Occasional review

Nitric oxide and lung disease

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Even five years ago few could have realised that the simple gas nitric oxide (NO) would be involved in the regulation of so many physiological functions and pathophysiological processes. There has been an explosion of information about nitric oxide which appears to be involved in an extraordinary range of functions including vascular regulation, neurotransmission, host defence, and cytotoxicity. An increasing volume of literature has shown the importance of nitric oxide in the regulation of various pulmonary functions and suggested its involvement in several pulmonary diseases. The purpose of this review is to highlight some of the recent advances in this rapidly moving field.

For over 20 years it has been recognised that the vasodilator responses to many agents are mediated by the release of a vasodilator substance from endothelial cells. The identity of endothelium derived relaxant factor (EDRF) remained elusive, largely because of its short half life, until 1987 when two groups independently suggested EDRF had the characteristics of nitric oxide. Many were surprised that such a simple molecule could account for all of the actions of EDRF, but extensive investigations in many species have now provided compelling supportive evidence. Parallel investigations established that nitric oxide was also important in macrophage mediated cytotoxicity.

Formation of nitric oxide

Nitric oxide is a highly reactive radical (more accurately designated NO) formed from the semi-essential amino acid L-arginine by the action of an enzyme nitric oxide synthase (NOS) (fig 1). The enzyme catalysing the reaction is stereospecific as L-arginine is a substrate but D-arginine is not. Nitric oxide is produced in neuronal and non-neuronal tissue as a product of the action of NOS catalysing the conversion of L-arginine to L-citrulline. Conversion of [H]arginine to [H]citrulline can be monitored in order to measure NOS activity as [H]citrulline is formed stoichiometrically with nitric oxide and is measured more easily than the short lived nitric oxide. Nitric oxide is rapidly oxidised to nitrite (NO2) which can also be used to monitor nitric oxide formation. The nitrogen of nitric oxide is derived from the terminal guanidino nitrogen of L-arginine and the oxygen is derived from molecular oxygen.

Nitric oxide is formed in the lungs and can be detected in the expired air of several species, including humans. Nitrite and S-nitrosothiols have also been detected in bronchoalveolar lavage fluid of normal human volunteers. The source of nitric oxide is not certain but it may be derived from the alveoli (epithelium or endothelium), from macrophages, or from airway epithelium.

NITRIC OXIDE SYNTHASES (NOS)

Several species of NOS have now been characterised and several distinct NOS genes have been identified. NOS exists as constitutive forms (cNOS) which are basally expressed in endothelial, neuronal and other cells and are Ca2+-calmodulin dependent. The purified enzyme migrates as a single 150 kDa band on gel electrophoresis, and the native enzyme appear to be a monomer. The messenger RNA is colocalised with NADPH diaphorase, and neuronal NOS and NADPH diaphorase appear to be one and the same enzyme. The histochemical location of NADPH diaphorase by the formation of blue formazans from tetrazolium can therefore be used to localise neuronal NOS. A form of cNOS distinct from the neuronal form is localised to vascular endothelial cells. cNOS from brain and endothelial cells have been cloned and these are distinct enzymes with about 60% homology, which are also homologous to cytochrome P-450.

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**Figure 1** Nitric oxide (NO) formation by NO synthase involves conversion of L-arginine to L-citrulline. Several cofactors are necessary for enzyme activity, including flavones (FAD, FMN) and tetrahydrobiopterin (BH4). NO synthase is inhibited competitively by arginine derivatives N6-monomethyl-L-arginine (L-NMMA), N7-nitroarginine methyl ester (L-NAME), and N7-nitrosoarginine (L-NOARG). Nitric oxide itself may be oxidised to nitrite (NO2), nitrate (NO3), and peroxynitrite (ONOO-) ions.
Nitric oxide activates soluble guanylyl cyclase (sGC) in target cells, resulting in an increase in cyclic guanosine 3′5′-monophosphate (cGMP) and smooth muscle relaxation.

The development of drugs that inhibit NOS has been the most important strategy in determining the physiological and pathophysiological roles of nitric oxide. Analogues of L-arginine act as false substrates for the enzyme and therefore block the formation of endogenous nitric oxide. This blockade can be overcome by adding back L-arginine, but not by adding D-arginine which is not a substrate for the enzyme. Several arginine analogues have been developed, including Nω-nitro-L-arginine (L-NMMA), Nω-nitro-L-arginine methyl ester (L-NAME), and Nε-iminoethyl-L-ornithine (L-NIO), all of which are extremely useful in revealing the role of endogenous nitric oxide in various processes. These inhibitors are stereospecific, the corresponding analogues of D-arginine being inactive. In addition, L-NMMA and L-NIO have been shown to inhibit L-arginine transport. Recently, hydroxyxycobalamin has been used as a tool to distinguish between the source of nitric oxide as it has a differential action in blocking responses to endothelium-derived nitric oxide in rat aortic rings but not neurogenic nitric oxide in the rat anococcygeus muscle. Alternatively, the NOS inhibitor L-NN-nitroarginine p-nitroanilide (L-NAPNA) appears to be a selective inhibitor of NOS in the brain, as shown by its antinociceptive activity in mice, with little effect on the endothelium-dependent relaxation of blood vessels. Selective inhibitors of iNOS have recently been reported, and aminoguanidine has a 10–100-fold selectivity for iNOS compared with cNOS and may be the basis for potentially important therapeutic agents in the future.

CELLULAR EFFECTS OF NITRIC OXIDE

Nitric oxide activates soluble guanylyl cyclase after binding to its haem moiety to initiate a three-dimensional conformation change in the shape of the enzyme which increases its activity and consequently the production of cyclic guanosine 3′5′-monophosphate (cGMP). The rise in cGMP results in relaxation of smooth muscle, but the mechanism by which this happens is unknown. Many possibilities have been suggested including the inhibition of inositol trisphosphate (IP3), inhibition of a cAMP phosphodiesterase, dephosphorylation of the light chain of myosin, activation of protein kinases, stimulation of membrane Ca2+ ATPase, opening of K+ channels, increased sequestration of cytosolic Ca2+, and inhibition of Ca2+ influx. This increase in cGMP underlies many of the neural and cardiovascular actions of nitric oxide. There have also been recent reports of effects of nitric oxide that are not dependent on cGMP. However, the second messenger systems involved in nitric oxide induced cell death are not as well understood. It had long been recognised that directly acting nitrovasodilators such as glycyltrinitrate are metabolised to nitric oxide within smooth muscle, endothelium, and plasma, and others like sodium nitroprusside liberate nitric oxide spontaneously. It has also been suggested that nitric oxide may form an intermediary complex with thiols such as cysteine, and it has been suggested that S-nitrosocysteine more
closely mimics the properties of EDRF than does nitric oxide itself. The formation of these sulphhydryl complexes may prolong the half-life of nitric oxide released from cells.

The fact that a simple gas such as nitric oxide is a physiological messenger has raised the possibility that other gases could have similar functions. Carbon monoxide is also an activator of guanylyl cyclase and is formed by the action of heme oxygenase. Localisation of the mRNA for the constitutive form of heme oxygenase has been found throughout the brain, and the localisation is similar to that for soluble guanylyl cyclase mRNA, suggesting that carbon monoxide may function as a neurotransmitter.

Role as a vasodilator

The observation that EDRF is nitric oxide immediately suggested that it may play a part in the regulation of the pulmonary circulation and this has been extensively investigated. NOS inhibitors reduce the vasodilator response to acetylcholine in animal and human pulmonary vessels in vitro and against endothelin-3 in rat pulmonary vessels. Endogenous nitric oxide appears to act as a braking mechanism against pulmonary vasoconstriction. Release of nitric oxide from endothelial cells in the pulmonary circulation appears to counteract hypoxic vasoconstriction and nitric oxide release is apparently decreased in hypoxia. In chronic experimental hypoxia there is a reduced response to nitrovasodilators which appears to result from impaired function of soluble guanylyl cyclase. There is circumstantial evidence that release of nitric oxide from pulmonary vessels may be impaired in patients with chronic obstructive pulmonary disease (COPD) and cystic fibrosis. Impaired endothelium dependent relaxation has also been observed in isolated pulmonary arteries obtained from patients undergoing heart-lung transplantation for end stage chronic lung disease, including cystic fibrosis.

Since nitric oxide is a potent pulmonary vasodilator, inhalation of nitric oxide might be effective as a selective pulmonary vasodilator in view of its short half life. Rapid combination of nitric oxide with haemoglobin contained in red blood cells would rapidly inactivate any nitric oxide reaching the systemic circulation, thereby limiting vasodilatation to pulmonary vessels. Inhaled nitric oxide (5–80 ppm) has been shown dose dependently to inhibit pulmonary vasoconstriction induced by either hypoxia or by infusion of a thromboxane analogue in conscious spontaneously breathing lambs and in normal volunteers with induced hypoxia without causing systemic hypotension. Inhaled nitric oxide reverses hypoxic pulmonary vasoconstriction in sheep without impairing gas exchange. Inhalation of nitric oxide (40 ppm) has also been shown to cause selective pulmonary vasodilatation in patients with pulmonary hypertension and in neonates with persistent pulmonary hypertension nitric oxide produces a significant improvement in oxygen exchange and arterial oxygen tension. Inhalation of nitric oxide has also been reported to cause selective pulmonary vasodilatation in patients with COPD. Inhalation of nitric oxide (18 ppm) has recently been reported to reduce pulmonary artery pressure in patients with severe adult respiratory distress syndrome (ARDS) and increase oxygenation by improving ventilation perfusion matching without systemic vasodilatation. Continuous nitric oxide inhalation was effective over a period of 3–53 days without evidence of tachyphylaxis, indicating that this might be an effective therapy for this condition although controlled trials are now required.

Nitric oxide is also a potent bronchial vasodilator in animal airways. Cigarette smoke contains a high concentration of nitric oxide which may account for the increase in airway blood flow after exposure of airways to cigarette smoke.

Role as a bronchodilator

Nitrovasodilators such as glyceryl trinitrate and sodium nitroprusside also relax airway smooth muscle in vitro, resulting in an increase in soluble guanylyl cyclase activity and an increase in cGMP. It is therefore to be expected that nitric oxide may act as a bronchodilator and this has been shown in canine airways in vitro. The effect of nitric oxide on isolated epithelium intact guinea pig tracheal strips contracted with carbachol was, however, considerably less than on epithelium denuded strips. The mechanism by which the epithelial layer reduces the effects of nitric oxide is unclear, but the epithelium may act as a metabolic or diffusional barrier for nitric oxide. High concentrations of inhaled nitric oxide (80 ppm) reduce the bronchoconstrictor effect of nebulised methacholine in anaesthetised rabbits. In anaesthetised guinea pigs methacholine induced bronchoconstriction is reduced, albeit transiently, by inhaled nitric oxide in a concentration dependent manner from 5 ppm to 300 ppm and by the nitric oxide releasing compound S-nitroso-N-acetylpenicillamine (SNAP). In addition, a high concentration of nitric oxide (300 ppm) causes a small degree of baseline bronchodilatation. There is no evidence for tolerance after prolonged administration and the bronchodilator effect of nitric oxide is additive with a β adrenoceptor agonist. Histamine release from bovine lung is also reduced by nitrovasodilators, probably by alterations in guanylyl cyclase activity, suggesting that nitric oxide may have mast cell stabilising effects.

Nitric oxide inhalation (at estimated doses of 0.5 mg and 10 mg) has no effect on airway resistance in normal subjects, although it is associated with a small fall in oxygen saturation, presumably as a result of changes in ventilation perfusion ratio. Inhaled nitric oxide at a concentration of 80 ppm similarly has no effect in normal individuals, but has a
small and inconsistent bronchodilator effect in asthmatic patients.\textsuperscript{66} This raises the possibility that nitric oxide inhalation or nitric oxide releasing compounds might have some therapeutic potential as alternative bronchodilators in obstructive airways diseases. An advantage of inhaled nitric oxide would be its lack of systemic effects since it would be rapidly inactivated by haemoglobin. There are, however, potential dangers of inhaling nitric oxide\textsuperscript{67} since, in the presence of oxygen, it is oxidised to nitrate and thence to nitrous and nitric acids\textsuperscript{56} which may increase airway responsiveness and, in high concentration, might cause pulmonary oedema.\textsuperscript{69,70} The interaction between nitric oxide and superoxide anions may lead to the formation of peroxynitrite (ONOO\textsuperscript{−}) that may generate tissue damaging hydroxyl radicals.\textsuperscript{68,71} There is also some evidence that high concentrations of nitric oxide may have effects on DNA and be both genotoxic and cytopoietic.\textsuperscript{72} Nitric oxide has been detected in the exhaled air of humans and is presumably derived from pulmonary capillary endothelial cells.\textsuperscript{73}

**Role as a non-adrenergic non-cholinergic (NANC) transmitter**

There is increasing evidence that nitric oxide may function as a neurotransmitter of inhibitory non-adrenergic non-cholinergic (iNANC) nerves, and "nitrergic" or "nitroxidergic" neurotransmission has been shown in the gut, bladder, and reproductive organs (fig 3).\textsuperscript{74} There is convincing evidence that nitric oxide is released from nerves themselves, since a particular form of cNOS has been localised to peripheral nerves\textsuperscript{74} and is activated by calcium entry when the nerve is depolarised. Nitric oxide accounts for approximately half of the iNANC (bronchodilator) response in guinea pig trachea in vitro.\textsuperscript{75,76} In pig tracheal smooth muscle there is a prominent iNANC response which is completely inhibited by NOS inhibition and reversed by L-arginine in a stereospecific manner.\textsuperscript{77,78} Similar results have been reported in cat and horse airways\textsuperscript{79,80} although a previous report in feline airways found no effect of NOS inhibitors.\textsuperscript{81} This indicates that the NANC response is evoked by release of nitric oxide and provides evidence that it is the principal iNANC neurotransmitter. Nitric oxide also appears to account for the bronchodilator NANC response in human airways in vitro in central and peripheral airways\textsuperscript{82-85} and, in contrast to guinea pig trachea, the neuropeptide vasoactive intestinal peptide (VIP) appears to play little or no part in this response.\textsuperscript{86} A similar mechanism seems to be responsible for iNANC responses in central and peripheral airways, as evidenced by similar kinetics and frequency dependency.\textsuperscript{86} The cellular source of nitric oxide is not completely certain, but it is most likely to be derived from intrinsic nerves within the airway rather than another transmitter substance inducing the release of nitric oxide from endothelial, epithelial, or smooth muscle cells. Epithelial removal has no effect on the iNANC response in guinea pig airways.\textsuperscript{87} However, nitric oxide is not stored in vesicles and electron microscopy studies have shown that NOS, at least in the myenteric plexus, is located in the cytoplasm. Immunocytochemical staining for neuronal cNOS and NADPH diaphorase shows localisation to nerves of guinea pig, ferret, and human airways.\textsuperscript{86-90} NOS immunoreactive nerves appear to supply airway vessels, airway smooth muscle, and submucosal glands. NOS is colocalised with VIP in these species. NOS immunoreactive neurones are found in parasympathetic ganglia of human but not guinea pig airways, and also in sympathetic and sensory ganglia supplying the airways.\textsuperscript{88,89} NOS immunoreactive nerves are more prominent in proximal than in distal airways.

Nitric oxide is probably synthesised on demand in a neurone and its release may involve simple diffusion. Alternatively, it can spontaneously combine with naturally occurring thiols such as cysteine in aqueous media to form nitrosothiols such as L-cysteine-NO\textsubscript{2} at acidic pH.\textsuperscript{91} Once formed, nitrosothiols may be stabilised at the low pH in secretory vesicles. If nitrosothiols are stored in secretory vesicles then release would expose these compounds to the higher pH of the extracellular fluid, with rapid breakdown to release nitric oxide. Another possibility is that nitric oxide is released by electrical field stimulation from glial cells that have been shown to contain L-arginine\textsuperscript{92} and voltage gated sodium channels, which could account for a tetrodotoxin sensitive iNANC response.\textsuperscript{93} In the gastrointestinal tract there is some evidence that VIP stimulates the release of nitric oxide from smooth muscle cells so that it acts as an indirect transmitter of relaxation.\textsuperscript{94,95} but this is unlikely in airways since VIP induced bronchodilatation is not reduced by NOS inhibitors.\textsuperscript{76,84,85}

![Inhibitory non-adrenergic non-cholinergic nerves in airways utilise nitric oxide (NO) as a neurotransmitter. Depolarisation of the nerve results in influx of calcium ions (Ca\textsuperscript{2+}) and activation of nitric oxide synthase. Nitric oxide diffuses from the nerve to activate soluble guanylyl cyclase in airway smooth muscle cells, resulting in increased cyclic GMP (cGMP) and relaxation. Nitric oxide may form an intermediate and more stable complex with cysteine.](image-url)
NOS inhibition markedly potentiates cholinergic neural responses in both human and guinea pig airways, suggesting that endogenous nitric oxide modulates cholinergic neural responses by acting as a functional antagonist to acetylcholine at airway smooth muscle. However, this effect is absent in human airways when the pulse width of the electrical field stimulation is reduced to 0-1 ms which suggests that iNANC responses are less evident under these conditions, so the chance of functional interactions between the NANC relaxation response and the cholinergic contractile response is reduced. There is also evidence to suggest that nitric oxide may modulate reflex bronchoconstriction in vivo. Endogenous nitric oxide modulates the atropine-sensitive component of bradykinin induced bronchoconstriction in the guinea pig. In addition, endogenous nitric oxide released in association with vagal nerve stimulation regulates the magnitude of NANC neurogenic bronchoconstriction (due to tachykinin release from sensory nerves) in guinea pigs in vivo.

Nitric oxide may be released from parasympathetic nerves as a cotransmitter with acetylcholine, although there is no direct evidence for this yet. In the gastrointestinal tract NOS immunoreactivity is colocalised with VIP, which may also be localised to cholinergic nerves, and NOS is colocalised with VIP in vascular nerves of guinea pigs. Some evidence that argues against the colocalisation of NOS in human airway cholinergic nerves is the observation that bronchodilator neural responses are virtually absent in extrinsically denervated airways obtained from heart-lung transplant recipients undergoing a second transplantation, while cholinergic neural responses are preserved. In an innervated guinea pig tracheal tube preparation a bronchodilator response to postganglionic nerve stimulation (via electrical field stimulation) is present which is reduced by NOS inhibitors as expected, but no bronchodilator response is observed after stimulation of the vagus nerve although both types of neural stimulation elicit a cholinergic bronchoconstrictor response. This suggests that the neurones which release nitric oxide are separate from cholinergic pathways and may arise from outside the airways. There is some evidence that the nerve cell bodies may be localised in the vicinity of the oesophagus.

Since bronchodilator NANC nerves are the only neural bronchodilator pathway in human airways, it is possible that there may be a defect in the function of these nerves in diseased airways. iNANC responses in tissues from patients with mild asthma were not altered, but the responses were significantly reduced in tissues from patients with cystic fibrosis compared with iNANC responses in donor tissue. This defect in iNANC function may be because nitric oxide released from either nerves or cells is degraded by inflammatory mediators. Airway inflammation may be associated with release of superoxide anions from activated inflammatory cells resulting in increased breakdown of nitric oxide. Alternatively, this could be the result of a malfunction at the level of the NOS or cyclic GMP activation.

Nitric oxide may also be involved in neurogenic vasodilator responses in the pulmonary circulation. Electrical field stimulation causes a vasodilator response in pulmonary vessels in vitro which is partly due to the release of nitric oxide from endothelial cells via ATP release from sympathetic nerves. Nitric oxide may also act as a neurotransmitter of vasodilator NANC responses in guinea pig pulmonary vessels. Selective destruction of adrenergic nerves with 6-hydroxydopamine has no effect on the NANC vasodilator response, suggesting that nitric oxide is likely to be released from some other type of nerve. Endothelial removal is similarly ineffective, indicating that nitric oxide is not of endothelial cell origin. Interestingly, superoxide anions appear to inhibit the NANC vasodilator response which may suggest that in inflammation neurogenic vasodilatation may be impaired.

NOS inhibition enhances adrenergic neural vasocostrictor responses in pulmonary vessels. This indicates that, as in the airways, endogenous nitric oxide appears to act as a functional antagonist or "braking" mechanism to vasoconstrictor responses in pulmonary vessels. Although the evidence is now convincing that endogenous nitric oxide mediates NANC relaxant effects, there are still some doubts as to whether nitric oxide itself acts as the transmitter or whether some more stable intermediate complex is formed. As for endothelial responses, nitrocysteine has been suggested as an intermediary compound and closely mimics NANC responses in the gut, and several S-nitrosothiols relax smooth muscle directly. Hydroquinone blocks relaxation induced by nitric oxide but not NANC nerve stimulation or other nitrovasodilators in guinea pig trachea in vitro, and the mechanism of action appears to be by free radical scavenging rather than superoxide anion generation. This differential effect may indicate that the NANC transmitter is not free nitric oxide but a nitric oxide containing or nitric oxide generating substance.

Role as an inflammatory mediator
Endothelial cells and nerves are not the only source of nitric oxide in airways and several types of inflammatory and structural cells have now been found to produce nitric oxide.

MACROPHAGES
There is convincing evidence that macrophages, including alveolar macrophages of some species, may synthesise nitric oxide after exposure to various cytokines and to endotoxin, and that nitric oxide is
important in host defence. Macrophages can perform many cytotoxic activities which include inhibition of mitochondrial respiration, aconitase activity and DNA synthesis, and are thought to be due to the inhibition of iron containing enzymes in target cells. Those activities, as well as NO$_2^-$ and NO$_3^-$ generation, are blocked by inhibitors of NOS. Macrophages express an inducible form of NOS that has recently been cloned. The induction of iNOS in alveolar macrophages can be blocked by glucocorticosteroids. Interleukin 10 also inhibits iNOS induced by interferon $\gamma$ in murine macrophages. Nitric oxide itself may act as a feedback inhibitor of iNOS in these cells, providing a mechanism to limit induced nitric oxide generation. Human alveolar macrophages, in contrast to rat and murine macrophages, do not appear to produce nitric oxide in vitro under the same experimental conditions, but it is possible that a suitable combination of cytokines and cofactors would allow the production of nitric oxide from these cells.

**MAST CELLS**

Rat serosal mast cells have been reported to synthesise nitric oxide from L-arginine, but whether nitric oxide is produced by activated human lung mast cells is not yet certain.

**SMOOTH MUSCLE CELLS**

Other cells also have an inducible form of NOS, including endothelial cells, neutrophils and vascular smooth muscle cells. A combination of cytokines (IL-1$\beta$, TNF$\alpha$, and IFN$\gamma$) and lipopolysaccharide potently induce iNOS in cultured rat pulmonary artery smooth muscle and this effect is blocked by an inhibitor of protein synthesis. Treatment of rats with in vivo lipopolysaccharide induced iNOS after six hours in the lung and induced the expression of mRNA for iNOS, but not cNOS, in rat lung. Induction of NOS by cytokines and lipopolysaccharide appears to result in much greater amounts of nitric oxide than is possible with cNOS. There is compelling evidence that nitric oxide in septic shock is a major contributor to the cardiovascular collapse.

**EPITHELIAL CELLS**

Airway epithelial cells are also a source of nitric oxide and this could be induced by exposure to cytokines such as TNF$\alpha$ and IFN$\gamma$ in the airway, although nitric oxide does not appear to be "epithelium derived relaxing factor." In guinea pigs inhaled L-NAME increases the bronchoconstrictor response to histamine in vivo and intraluminal application in a tracheal tube preparation also enhances the contractile response to histamine in vitro. This effect is mimicked by removal of airway epithelium, suggesting that airway epithelium releases nitric oxide which counteracts the bronchoconstrictor effect of spasmogens. Cultured bovine epithelial cells metabolise L-arginine to L-citrulline, an effect blocked by NOS inhibitors, indicating that airway epithelial cells have the capacity to produce nitric oxide. A cultured human epithelial cell line also produces nitrite when incubated with L-arginine, indicating the capacity of human airway epithelial cells to produce nitric oxide. Immunocytochemical staining has demonstrated the expression of iNOS in human and rat airway epithelial cells and there is evidence for increased iNOS expression in epithelial cells of bronchial biopsy specimens from asthmatic patients compared with normal controls, whereas cNOS is not expressed in either group. This suggests that iNOS may be induced by cytokines produced in asthmatic inflammation, including IFN$\gamma$, TNF$\alpha$, and IL-1$\beta$. Endogenous nitric oxide may be an important modulator of mucociliary clearance and is reported to increase ciliary beat frequency in bovine ciliated epithelial cells.

**FIBROBLASTS**

Rat lung fibroblasts are capable of producing nitric oxide after stimulation with IFN$\gamma$, and this effect is enhanced by lipopolysaccharide and IL-1$\beta$. The role of endogenous nitric oxide in fibrogenesis is not yet certain, however.

**IMMUNOLOGICAL EFFECTS**

Alveolar macrophages have a suppressive effect on T lymphocyte proliferation both in rats and humans. There is some evidence that at least part of this immunosuppressive effect is mediated by nitric oxide in rats. This suggests that endogenous nitric oxide may have an immunomodulatory role in the airways.

**ROLE IN LUNG INFLAMMATION**

Endogenous nitric oxide may be a double edged sword. Nitric oxide produced in small amounts locally by cNOS activation may be beneficial in relaxing airway smooth muscle in airways, but may have deleterious effects when produced in much higher concentrations from iNOS. Nitric oxide is a potent vasodilator and might contribute to the hyperaemia of asthmatic airways. This may also increase exudation of plasma from leaky post capillary venules in the airways. Indeed, inhibition of endogenous nitric oxide production significantly reduces plasma exudation and inflammation, both in skin and in airways. High concentrations of nitric oxide may have cytotoxic effects in the airway and could conceivably contribute to the epithelial shedding observed in asthmatic patients. Ozone inhalation increases nitric oxide release from rat alveolar macrophages and this could contribute to the damaging effect of ozone on airway epithelial cells. Nitric oxide may be involved in acute and chronic lung injury. L-NAME protects rats against lung injury induced by injection of immune complexes and the high levels of nitrite recovered from bronchoalveolar lavage fluid of these animals provide further evidence for the involvement of nitric oxide in this process.
EFFECT OF CORTICOSTEROIDS
Corticosteroids potently inhibit the expression of the inducible, form of NOS26 27 and this may contribute to their anti-inflammatory action since massive nitric oxide formation may be detrimental as in the case of endotoxin shock. If NOS is induced in airway epithelial cells in asthma as a result of exposure to cytokines released from inflammatory cells, inhaled steroids may act to reduce the formation of nitric oxide and thus downregulate the vascular components of the inflammatory response. Steroids would not be expected to affect the release of nitric oxide from bronchodilator nerves since the neural constitutive form of the enzyme is not steroid sensitive.

CLINICAL PROSPECTS
It is clear that nitric oxide may have a very important regulatory role in pulmonary function and may be implicated in the pathophysiology of several lung diseases.

PULMONARY VASCULAR DISEASE
Inhaled nitric oxide is clearly effective in counteracting hypoxic pulmonary vasoconstriction and has the major advantage that there are no systemic vasodilator effects, which are the major limitation to the use of other vasodilators (such as prostacyclin) for the treatment of pulmonary hypertension. Furthermore, inhaled nitric oxide selectively dilates the pulmonary vasculature to which it is delivered thereby improving ventilation perfusion matching, whereas systemically administered vasodilators may increase ventilation perfusion mismatch. Inhaled nitric oxide is also effective in improving the oxygenation in ARDS.56 These preliminary clinical studies suggest that nitric oxide may be useful as an acute selective pulmonary vasodilator in critical care and may be administered safely over a period of several days.

BRONCHODILATATION
Recent studies with nitric oxide have revived interest in nitrovasodilators as alternative bronchodilators which work by a different molecular pathway from β agonists and theophylline. Previous studies of such drugs in asthma have not been impressive48 but new nitric oxide donors such as S-nitrosothiols may have advantages.49 In particular, inhalation of nitric oxide gas or nitric oxide releasing compounds may present a useful approach in the treatment of diseases such as asthma, and rapid inactivation of inhaled nitric oxide by red blood cells will prevent any systemic haemodynamic side effects. Nitric oxide itself, however, may not reach airway smooth muscle in adequate concentrations unless large amounts are inhaled when the risk of vasodilator effects is increased. Inhaled nitric oxide would probably be suitable only as an acute bronchodilator in severe asthma, but there is likely to be no advantage (and several disadvantages) compared with nebulised β agonists. Studies of the effects of inhaled nitric oxide in human airway are limited, but inhalation of >15 000 ppb in healthy subjects causes a small fall in arterial oxygen tension and an increase in airways resistance at concentrations of >20 000 ppb.49 Another approach would be to enhance the release of neuronal nitric oxide which should give selective bronchodilatation.

TOXICITY
High concentrations of nitric oxide are likely to have toxic effects resulting from the formation of nitrites and peroxy nitrites radicals which may have cytotoxic and genotoxic effects.71 72 Formation of nitrite may lead to methaemoglobinemia. Nitric oxide is a recognised air pollutant derived from car exhaust and from domestic gas cookers, but little is known about its effects on airway function after long term exposure. It is a major constituent of the gaseous phase of cigarette smoke, derived from nitrates in tobacco, and concentrations of >50 ppm are inhaled in mainstream smoke.56 The nitric oxide content is particularly high in French and American blends because of the high nitrate content.

L-ARGININE
Since L-arginine is the necessary precursor for nitric oxide formation, it is possible that this amino acid may increase endogenous nitric oxide production, particularly when there is a possibility that endogenous L-arginine is the limiting factor. L-Arginine infusion has been studied in patients with primary pulmonary hypertension, without any effect, however.49

NOS INHIBITORS
The production of large amounts of nitric oxide by induction of NOS in response to cytokines may be deleterious and contribute to the inflammatory response in both airways and in lung parenchyma. This has suggested that NOS inhibitors might have therapeutic potential. However, non-selective NOS inhibitors such as L-NMMA and L-NAME lead to hypertension by blocking cNOS in endothelial cells.49 What is required is a selective inhibitor of iNOS and this is achieved by corticosteroids, as discussed above. A selective iNOS inhibitor might have the beneficial anti-inflammatory effects of steroids, without the systemic side effects which limit their usefulness when systemic administration is required. There is evidence that some NOS inhibitors may have selectivity for particular types of NOS; aminoguanidine appears to have a 10–100-fold selectivity for inhibition of iNOS compared with cNOS indicating that selective inhibition is a realistic possibility in the future. Cloning of NOS has revealed several distinct enzyme genes, each of which may have subtypes resulting from differential splicing, and this
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raises the prospect that even more selective NOS inhibitors may be developed.

Conclusions
Endogenous nitric oxide may have a key role in many physiological and pathophysiological events in the lung. It appears to be important in neural bronchodilator and vasodilator mechanisms, in the regulation of airway and pulmonary blood flow, and in immune defence. When produced in low concentrations it appears to have a generally beneficial role, but when produced in large amounts by inducible NOS it may result in increased inflammatory responses and tissue damage, suggesting that nitric oxide may be involved in the pathophysiology of several pulmonary diseases. This raises the prospects for new approaches to the treatment of airway inflammation, ARDS, and pulmonary vascular disease in the future.

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