Early View

Research letter

Pneumothorax as a complication of dry needling technique

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Pneumothorax as a complication of dry needling technique. Jan S.B. Bontinck, MD, Emergency Department, University Hospital, Ghent, Belgium Cathelijne Lyphout, MD, Emergency Department, University Hospital, Ghent, Belgium Thomas L.A. Malfait, MD, Department of Respiratory Medicine, University Hospital, Ghent, Belgium Correspondence address: Dr. Jan Bontinck Emergency Department Universitair Ziekenhuis

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To the editor:

Dry needling involves the insertion of a solid needle into palpably taut muscle bands or muscle knots known as 'myofascial trigger points'. After insertion the needle may be redirected in several planes or electrically stimulated to cause muscle contraction which should relieve pain and improve range of motion. The technique is referred to as 'dry' because no medication is injected. The underlying mechanism of action is subject to debate, as is the exact pathogenesis of 'myofascial trigger points'. Dry needling is used in the treatment of a variety of musculoskeletal disorders e.g. tendinopathy, shoulder impingement, non-radicular neck pain or chronic low back pain. Being considered as its Western counterpart, dry needling is similar to acupuncture, which originates from traditional Chinese medicine. The latter technique however focuses on 'flows of life force (chi)' and aims to treat conditions in multiple organ systems whereas the former is used mainly in the musculoskeletal field.

Evidence regarding the effectiveness of dry needling is conflicting. ²⁻⁶ However not robust, literature suggest dry needling may be a useful addition in the management of muscle pain. The technique requires training to perform properly and results may vary depending on clinician skill. In Belgium procedures are usually performed by physiotherapists who have completed an additional training in private academies. The duration of these courses ranges from 14 hours to 2 years, with some requiring no medical background at all. The lack of regulation may lead to a great variety in quality of care.

High-quality data regarding adverse events associated with dry needling are lacking. Most literature on this topic is based on patient or physician surveys and case reports. Being practiced for centuries and thus more widespread, acupuncture is the topic of a vast body of literature, including publications regarding safety. Dry needling being considered a low-risk intervention is partly due to extrapolation of the presumed safety of acupuncture. The most common adverse events after small diameter needle insertion are mild bleeding, bruising and post-needling soreness. Albeit rare, more serious adverse events like nerve injury, hematoma, infection and cardiac tamponade have been described. In a prospective study Witt et al found that 2 in nearly 230.000 patients reported pneumothorax after acupuncture treatment. A systematic review of the Chinese literature found that pneumothorax was the most frequent serious acupuncture-associated adverse event with a total of 201 cases, 4 of which were fatal. To our knowledge only two cases of pneumothorax after dry needling have been described. Under the present a series of four cases in one Belgian university hospital over a period of 15 months.

Four patients were seen at the emergency department (ED) with post-dry needling pneumothorax between September 2022 and December 2023. All were women aged 28 to 35. One patient was a smoker. None of the patients had any relevant medical history. All had been treated for pain located in the left shoulder, trapezius muscle or neck region in outpatient physiotherapist practices. At least three different physiotherapists were involved, one could not be identified. One patient presented to the ER on the same day as the dry needling procedure, the others presented the day after. All mentioned thoracic pain (4/4) and dyspnoea (4/4). Clinical examination in all of these patients was unremarkable, as were their vital signs. Diagnosis was confirmed with ultrasound (US) and chest X-ray (CXR) in all patients. The latter exam showed left-sided apical pleural detachment with a median of 3.65 cm in expiration.

Two patients were managed conservatively. One patient (initial pneumothorax 2.5 cm) was discharged from the ED and US two days later displayed a normally expanded lung. One patient with an initial apical size of 2.8 cm was admitted with 2 litres of oxygen through a nasal canula and discharged from the hospital the next day after US had shown no increase in size. Her control CXR 4 days later showed only minimal pleural detachment measuring 6 mm. The two other patients were treated with US guided needle aspiration in the ED. One patient with detachment initially being 4.5 cm showed decreased size of the pneumothorax immediately after aspiration. She was admitted to the respiratory medicine ward and discharged the next day. Control US and CXR after 1 week showed no more signs of pneumothorax. In the other patient, with detachment initially being 5.5 cm, needle aspiration resulted in complete deployment on US immediately after the procedure, but control CXR showed a totally collapsed lung 3 hours later. A small bore chest drain was placed but persistent air leakage was seen. Several trials of clamping the drain resulted in recurrent collapsing of the lung. After CT-scan had shown no structural deformities of the lung, suction was gradually reduced and the drain was successfully removed on the sixth day after placement. Patient was discharged home. Control CXR 3 weeks later was normal.

This case series describes four young women suffering a pneumothorax after dry needling of the shoulder- and neck region. In July 2023 the British Thoracic Society (BTS) published updated guidelines for the management of spontaneous pneumothorax¹³. Although it does not deal with traumatic pneumothoraxes, the BTS treatment algorithm guided our approach to these four patients. Since all were symptomatic but none exhibited high-risk characteristics (e.g. hemodynamic compromise or significant hypoxia), size of the pneumothorax and patient preference were decisive for treatment choice. This led to conservative treatment in two, needle aspiration in two, and hospital admission in three out of four cases with a mean length of stay of 2.7 (range 1-6) days and chest drain placement in one patient. Besides the psychological and financial burden, all patients dealt with incapacity for work.

Our series suggests post-dry needling pneumothorax is, contrary to numbers cited in literature, not extremely rare. With rising popularity of the technique we expect complications to occur more often. Patients and referring doctors should be aware of this. In their informed consent practitioners should mention pneumothorax as a considerable risk of dry needling procedures in the neck, shoulder or chest region.

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