Long term imatinib treatment in pulmonary arterial hypertension

Pulmonary arterial hypertension (PAH) is a life-threatening condition characterised by progressive obliteration of the small pulmonary arteries leading to increased pulmonary arterial resistance and right heart failure. Treatment for PAH has developed in the last few years; hence the description of new pathways related to the disease. Recently, short-term (6 months) use of imatinib, a platelet derived growth factor (PDGF) receptor antagonist, in combination with maximal PAH treatment (prostacyclin derivative, endothelin receptor antagonist, and type 5 phosphodiesterase inhibitor) has been shown to improve the haemodynamics and functional capacity in a single case of severe PAH.1 We here report the first two cases of the long term (3 years or more) use of imatinib, as monotherapy or in combination with bosentan, a dual endothelin receptor antagonist.

Case 1 was a 34 year old man with PAH associated with type 1 glycogen storage disease. Until 1999 the patient had remained stable with functional class II PAH (New York Heart Association classification) without specific treatment. In 2000 he presented with increased white blood cell count at his routine evaluation resulting in the diagnosis of chronic myeloid leukemia, a known late complication of type I glycogen storage disease. The patient remained in functional class II and the haemodynamic pattern showed a trend to worsening without a significant change at the 6 minute walk test (fig 1A). Imatinib was started as first line treatment for leukemia without any associated PAH treatment. During 3 years of imatinib use his leukemia was adequately controlled, functional capacity was sustained, and the haemodynamic profile was improved.

Case 2 was a 65 year old woman with a known diagnosis of chronic myeloid leukemia since 1994 which was satisfactorily controlled with hydroxyurea/cytemamine, cytarabine, and Imuran. In 1996 she reported dyspnoea on exercise with insidious progression during the next 4 years. In 2000 an echocardiogram showed a right ventricular systolic pressure of 65 mm Hg. In 2002 she presented with functional class III PAH and was referred. The investigation showed no other condition associated with PAH but significant haemodynamic impairment (fig 1B). Treatment with bosentan was initiated with a good haemodynamic and functional response after 3 months. At that time interferon was withdrawn and imatinib was started as treatment for leukemia. Since then, after more than 3 years of treatment with imatinib and bosentan, there has been functional and haemodynamic improvement (fig 1B).

In these two cases of long term use of imatinib (alone or in combination with bosentan), functional and clinical stabilisation or improvement in PAH were observed. Of note is the progressive increase in cardiac index during the 3 years of treatment with imatinib. Such a progressive increase in cardiac index is certainly an indicator of a good prognosis, as previously discussed in PAH.1 PDGF is a potent mitogen that has been related to the chemotaxis and proliferation of pulmonary vascular smooth muscle cells, and inhibition has been shown to prevent and reverse pulmonary hypertension in experimental models, raising the potential of its use in clinical practice.1 Imatinib is approved for the treatment of chronic myeloid leukemia, which was the reason for using it in the cases leading to haematological remission. In parallel, the haemodynamic response to imatinib alone or in combination with bosentan was significant. Although many side effects have already been described with the long term use of imatinib,1 only mild anaemia was observed in our first case. We conclude that PDGF inhibition should be tested in future PAH clinical trials in order to establish its safety and efficacy.

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References

Reduced exercise capacity in a mouse model of asthma

One of the important clinical features of asthma is the exercise intolerance due to an exacerbation.1 Yet, to our knowledge, this end point has never been assessed in animal models of asthma. In a mouse model of chemical induced asthma we found early and late alterations in ventilatory function that were retained whole body plethysmography.2–4 Partly because this technique has been heavily criticised,2 we sought to provide another measure of functional impairment in asthma: swimming. This was done by a simple exercise test in which mice were forced to swim against a limited downwards current,2 if they stopped swimming the current pulled them under the water. Swimming endurance time is measured as a proxy of exercise tolerance. Task failure is defined as a period of 5 seconds under water.

The protocol for sensitising and challenging mice was similar to that used previously,2 with small modifications. BALB/c mice (± 20 g, 6 weeks old) received dermal applications of 20 μl vehicle (2:3 acetonitrile:0.5% aqueous 2,4-diso- cyanate (TDI) on each ear on days 1 and 8. On day 15 they received intranasal instillation of 10 μl vehicle or 0.1% TDI in each nostril. Treatment with TDI is indicated as 1, while treatment with vehicle is indicated as 0. Thus, the 1/1/1 type consists of mice that received dermal applications of TDI (days 1 and 8) and an intranasal instillation of TDI (day 15), while the 0/0/0 control group consists of mice that received the vehicle on all occasions. All mice had a 5 minute swimming training session on day 13. All swim tests were done in water at 34°C.

On day 15 resting ventilatory function (enhanced pause (Penh)) of each mouse was recorded by whole body plethysmography (EMKA Technologies, Paris, France) for 5 minutes before and 40 minutes after the intranasal instillation. One hour after instillation the mice were made to swim in groups of 2 mice until exhaustion (fig 2A). On the day 16 (that is, 22 hours after intranasal instillation) the mice had a second swim test. Two hours later methacholine reactivity was assessed by whole body plethysmography. The measurement of the swimming time was not conducted in a blinded manner, but we are confident that this did not influence the results. All experimental procedures were approved by the local Ethical Committee for Animal Experiments.

Figure 1 shows the early increase in Penh (panel A) and late increase in methacholine reactivity (panel B) in mice sensitised and challenged with TDI (group 1/1/1) compared with non-sensitised mice (1/0/0) and control mice (0/0/0). These changes are similar to those of our previous experiments.2 The outcomes of the swimming test paralleled those of ventilatory function: in the 1/1/1 group endurance was reduced by 13 minutes (panel C) and by 5 minutes (panel D) at the early and late time points, respectively, compared with the 0/0/0 group whose swimming times did not differ from those of the 0/0/0 control group. The swim time of the control mice was comparable to those of Matsumoto et al.2 When looking at responses in mice, the magnitude of the impairment in the swimming test correlated well with the magnitude of the early increase in Penh (r = 0.84, www.thoraxjnl.com
A Wallis test, Dunn’s multiple comparison post hoc test).

PostScript

Figure 1 (A) Ventilatory response (Penh) before and after intranasal instillation (arrow) with TDI or vehicle. (B) Methacholine responsiveness (Penh) 24 hours after instillation of TDI or vehicle. (C) Swim test 1 hour after instillation with TDI or vehicle. (D) Swim test 22 hours after instillation with TDI or vehicle. Experimental groups are identified by three symbols. 0 and 1 represent administration of vehicle (acetone:olive oil) or TDI, respectively; the first two symbols identify the agent applied dermally on days 1 and 8, and the third symbol identifies the agent instilled intranasally on day 15. n = 5 per group. **p<0.01 compared with the 0/0/0 control group (non-parametric Kruskal-Wallis test, Dunn’s multiple comparison post hoc test).

p<0.001) and the late increase in methacholine reactivity (r = −0.72, p<0.01). The second swim test correlated also with the early increase in Penh (r = −0.85, p<0.001) and the increased methacholine reactivity (r = −0.83, p<0.001).

This is the first study to use an exercise test to measure the physiological response in a mouse model of asthma. In mice that had been dermally sensitised and then intranasally challenged with TDI, there was an early decrease in endurance 1 hour after challenge and this was still present 1 day later, though to a lesser degree. As in our previous experiments,3 the physiological response depended on the prior dermal administration of TDI, thus excluding a non-specific toxicity of TDI. Although our findings provide further evidence of a relevant functional response in our asthma model, it remains to be verified whether the changes in Penh and in exercise capacity found here are due to reduced nasal patency, bronchoconstriction, or other effects in small airways.

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References