Chest physiotherapy (CPT) with bronchial drainage is the standard treatment for mobilization and removal of airway secretions in many types of respiratory dysfunction especially in chronic lung disease, such as cystic fibrosis, bronchiectasis, bronchitis, bronchial asthma, primary ciliary dyskinesia syndrome. CPT has been shown to be effective in maintaining pulmonary function and prevention or reduction of respiratory complications in patients with chronic respiratory diseases. However, standard CPT is very labor-intensive and time-consuming both for hospitalized and non-hospitalized patients with impaired airway clearance. For this reason many patients refuse to do daily physiotherapy and interrupt it with all bad consequences.

In recent years, devices of respiratory physiotherapy have emerged which offer alternatives to standard CPT which are less time-consuming and offer greater independence to the patient with chronic lung disease. According to recent literature, devices of respiratory physiotherapy are introduced as alternative therapy methods in order to facilitate and improve mobilization of mucus from airways, through which better lung ventilation and improved pulmonary function can be achieved. These devices are safe and offer acceptable airway clearance to conventional CPT.

Patients with chronic respiratory diseases prefer to use devices of respiratory physiotherapy because of their benefits, such as the independent application and the reduced cost of therapy. Aerosol treatments may be given while the patient is using these devices, if needed. The current devices of respiratory physiotherapy are: Positive Expiratory Pressure, High Frequency Chest Wall Oscillation, Oral High Frequency Oscillation, Intrapulmonary Percussive Ventilation, Incentive Spirometry, the Flutter and the Acapella. Current devices seem to be effective in terms of mucus expectoration and pulmonary function improvement, as it is shown by published studies. The choice of the suitable device for each patient is a challenge for the physiotherapist in order to achieve better compliance in daily treatment. More controlled studies are needed due to the fact that the number of published studies is limited.

Positive Expiratory Pressure (PEP) device
Positive expiratory pressure (PEP) therapy was developed in the 1970s and has been introduced in the United States as an alternative to conventional physiotherapy. The device consists of (Figures 1, 2): a face-mask or mouthpiece, a one-way valve, to which expiratory resistances can be attached and a manometer between the valve and the resistance, to monitor the actual value of pressure, which should be between 10 and 20 cm H$_2$O during mid-expiration.

This device is considered to allow more air to enter peripheral airways via collateral channels, to allow pressure air to go behind secretions, moving them towards larger airways where they can easily be expelled and to prevent the alveoli from collapsing.

Application: Patient must be able to perform certain manoeuvres in an upright or sitting position. The patient slowly inspires to vital capacity and then holds his breath for about 3 seconds. Then slowly exhales through the
mouthpiece with the fixed orifice resister that is usually set to create an expiratory pressure resistance between 10-20 cm H₂O. This slow deep breathing manoeuvre is repeated anywhere from 10 to 20 times. Then a “huff” or a forced exhalation should follow to clean secretions that have been mobilized. Periods for relaxation and breathing control for about 1-2 minutes are necessary. Full expiration should be avoided. Therapy duration and frequency are adjusted according to individual’s needs.

PEP device may give independence to patients with chronic respiratory diseases, as it can be carried out without an assistant and is easy and convenient in use. Small clinical studies have reported improved tracheobronchial clearance and patient comfort with PEP devices compared to standard CPT. Reduction in pulmonary infections/antibiotic courses and improved bronchodilation is also reported. In addition, there has also been reported improvement in compliance and shorter hospital stays. Other studies report PEP as an acceptable and effective treatment regimen to lung function.

High Frequency Chest Wall Oscillation (HFCWO) device

HFCWO is performed with a mechanical device (ThAIRaphy® bronchial drainage system, Hayek Oscillator). During HFCWO, positive pressure air pulses are applied to the chest wall, for example by means of an inflatable vest and an air pulse generator (Figures 3, 4). The generator produces pressures of about 50 cm H₂O at a frequency of around 525 Hz delivered via a pneumatic vest which surrounds the thorax. These air pulses oscillate the chest and the vibrations reportedly cause transient flow increases in the airways, loosening mucus and producing cough like sheer forces.
It is hypothesized that increases in cough clearing ability may be due to an increase in mucus/airflow interaction and/or a shearing mechanism leading to a decrease in the viscoelasticity of mucus. Several studies report that the high frequency chest wall oscillation device: a) provides an adequate physiotherapy method to conventional physiotherapy, b) helps sputum expectoration, c) contributes in stabilization or improvement of respiratory function and d) increases airflow in low lung volumes.

Application: The manufacturers recommend that the patient must undergo this therapy while on upright or sitting position and concomitantly receiving aerosol bronchodilator therapy. The HFCWO treatment is started at low pressures and frequency and then increased to the recommended pressure/frequency as per patient tolerance.

These therapy sessions generally last about 30 minutes and should not be performed on patients with significant hemoptysis. Despite these restrictions, these devices have been shown to mobilize more secretions than standard CPT in small clinical trials. More research is needed to determine its efficacy, cost benefits and optimum treatment strategies.

Oral High Frequency Oscillation (OHFO) device
Oral high frequency oscillation has been developed from the technique of high frequency jet ventilation. Oscillation of the air within the lungs at high frequency is associated with an increased clearance of CO₂. Because of the high frequency and low volume of these oscillations, spontaneous breathing is unhindered and the technique has potential value as a supplement to ventilation. Sine wave oscillations are produced by an eccentric cam piston or the diaphragm of a loudspeaker and can be superimposed on normal tidal breathing. The low volumes of approximately 48 ml and pressures approximately 0.2 cm-2.0 cm H₂O with a mean pressure of zero allow patients to breathe spontaneously.

This device provides a practical and simple method of supplementing breathing in conscious subjects, and it may also have application in the management of patients with acute or chronic respiratory failure, where intubation and conventional ventilation might be avoided.

In normal subjects under laboratory conditions OHFO has been shown that increases mucociliary clearance and reduces minute ventilation.

Data on the effect on mucociliary clearance of oral high frequency oscillation is conflicting. Nevertheless this device has not been demonstrated to be more effective than other techniques in patients with cystic fibrosis or chronic bronchitis, it may be an alternative for some patients. Increase in mucociliary clearance in normal man induced by oral high frequency oscillation OHFO has considerable potential in the management of patients with chronic airflow obstruction (CAO), where it may be of value as an assistance to breathing and in the relief of breathlessness.

This device of chest wall oscillation is widely used in the USA. Direct comparisons with other airway clearance treatments more commonly used in other countries, are difficult to make.

Intrapulmonary Percussive Ventilation (IPV) device
Intrapulmonary percussive ventilation (IPV) combines aerosol inhalation and internal thoracic percussion applied via a mouthpiece. The IPV is the delivery of a pulsatile flow of gas released with each pulse that can be preset and the pulsation frequency adjusted to each individual.

Application: The patient initiates the flow of gas and during inspiration the pulsatile flow results in an internal percussion. Interruption of the respiratory flow allows for passive expiration. According to some researchers the device of intrapulmonary percussive ventilation is as effective as «standard» respiratory physical therapy, assists mucus clearance and may be an alternative for some patients.

Further studies are need to determine the short-term and long-term goals and to establish the rank of IPV in airway clearance in patients with chronic diseases.

Incentive Spirometry (I.S.) device
Incentive spirometry provides feedback at a preset inspiratory flow of volume of air (Figure 5). Incentive Spirometry is designed to mimic natural sighing or yawning by encouraging the patient to take slow, deep breaths. A device called Incentive Spirometer is used to provide information about patient’s inspiratory effort by measuring the air flow (FEV₁) and the air volume (FVC).

The use of incentive spirometer has been supported to increase or maintain inhaled lung volume, improve sputum expectoration and to avoid serious lung infection, especially after surgery.
**Application:** Holding the incentive spirometer upright, patient is encouraged to take a slow and deep breath with his lips sealed around the mouthpiece and is motivated to achieve a preset volume by visual feedback, such as a piston rising to a preset marker. Holding breath for 2-3 sec at full inspiration is very important. Expiration is slow and calm. After each set of 10 breaths, cough should be encouraged in order to clear the lungs from mucus. It is recommended a repetition of at least ten breaths every hour on the incentive spirometer.

The pattern of breathing while using incentive spirometry is important. Emphasis should be given in the expansion of the lower chest during full inspiration, rather than the use of the accessory muscles of respiration, which would encourage expansion of the upper chest. IS has been compared with few of the airway clearance regimens and it is difficult to ascertain its effectiveness. There is little evidence to support the use of IS in airway clearance but it is thought to be an important factor in the reduction or prevention of postoperative pulmonary complications and its use in those patients is frequent. Using incentive spirometer after surgery help keeping lungs clear. Deep inhalations promote the mobilization of secretions and the opening up of lung areas that may have become collapsed. Also it exercises the lungs, keeping them active, especially during the recovery from surgery, as if the patient was at home performing daily activities. The use of IS appears to improve arterial blood gases and health-related quality of life in patients with COPD exacerbations, although it does not alter pulmonary function parameters.

**Flutter Device**

The Flutter device was developed in Switzerland and combines positive expiratory pressure therapy with high-frequency oscillations within the airway. It is a controlled vibration system which produces positive expiratory pressure and cyclic oscillation of the airways during expiration. The Flutter device is a portable device designed to help clear mucus in patients with lung disorders.

The device consists of a tube based on oscillations of a steel ball during expiration through a pipe-type device. The principle behind this device is that exhalation into the Flutter valve causes a steel ball-bearing to oscillate at a high frequency, resulting in vibration of the airways and intermittent positive expiratory pressure, to facilitate mucus expectoration (Figures 6,7).

Exhalation through the Flutter results in oscillations of expiratory pressure and airflow, which vibrate the airway walls (loosening mucus), decrease the collapsibility of the airways and accelerate airflow facilitating movement of mucus up the airways and improving lung function and oxygenation.

**Application:** Flutter device must be used in the sitting or supine lying position. The patient is instructed to inhale deeply and hold his breath for 2 to 3 sec. Expiration should be slow through the Flutter valve, causing oscillations of the steel ball inside the cone of the Flutter (Figure 8). Patients apply repeated exhalations through the Flutter valve. Routinely, three sets of 15 exhalations are performed over 12–20 min. After each series of exhalations, patients were instructed to “huff” and cough, thereby aiding expectoration.

The frequency of the oscillations can be modulated by changing the inclination of the Flutter device slightly up or down from its horizontal position (Figure 9). The patient selects the position that results in the best transmission of vibration to chest wall, optimizing the mobilization of mucus.

Effective use of the Flutter device requires training, concentration, and appropriate positioning of the mouthpiece. The Flutter device is simple to use inexpensive and easily portable and once the patient and his family are instructed in its use it does not require the assistance of a caregiver.
Additionally, patients with severe obstruction may not be able to generate sufficient airflow to cause vibration of the steel ball housed in the pipe-like extension of the Flutter valve, thus limiting the effectiveness of this device in these patients.

Long-term studies of the use of the Flutter seem justified to determine its effects on pulmonary function and outcome. For hospitalized patients elimination of the need for a therapist could reduce health care costs. The Flutter therapy is an acceptable alternative to standard RPT during in-hospital of patients with CF.

Other devices producing similar effect such as Flutter device, based in different mechanism of oscillations and vibration effect within chest wall can be used by patients with chronic respiratory diseases. These devices are Acapella and Cornet.

**Acapella device**

The Acapella (Smiths Medical Inc, Carlsbad, California, USA) is a handheld airway clearance device (Figure 10) that operates on the same principle as the Flutter, i.e. a valve interrupting expiratory flow generating oscillating PEP. Utilizing a counterweighted plug and magnet to achieve valve closure, the Acapella is not gravity dependent like the Flutter. The Acapella comes in three models, a low flow (<15 L/min), high flow (>15 L/min) and the Acapella Choice. The high and low flow models have a dial to set expiratory resistance while the Choice model has a numeric dial to adjust frequency. All models can be used with a mask or mouthpiece and can be used in line with a nebulizer. While these attributes may offer the Acapella some advantage over the Flutter, no long-term studies have been done in CF patients. A bench study of the performance characteristics of the two devices showed a slight advantage for the Acapella, with more stable wave form and a wider range of PEP at low air flow.
**Cornet device**

The Cornet device (R. Cegla, Montabaur, Germany) consists of a semi-circular tube containing a flexible latex-free hose (Figure 11). Expiration through the Cornet causes the hose to flex, buckle and unbuckle, causing oscillating positive pressure in the airways which fluctuates many times per second. The mouthpiece can be adjusted to produce the optimal effect. Operating principle and use are similar to the Flutter valve, although the Cornet is not gravity dependent and can be used in any position. Like the Flutter the Cornet cannot be used in line with a nebulizer. No studies showing the long-term effectiveness of the Cornet in CF patients are available yet. The Cornet is available in Europe but not in the USA.

![Figure 11: Cornet device.](image)

![Figure 12: Representation of oscillations of the latex-free hose.](image)

**Critical analysis of Research studies**

Sixty eight studies regarding devices of respiratory physiotherapy published to PubMed the last twenty years were studied in order to define their effectiveness. The effectiveness of standard chest physiotherapy (CPT) has been confirmed by many studies. CPT is considered the base of respiratory physiotherapy and is characterized as ‘gold standard’ of physiotherapy.

A great number of the published studies (36 studies- percentage 53%) compare the devices of respiratory physiotherapy with standard CPT. Eighteen studies (percentage 26%) are referred to the effectiveness of each device separately, while a small number of reports (8 reports- percentage 12%) compare the devices between them. Limited number of studies compares the devices with other active techniques of respiratory physiotherapy (6 studies- percentage 9%) (Table 1, Figure 13).

<table>
<thead>
<tr>
<th>Table 1: Published studies.</th>
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<tr>
<td><strong>Comparison devices vs CPT</strong></td>
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<td><strong>Effectiveness of each device</strong></td>
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<td><strong>Comparison of devices between them</strong></td>
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<td><strong>Comparison devices vs active techniques</strong></td>
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![Figure 13: Percentages of published studies.](image)

The greater number of published studies (63 studies- percentage 93%) is concerned the short-term effects, while only 5 studies (percentage 7%) were referred to long-term results of devices (Table 2, Figure 14).

<table>
<thead>
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<th>Table 2: Results of studies.</th>
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<tr>
<td><strong>Short-term results</strong></td>
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<tr>
<td><strong>Long-term results</strong></td>
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A great number of studies (25 studies- percentage 73%) determine the results of devices in patients with cystic fibrosis. Eight reports (percentage 24%) refer to patients with COPD and only 1 study (percentage 3%) to patients with primary ciliary dyskinesia. (Table 3, Figure 15)
The devices used more often according to published researches are Flutter device (23 studies) percentage 35%, PEP device (18 studies) percentage 27% and HFCWO device (11 studies) percentage 16%. Following the IPV device (9 studies) percentage 13% and in the same number of reports the IS device and Cornet device (3 studies) percentage 4%, while the less used device is Acapella device (1 study) percentage 1% (Table 4, Figure 16).
Conclusively, according to the results of published studies we support that more research is needed to define the long-term outcomes of the devices of respiratory physiotherapy, to evaluate patients’ compliance to therapy and to underline their effect to quality of life. It is difficult to determine which device of respiratory physiotherapy is better since there are no long-term trials that compare the devices between them. Flutter and PEP device are the most popular used devices, even though this is not confirmed by long-term studies, while for the other devices the literature is limited and ambiguous.

Discussion

Stasis of secretions in respiratory diseases leads to chronic infection, inflammation and lung destruction\textsuperscript{44}. Respiratory physiotherapy has been used for many years to help in removal of secretions. However the denial of the patients to do daily physiotherapy especially in chronic obstructive diseases leads to the creation of regimens which provide independent application.

The current respiratory devices have been designed to enhance patients’ compliance and independence.

According to the research studies the current devices of respiratory physiotherapy are effective in improving pulmonary function, lung oxygenation, clearing mucus from bronchi and making feasible better compliance of patients in treatment. Also they decrease the respiratory complications\textsuperscript{4,4,8,35,38,49-51}. Furthermore these devices are easy in use and they reduce cost of therapy\textsuperscript{8}.

The number of published reports is limited, though more research is needed to define the effectiveness of the devices of respiratory physiotherapy and their place among the current techniques available.

Physiotherapists should be informed about current devices of respiratory physiotherapy in order to choose the appropriate device for each patient, according to patient’s age and clinical condition. Also they should teach patient the direct use of the device and give practical advices. Also the patient must be informed that these devices must not replace the programme of respiratory physiotherapy, because they have a supplemental role to respiratory physiotherapy, contributing to better results in pulmonary function. Alternation of using the devices, according to the patient’s condition will give a motivation to the patients in order to continue their treatment.

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