

Nebulisers are used to deliver bronchodilators to people with asthma or COPD or antibiotics to those with bronchiectasis.

There is now substantial evidence that the use of a metered dose inhaler (MDI) with a spacer is just as effective as a nebuliser, even in acute asthma.

Indications for nebuliser use

For acute deteriorations in asthma or COPD

- If the patient is unable to use a MDI and spacer e.g. very young children, intellectual disability, anxiety.
- If the episode is so severe that the rate of breathing is above 30 per minute at rest (in an adult), and the patient is unable to inhale from a spacer properly.
- The patient has a condition e.g. rheumatoid arthritis, which impairs use of the MDI or spacer.
- If the patient has a history of "crashing" or lifethreatening asthma.

For chronic asthma or COPD

- For a patient with COPD whose lung function is so poor that they cannot inhale properly from any hand held delivery device.
- For a patient with asthma or COPD with another disability which prevents them using a MDI and spacer.

For cystic fibrosis or bronchiectasis

- For delivery of prophylactic antibiotics.
- For delivery of hypertonic saline.
- For delivery of DNA-ase.

What are the potential harmful effects of nebuliser therapy?

Regular nebulised bronchodilator therapy (i.e. for more than a few days) should be avoided. The risk of adverse effects is based on the following:

- The delivered dose using a nebuliser is high. The dose in a single nebule may be equivalent to either 25 or even 50 puffs of bronchodilator from a MDI.
- Beta-agonists have a pro-inflammatory effect when given at high doses.
- Beta-agonists result in an increase in airway hyper-responsiveness (AHR) when given regularly or at high doses.
- In some patients the increase in AHR is associated with rebound bronchconstriction at the end of the dosing interval. Therefore, a vicious circle develops in some patients because with increased inflammation and AHR, symptoms worsen and there is a perception that even more "reliever" is required.

There are other pitfalls. Using a nebuliser may create a false sense of security, especially during an acute episode of asthma. The temporary relief from a high dose of nebulised bronchodilator may result in a delay in starting steroid therapy – or in ringing the ambulance. Also, the "rush" which occurs with nebulised beta-agonist - often associated with a fast heart rate or tremor – is enjoyable for some people. "Reliever" beta-agonists are modified forms of adrenaline and, because the delivered dose of nebulised beta-agonist is so high, it affects not only the airways but other organs also. The "adrenaline rush" is therefore real, and psychological dependence may occur.



The Asthma and Respiratory Foundation of New Zealand (Inc.) Te Taumatua Huango, Mate Ha o Aotearoa

These problems may be encountered in people with COPD just as much as in people with asthma, although the evidence is less clear cut. In people with predominantly emphysema, there is an additional problem – the smooth muscle relaxation which occurs with nebulised bronchodilator may actually cause "floppy" airways to collapse. Paradoxically the patient gets worse, not better. Some patients may benefit from beta-agonist withdrawal.

Psychological dependence on beta-agonists is a recognised problem. One of the ways of identifying psychological dependence is to measure peak flow at times when the patient feels the need to use the nebuliser and then afterwards. If there is no fall in peak flow associated with symptoms, or no increase with bronchodilator, then it may be that symptoms are not due to bronchospasm but something else, including vocal cord dysfunction or anxiety hyper-ventilation. This scenario may not apply to patients with COPD in whom the relationship between airway spasm and symptoms is less clear.

A very carefully constructed plan of action to wean someone off excessive beta-agonist therapy may be required. Sometimes it even requires admission to hospital for 3–4 days so that the patient can be weaned off safely. If this situation arises, consultation with a respiratory specialist is advised.

Professor D. Robin Taylor,

Medical Adviser,

Asthma and Respiratory Foundation of New Zealand. July, 2008



Otago Department of General Practice Dunedin School of Medicine

Postgraduate Diploma in Rural & Provincial Hospital Medicine (PGDipRPHP)

GENX 721 Rural Hospital Medical Medicine 30 pts (full year) 2009 Maximum of 10 participants

NEW

GENX 708 Special Topic Echocardiography and GENX 713 Medical Ultrasound 30 pts each 2009

Taught co-requisitely. Maximum of 8 participants

Short Course in Rural Hospital Cardiology

For more information please contact:

Raelene Abernethy

Tel 03 479 9186 or 021 263 2635

Email raelene.abernethy@stonebow.otago.ac.nz

Postgraduate Diploma in General Practice (PGDipGP)

GENX 820 Core Studies in General Practice 30 pts (full year) 2009

Semester One 2009 GENX 825 Medical Anthropology 15 pts (semester one) 2009

Semester Two 2009 GENX 826 Special Topic - Complementary Medicine 15 pts (semester two) 2009

For more information about the Postgraduate Diploma in General Practice (PGDipGP) contact:

Anita Fogarty

Tel 03 479 7424 or 027 2823 009 Email anita.fogarty@otago.ac.nz

