lung maturation OR lung development OR respiratory distress syndrome OR fetal lung OR prematurity OR bronchopulmonary dysplasia). No additional titles were found using the MeSH terms or by search in the Cochrane Database. Additional studies were identified by manual search of reference lists of all full-text publications selected. Human, animal, and laboratory studies in English written language were included without time limit. The last search was made on January 12, 2015.

Exclusion criteria were as follows: 1) non-English language, 2) reviews or commentaries, and 3) topics not related to the review (studies without data on vitamin D in newborns, fetal and/or neonatal lung maturation or development, surfactant production, or respiratory problems attributable to altered lung function in newborns).

Broader nonsystematic searches were performed to serve as background material, including vitamin D AND (prematurity OR respiratory distress syndrome OR fetal lung OR prematurity OR bronchopulmonary dysplasia). No additional titles were found using the MeSH terms or by search in the Cochrane Database. Additional studies were identified by manual search of reference lists of all full-text publications selected. Human, animal, and laboratory studies in English written language were included without time limit. The last search was made on January 12, 2015.

Decision making. Search results were imported into Excel. The screening, selection, data extraction, and risk of bias assessment were performed by two independent reviewers (S. Lykkedegn and H. Christesen) with a third reviewer (T. Ussing) as a tiebreaker. Disagreements were resolved by discussion until consensus was achieved.

Risk of bias. To access the risk of bias of RCTs, the Cochrane Collaboration has developed the Cochrane RoB Tool (41). Systematic reviews of experimental animal studies are not yet common practice, and just recently the SYstematic Review Centre for Laboratory animal Experimentation has presented a SYRCLE’s RoB Tool for animal studies (42). The SYRCLE’s RoB Tool is based on the Cochrane RoB Tool and has been adjusted for the aspects of bias known to play a role in animal intervention studies. We assessed the risk of bias in the included RCTs and animal intervention studies using the above-mentioned tools (Table 1).

Results

Our search identified 1,324 records, of which 246 were duplicates (Fig. 2). After application of the exclusion criteria, 16 publications remained. The study characteristics are summarized in Tables 2, 3, and 4. Only three human studies were identified through the search, an RCT and two observational cohort studies. In addition, ten animal studies, two laboratory studies, and one combined animal and laboratory study were obtained.

Human studies. One human RCT and two observational studies were identified in the systematic search on human data. In 1999, Backström et al. (5) randomized 39 premature children born before the 33rd gestational week to vitamin D 200 IU/kg per day (maximum 400 IU/day) or 960 IU/day until 3 mo of age. Gestational age, birth weight, and birth length were comparable between the groups. At birth, the mean serum 25(OH)D concentrations were 29.8 vs. 29.2 nmol/l. At 6 wk of age, 25(OH)D was significantly higher in the high-dose vitamin D group (mean 66.7 vs. 45.7 nmol/l). The authors focused on infant bone mineralization obtained by dual-energy X-ray absorptiometry scans but noted, as the only significant finding, a reduced need for assisted ventilation in the high-dose vitamin D group (median 0 vs. 4 days, \( P = 0.01 \)). Moreover, a trend toward a lower duration of oxygen supplementation was recorded (median 2 vs. 14 days, \( P = 0.06 \)). Respiratory acidosis was more prevalent in the low-dose vitamin D group.

In 2013, Ataseven et al. (4) performed an observational cohort study investigating vitamin D deficiency as a risk factor for RDS. A total of 152 infants with a gestational age of 29 to 35 wk was included. Data on sex, gestational age, body weight, Apgar scores, antenatal steroid use, type of delivery, and antenatal problems were recorded, and serum 25(OH)D and calcium levels were measured. All 152 infants had 25(OH)D below 75 nmol/l (64% below 25 nmol/l, 33% 25–50 nmol/l, and 3% 50–75 nmol/l). No correlation between gestational age and vitamin D status was found. In unadjusted analysis, RDS was seen in 28% of the group with 25(OH)D <25 nmol/l compared with 14% in those with higher 25(OH)D. RDS was reduced 3.34 times in newborns with higher vitamin D levels, but no multivariate analysis was done.

In an observational study from 2014, Koroglu et al. (50) found an increased adjusted odds ratio (OR = 4.11, 95% CI 1.08–15.68; \( P = 0.038 \)) for BPD among 109 preterm babies with the variant VDR Fok 1 polymorphism Ff or ff, independent of patent ductus arteriosus, sepsis, mechanical ventilation, and surfactant treatment. However, when controlling for gestational age and birth weight, they saw no effect of variant Fok 1.

Animal studies. The ten animal studies (23, 59–61, 68–71, 85, 108) all showed a positive impact of vitamin D on fetal and neonatal lung development and maturation. These effects included a VDR-dependent increase in the synthesis and secretion of surfactant phospholipids in ATII cells and a postnatal vitamin D-dependent alveolar growth. The presence of the VDR in the fetal lung was restricted to the late period of gestation, corresponding to the onset of the surfactant synthesis and the differentiation of the ATII cells.

In 1987, Nguyen et al. (68) identified a significant amount of VDR related to pneumocytes in the fetal rat lung during the last quarter of gestation (days 19–21 of gestation). Crude and partially purified lung cytosols were incubated in the absence or presence of either 1,500 nM unlabeled 1,25(OH)2D3 in increasing concentrations or a 50-fold excess of radioinert 1,25(OH)2D3. The amount of VDR was either calculated by Scatchard analysis or a result of measured radioactivity using a liquid scintillation spectrometer. This study suggested that pneumocytes might be a major target tissue for 1,25(OH)2D3 during fetal life.

In a subsequent study from 1990 (69), the group investigated which cell types were directly responsive to 1,25(OH)2D3. VDR-dependent alveolar growth. The presence of the VDR in the fetal lung was restricted to the late period of gestation, corresponding to the onset of the surfactant synthesis and the differentiation of the ATII cells.

In vivo, lung tissue from rat fetuses (day 21) and newborn rats (3 days after birth) were prepared for immunohistochemical staining with a monoclonal antibody against VDR (9A7γ). Furthermore, the level of VDRs was estimated in cytosols from...
fresh lung tissue using a 50-fold excess of unlabeled 1,25(OH)₂D₃, as previously described by the group (68). The number of specific VDRs was calculated as the difference between total binding and nonspecific binding. The highest levels of VDR were located in cells corresponding to ATII cells at the end of pregnancy (days 20–21), whereas little or no immunostaining was observed in the tissue from the 3-day-old pups. The level of VDRs decreased a few hours before delivery and remained low during the first 5 days of life.

In vitro, the level of VDRs was estimated in cytosols from fetal lung explants taken on days 20–21 of gestation and cultured for 48 h using the same method as in vivo. The number of VDRs in vitro was different from the in vivo pattern, as the number did not decrease during the culture period (48 h). In addition, the group used the fetal lung explants to test the effect of factors known to affect lung maturation, including 1,25(OH)₂D₃, dexamethasone, thyroxine, prolactin, terbutaline, retinoic acid, insulin, and oxytocin.
Prolactin, thyroxine, 1,25(OH)₂D₃, and to a lesser extent dexamethasone increased the capacity of ATII cells to bind 1,25(OH)₂D₃ by inducing an increase in the number of VDRs without altering the receptor-binding affinity.

The same year Marin et al. (61) demonstrated that 1,25(OH)₂D₃ (10⁻⁹ M) treatment of 18-day-old immature fetal rat lung significantly increased the levels of surfactant-related phospholipids ex vivo. Lipids were extracted from fetal lung explants, and the levels of phospholipids were visualized by thin-layer chromatography. An increase in the surfactant-related phospholipids was also seen in dexamethasone-treated explants, but not to the same extent. Dexamethasone (10⁻⁷ M) did not have any additive effect when present together with 1,25(OH)₂D₃ in the culture medium. Morphological observations were studied by light and electron microscopy. Whereas surfactant mainly was accumulated intracellularly in the dexamethasone-treated explants causing abnormally large OLBs, it was mainly found extracellularly in the luminal space in the 1,25(OH)₂D₃-treated explants. Thus 1,25(OH)₂D₃, not only stimulated the synthesis of phospholipids, but also triggered the surfactant secretion.

The same group used a similar setup in 1993 to analyze the morphological changes in surfactant synthesis and release induced by 1,25(OH)₂D₃ (10⁻⁹ M) and dexamethasone (10⁻⁷ M) (60). Changes were visualized by both light and electron microscopy. They showed no significant effect of 1,25(OH)₂D₃ on the number of immature pneumocytes entering differentiation and no significant effect on structural changes (increase of

Table 2. Study characteristics, human studies

<table>
<thead>
<tr>
<th>Source</th>
<th>Population</th>
<th>Study Design</th>
<th>Intervention</th>
<th>Outcome</th>
<th>Results</th>
<th>Statistics</th>
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</thead>
<tbody>
<tr>
<td>Backström et al. 1999</td>
<td>39 preterm infants</td>
<td>RCT</td>
<td>200 IU/kg body wt per day ≤400 IU/day (a) vs. 960 IU/day (b)</td>
<td>Gestational age, Birth weight, Birth length, Duration of assisted ventilation, Duration of oxygen supplement,</td>
<td>30 ± 3 (a) vs. 30 ± 6 (b)</td>
<td>P = 0.19</td>
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<td>Birth weight, Birth length, Duration of assisted ventilation, Duration of oxygen supplement,</td>
<td>1.365 g (a) vs. 1.510 g (b)</td>
<td>P = 0.45</td>
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<td>Birth weight, Birth length, Duration of assisted ventilation, Duration of oxygen supplement,</td>
<td>39 cm (a) vs. 40 cm (b)</td>
<td>P = 0.65</td>
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<td>Birth weight, Birth length, Duration of assisted ventilation, Duration of oxygen supplement,</td>
<td>4 (a) vs. 0 (b)</td>
<td>P = 0.01</td>
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<td>Birth weight, Birth length, Duration of assisted ventilation, Duration of oxygen supplement,</td>
<td>14 (a) vs. 2 (b)</td>
<td>P = 0.06</td>
</tr>
<tr>
<td>Ataseven et al. 2013</td>
<td>152 preterm infants</td>
<td>Observational cohort study</td>
<td>No intervention</td>
<td>Gestational age, Birth weight, Antenatal corticosteroids, 25(OH)D₃, ng/ml</td>
<td>18 vs. 63</td>
<td>P = 0.01</td>
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<td>Birth weight, Antenatal corticosteroids, 25(OH)D₃, ng/ml</td>
<td>1.667 ± 505 vs. 1.974 ± 585</td>
<td>P = 0.00</td>
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<td>Birth weight, Antenatal corticosteroids, 25(OH)D₃, ng/ml</td>
<td>23 vs. 32</td>
<td>P = 0.40</td>
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<td></td>
<td>Birth weight, Antenatal corticosteroids, 25(OH)D₃, ng/ml</td>
<td>7.5 ± 4.9 vs. 9.6 ± 5.7</td>
<td>P = 0.06</td>
</tr>
<tr>
<td>Koroglu et al. 2014</td>
<td>109 preterm infants</td>
<td>Observational cohort study</td>
<td>No intervention</td>
<td>Gestational age, Birth weight, Surfactant treatment, Duration of assisted ventilation, Duration of oxygen therapy, Survival</td>
<td>30.17 vs. 27.19</td>
<td>P &lt; 0.001</td>
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<td>Gestational age, Birth weight, Surfactant treatment, Duration of assisted ventilation, Duration of oxygen therapy, Survival</td>
<td>1,523.79 vs. 980.04</td>
<td>P &lt; 0.001</td>
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<td>Gestational age, Birth weight, Surfactant treatment, Duration of assisted ventilation, Duration of oxygen therapy, Survival</td>
<td>29 vs. 83</td>
<td>P &lt; 0.001</td>
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<td>Gestational age, Birth weight, Surfactant treatment, Duration of assisted ventilation, Duration of oxygen therapy, Survival</td>
<td>3.59 vs. 41.95</td>
<td>P &lt; 0.001</td>
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<td>Gestational age, Birth weight, Surfactant treatment, Duration of assisted ventilation, Duration of oxygen therapy, Survival</td>
<td>3.37 vs. 79.02</td>
<td>P &lt; 0.001</td>
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<td>Gestational age, Birth weight, Surfactant treatment, Duration of assisted ventilation, Duration of oxygen therapy, Survival</td>
<td>62 vs. 43</td>
<td>P = 0.032</td>
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</tbody>
</table>

RCT, randomized controlled trial. (a), Group receiving 200 IU/kg body wt per day ≤400 IU/day intervention; (b), group receiving 900 IU/day intervention.
<table>
<thead>
<tr>
<th>Source</th>
<th>Animals</th>
<th>Endpoints</th>
<th>Analysis</th>
<th>Results</th>
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<tbody>
<tr>
<td>Marin et al. 1990 (61)</td>
<td>Sprague-Dawley rats</td>
<td>Content of surfactant-related phospholipid content (a) Morphology (b)</td>
<td>Thin-layer chromatography (a) Electron microscopy (b) Light microscopy (b)</td>
<td>1,25(OH)<em>{2}D</em>{3} significantly increased the levels of phospholipids. Dexamethasone also increased the levels of phospholipids but not to the same extent, and no additive effects of the 2 treatments were shown.</td>
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<tr>
<td>Nguyen et al. 1993 (60)</td>
<td>Pregnant Sprague-Dawley rats</td>
<td>Morphology</td>
<td>Electron microscopy Light microscopy</td>
<td>1,25(OH)<em>{2}D</em>{3} had no significant effect on either the number of immature pneumocytes entering differentiation or the structural changes in the ATII cells but stimulated the release of surfactant. Dexamethasone slowed down the enlargement of the luminal space significantly.</td>
</tr>
<tr>
<td>Edelson et al. 1993 (23)</td>
<td>Pathogen-free Wistar rats Male adult and neonatal Sprague-Dawley rats</td>
<td>Influence of 1,25(OH)<em>{2}D</em>{3} on alveolar epithelial proliferation</td>
<td>Thymidine incorporation Cell number Autodradiography Flow cytometry</td>
<td>1,25(OH)<em>{2}D</em>{3} increased thymidine incorporation into DNA in the late neonatal and adult ATII cells but not in fetal or early neonatal ATII cells. The increase in the thymidine incorporation was accomplished by an increase in cell number.</td>
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<tr>
<td>Nguyen et al. 1996 (70)</td>
<td>Sprague-Dawley rats</td>
<td>Localization of VDR (a) Lung cell characterization (b) Binding of 1,25(OH)<em>{2}D</em>{3} (c) Phospholipid synthesis and release (d) Metabolism of 1,25(OH)<em>{2}D</em>{3} (e) VDR labeling</td>
<td>Immunohistochemistry (a) Light and electron microscopy (b) Binding studies (c) Thin-layer chromatography (d) HPLC (e)</td>
<td>Whereas HPLC was used to demonstrate a paracrine system during the last 3 days of pregnancy (days 19–21 of gestation), immunostaining showed that ATII express VDR and fibroblasts do not. Thin-layer chromatography showed that 1,25(OH)<em>{2}D</em>{3} stimulated the production and release of phospholipids by ATII cells.</td>
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<td>Nguyen et al. 2004 (71)</td>
<td>Sprague-Dawley rats</td>
<td>VDR labeling</td>
<td>Electron microscopy</td>
<td>VDRs were located to the ATII cell nucleus, cytoplasm, and endoplasmic reticulum at 21 days of gestation. 1,25(OH)<em>{2}D</em>{3} was especially active in the intermediate stage of ATII cell differentiation. In vitro, incubation with 1,25(OH)<em>{2}D</em>{3} or 1,25(OH)<em>{2}D</em>{3}-3-epi-D_{3} and key markers for alveolar epithelial-mesenchymal interactions caused increases in PTHrP receptor, PPAR-γ and ADRP in lipofibroblasts and in SP-B and leptin receptor in ATII cells. Furthermore, both treatments caused a dose-dependent increase in the proliferation of both lipofibroblasts and ATII cells by inhibiting apoptosis. In vivo, immunoprecipitation administration with either 1,25(OH)<em>{2}D</em>{3} or 1,25(OH)<em>{2}D</em>{3}-3-epi-D_{3} during the first 2 wk of postnatal life led to an increase in the expression of key markers for both lipofibroblast (PPAR-γ and ADRP) and ATII cell (SP-B and SP-C) differentiation. In addition, immunohistochemical staining showed a significant increase in both alveolar count and alveolar septal thickness in vitamin-treated animals compared with controls. TGV was significantly smaller in vitamin D-deficient mice compared with replete controls, and lung mechanics was significantly higher. Stereology also showed significantly smaller lung volume in the vitamin D-deficient mice compared with controls but no difference in either surface area or septal thickness between the groups. The number of alveoli was lower in the vitamin D-deficient group of female mice but not in the male mice.</td>
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Continued
Primary cultures of ATII cells were exposed to 1,25(OH)2D3 neonatal (7 and 18 days postpartum), and adult rat lung. epithelial cells from fetal (gestational
panied by an increase in cell number, demonstrating that 1,25(OH)2D3 acts as a growth factor for ATII cells in postnatal
incorporation into DNA in primary cultures of late neonatal
They showed that 1,25(OH)2D3 acts to increase thymidine
expression a functional VDR that can be upregulated by
1,25(OH)2D3. To study the ability of fibroblasts and ATII cells to convert 25(OH)D3 into 1,25(OH)2D3 during the last days
VDRs were located in the nucleus, cytoplasm, and endoplas-
the total pulmonary compliance was not significantly
different in the 4 groups, but the tracheal
contractility response to acetylcholine showed
increased dose of cholecalciferol in the diet.
levels of calcium were unaffected by both
decrease was seen. Perinatal
appeared to block the altered airway contractility and
VDR, vitamin D receptor; ATII, alveolar type II cell; HPLC, high-performance liquid chromatography; PTHrP, parathyroid hormone-related protein; PPAR-γ, peroxisome proliferator-activated receptor-γ; ADRP, adipocyte differentiation-related protein; SP-B, surfactant protein B; TGV, thoracic gas volume. Letters in parentheses connect the endpoints in the analysis.
the surface area of the epithelial structures and the numbers of cells) in the ATII cells. In contrast, the presence of dexameth-
asone slowed down the enlargement of the luminal space
significantly. 1,25(OH)2D3 stimulated the release of surfactant
in the mic reticulum of ATII cells at 21 days of gestation in fetal rat
8 years later in 2004, Nguyen et al. (71) discovered that
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Table 4. Study characteristics, laboratory studies

<table>
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<td>Rehan et al. 2002 (79)</td>
<td>NCI-H441</td>
<td>Identification of metabolite M (a) Surfactant phospholipid synthesis (b)</td>
<td>HPLC (a) GC/MS (a) Liquid scintillation</td>
<td>HPLC and GC/MS identified the metabolites of 1,25(OH)2D3, and both 1,25(OH)2D3 and 1,25(OH)2D3-3-epi-D3 were found to be significant stimulators of the synthesis of surfactant phospholipids in ATII cells. 1,25(OH)2D3-3-epi-D3 also increased mRNA expression and synthesis of SP-B.</td>
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<tr>
<td>Nguyen M et al. 2004 (71)</td>
<td>NCI-H441</td>
<td>Expression of F1,6-BP mRNA (a) VDR binding (b)</td>
<td>Northern blot (a) Binding study (b)</td>
<td>Incubation with 1,25(OH)2D3 led to a 1.4-fold increase in the ability of ATII cytosol to bind 1,25(OH)2D3 and increased the expression of F1,6-BP mRNA. VDR was barely detectable in human fetal lung and human ATII cells in the absence of 1,25(OH)2D3, but increased dramatically in the presence of the hormone. In general, 1,25(OH)2D3 decreased the expression of SP-A mRNA in human fetal lung tissue and reduced SP-A mRNA and protein levels in isolated ATII cells. No significant effect of 1,25(OH)2D3 on SP-B and SP-C mRNA levels was observed in human fetal lung tissue, but the levels of both SP-B mRNA and SP-B protein in ATII cells were increased in the presence of 1,25(OH)2D3.</td>
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<tr>
<td>Phokela et al. 2005 (75)</td>
<td>NCI-H441</td>
<td>Expression of SP-B mRNA (c) SP-B protein synthesis (d)</td>
<td>SP-B protein synthesis (d)</td>
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<td>Expression of SP-B mRNA (c) SP-B protein synthesis (d)</td>
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<td>culture</td>
<td>Expression of SP-B mRNA (c) SP-B protein synthesis (d)</td>
<td>Western blot (c)</td>
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In 2009, Sakurai et al. (85) determined a role of both 1,25(OH)2D3 and its metabolite 1,25(OH)2D3-3-epi-D3 in key alveolar epithelial-mesenchymal interactions.

Embryonic day 19 primary rat lung lipofibroblasts and ATII cells were incubated with 1,25(OH)2D3 (10^{-11} M) or 1,25(OH)2D3-3-epi-D3 (10^{-8} M) in vitro, and key markers for alveolar epithelial-mesenchymal interactions were visualized by Western blotting. Both treatments caused increases in the expression of those markers. Whereas treatment of lipofibroblasts caused increases in parathyroid hormone-related protein receptor, peroxisome proliferator-activated receptor-γ (PPAR-γ), and adipocyte differentiation-related protein (ADRP), treatment of ATII cells resulted in an increase in both SP-B and leptin receptor. Furthermore, both 1,25(OH)2D3 and 1,25(OH)2D3-3-epi-D3 treatments caused a dose-dependent increase in the proliferation of both lipofibroblasts and ATII cells by inhibiting apoptosis.

Newborn rat pups were administrated with 1,25(OH)2D3 (10 ng/kg body wt) or 1,25(OH)2D3-3-epi-D3 (50 ng/kg body wt) intraperitoneally during the first 2 wk of postnatal life, and the effects on markers of spontaneous lung maturation were determined by Western blotting. Compared with controls, both treatments led to an increase in the expression of key markers for both lipofibroblast (PPAR-γ and ADRP) and ATII cell (SP-B and SP-C) differentiation. In addition, immunohistochemical staining of lungs from 1,25(OH)2D3-treated animals showed a significant increase in both alveolar count and alveolar septal thickness compared with controls. Altogether, this study implied an essential physiological role of vitamin D in perinatal pulmonary maturation.

In 2011, Zosky et al. (108) showed a connection between vitamin D deficiency and altered postnatal lung development. Female BALB/c mice were assigned to either a vitamin D-deficient or vitamin D-replete (2.195 IU/kg) diet. Pups of both sexes were studied at 2 wk of age. Whereas lung volume (thoracic gas volume [TGV]) and lung mechanics were assessed through respectively plethysmographic measurements and modified low-frequency forced-oscillation technique, respectively, lung structure was assessed after fixation with 2,5% glutaraldehyde and visualized by stereology. The results showed significantly smaller TGV and significantly higher airway resistance and lung mechanics (tissue damping and elastance) in vitamin D-deficient mice compared with replete controls, which still after adjustment for body length suggest that vitamin D has an effect on postnatal lung growth. Stereology also showed significantly smaller lung volume in the vitamin D-deficient mice compared with controls but no difference in either surface area or septal thickness between the groups. However, the number of alveoli was lower in the vitamin D-deficient group of female mice but not in the male mice.

In a study from 2014, Mandell et al. (59) examined the effects of vitamin D on neonatal rat pup survival after antenatal endotoxin injection and on isolated endothelial and ATII cells. Pregnant Sprague-Dawley rats received intra-amniotic injections at 20 days of gestation. The animals were assigned to a saline control (50 μl), endotoxin alone (10 μg endotoxin/50 μl saline), or endotoxin + vitamin D (50 pg vitamin D/50 μl saline). Cesarean sections were performed on day 22 of gestation. Newborn pups that were exposed to endotoxin alone
received daily intraperitoneal injections of vitamin D (1 ng/g) or saline for 14 days. Oxygen saturation was measured at delivery, and survival was monitored daily throughout the observation period. Survival rate was calculated as the number of survived pups divided by the number of placental sacs that received intra-amniotic injections.

Antenatal vitamin D treatment improved oxygenation (78% vs. 87%) and survival (84% vs. 57%) significantly after endotoxin exposure compared with endotoxin alone. Furthermore, the study also included a description of the alveolar growth. Fetal sheep ATII cells were isolated, and the viability of the ATII cells was assayed using Trypan blue exclusion and counted on a hemacytometer (viability >90%). The study concluded that vitamin D has a proliferative and protective effect on fetal ATII cells and suggested that early vitamin D therapy might be a potential strategy for reducing the risk of acute respiratory distress.

In the most recent study from 2014, Yurt et al. (107) studied the effects of vitamin D deficiency on lung molecular and structural alterations in rat pups. Sprague-Dawley dams were assigned to four different dietary groups 4 wk before mating: 0, 250, 500, or 1,000 IU/kg cholecalciferol. All diets contained 4.5 g/kg calcium. The assigned dietary regimens were continued throughout pregnancy and lactation until the pups were delivered, and survival was monitored daily throughout the observation period. Survival rate was calculated as the number of survived pups divided by the number of placental sacs that received intra-amniotic injections.

Vitamin D deficiency has been shown to alter lung development and function in preterm and term infants. This is because vitamin D deficiency inhibits the synthesis of surfactant proteins and affects the expression of genes involved in lung development. The study by Yurt et al. (107) demonstrated that antenatal vitamin D treatment improved oxygenation and survival in rat pups, providing evidence that early vitamin D therapy might be a potential strategy for reducing the risk of acute respiratory distress.

In a study by Rehan et al. (79), human ATII cells were able to convert 1,25(OH)2D3 into its metabolite 1,25(OH)2D3-3-epi-D3. The study compared the effects of 1,25(OH)2D3 and 1,25(OH)2D3-3-epi-D3 on mRNA expression and protein levels of surfactant proteins and pro-inflammatory cytokines. The study found that 1,25(OH)2D3-3-epi-D3 was equipotent to 1,25(OH)2D3 in promoting surfactant protein expression and suppressing pro-inflammatory cytokine expression.

To determine the effects of vitamin D on surfactant synthesis, NCI-H441 cells were incubated with either 1,25(OH)2D3 (10−9 M) or 1,25(OH)2D3-3-epi-D3 (10−9 M) and the metabolites were analyzed using gas chromatography mass spectrometry. The study found that both vitamin D compounds increased the synthesis of surfactant, the 1,25(OH)2D3-3-epi-D3 group compared with the supplemented groups. Furthermore, a dose-dependent increase in choline incorporation into saturated phosphatidylcholine and triolein uptake was observed, indicating that vitamin D deficiency inhibited both triglyceride uptake and de novo surfactant phospholipid synthesis. In conclusion, perinatal supplementation with 500 IU/kg cholecalciferol appears to block the altered airway contractility and alveolar epithelial-mesenchymal signaling caused by vitamin D deficiency.

The study by Rehan et al. (79) also investigated the effects of vitamin D on surfactant protein expression. NCI-H441 cells were incubated with either 1,25(OH)2D3 (10−9 M) or 1,25(OH)2D3-3-epi-D3 (10−9 M) and the metabolites were analyzed using gas chromatography mass spectrometry. To study the effects of vitamin D on surfactant synthesis, NCI-H441 cells were incubated with either 1,25(OH)2D3 (10−9 M) or 1,25(OH)2D3-3-epi-D3 (10−9 M) and prepared for liquid scintillation spectrometry. The study found that both vitamin D compounds increased the synthesis of surfactant, the 1,25(OH)2D3-3-epi-D3 group compared with the supplemented groups.

Nguyen et al. (71) demonstrated in 2004 a 1.4-fold increase in the ability of ATII cytosol to bind 1,25(OH)2D3 in human ATII cells. Human NCI-H441 type II cells were incubated in the presence of 1,25(OH)2D3 (10−8 M) or ethanol in the absence or presence of radioinert hormone, and the binding capacity of the ATII cells was estimated. The cells were also used to study the expression of F1,6-BP, a regulatory enzyme in glucose synthesis and degradation, by Northern blot analysis. The F1,6-BP mRNA expression was increased by 1.25(OH)2D3 action on surfactant synthesis via the gluconeogenesis pathway.

One year later in 2005, Phokela et al. (75) investigated the effects of 1,25(OH)2D3 on NCI-H441 cells, human fetal lung tissue in organ culture, and isolated ATII cells in primary culture. To study the expression of VDR in human fetal lung and primary ATII cells, cells were incubated in the absence or presence of 1,25(OH)2D3 (10−7 M), and lysates were analyzed by Western immunoblot analysis. VDR was barely detectable in the absence of 1,25(OH)2D3 but increased dramatically in the presence of the hormone. Expression of SP-A was visualized...
by Northern blot and Western blot analysis in human fetal lung, primary ATII cells, and NCI-411 cells after incubation with various concentrations of 1,25(OH)2D3 (10−10 M or 10−7 M) in the absence or presence of dibutyryl cyclic AMP. Expression of SP-B was studied in human fetal lung and primary ATII cells and SP-C only in human fetal lung after the same procedure as described above. In general, 1,25(OH)2D3 decreased the expression of SP-A mRNA in human fetal lung tissue and reduced SP-A mRNA and protein levels in isolated ATII cells. No significant effect of 1,25(OH)2D3 on SP-B and SP-C mRNA levels was observed in human fetal lung tissue, but the levels of both SP-B mRNA and SP-B protein in ATII cells were increased in the presence of 1,25(OH)2D3.

Discussion

The evidence of an impact of vitamin D on human fetal and neonatal lung diseases is sparse. Our systematic search identified only one small human RCT and two observational studies, of which only one had adjusted the association analysis for confounders. Regarding lung development and maturation, several studies in rodents showed a positive effect of vitamin D on the proliferation of ATII cells and fibroblasts, surfactant synthesis, and upregulation of VDR in the lungs. These findings were supported by laboratory studies.

Human studies. The RCT of Backström et al. (5) showed a decreased need of ventilation and a trend toward a shorter duration of oxygen supplementation in the high-dose vitamin D group, implying an improved lung function. The RCT was not designed primarily to investigate the effect of vitamin D on mechanical ventilation days or days with oxygen supplementation, and no statistical power calculation was performed for the association. An increased serum 25(OH)D level in the high-dose vitamin D group was not documented until 6 wk of age (~42 days). However, the number of days on mechanical ventilation ranged up to 50 days and 60 days in the low- and high-dose vitamin D group, respectively, which was within the time period of documented increased serum 25(OH)D levels. A decreased need of mechanical ventilation is an important finding, as even modern invasive ventilator treatment for preterm infants born at gestational age <32 wk and 25(OH)D was not determined. Most probably, the rachitic respiratory distress syndrome was due to postnatal softening and fracture of the ribs as well as weakness of the respiratory muscles.

Hypovitaminosis D is frequent, not only in pregnancy, but also in preterm neonates. In Ireland, 78% of very preterm neonates born (gestational age <32 wk, or birth weight <1,500 g) had a serum 25(OH)D ≤50 nmol/l at 18 days of age despite vitamin D supplementation (65). Other studies have reported a mean cord blood 25(OH)D3 in preterm infants between 14.5 and 29.2 nmol/l (19, 22). This high prevalence implies that hypovitaminosis as a risk factor of RDS or BPD would be frequent and therefore detectable in larger cohort studies or matched case-control studies. Well-designed cohort or case-control studies on the impact of vitamin D on RDS or BPD in very and extremely preterm neonates are encouraged. To date, the sparse existing human data on the impact of vitamin D on RDS or BPD do, however, in conjunction support a hypothesis of a risk factor to be tested in RCTs.

Animal and laboratory studies: fetal lung development. The animal studies (23, 59–61, 68–71, 85, 108) demonstrated the presence of VDR in the lung primarily during the last period of gestation (days 19–22) when ATII cells differentiate, surfactant biosynthesis and secretion begin, and the glycogen content...
decreases. Incubation with 1,25(OH)2D3 induced an upregulation of the receptor and stimulated differentiation and proliferation in both fibroblasts and ATII cells.

According to Marin et al. (60, 61) 1,25(OH)2D3 accelerated the decrease in glycogen content and increased the synthesis of surfactant-related phospholipid and the secretion of surfactant by ATII cells isolated from fetal rat lung. Of clinical interest, dexamethasone had no additional effects on surfactant concentration in the presence of 1,25(OH)2D3, and dexamethasone alone did not induce surfactant exocytosis in contrast to 1,25(OH)2D3 (60, 61). Antenatal prevention of RDS with dexamethasone or other glucocorticoids to the pregnant mother is widely used and evidence based in threatening premature birth (13, 82). The animal data suggest that vitamin D may be an additional, if not alternative, option to glucocorticoids in the prevention of RDS in mothers with threatening premature birth.

Nguyen et al. (71) brought further evidence of a physiological role of 1,25(OH)2D3 by presenting an association between the expression of VDR and the different stages of lung maturation in fetal rats. Their findings of an increased F1,6BP mRNA expression by 1,25(OH)2D3 in human ATII cells indicates that activation of F1,6-BP may play a role in the actions of 1,25(OH)2D3 on surfactant synthesis through the pathway of gluconeogenesis. Besides the effects on gluconeogenesis, F1,6-BP might facilitate surfactant synthesis in the postnatal lung (30). The F1,6BP gene is known to bear VDRE, and both RA and 1,25(OH)2D3 induce F1,6BP gene expression in human differentiated monocytes (28).

Whereas Sakurai et al. (85) illustrated the ability of 1,25(OH)2D3 to stimulate the differentiation of both fibroblasts and ATII cells, others (53) have shown that RA also is a potential stimulator of lung maturation. Both 1,25(OH)2D3 and RA exert their effects by binding as ligands to their respective intracellular receptors, RARs and VDRs, which both can form heterodimers with RXR and thereby transduce the hormonal signal into altered gene transcriptional events. Furthermore, interactions between RA and vitamin D signaling pathways have been described in lungs (53) as well as in other organ systems (87). The laboratory studies (71, 75, 79) supported that vitamin D has an effect on several physiological lung maturation processes in rodents. Rehan et al. (79) demonstrated the ability of vitamin D to stimulate the production and secretion of surfactant-related phospholipids in human ATII cells. Phokela et al. (75) demonstrated how 1,25(OH)2D3 increases SP-B mRNA expression in human ATII cells and reduces the expression of SP-A mRNA in human fetal lung tissue and isolated ATII cells. The increase in SP-B mRNA and reduction in SP-A mRNA caused by 1,25(OH)2D3 are similar to observations made in the presence of both glucocorticoids and RA (11, 75, 95), implying a complex regulation of fetal lung maturity and surfactant synthesis. Phokela et al. (75) also demonstrated how 1,25(OH)2D3 upregulates VDR in human fetal lung tissue and human isolated ATII cells, confirming findings by Nguyen et al. (71) in human ATII cells.

Taken together, the animal and laboratory studies have given detailed insights in the mechanistic actions through which vitamin D stimulates maturation of the fetal lung. Still, no studies have to our knowledge clarified the relationship between the production of 1,25(OH)2D3 in the fibroblasts, the upregulation of VDR, and the biosynthesis of surfactant in ATII cells during the late period of gestation. Further investigation of these relationships is warranted to elucidate the physiological role of 1,25(OH)2D3 in the fetal lung.

**Animal and laboratory studies: postnatal lung development.**

In the rat lung, the widespread expression of VDR decreases before term delivery and remains low the first days of life (69–71). VDR is present after birth in pulmonary macrophages as in almost all immune cells (55, 64, 68), and alveolar macrophages convert 25(OH)D to 1,25(OH)2D (23, 68, 80).

Besides the stimulation of differentiation and proliferation in ATII cells and fibroblasts, 1,25(OH)2D3 inhibits apoptosis in both cell types during the most active period of alveolarization and increases the alveolar count postnatally (23, 59, 60, 85). These findings suggest a local alveolar modulation of the postnatal alveolar growth by vitamin D.

Both Zosky et al. (108) and Mandell et al. (59) showed a link between vitamin D deficiency and altered postnatal lung development and maturation in rodents. Vitamin D improved oxygenation and survival after antenatal endotoxin exposure (59).

Even though the risk of bias analysis weakens the results by Sakurai et al. (85), others have found similar results (23, 59, 60, 71), which especially supports their postnatal findings. In the paper by Mandell et al. (59), blinding of the caretakers and investigators is not well described, but their findings are supported by Zosky et al. (108), whose risk of bias analysis does not raise any concerns.

Taken together, fetal and postnatal effects of vitamin D on lung maturation are well documented and suggest that vitamin D therapy may be a preventive or therapeutic option in preterm neonates against, not only RDS, but also BPD.

Future animal studies should be designed to identify optimal time windows and optimal doses for intervention with vitamin D treatment in vitamin D-depleted models at different stages in early pregnancy, late pregnancy, and in preterm offspring. However, the present knowledge from animal and laboratory data should encourage researchers to perform RCTs on the effect of vitamin D in the prevention and treatment of RDS and BPD. Population-based RCTs with vitamin D supplementation in pregnancy as a primary prevention must be very largely dimensioned to overcome the rareness of extreme or very premature delivery. Moreover, a control group without vitamin D supplementation would not be feasible in countries with recommended vitamin D supplementation. RCTs restricted to women with documented vitamin D deficiency or insufficiency must take the time delay of 25(OH)D determination into account.

RCTs targeted to women with threatening preterm birth or to the preterm neonates from the earliest possible time are more feasible options but with a short time interval between intervention and outcome, possibly requesting a high, single, or repeated vitamin D dose analogous to the well-established antenatal steroid therapy to women with threatening preterm delivery. The RCTs should investigate the effect of different vitamin D doses with determination of 25(OH)D to establish cut-off values for eventual effects and toxicity.

Animal data have shown an effect of vitamin D on respiratory airways. Foong et al. (27) showed a decreased expression of transforming growth factor (TGF)-β1 and TGF-β receptor I in vitamin D-depleted female mice fetuses on day E17.5. At the
postnatal age of 8 wk, airway resistance and airway smooth muscle mass were significantly increased in the large airways; lung volume, volume of parenchyma, and alveolar septa were smaller, and TGF-β levels were reduced in bronchoalveolar lavage fluid. Male mice only showed the latter change.

Yurt et al. (107) recently demonstrated both proximal and distal airway molecular and functional alterations caused by perinatal vitamin D deficiency and dose-dependent prevention of these alterations with cholecalciferol supplementation, indicating that vitamin D supplementation may have a clinical effect in preventing childhood asthma.

In keeping with the animal data, a recent human study (109) showed that lower maternal vitamin D concentrations in pregnancy were after confounder control associated with current wheeze (both sexes, questionnaire data, \( P = 0.05 \)), current asthma (boys, doctor’s diagnosis, \( P = 0.04 \)), decreased functional vital capacity (FVC)-Z-score (both sexes, \( P = 0.02 \), largely driven by decreased FVC in girls), a trend toward lower forced expiratory volume in 1 s (FEV1)-Z-score at 6 yr (girls, \( P = 0.09 \)), and decreased FEV1/FVC Z-score in 14-yr-old girls (\( P = 0.05 \)). Whereas some other studies support an inverse relation between vitamin D status in pregnancy and offspring wheeze or asthma, others have found no associations, or U-shaped or direct associations, although many of these studies were of minor quality, e.g., in the diagnosis of childhood asthma (62, 63). Hopefully, ongoing RCTs like the Vitamin D Antenatal Asthma Reduction Trial (VDAART) (54) will provide high-quality, high-level evidence on this question.

Strengths of our study included the systematic review method without time limit in the search and the inclusion of human, animal, and laboratory data. Furthermore, a risk of bias analysis was performed.

Limitations included the paucity of high-evidence human data, which severely hampered conclusions for the clinical use of vitamin D. Extrapolation of rodent data to humans may be hampered by differences in lung development and maturation. The delayed surfactant synthesis and lack of prenatal alveolar phase in rodents provide suitable models for studies on lung diseases of the preterm but less suitable for studies on lung diseases of infants and children.

Conclusions on VDR expression were constrained by methodological issues on VDR detection by immunohistochemistry (69–71). New data show that the widely used VDR monoclonal antibody 9A7γ especially in rat tissue, not only binds to VDR, but also possesses nonspecific interactions with unidentified proteins (102, 103). On the basis of a parallel comparison of a large selection of VDR antibodies, the mouse monoclonal antibody D-6 possesses the highest specificity, sensitivity, and versatility. The antibody is capable of binding VDRs from human, monkey, pig, chicken, rat, and mouse and can be used for multiple immunoassays (102). Future studies on the VDR expression should use a more specific VDR antibody, e.g., D-6, which is highly sensitive, specific, and versatile.

Conclusion. Our systematic review on human data showed sparse evidence, allowing no conclusions on the potential role of vitamin D in the prevention or treatment of RDS or BPD in preterm neonates. Animal and laboratory data showed substantial evidence of multiple physiological actions through which vitamin D stimulates maturation of the fetal lung including ATII cell maturation and the alveolarization. These data give support to a hypothesis of vitamin D deficiency or insufficiency as a frequent, modifiable risk factor of RDS and BPD, which should be investigated in cohorts or case-control studies and tested in RCTs on pregnant women, especially with threatening preterm delivery, or in the preterm neonates themselves. Moreover, the effect of vitamin D on RDS or BPD in preterm neonates without vitamin D deficiency or insufficiency and potential adverse effects of high doses, including long-term outcomes as asthma, should be specifically addressed. Future experimental and human studies should aim to identify optimal time windows, vitamin D doses, and cut-off levels for 25(OH)D for the intervention against RDS, BPD, and later respiratory outcomes.

DISCLOSURES

No conflicts of interest, financial or otherwise, are declared by the authors.

AUTHOR CONTRIBUTIONS

Author contributions: S.L. and H.T.C. conception and design of research; S.L. and H.T.C. analyzed data; S.L. interpreted results of experiments; S.L. prepared figures; S.L., draft manuscript; S.L., G.L.S., S.S.B-N., and H.T.C. edited and revised manuscript; S.L., G.L.S., S.S.B-N., and H.T.C. approved final version of manuscript.

REFERENCES


VITAMIN D, FETAL LUNG, AND SURFACTANT


