Sildenafil: from angina to erectile dysfunction to pulmonary hypertension and beyond

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Abstract | In less than 20 years, the first selective type 5 phosphodiesterase inhibitor, sildenafil, has evolved from a potential anti-angina drug to an on-demand oral treatment for erectile dysfunction (Viagra), and more recently to a new orally active treatment for pulmonary hypertension (Revatio). Here we describe the key milestones in the development of sildenafil for these diverse medical conditions, discuss the advances in science and clinical medicine that have accompanied this journey and consider possible future indications for this versatile drug.

However, the therapeutic potential of nitrates is limited by the rapid induction of tachyphylaxis with prolonged administration. This limitation led some scientists at Pfizer to consider alternative approaches to modulate NO signalling.

PDE5 is selected as a target

Although the precise mechanism of tolerance to nitrates is not clear, any treatment that does not directly increase NO levels might circumvent this problem. The scientific team at Pfizer therefore proposed that a downstream target in the NO/cGMP pathway could be modulated. Cyclic nucleotides (cAMP and cGMP) are degraded by cyclic nucleotide phosphodiesterases (PDEs). Five subtypes of PDEs were identified. PDE3 and PDE4 specifically catalyse the breakdown of cAMP, whereas PDE1 and PDE2 catalyse the breakdown of both cAMP and cGMP. The fifth member of this group, PDE5, is present in the smooth muscle of the systemic vasculature, and in platelets. Studies by Corbin et al. revealed that the regulatory domain in the amino-terminal portion of PDE5 contains a phosphorylation site (serine 92), two allosteric cGMP-binding sites, a and b, and at least a portion of the dimerization domain. The catalytic domain in the carboxy-terminal portion of the protein contains the two Zn+2-binding motifs A and B, and a cGMP substrate-binding site (Fig. 2). In 1996 Pfizer formed a project team of scientists at Sandwich, with the aim of developing a selective inhibitor of PDE5 and evaluating its preclinical pharmacology (timeline). Eventually the team succeeded in synthesizing novel pyrazolopyrimidines that were highly potent inhibitors of PDE5. A compound...
designated chemically as 1-[[3-(6,7-dihydro-1-methyl-7-oxo-3-propyl-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-ethoxyphenyl]sulphonyl]-4-methylpiperazine, was chosen for further profiling (FIGS 3, 4). It was initially named UK-92,480, but is now better known as sildenafil, of which the citrate salt is marketed under the trade name Viagra. The compound was demonstrated to have very good potency (an IC$_{50}$ of 3.5 nM against PDE5 derived from human platelets), and excellent selectivity over PDEs 1–4 (REF. 7). Moreover, in preclinical studies, this compound displayed vasodilatory effects, abrogated platelet aggregation, and inhibited thrombus reformation in a damaged carotid artery (modified Folt's model). Therefore, in 1989 UK-92,480 was selected to undergo further development and eventually entered clinical trials in 1991 (REF. 9).

**Clinical development programme for angina**

In 1991 studies in healthy volunteers commenced, in which single doses of UK-92,480 up to 200 mg were administered. UK-92,480 had a relatively short plasma half-life (~4 hours)$^{10}$. The doses were generally well tolerated and, at moderate and higher doses, modest reductions in systemic blood pressure and side effects associated with vasodilatation (such as headache and flushing) were observed. At doses of 150 mg and higher, transient disturbances in colour vision perception were reported (later in the development programme, these were demonstrated to be caused by a weak inhibitory effect of UK-92,480 on PDE6 in photoreceptors)$^{11}$.

Subsequently, in a study involving intravenous administration of UK-92,480, a moderate vasodilatory effect was observed in patients with angina. A later study in healthy volunteers that was designed to investigate a potential pharmacodynamic interaction between UK-92,480 and glyceryl trinitrate (GTN) demonstrated that UK-92,480 augmented the vasodilatory and antihypertensive effects of GTN$^{12}$. Although this finding was positive in confirming the mode of action of sildenafil in a clinical setting, it was also potentially problematic. UK-92,480 would have to be contraindicated in patients taking nitrates or, alternatively, a safe way of co-administering the two different agents needed to be developed and demonstrated.

In 1992, several multiple-dose, healthy volunteer studies were undertaken to investigate the pharmacokinetics, pharmacodynamics and tolerance of UK-92,480. When administered at doses of up to 75 mg three times per day for 10 consecutive days, some volunteers reported headaches, flushing, indigestion and muscle aches. Some volunteers also reported penile erections as a side effect. Initially this was not considered to be of major significance, because the volunteers were reporting these effects after a mere several days of UK-92,480 administration. Therefore, cardiovascular indications remained the primary focus of ongoing clinical investigations during 1992 and 1993. However, by mid-1993, UK-92,480 was looking less promising as a new treatment for angina pectoris. The relatively short half-life indicated that treatment would have to be administered at least three times per day for the chronic treatment of...
thoughts to erectile dysfunction

During the 1980s, advances were made in the recognition and treatment of erectile dysfunction (ED). Prior to then, many clinicians considered ED to be either a relatively trivial issue and/or a condition that was predominantly attributable to psychological causes. However, urologists interested in ED had started treating their patients with intracavernosal injections of vasodilating drugs (for example, papaverine and prostaglandin E1) that functioned by modulating levels of cAMP. This pharmacological approach worked for many patients, but drawbacks included the invasive nature of the treatment, the induction of an ‘artificial’ erection that would often last long after intercourse had finished, with the added risks of local bleeding, bruising and priapism (a prolonged painful erection lasting longer than 6 hours). There was no doubt that an effective oral agent would be a major breakthrough in the treatment of this distressing condition, although at the time few researchers and clinicians in the field thought this was possible.

The decision to undertake pilot studies with sildenafil in ED was supported by the observation that penile erections were a common side effect in the multiple-dose sildenafil Phase I study. Furthermore, emerging data implicated NO as a key mediator of the neural and haemodynamic effects that lead to penile erection in men. In particular, in the early 1990s, Ignarro and others reported that NO is the neurotransmitter that is released from cavernous nerves during sexual stimulation. The NO diffuses into vascular smooth muscle cells of the penis, stimulating the production of cGMP and leading to corpus cavernosum smooth muscle relaxation, vasocongestion, veno-occlusion (by constriction of the venous outflow from the penis against the tunica albuginea) and, ultimately, erection13–15. Local neural production of high quantities of NO in the presence of sexual stimulation enables selective vasodilatation of the penile vasculature. The Pfizer team postulated that the administration of an inhibitor of cGMP breakdown would enhance and prolong the vasodilatory response, but only during sexual stimulation. The prospect of an oral agent that could work naturally with sexual stimulation to facilitate and maintain erections, without causing excessive vasodilatation in the systemic vasculature, was indeed an exciting prospect.

Clinical studies in erectile dysfunction

In late 1993 the first clinical study in patients with ED was undertaken using a very novel trial design. In order to overcome the practical problem of confirming an erectogenic effect while allowing the patients to be sexually stimulated, a Rigiscan device that contains two loops that are placed round the base and tip of the penis was used to monitor and record the girth and hardness of the penis during sexual stimulation. In order to maximize the chances of detecting a signal, the patients selected for the first two studies were relatively healthy men with ED but no clinically apparent cardiovascular diseases or other underlying risk factors. By mid-1994, after completion of two separate clinical studies, single doses of sildenafil were shown to enhance erectile responses to sexual stimulation and to be well tolerated16; moreover, a clear dose–response relationship was observed17.

After this initial success, a full development programme was designed. Again, novel clinical trial designs (including the development of a new instrument, the International Index of Erectile Function17) were used. Sildenafil was effective in almost all types of patients with ED18–21, including those with diabetes mellitus22,23, cardiovascular disease24, multiple sclerosis25, spinal cord injury26 and even in patients who had undergone radical prostatectomy27. Efficacy response rates of 70% or higher were consistently observed in clinical trials, although response rates were somewhat lower in post-prostatectomy patients (~40–50%) and those with diabetes mellitus (~60%). Sildenafil was very well tolerated when taken on demand (usually about twice per week) and muscle aches were very rarely reported in the Phase IIb/III programme27.

During and after the full development programme, a further six families of PDE (PDEs 6–11) were identified (Table 2). The activity of sildenafil was assessed against representative members of all of these PDE families. Sildenafil exhibited excellent selectivity over PDEs 7–11, but only about tenfold selectivity for PDE5 versus PDE628. Subsequent in vitro and in vivo investigations confirmed that the transient visual side effects observed with sildenafil treatment were almost certainly due to weak inhibition of PDE6, a cGMP-metabolizing enzyme that is present exclusively in photoreceptors. Sildenafil was shown to have similar potency against PDE6 in all species routinely evaluated in toxicological screens, and it was further demonstrated that long-term sustained inhibition of PDE6 by sildenafil (even at doses many times greater than those administered to humans) did not lead to any lasting damage to the structure and function of the eye29–32.
Indeed there are now multiple scientific papers suggesting a potential utility of sildenafil in protecting the ischaemic myocardium and in treating stroke\(^{36-45}\).

There have also, since 2000, been occasional case reports\(^{46-49}\) of non-arteritic anterior ischaemic optic neuropathy (NAION) in patients taking sildenafil. Although NAION is the most common acute optic neuropathy in people older than 50 years it is a relatively rare event causing partial visual loss in one eye, and is associated with various risk factors, including cardiovascular disease and a small cup:disk ratio. In a recently published review of clinical trial data\(^6\), Gorkin et al. estimated an incidence of 2.8 cases of NAION per 100,000 patient-years of sildenafil exposure, which is similar to estimates reported in the general US population (2.5–1.8 cases per 100,000 men aged >50 years)\(^{51,52}\). Very recently the original authors of many of these case reports published a further review\(^6\) and concluded that most of the case reports of NAION might be an expected coincidence because sildenafil is a top-selling medication and patients who receive the drug are frequently older, vasculopathic and already at risk of NAION. They further conclude that the only patients who need to avoid PDE5 inhibitors for visual reasons are those who have previously suffered NAION in one eye.

At the time of preparing this paper, sildenafil has been on the market as a treatment for ED for more than 7 years. It is now estimated that more than 750,000 physicians have prescribed sildenafil to over 23 million men, and sildenafil remains by far the most widely used mode of action of sildenafil in treating ED, the full scope of ED patients amenable to treatment with sildenafil and the potential for the management of other indications.

Post-marketing studies have revealed that conjunctival injection and epistaxis are relatively rare side effects of sildenafil administration. Contraindications have been extended to include left ventricular outflow obstruction, multiple system atrophy, and additional caution is necessary if patients are receiving alpha-blockers.

**Thoughts turn to pulmonary hypertension**

After the approval of sildenafil for the treatment of ED, thoughts started to turn to other potential indications for this drug. Sanchez et al.\(^5\) observed upregulation of PDE5 gene expression in pulmonary hypertensive lungs. Furthermore, it was observed that zaprinast (M&B 22948), E4021 and dipyridamole (a relatively non-selective PDE inhibitor with PDE5-inhibitory activity) could ameliorate pulmonary pressure in experimental pulmonary hypertension models\(^{56-60}\). With the availability of the more potent and selective PDE5 inhibitor sildenafil, a series of preclinical and clinical investigations were conducted to examine the therapeutic potential of this approach in pulmonary vascular diseases. The first intravenous placebo control study (Pfizer study 1024) was conducted to evaluate the effect of intravenous administration of sildenafil at various doses. The study was conducted between 1998 and 2000, and showed that sildenafil selectively reduced pulmonary pressure and
pulmonary vascular resistance in more than 80 patients with pulmonary arterial hypertension, pulmonary venous hypertension and pulmonary hypoxic hypertension. It was also observed that the effect reached a plateau at a plasma concentration of 100 ng per ml of sildenafil (G. Butrous, personal communication). During this period, interest in the role of sildenafil in pulmonary hypertension gained significant momentum.

**Pulmonary hypertension: background.** Pulmonary hypertension is a progressive disease of various origins which has a poor prognosis and results in right heart dysfunction\(^{41}\). According to the new classification for pulmonary hypertension from the Third World Conference on Pulmonary Hypertension in Venice in 2003 (REF 62), five subclasses of chronic pulmonary hypertension can be defined. Among these groups, class 1 represents the so-called pulmonary arterial hypertension (PAH) group, which includes a mixture of diseases that have some pathophysiological, histological and prognostic features in common. Idiopathic pulmonary arterial hypertension (iPAH, formerly known as primary pulmonary hypertension (PPH)) is the most prominent, although exceptionally rare, representative of this class, and is a disease for which loss-of-function mutations in the bone morphogenetic protein type 2 receptor (BMPR2) have been identified as one underlying mechanism\(^{63-65}\). Recent randomized controlled trials for drug development in this area have focused on patients suffering from iPAH as well as associated forms including connective tissue disease-associated pulmonary hypertension and pulmonary hypertension secondary to congenital heart disease\(^{66}\).

**Previous treatments for pulmonary arterial hypertension.** Continuous infusion of prostacyclin was the first licensed treatment for severe pulmonary arterial hypertension, and this approach has been shown to be life-saving\(^{67}\) and to improve exercise capacity\(^{68}\) in controlled clinical trials. However, there are also drawbacks of this therapy inherent both to the nature of the drug, and to the necessity of implanting a continuous central intravenous line. Preserving the advantageous effects of prostacyclin, while avoiding several of its limitations, the concept of aerosolized iloprost (a long-acting prostacyclin analogue) for the treatment of PAH was developed\(^{69,70}\). This therapy is currently approved in the US, Europe and many other countries for the treatment of PAH\(^{71}\); however, its use is cumbersome because of the need for frequent inhalation manoeuvres (six to nine times daily) and elaborate nebulization techniques\(^{72}\). The first oral drug approved for the treatment of pulmonary arterial hypertension was the non-selective oral endothelin receptor antagonist bosentan. The efficacy of this therapeutic approach has been proven in randomized controlled trials and led to approval in the US and Europe\(^{73,74}\). However, the long-term use of this vasodilator might be limited by liver toxicity, which affects up to 10% of all patients treated. Therefore the search for an ‘ideal’ pulmonary vasodilator that combines high treatment efficacy, pulmonary selectivity, simplicity of administration and reduced side effects continued until the PDE5 inhibitor sildenafil became the focus of investigation.

**Role of PDE5 in the pulmonary vasculature.** Nitric oxide is constitutively produced in the lung by NO synthases (NOS). The main enzymatic source of lung NO production is endothelial (or constitutive) NOS (eNOS), which is located in the vascular endothelium and the airway epithelium\(^{75,76}\). eNOS-derived NO activates soluble guanylate cyclase, which in turn increases intracellular cGMP levels. Adaptation of the perfusion distribution to well-ventilated areas of the lung (ventilation/perfusion (V/Q) matching) is regulated primarily by local NO production\(^{77,78}\), because the most prominent stimulus for
Alveolar hypoxia
Reduced oxygen levels (<80 mm Hg) in the lung alveoli caused by impaired ventilation (for example, in chronic lung disorders) or reduced oxygen content in the inspired air (for example, at high altitudes).

Hypoxic pulmonary vasoconstriction
Constriction of pulmonary vessels in the presence of alveolar hypoxia, which prevents the perfusion of non-ventilated areas of lung and maintains optimized gas-exchange properties (also known as the von Euler–Liljestrand mechanism).

In 1991, Haynes and colleagues demonstrated that the PDE5 inhibitor zaprinast caused a reduction in the vasoconstrictor response of isolated rat lungs to acute alveolar hypoxia80. The same compound was shown to induce that it was not only eNOS-derived NO that contributed to the effects of the PDE5 inhibitor85. The authors suggested the involvement of natriuretic peptides — which increase intracellular cGMP levels via receptor–linked activation of the particulate guanylate cyclase — by showing that in natriuretic peptide receptor A (NPR-A) knockout mice, the anti-remodelling effects of sildenafil on the pulmonary vasculature and the right ventricle were reduced89. Although all of the investigations described above involved initiating the treatment of chronically hypoxic animals at the onset of hypoxia, Sebkhi et al. demonstrated that even curative application of sildenafil reduces pulmonary artery pressure and vascular muscularization in the lungs of chronically hypoxic rats84.

In essence, these investigations demonstrate that PDE5 inhibition has anti-pulmonary hypertensive effects, with selective effects on the pulmonary vascular resistance. Therefore the selective pulmonary effects of PDE5 inhibitors are most probably attributable to a generally high level of PDE5 in the pulmonary circulation compared with the systemic circulation110,111,112 and the fact that NO production in the lung is high, akin to the situation in the corpus cavernosum. In a recent publication, Itoh et al. investigated the effect of sildenafil alone, or in combination with beraprost, a prostacyclin analogue, on the development of experimental PAH. Long-term administration of sildenafil to rats in a monocrotaline model of pulmonary hypertension resulted in a decrease in right ventricular systolic pressure, right heart hypertrophy and medial wall thickness. The curative properties of sildenafil after the development of pulmonary hypertension yielded similar results. Sildenafil reduced pulmonary artery pressure and vascular muscularization in lungs from chronically ill rats, and reduced the expression of matrix metalloproteinases (MMP) 2 and 9. Additionally, the degree of fully muscularized small (<50 μm) pulmonary arteries was decreased.

Clinical experience with sildenafil for the treatment of chronic pulmonary hypertension. The vasodilatory effects of inhaled NO are restricted to the pulmonary vasculature. In addition, NO has a very short half-life, is used as a screening agent for lung vasoreactivity113, and is effective in improving gas exchange in selected patients with adult respiratory distress syndrome (ARDS). Weaning from chronic NO treatment in patients with ARDS was facilitated by oral sildenafil. An initial case report of an adult patient suffering from severe pulmonary arterial hypertension who was treated chronically with very high doses of oral sildenafil indicated that this approach might be effective. In paediatric patients, the administration of intravenous prostacyclin is hampered to a greater extent by problems associated with the mode...
of administration than it is in adult patients. An early study reporting the successful use of oral sildenafil in a child with severe pulmonary hypertension therefore attracted a lot of attention, not only within the medical community but also in the media104–106. Trials addressing the characterization of the acute effects of sildenafil on pulmonary and systemic haemodynamics in a larger number of patients with pulmonary arterial hypertension showed that sildenafil effectively reduced pulmonary vascular resistance in a dose-dependent manner107. Notably, the vasodilatory effects were restricted primarily to the pulmonary circulation, and were significantly stronger than the effects seen with inhaled NO. In combination with inhaled iloprost, augmentation of the pulmonary vasodilatory effect of each single agent was observed107,108. Long-term treatment of patients with pulmonary arterial hypertension was investigated in a number of single-centre studies, all confirming the high efficacy and excellent tolerance of chronic oral sildenafil treatment109–111. In patients with deteriorating severe PAH despite ongoing prostanoid treatment, additional long-term administration of oral sildenafil improved exercise capacity and pulmonary haemodynamics111. The combination of prostanoids and sildenafil therefore has potential as a possible future treatment for pulmonary hypertension and numerous reports on the clinical use of sildenafil in pulmonary arterial hypertension in uncontrolled trials have been published to date109,109,110,113–118.

Interestingly, sildenafil seems to also be effective for treating patients with pulmonary hypertension of origins other than primary pulmonary hypertension. In patients suffering from human immunodeficiency virus (HIV)-related pulmonary hypertension, sildenafil was effective in reducing pulmonary vascular resistance, as it was in PPH118,119. Recent data also suggest that long-term oral sildenafil treatment in patients with non-operable chronic thromboembolic pulmonary hypertension is beneficial120. The importance of this finding lies in the fact that there is currently little to offer these patients in the way of therapeutic options, with the exception of lung transplantation.

**Pivotal trial and approval of sildenafil for the treatment of PAH (SUPER-1 study).** The growing body of evidence from various studies between 1998–2001, which had already demonstrated the efficacy of sildenafil in the treatment of PAH, led to the design of a large, randomized, controlled, multinational trial to provide final proof of this new treatment concept, and to obtain legal approval for sildenafil as a new treatment for pulmonary arterial hypertension. The SUPER-1 (Sildenafil Use in Pulmonary HypERTension) study started in 2002 and included 278 patients with symptomatic pulmonary arterial hypertension who were treated either with placebo or sildenafil (20, 40 or 80 mg) orally three times daily (TID) for 12 weeks. The primary endpoint in this trial—as in many previous trials with other medications similarly indicated—was the change from baseline to week 12 in the 6-minute walk test. Sildenafil, at all of the applied doses, improved exercise capacity (up to 50 metres [placebo-corrected value] in the 80 mg TID group), functional class and haemodynamics, as compared with placebo-treated patients, and was very well tolerated121. Additionally, patients completing the double-blind phase were able to enter a long-term extension trial, which was conducted over a 2-year period with 80 mg sildenafil TID. The increase in the 6-minute walk distance achieved after 3 months in the placebo-controlled phase was maintained even after a year of therapy, as were the improvements in functional class, which are both strongly indicative of the maintenance of the effect in spite of the severity of the disease.

Based on the very favourable mid- and long-term effects of this new oral treatment, sildenafil was approved by the FDA and the EMEA in 2003 for the treatment of patients suffering from PAH. Both agencies decided to approve only the 20 mg TID dose, as only a flat (non-significant) dose–effect relationship between 20–80 mg TID was observed regarding the primary endpoint of the study, the change in 6-minute walking distance over 12 weeks of treatment. In addition, analysis of sildenafil plasma levels in the SUPER-1 study showed no dose–effect relationship with the doses studied (Pfizer Chemical Research Group, unpublished data). There is evidence from some clinical and experimental settings that the duration of action of sildenafil might not be accurately reflected by plasma levels and the applied dosage122. It has been shown that the affinity of sildenafil for PDE5 is even increased after intracellular phosphorylation of the enzyme123. In addition, conformational changes to PDE5 and the slow dissociation rate of sildenafil from the enzyme could contribute to the flat dose–effect relationship124–127. One possible explanation is that sildenafil binding to the catalytic site of PDE5 could occur at higher affinity intracellularly than estimated.
previously, which might retard clearance of the inhibitor from the cells (G. Butrous, personal communication). On the other hand, however, in the SUPER-1 trial there were clear trends in some secondary endpoints (some showed statistically significant differences between the three applied doses) indicating that, for a subgroup of patients, higher doses might be more efficacious than the approved 20 mg TID dosage. Moreover, in the majority of preceding short- and long-term studies, daily doses of 100–300 mg were investigated and reported to be efficacious and well tolerated. Future studies are therefore warranted that address the long-term efficacy of 20 mg TID or even lower doses of sildenafil for the treatment of PAH.

**Future indications for PDE5 inhibitors**

*Raynaud's phenomenon and digital ulcers in collagen vascular diseases.* Patients with pulmonary arterial hypertension often complain about intermittent, temperature-dependent peripheral vasospasms resulting in perfusion deficiencies in their fingers and toes (so-called Raynaud's phenomenon) even in the absence of a proven collagen vascular disease. When treated with vasodilators, these symptoms can improve in parallel with, but also independent of, improvements in pulmonary haemodynamics. Raynaud's phenomenon and digital ulcers are of an even higher prevalence, and are clinically more important, in patients with systemic sclerosis, CREST syndrome (which comprises calcinosis cutis, Raynaud's phenomenon, oesophageal dysmotility, sclerodactyly and telangiectasia), and systemic lupus erythematosus. Treatment currently includes the application of calcium-channel blockers, infused prostanoids and alpha-2 blockade. However, the clinical efficacy of these therapies is often modest at best. A growing number of uncontrolled trials indicated that the efficacy of sildenafil for the treatment of digital ulceration and Raynaud's phenomenon in patients with scleroderma with or without pulmonary hypertension warranted attention. In an initial randomized controlled trial, Fries et al. investigated the effects of sildenafil (50 mg given twice daily) on the symptoms and capillary perfusion in patients with Raynaud's phenomenon. Most notably, only patients previously treated with other vasodilators that showed insufficient improvement were included in this trial. In contrast to the effects of placebo, chronic sildenafil treatment for 4 weeks reduced the mean frequency and duration of Raynaud attacks, and lowered the

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<thead>
<tr>
<th>PDE family</th>
<th>Role(s)</th>
<th>Evidence*</th>
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<tbody>
<tr>
<td>1</td>
<td>Vascular smooth muscle proliferation; Ca2+ modulation of olfaction</td>
<td>Broad distribution, but highest levels in proliferating vascular smooth muscle cells, testes, heart and neural tissues (for example, olfactory epithelial cells). Binding and inactivation by Ca2+/calmodulin</td>
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<tr>
<td>2</td>
<td>Regulation of Ca2+ channels, olfaction, platelet aggregation and aldosterone secretion</td>
<td>Broad distribution, but highest levels in brain and adrenal cortex</td>
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<tr>
<td>3</td>
<td>Cardiac contractility, insulin secretion and lipolysis</td>
<td>Broad distribution, but particular abundance in adipose tissue, liver, cardiac muscle, vascular smooth muscle and platelets; inhibited by drugs with cardiotonic, vasodilatory, thrombolytic, and antiplatelet aggregation properties. Stimulated by insulin, leptin and insulin-like growth factor</td>
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<tr>
<td>4</td>
<td>Immunological and inflammatory signalling processes; smooth muscle tone; depression</td>
<td>Broad distribution, highest levels in neural and endocrine tissue. Inflammatory cells thought to participate in the pathogenesis of inflammatory diseases (such as asthma and chronic obstructive pulmonary disease) preferentially express PDE4</td>
</tr>
<tr>
<td>5</td>
<td>Penile erection; smooth muscle tone of vasculature, airways and gastrointestinal tract</td>
<td>Abundant distribution in smooth muscle. The PDE5-specific inhibitor sildenafil has clinical efficacy for the treatment of erectile dysfunction</td>
</tr>
<tr>
<td>6</td>
<td>Vision</td>
<td>Distribution in rod and cone photoreceptor cells. Some visual defects are related to PDE6 mutations</td>
</tr>
<tr>
<td>7</td>
<td>T-lymphocyte activation and proliferation; skeletal muscle metabolism</td>
<td>Distribution is predominantly in T lymphocytes (PDE7A1). PDE7 mRNA is abundant in skeletal muscle tissue, T lymphocytes and B lymphocytes, but protein and activity are readily measurable only in T lymphocytes</td>
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<tr>
<td>8</td>
<td>T-cell activation</td>
<td>PDE8A mRNA is widely expressed (highest in testis). PDE8B is unique to the thyroid gland</td>
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<td>9</td>
<td>Possibly maintains basal intracellular cGMP levels or natriuresis and vascular tone</td>
<td>mRNA is widely expressed, particularly in spleen, intestine, kidney, heart and brain</td>
</tr>
<tr>
<td>10</td>
<td>Unknown</td>
<td>Human PDE10 is widely distributed</td>
</tr>
<tr>
<td>11</td>
<td>Sperm capacitation; other functions unknown</td>
<td>mRNA occurs at highest levels in skeletal muscle, prostate, kidney, liver, pituitary and salivary glands, and testis. Protein is localized to vascular smooth muscle cells, cardiac myocytes, corpus cavernosum of the penis, prostate and skeletal muscle</td>
</tr>
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*See REFS 191–196, cGMP, cyclic guanosine monophosphate; PDE, phosphodiesterase.
Figure 5 | Adaptation of blood flow to ventilation in the pulmonary circulation.

Blood flow (Q) in the pulmonary circulation must, ideally, be directed to well-ventilated areas (symbolized by big V (ventilation) and O2 (oxygenation) in the largest alveolus (blue circle at bottom of figure)) to ensure optimized gas exchange (‘matching’), whereas only a small amount of blood should flow through areas of minor or no ventilation (midsize and small alveolus, respectively) (left panel). Lung vessel dilatation is mainly regulated by the compartmentalized production of nitric oxide (NO) and subsequent intracellular cGMP formation, where alveolar distension and oxygenation represent the most potent stimuli for this local NO release. Similarly, less NO/cGMP is produced in non-ventilated areas of the lung, resulting in hypoxic vasoconstriction (the so-called von Euler–Liljestrand mechanism). During application of non-selective vasodilators and/or under disease conditions (for example, chronic obstructive lung disease, lung fibrosis, sepsis or acute respiratory distress syndrome), vasodilatation is induced in poorly or non-ventilated areas of the lung resulting in venous admixture and worsening of gas-exchange (‘mismatch’, right panel). There is strong evidence that oral sildenafil preferentially dilates vessels in well-ventilated areas of the lung, thereby both reducing overall vascular resistance and improving overall oxygenation (re-matching’ drug)\textsuperscript{143,144}.

Exercise tolerance

Ability to perform physical strain until limited by occurrence of peripheral (muscular) exhaustion, shortness of breath and/or insufficient blood supply to the myocardium (due, for example, to coronary heart disease).

Heart failure.

Chronic heart load leads to ventricular hypertrophy as an initial process of adaptation and can ultimately result in ventricular dilatation and failure if not treated chronically\textsuperscript{145}. Although cardiac hypertrophy applies both to disorders that lead to left ventricular loading as well as those that lead to right ventricular loading, there are important differences with respect to the reversibility of muscular hypertrophy of both ventricles. Although the right ventricle — even at advanced stages of dilatation and decompensation — can return to almost normal structure and function once the load is effectively reduced, left ventricular hypertrophy is only partly reversible once a certain degree of hypertrophy has been exceeded\textsuperscript{145,146}. In chronic pulmonary hypertension, right ventricular dysfunction is the ultimate cause of death over the course of the disease; however, effective reduction of the pulmonary vascular resistance — after lung transplantation, for example — might reverse right ventricular hypertrophy\textsuperscript{147}. Wilkins et al. have elegantly shown that effective treatment of pulmonary arterial prostanoid, selectivity for well-ventilated lung areas was demonstrated for sildenafil, resulting in an improvement, rather than deterioration, of gas exchange\textsuperscript{142}. Pulmonary hypertension impairs right ventricular performance because it increases right-heart afterload. However, it is still unclear to what extent exercise tolerance is limited by this mechanism. In a recent investigation, this issue was addressed under conditions of acute hypoxia at sea level, and prolonged hypoxia at the altitude of Mount Everest Base Camp\textsuperscript{144}. These investigations were performed in healthy volunteers to exclude other confounding factors that might have added to the limitation of exercise tolerance in patients suffering from chronic hypoxia (such as muscle wasting or chronic immobilization). In essence, acute and prolonged hypoxia induced significant pulmonary hypertension in the study subjects. As expected, exercise tolerance was dramatically reduced as a consequence of severe hypoxaemia and significant pulmonary hypertension. Sildenafil significantly reduced pulmonary hypertension under resting conditions, as well as during exercise. The most interesting finding of this study was that the reversal of pulmonary hypertension resulted in an immediate improvement of exercise tolerance, irrespective of improvements in oxygenation. Further studies investigating the effects of acute and chronic sildenafil administration in hypoxic pulmonary hypertension confirmed the anti-pulmonary hypertensive potential, and the beneficial effects of sildenafil on exercise performance, under these conditions\textsuperscript{143,144}. The results of these studies stimulated further investigations that addressed the therapeutic potential of this drug in patients suffering from chronic hypoxic pulmonary hypertension as it occurs in various chronic diseases (such as chronic obstructive pulmonary disease (COPD), interstitial lung disease and obstructive sleep apnoea)\textsuperscript{147–150}. In fact, very recent work supports the possibility of effective treatment of pulmonary hypertension in patients suffering from advanced COPD\textsuperscript{151}. Based on the significant impact of COPD on public health, studies in this field are warranted.
An interesting further insight into the mode of action of sildenafil comes from several preclinical investigations, including a recent publication by Musicki et al., which investigated the effects of long-term treatment of sildenafil in young and aged rats. In aged rats, but not in their young counterparts, sildenafil prolonged erection and increased the protein expression of phosphorylated endothelial nitric oxide synthase (eNOS) and phosphorylated Akt. Therefore the description of the mode of action of sildenafil in erectile dysfunction, and possibly also in pulmonary vascular diseases, can be further elaborated. It can be proposed that the initial increase in NO levels — due to neurogenic stimulation or a result of alveolar distension, for example — is followed by subsequent maintenance of vasodilation by increased endothelial generation of NO via activated eNOS. It can also be speculated that the resulting increase in blood flow, and consequently shear stress, across the vascular endothelial cell surface activates phosphatidylinositol 3-kinase (PI3K)/Akt, which phosphorylate and thereby further activate eNOS. The activated eNOS can then lead to a sustained further release of NO, which acts to maintain the smooth muscle relaxation, vascular engorgement and increased blood flow (see figure). As sildenafil is being explored for other potential indications, and provided there is a dual effect of chronic administration on activation of eNOS and inhibition of the phosphodiesterase type 5 (PDE5), the therapeutic potential for various diseases involving vascular pathology might therefore be greater than originally envisaged.

**Cerebral circulation.** Various studies have provided compelling evidence for a functional role of the NO-cGMP pathway in the brain[173,174]. Inhibition of hippocampal NOS resulted in a state-dependent impairment in object recognition[175]. As no direct influence of PDE5 inhibitors on the overall cerebral blood flow was found[173,175,177], the functional changes described in response to these agents were attributed to neuronal alterations[178]. Static flow-measures might, however, not be the appropriate parameter to characterize changes in microcirculatory adaptation to demand, where this regulation is mediated via neurovascular coupling in a fast and fine-tuned manner[179-181]. Concerning the cerebral vasculature, numerous studies have addressed the distribution of regional brain perfusion in health and disease states[176,182-184]. Investigations addressing the cerebral effects of sildenafil showed no changes in overall cerebral blood flow[173,175,177] after administration of this agent. On the other hand, despite the absence of changes in overall cerebral blood flow, an effect of the PDE5 system on cognitive function[185], repair mechanisms after stroke and expression profiles of second messenger levels in distinct brain areas have been proposed[153,186,187]. This discrepancy might well be reconciled by a recent finding suggesting that sildenafil significantly improves neurovascular coupling while leaving overall cerebral blood flow unchanged[188]. Furthermore, this finding is in line with previous observations that the initial blood flow response to cortical activity is mainly governed by stimulation of the NO system[189]. It has also been proposed that there might be therapeutic potential for sildenafil in cerebral ischaemic disorders, because cGMP levels were shown to be inversely correlated with cardiac hypertrophy in an isoproterenol-induced cardiac hypertrophy model in rats[190].

However, the potential usefulness of sildenafil in chronic heart failure might result from a variety of actions in addition to an effect on ventricular hypertrophy. A variety of investigations have indicated that sildenafil can be cautiously administered to selected patients with left heart failure[163-172]. Endothelial function (the capacity of arteries to enable increases in regional blood flow in response to appropriate stimuli such as ischaemia) is severely limited in chronic congestive heart failure. In an elegant study Katz et al. showed that sildenafil could improve endothelial function in such patients. Further studies investigating the long-term effects of PDE5 inhibitors in patients with these diseases would therefore be logical. Also, by virtue of its effect on vasodilation, arterial stiffness and wave reflection, sildenafil has been shown to reduce aortic pressure and the augmentation index, and could therefore have a role in the management of systemic hypertension[191].

**Box 1 | Further insights into the mechanism of action of sildenafil**

An interesting further insight into the mode of action of sildenafil comes from several preclinical investigations, including a recent publication by Musicki et al., which investigated the effects of long-term treatment of sildenafil in young and aged rats. In aged rats, but not in their young counterparts, sildenafil prolonged erection and increased the protein expression of phosphorylated endothelial nitric oxide synthase (eNOS) and phosphorylated Akt. Therefore the description of the mode of action of sildenafil in erectile dysfunction, and possibly also in pulmonary vascular diseases, can be further elaborated. It can be proposed that the initial increase in NO levels — due to neurogenic stimulation or a result of alveolar distension, for example — is followed by subsequent maintenance of vasodilation by increased endothelial generation of NO via activated eNOS. It can also be speculated that the resulting increase in blood flow, and consequently shear stress, across the vascular endothelial cell surface activates phosphatidylinositol 3-kinase (PI3K)/Akt, which phosphorylate and thereby further activate eNOS. The activated eNOS can then lead to a sustained further release of NO, which acts to maintain the smooth muscle relaxation, vascular engorgement and increased blood flow (see figure). As sildenafil is being explored for other potential indications, and provided there is a dual effect of chronic administration on activation of eNOS and inhibition of the phosphodiesterase type 5 (PDE5), the therapeutic potential for various diseases involving vascular pathology might therefore be greater than originally envisaged.
with an increase in the phosphorylation of glycogen synthase kinase 3 (GSK3), a downstream target of Akt. Co-incubation of neurospheres with sildenafil and LY 294002, a pharmacological inhibitor of phosphatidyl-
inositol 3-kinase (PI3K)/Akt, abolished sildenafil-
induced phosphorylation of Akt and GSK3, suggesting that sildenafil enhanced neurogenesis through activation of the PI3K/Akt/GSK3 pathway.

**Conclusion**

Sildenafil started clinical development as an agent for the treatment of hypertension and angina, and subsequently evolved into a revolutionary new oral treatment for erec-
tile dysfunction. Sildenafil was then further developed to become a much-needed new oral treatment for pul-
monary arterial hypertension, and has also been shown to be effective in treating severe Raynaud’s phenomenon associated with systemic sclerosis and digital ulceration. Later investigative studies have suggested that sildenafil also has promise in the treatment of respiratory disor-
ders with ventilation/perfusion mismatch, congestive cardiac failure, hypertension and even stroke. Moreover, many years after the original indication was abandoned, single-dose sildenafil was shown to prolong exercise time in men with angina. Although these findings might seem to be quite disparate and unrelated, all of the disorders described above are in fact characterized by regional deficiencies in blood supply. The successful application of a PDE5 inhibitor (as opposed to non-selective vaso-
dilators) to treat these conditions can be understood in terms of the capacity of sildenafil to reverse endothelial dysfunction, and to selectively improve regional blood flow in areas of greatest need. Further applications of PDE5 inhibitors may arise from new insights into the mode of action of chronic sildenafil administration (BOX 1). Many patients with erectile dysfunction and pulmonary arterial hypertension are now benefiting from advances in our understanding of vascular biology and pathophysiology, and the advent of selective inhibitors of PDE5 (TABLE 5). It is hoped that the clinical potential of this mechanism to treat the other serious medical condi-
tions described above (and others such as pre-eclampsia and certain forms of intrauterine growth retardation) will soon be realized, so that many more patients can benefit from these amazing breakthroughs in science, technology and medicine.

### Table 3 | PDE inhibition and selectivity

<table>
<thead>
<tr>
<th>Drug</th>
<th>Geometric mean IC₅₀ values (µM) [fold selectivity versus PDE5 in parentheses]</th>
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<tbody>
<tr>
<td>Vardenafil</td>
<td>0.070 [500] 6.20 [44,290] &gt;1.0 [7,140] 6.10 [43,570] 0.00014 [0.035] [4] 0.0006 [30 &gt;214,000] &gt;30 &gt;214,000 0.581 [4,150] 3.0 [21,200] 0.162 [1,160]</td>
</tr>
</tbody>
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IC₅₀ values were determined using either native enzyme purified from human tissue (PDE1, heart; PDE2, 3 and 5, corpus cavernosum; PDE4, skeletal muscle; PDE6, retina) or using recombinant human enzymes expressed in Sf9 cells (PDE7-11) and purified by anion-exchange chromatography. Adapted from REF. 29. IC₅₀ drug concentration necessary to inhibit 50% of enzyme activity; PDE, phosphodiesterase.

18. Important review of the International Index of Erectile Function (IIEF), the current ‘gold standard’ measure for efficacy assessment in clinical trials of erectile dysfunction.
21. First publication of the results of two sequential double-blind studies proving the efficacy and safety of sildenafil, administered as needed in men with erectile dysfunction.


29. Very informative data analysis on safety and tolerability of sildenafil in the treatment of erectile dysfunction from a series of double-blind, placebo-controlled studies and from ten open-label extension studies.

30. Cleker, E., Bethell, S., Fancett, L., Mount, N. & Phillips, S. Phosphodiesterase 5 inhibitor profiles from controlled studies and from ten open-label controlled studies and from ten open-label extension studies.


41. First systematic clinical trial proving that sildenafil had no adverse cardiovascular effects in men with severe coronary artery disease.


47. Interesting experimental work addressing the importance of lung nitric oxide production in the regulation of hypoxic vasoconstriction.


The first clinical study to show that an oral vasodilator (sildenafil) can improve gas exchange, with comparable effects to those achievable with inhaled nitric oxide.}


