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Determinants of diaphragm inspiratory motion, diaphragm thickening and its performance for predicting respiratory restrictive pattern in Duchenne muscular dystrophy

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Abstract

Introduction/Aims

Respiratory status is a key determinant of prognosis in patients with Duchenne muscular dystrophy (DMD). We aimed to evaluate the determinants of diaphragm ultrasound and its performance in predicting restrictive respiratory patterns in DMD.

Methods

We included DMD patients who experienced diaphragm ultrasound and pulmonary functional tests.

Results

This study included 74 patients with DMD. The right diaphragm thickening fraction (TF) was significantly associated with age ($p = 0.001$), Walton score ($p = 0.012$), inspiratory capacity (IC) ($p = 0.004$), upright FVC ($p < 0.0001$), supine FVC ($p = 0.038$), and maximal inspiratory pressure (MIP) ($p = 0.002$). Right diaphragm excursion was significantly associated with age ($p < 0.0001$), steroid use ($p = 0.008$), history of spinal fusion ($p < 0.0001$), BMI ($p = 0.002$), Walton score ($p < 0.0001$), IC ($p < 0.0001$), upright FVC ($p < 0.0001$), supine FVC ($p < 0.0001$), and MIP ($p < 0.0001$). A right diaphragm TF $> 28\%$ was associated with a FVC $> 50\%$, with an area under the curve (AUC) at 0.95 ($p = 0.001$). A right diaphragm excursion > 25.4 mm was associated with an AUC at 0.93 ($p < 0.001$). A left diaphragm TF $> 26.8\%$ was associated with upright FVC $> 50\%$, with an AUC at 0.95 ($p = 0.011$). A left diaphragm excursion > 21.5 mm was also associated with a pulmonary upright FVC $> 50\%$, with an AUC at 0.97 ($p < 0.001$).

Discussion

DIM and diaphragm TF can predict restrictive pulmonary insufficiency in DMD.

Key words: Duchenne muscular dystrophy, pulmonary, diaphragm, ultrasound

Clinical trial NCT 04045158

MANUSCRIPT

INTRODUCTION

Respiratory dysfunction in Duchenne muscular dystrophy (DMD) is characterized by a progressive reduction in lung volume with the onset of a restrictive pulmonary pattern and nocturnal followed by diurnal hypoventilation (1). Non-invasive mechanical ventilation is recommended in patients with an upright forced pulmonary vital capacity (FVC) of less than 50% of the predicted value (1). Respiratory status is a key determinant of prognosis in patients with DMD, as patients are at risk of acute respiratory failure and pneumonia (1). It is recommended to perform pulmonary respiratory tests regularly in DMD patients after the age of 6–7 years (1). Ultrasound has emerged as a recent technique that can help physicians to assess and monitor the diaphragm, which is the main inspiratory muscle. Diaphragm ultrasound is a single, reproducible technique without radiation exposure, which can be performed at the bedside (2, 3). Diaphragm ultrasound is used in the intensive care unit (ICU) during the weaning trial process in patients on mechanical ventilation to search for predictive factors of weaning trial failure (4, 5). In the field of neuromuscular disorders, this technique has emerged as a non-invasive test to assess diaphragm function (6). The diaphragm thickening fraction (TF) and diaphragm excursion are ultrasound parameters measured in radiology. The diaphragm TF was defined as $(\text{end inspiratory thickness} - \text{end expiratory thickness}) / \text{end expiratory thickness}$ (7). The diaphragm TF can be used as an index of respiratory effort and is correlated with inspiratory muscle pressure (7, 8). We aimed to evaluate patients with DMD and identify the determinants of the hemidiaphragm excursion and diaphragm TF, and accuracy of diaphragm ultrasound parameters to predict a FVC > 50% of predicted value.

METHODS

Study population

From 2015 to 2018, we included, retrospectively, DMD patients followed in our center and admitted for annual checkup including cardiopulmonary evaluation and diaphragm ultrasound. Patients with DMD who were ventilated 24/24 h invasively were excluded. From the cardiac ultrasound database, we noted the left ventricular ejection fraction (LVEF), the systolic pulmonary artery pressure, and diaphragm ultrasound parameters available. The Gardner Medwin Walton scale was used to evaluate the peripheral skeletal muscle disability (0, normal; 1, unable to run; 2, altered gait; 3, climbs stairs with hand support; 4, unable to climb stairs; 5, unable to rise from a chair; 6, walk only with aids; 7, cannot walk; 8, unable to move in a manual wheelchair; 9, unable to sit unsupported; 10, bedridden) (9). This study was performed in compliance with the ethical principles formulated in the Declaration of Helsinki and was approved by the French Regulatory Board (*Commission Nationale de l'Informatique et des libertés, CNIL*). Informed consent was obtained from all participants. This study was registered in a clinical trial (NCT 04045158).

Diaphragm ultrasound

Diaphragm ultrasound was performed simultaneously by an experienced operator (A.F.) during echocardiographic examination. Diaphragm thickness was measured at the apposition zone using a transducer placed perpendicular to the chest wall, between the anterior and the mid-axillary lines (**supplemental figures**). The patients were placed in a semi-recumbent position. For this measurement, we used a linear transducer probe with a higher frequency 9 L-D (2.4–10 MHz). Using the B mode, the diaphragm was visualized as a hypoechoic layer of muscle tissue surrounded by two hyper-echoic lines: the pleural and peritoneal lines. Diaphragm thickness was measured at the end of the expiratory phase and at the end of deep inspiration. The diaphragm TF was calculated using the following formula: $TF = (\text{end inspiratory thickness} - \text{end expiratory thickness}) / \text{end expiratory thickness}$ (10).

To record diaphragm excursion, a patient was placed in a semi-recumbent position, and the cardiac probe M5Sc-D (1.5–4.6 MHz) was positioned at the subcostal area between the mid-clavicular and

anterior axillary lines (**supplemental figures**); the liver and spleen were used as an acoustic window for the right and left hemidiaphragm, respectively. From the subcostal approach, M-mode tracing was recorded, and the ultrasound beam reached the posterior third of the hemidiaphragm. During inspiration, the diaphragm moves toward the probe. The diaphragm is recognized as an echogenic line that covers the liver. We recorded the diaphragm excursion during a deep inspiratory maneuver.

Pulmonary functional tests

Pulmonary functional tests were performed according to the ATS/ERS recommendations (11) using a Vmax 229 SensorMedics System (Yorba Linda, USA) in the upright position. Forced vital capacity (FVC) was also measured in the supine position. Maximal inspiratory pressure (MIP) was measured from the functional residual capacity in the upright position. Maximal expiratory pressure (MEP) was measured at total lung capacity. MIP was measured using a flanged mouthpiece with the maneuvers repeated at least three times or until two identical readings were obtained. Once the operator was satisfied, the maximum value of the three maneuvers that varied by less than 20% was recorded (12).

STATISTICAL ANALYSIS

Categorical and continuous variables were presented as numbers (percentages) and medians (interquartile ranges), respectively, due to the non-normal distribution of data. Non-parametric tests were performed in accordance with each other. Associations were assessed using non-parametric correlations. Finally, the discriminative characteristics of diaphragm echocardiographic parameters were assessed using the area under the curve (AUC) of receiver operator characteristics. Alpha risk was 5%. All calculations were performed using SPSS version 23.0 (IBM, Armonk, NY, USA).

RESULTS

Study population, diaphragm ultrasound, and respiratory function

From the 93 eligible patients with DMD, we included 74 patients who underwent diaphragm ultrasound to measure their diaphragm excursion (see **Figure 1**, flowchart), 68 (92%) patients (92%) were nonambulatory. Genetic tests did not find any mutation in one patient. Genetic findings in the 73 other patients were distributed as follows: exonic deletion, 60%; exonic duplication, 10 %; nonsense mutation, 11%; small deletion, 3%, small mutation, 12%; and intronic mutation, 4%. The median upright pulmonary FVC decreased. Non-invasive ventilation was used by most patients. The respiratory muscle strength and the right and left diaphragm TF were reduced. In addition, the right and left deep diaphragm excursion were reduced. **Table 1** summarizes the clinical and laboratory findings in patients with DMD.

Determinants of right diaphragm thickening fraction

The TF of the right diaphragm was significantly associated with that of the left diaphragm and it was inversely and significantly associated with age and Walton score. The right diaphragm TF was positively associated with LVEF, inspiratory capacity, upright FVC, supine FVC, MIP, peak expiratory flow (PEF), and MEP. **Figure 2 panel A** shows the relationship between the right hemi-diaphragm TF and MIP.

Determinants of the right diaphragm excursion

Right diaphragm excursion was significantly associated with left diaphragm excursion, and it was inversely and significantly associated with age and Walton score. The right diaphragm excursion was significantly positively associated with steroid use ($p = 0.008$), BMI, inspiratory capacity, upright FVC, supine FVC, MIP, PEF, and MEP. **Figure 2 panel B** shows the relationship between the right diaphragm excursion and MIP. **Figure 2 panel C** shows the relationship between pulmonary FVC and right diaphragm excursion, while **Figure 2 panel D** shows the relationship between the right diaphragm excursion and inspiratory capacity. The **Table 2** summarizes relationship between diaphragm ultrasound parameters and spirometric parameters.

Diaphragm ultrasound accuracy for predicting respiratory involvement in DMD

In DMD, a right diaphragm TF > 28% is associated with a pulmonary upright FVC > 50% with a sensitivity at 100%, a specificity at 87%, and an AUC at 0.95 ($p = 0.001$). A right diaphragm excursion >25.4 mm was also associated with a pulmonary upright FVC > 50%, with a sensitivity of 100%, a specificity of 85.2%, and an AUC at 0.93 ($p < 0.001$). For the left hemidiaphragm, a left diaphragm TF > 26.8% is associated with upright FVC > 50%, with a sensitivity at 100%, a specificity at 90%, and an AUC at 0.95 ($p = 0.011$). A left diaphragm excursion >21.5 mm was also associated with a pulmonary upright FVC > 50%, with a sensitivity of 100%, a specificity of 93.9%, and an AUC at 0.97 ($p < 0.001$).

Figure 3 summarizes the ROC curves for the different diaphragm ultrasound parameters.

Equation for predicting pulmonary forced vital capacity

Using a regression that included right diaphragm excursion (RDE) and right inspiratory diaphragm thickness (RIDT), the FVC can be predicted using the following equation: $FVC (ml) = Constant + 295.9 \times RIDT (mm) + 52.4 \times RDE (mm)$.

DISCUSSION

In this study, we found that diaphragm ultrasound parameters were significantly correlated with lung volumes and respiratory inspiratory muscle strength in patients with DMD. We found a significant association between diaphragm TF and inspiratory capacity, FVC, and MIP, and a significant association between diaphragm excursion and inspiratory capacity, FVC, and MIP.

In addition, we suggest that diaphragm ultrasonography may predict respiratory involvement in DMD. Diaphragm ultrasound is a free-risk and reproducible technique that can be used at the bedside (13). Diaphragm ultrasound is used in the ICU during the weaning process to predict successful extubation (5). The diaphragm TF is associated with trans-diaphragmatic pressure and reflects the work of breathing in ICU patients (8, 13). In healthy individuals, diaphragm ultrasound has been found to be significantly associated with inspiratory muscle strength and pulmonary

function (14). In healthy subjects, the diaphragm TF is also significantly associated with gastric pressure at inspiration, and diaphragmatic deep inspiratory motion is also significantly related to trans-diaphragmatic pressure following the sniff maneuver (15). This significant association between diaphragm excursion and trans-diaphragmatic pressure has been observed in patients undergoing postoperative cardiac surgery (16). In ICU patients, diaphragm TF has already been found to be significantly associated with inspiratory muscle pressure (17).

Our results are similar to findings in previous studies of various other neuromuscular diseases such as facio-scapulo-humeral muscular dystrophy (18), late-onset Pompe disease (LOPD) (19), and amyotrophic lateral sclerosis (ALS) (20). In fact, Henke et al. (18) reported a significant association between diaphragm thickening and MIP and between diaphragm excursion and MIP. In LOPD patients, Ruggeri et al. (19) reported a significant association between vital capacity and diaphragm excursion, TF, and inspiratory muscle strength. Similar results were reported by Fantini et al. (20) in patients with ALS. In DMD, diaphragm thickness is lower than in healthy subjects (6), and diaphragm function decreases with age (21). The present study highlights the usefulness of diaphragm ultrasonography in predicting respiratory involvement in DMD.

We found that a right diaphragm TF > 28% could predict a pulmonary upright FVC > 50% in DMD. This diaphragm cut-off is similar to that in ICU patients and is used to predict weaning trials (22). A diaphragm TF < 29% has been reported to identify diaphragm dysfunction, defined by a change in endotracheal tube pressure less than 11 cm H₂O, after a bilateral anterior magnetic phrenic nerve stimulation (22). The required diaphragm TF *cut-off* required for extubation success varies among authors and extends from 30% to 36% (8, 17).

We found that a right diaphragm excursion > 25.4 mm can predict a pulmonary upright FVC > 50% using the subcostal approach. This result is similar to cardiac surgery findings previously reported (16). In fact, Lerolle et al. (16) found a cut-off value of 25 mm to identify diaphragm dysfunction. Mechanisms involved in diaphragm dysfunction in DMD are multifactorial and include muscle

dystrophy, mechanical ventilation, and diaphragm loading. In the ICU, mechanical ventilation may expose patients to muscle trauma, which may be due to excessive ventilation (23, 24). Because some patients with DMD experience cognitive impairment and disclosed inability to maintain an adequate mouth seal during spirometry, diaphragm ultrasound may be proposed to assess respiratory function. Magnetic resonance imaging can be used to assess diaphragm function and respiratory muscle fatty infiltration (25); however, this technique requires a supine position that can be very difficult to maintain in DMD patients with respiratory insufficiency and spinal fusion.

Finally, the FVC depends on the inspiratory muscles, expiratory muscles, and mechanical properties of the thoracoabdominal wall. Thus, the significant association between diaphragm ultrasound parameters and FVC, compared with the significant association between inspiratory capacity and diaphragm ultrasound, suggests that diaphragm ultrasound parameters reflect not only inspiratory muscle function but also global respiratory function.

Our study has some limitations owing to its retrospective design. In addition, diaphragm motion is pertinent, particularly in patients on unassisted breaths; in patients on mechanical ventilation, the diaphragm downward displacement is the combined diaphragm displacement induced by the ventilator and the diaphragm displacement due to patient inspiratory efforts (26). In addition, most DMD patients are on nocturnal noninvasive mechanical ventilation, which may affect diaphragm function over time. Finally, external validation of our data was not performed. Prospective longitudinal studies that focus on the evolution of DMD patients before and after ventilation will help to better stratify patients.

Conclusion

DIM and diaphragm TF were significantly associated with spirometric parameters and inspiratory muscle strength. These diaphragmatic ultrasound parameters may predict the respiratory involvement in DMD. Diaphragm ultrasound can be used to assess and monitor respiratory status in DMD and may be appropriate for assessing DMD-like myopathies such as sarcoglycanopathies.

Findings: none

List of abbreviations

ALS: amyotrophic lateral sclerosis

AUC: area under the curve

DMD: Duchenne muscular dystrophy

DIM: diaphragm inspiratory motion

FVC: forced vital capacity

GMW: Gardner Medwin Walton score

IC: inspiratory capacity

ICU: Intensive Care unit

LOPD: late-onset Pompe disease

LVEF: left ventricular ejection fraction

MIP: maximal inspiratory pressure

MEP: Maximal expiratory pressure

PEF: Peak expiratory flow

RDE: right diaphragm excursion

RDIM: right diaphragm inspiratory motion

RDIT: right inspiratory diaphragm thickness

TF: thickening fraction

REFERENCES

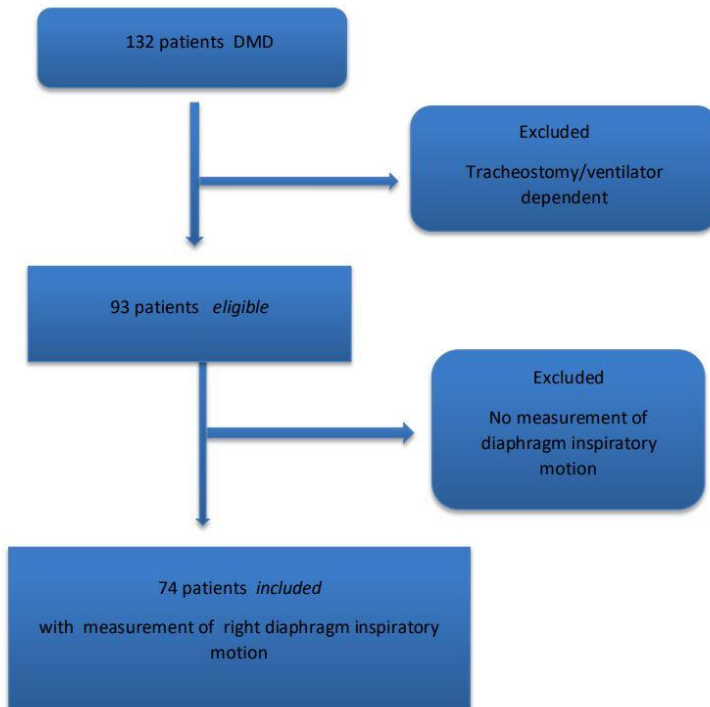
- 1) Birnkrant DJ, Bushby K, Bann CM, Alman BA, Apkon SD, Blackwell A, Case LE, Cripe L, Hadjiyannakis S, Olson AK, Sheehan DW, Bolen J, Weber DR, Ward LM, Care DMD, Considerations Working Group. Diagnosis and management of Duchenne muscular dystrophy, part 2: respiratory, cardiac, bone health, and orthopaedic management. *Lancet Neurol.* 2018 Apr;17(4):347-361.
- 2) Boussuges A, Gole Y, Blanc P. Diaphragmatic motion studied by m-mode ultrasonography: methods, reproducibility, and normal values. *Chest.* 2009 Feb;135(2):391-400.
- 3) Cappellini I, Picciafuochi F, Bartolucci M, Matteini S, Virgili G, Adembri C. Evaluation of diaphragm thickening by diaphragm ultrasonography: a reproducibility and a repeatability study. *J Ultrasound.* 2020 May 1.
- 4) Kim WY, Suh HJ, Hong SB, Koh Y, Lim CM. Diaphragm dysfunction assessed by ultrasonography: influence on weaning from mechanical ventilation. *Crit Care Med.* 2011 Dec;39(12):2627-2630.
- 5) DiNino E, Gartman EJ, Sethi JM, McCool FD. Diaphragm ultrasound as a predictor of successful extubation from mechanical ventilation. *Thorax.* 2014 May;69(5):423-427.
- 6) Laviola M, Priori R, D'Angelo MG, Aliverti A. Assessment of diaphragmatic thickness by ultrasonography in Duchenne muscular dystrophy (DMD) patients. *PLOS ONE.* 2018 Jul 26;13(7):e0200582.
- 7) Umbrello M, Formenti P, Longhi D, Galimberti A, Piva I, Pezzi A, Mistraletti G, Marini JJ, Iapichino G. Diaphragm ultrasound as indicator of respiratory effort in critically ill patients undergoing assisted mechanical ventilation: a pilot clinical study. *Crit Care.* 2015 Apr 13;19(1):161.
- 8) Vivier E, Mekontso Dessap A, Dimassi S, Vargas F, Lyazidi A, Thille AW, Brochard L. Diaphragm ultrasonography to estimate the work of breathing during non-invasive ventilation. *Intensive Care Med.* 2012 May;38(5):796-803.
- 9) Walton JN, Gardner-Medwin D. Progressive muscular dystrophy and the myotonic disorders. In: Walton JN, editor. *Disorders of the voluntary muscle.* 4th ed. London: Churchill Livingstone; 1981.

- 10) Carrillo-Esper R, Pérez-Calatayud AA, Arch-Tirado E, Díaz-Carrillo MA, Garrido-Aguirre E, Tapia-Velazco R, Peña-Pérez CA, Espinoza-de Los Monteros I, Meza-Márquez JM, Flores-Rivera OI, Zepeda-Mendoza AD, de la Torre-León T. Standardization of sonographic diaphragm thickness evaluations in healthy volunteers. *Respir Care*. 2016 Jul;61(7):920-924.
- 11) Quanjer PH, Tammeling GJ, Cotes JE, Pedersen OF, Peslin R, Yernault JC. Lung volumes and forced ventilatory flows. Report Working Party Standardization of lung Function Tests, European Community for steel and coal. Official Statement of the European Respiratory Society. *Eur Respir J Suppl*. 1993;16:5-40.
- 12) Wilson SH, Cooke NT, Edwards RH, Spiro SG. Predicted normal values for maximal respiratory pressures in Caucasian adults and children. *Thorax*. 1984;39(7):535-538.
- 13) Goligher EC, Laghi F, Detsky ME, Farias P, Murray A, Brace D, Brochard LJ, Bolz SS, Rubenfeld GD, Kavanagh BP, Ferguson ND. Measuring diaphragm thickness with ultrasound in mechanically ventilated patients: feasibility, reproducibility and validity. *Intensive Care Med*. 2015 Apr;41(4):642-649.
- 14) Cardenas LZ, Santana PV, Caruso P, Ribeiro de Carvalho CR, Pereira de Albuquerque AL. Diaphragmatic ultrasound correlates with inspiratory muscle strength and pulmonary function in healthy subjects. *Ultrasound Med Biol*. 2018 Apr;44(4):786-793.
- 15) Spiesshoefer J, Herkenrath S, Henke C, Langenbruch L, Schneppe M, Randerath W, Young P, Brix T, Boentert M. Evaluation of Respiratory Muscle Strength and Diaphragm ultrasound: normative Values, Theoretical Considerations, and Practical Recommendations. *Respiration*. 2020;99(5):369-381.
- 16) Lerolle N, Guérot E, Dimassi S, Zegdi R, Faisy C, Fagon JY, Diehl JL. Ultrasonographic diagnostic criterion for severe diaphragmatic dysfunction after cardiac surgery. *Chest*. 2009 Feb;135(2):401-407.
- 17) Ferrari G, De Filippi G, Elia F, Panero F, Volpicelli G, Aprà F. Diaphragm ultrasound as a new index of discontinuation from mechanical ventilation. *Crit Ultrasound J*. 2014 Jun 7;6(1):8.
- 18) Henke C, Spiesshoefer J, Kabitz HJ, Herkenrath S, Randerath W, Brix T, Görlich D, Young P, Boentert M. Respiratory muscle weakness in facioscapulohumeral muscular dystrophy. *Muscle Nerve*. 2019 Dec;60(6):679-686.

- 19) Ruggeri P, Lo Monaco L, Musumeci O, Tavilla G, Gaeta M, Caramori G, Toscano A. Ultrasound assessment of diaphragm function in patients with late-onset Pompe disease. *Neurol Sci.* 2020 Aug;41(8):2175-2184.
- 20) Fantini R, Mandrioli J, Zona S, Antenora F, Iattoni A, Monelli M, Fini N, Tonelli R, Clini E, Marchioni A. Ultrasound assessment of diaphragmatic function in patients with amyotrophic lateral sclerosis. *Respirology.* 2016 Jul;21(5):932-938.
- 21) Fayssoil A, Chaffaut C, Ognia A, Stojkovic T, Lamothe L, Mompoin D, Meng P, Prigent H, Clair B, Behin A, Laforet P, Bassez G, Carlier R, Orlikowski D, Amthor H, Quijano Roy S, Crenn P, Chevret S, Eymard B, Lofaso F, Annane D. Echographic assessment of diaphragmatic function in Duchenne muscular dystrophy from childhood to adulthood. *J Neuromuscul Dis.* 2019;6(1):55-64.
- 22) Dubé BP, Dres M, Mayaux J, Demiri S, Similowski T, Demoule A. Ultrasound evaluation of diaphragm function in mechanically ventilated patients: comparison to phrenic stimulation and prognostic implications. *Thorax.* 2017 Sep;72(9):811-818.
- 23) Schepens T, Dianti J. Diaphragm protection: what should we target? *Curr Opin Crit Care.* 2020 Feb;26(1):35-40.
- 24) Goligher EC, Brochard LJ, Reid WD, Fan E, Saarela O, Slutsky AS, Kavanagh BP, Rubenfeld GD, Ferguson ND. Diaphragmatic myotrauma: a mediator of prolonged ventilation and poor patient outcomes in acute respiratory failure. *Lancet Respir Med.* 2019 Jan;7(1):90-98.
- 25) Barnard AM, Lott DJ, Batra A, Triplett WT, Forbes SC, Riehl SL, Willcocks RJ, Smith BK, Vandenberghe K, Walter GA. Imaging respiratory muscle quality and function in Duchenne muscular dystrophy. *J Neurol.* 2019 Nov;266(11):2752-2763.
- 26) Schepens T, Fard S, Goligher EC. Assessing diaphragmatic function. *Respir Care.* 2020 Jun;65(6):807-819.

FIGURES

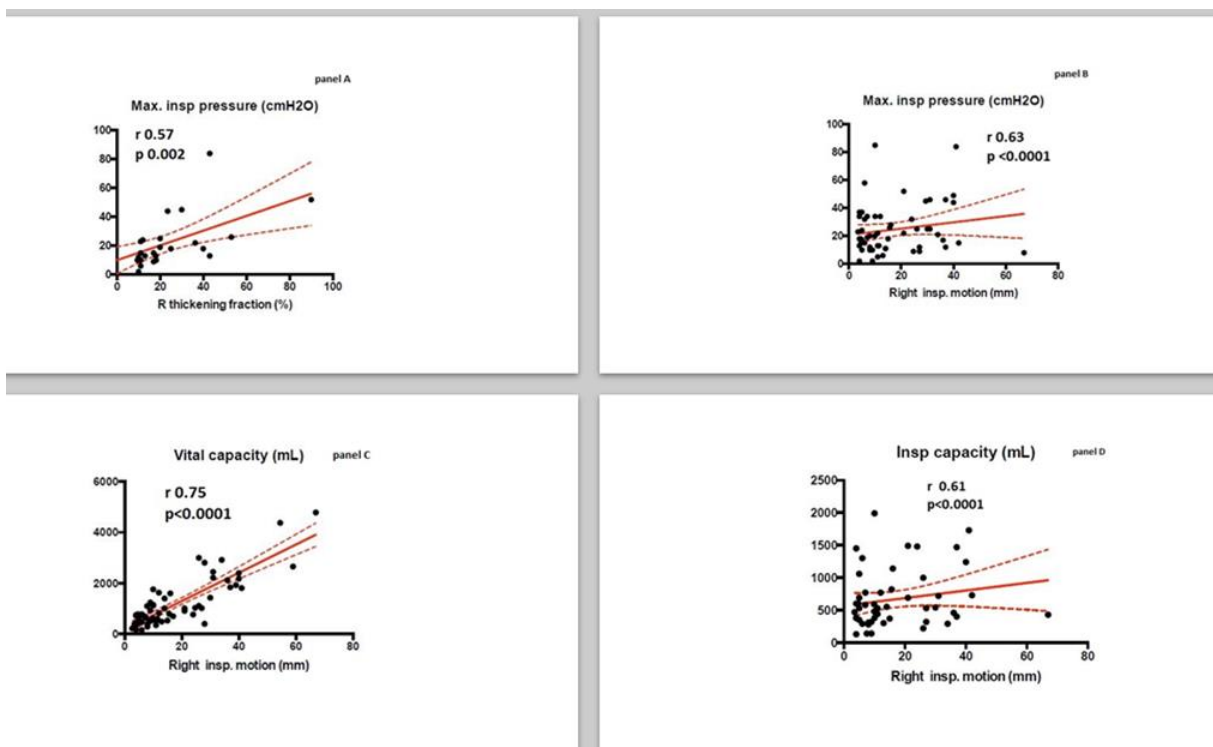
Figure 1: *flow chart* diagram of patients included in the study



DMD: Duchenne muscular dystrophy

Figure 2:

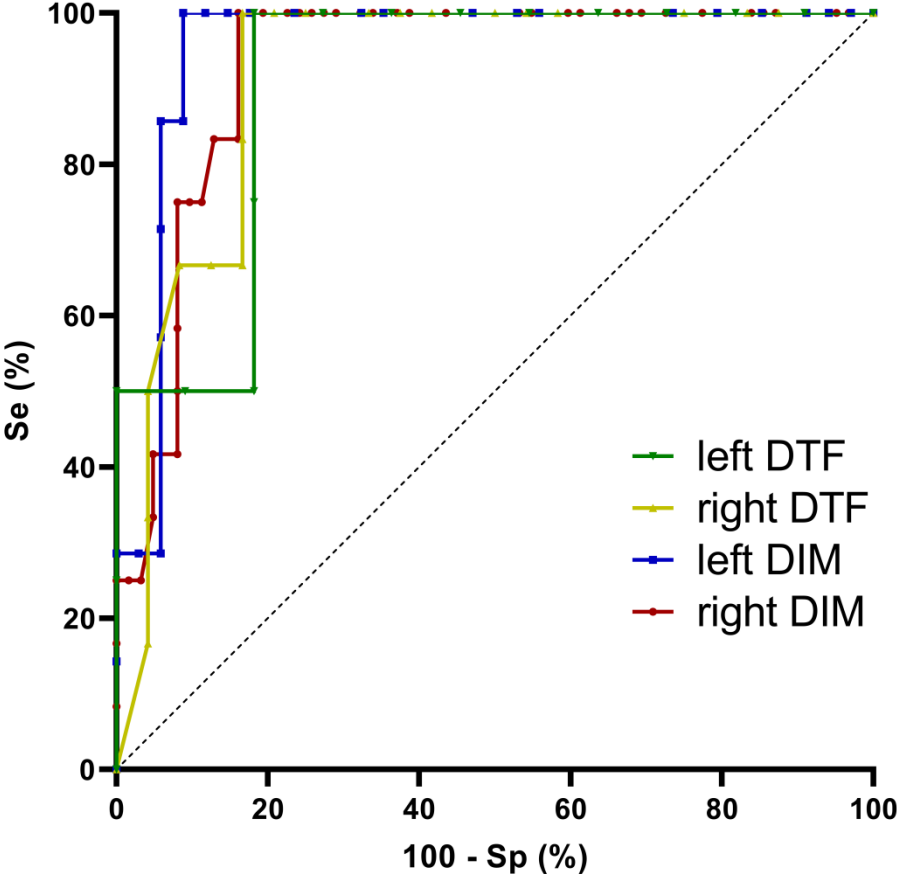
Panel A: Relationship between diaphragm R (right) thickening fraction and max inspi pressure (maximal inspiratory pressure). **Panel B:** Relationship between right diaphragm inspi (inspiratory) motion and max inspi pressure (maximal inspiratory pressure). **Panel C:** Relationship between pulmonary forced vital capacity (ml) and diaphragm right inspi (inspiratory) motion (mm). **Panel D:** Relationship between right diaphragm inspiratory motion and inspiratory capacity



Right diaphragm inspi motion= right diaphragm excursion

Diaphragm inspiratory motion= diaphragm excursion

Figure 3: ROC curves of diaphragm ultrasound parameters to predict pulmonary upright FVC>50%.



ROC: Receiver Operator Characteristic. FVC: forced vital capacity. DIM: diaphragm inspiratory motion= diaphragm excursion. DTF: diaphragm thickening fraction

TABLES

Table 1: clinical, spirometric and ultrasound parameters of patients included in the study

| Parameters | N (number of patients included) | Median [range] or n (%) |
|---|--|--------------------------------|
| Age (years) | 74 | 22[19-25] |
| BMI (kg/m ²) | 61 | 19.5[15.4-25] |
| Wheelchair bound | 74 | 68 (92%) |
| GMW score | 73 | 7 [7-8] |
| Steroids*** | 74 | 14 (19%) |
| Spinal fusion | 55 | 39(71%) |
| Nocturnal non-invasive ventilation | 74 | 52 (70%) |
| Nocturnal and diurnal non-invasive ventilation | 74 | 13 (17.6%) |
| Upright FVC (ml) | 63 | 780[500-1620] |
| Upright FVC (%) | 63 | 17[11-35] |
| Supine FVC (%) ** | 37 | 16[11-30] |
| FEV1 (ml) | 48 | 590[460-1090] |
| IC(ml) | 50 | 530[370-770] |
| IC (%) | 49 | 16[11-22] |
| PEF (l/s) | 51 | 2.3[1.1-3.4] |
| RV(ml) | 33 | 1400[1240-1860] |
| MIP (cmH2O) | 56 | 20[12-34] |
| Bicarbonate (mmol/l) | 46 | 25.6[23-28] |
| LVEF (%) | 74 | 50[42-60] |
| sPAP (mmHg) | 26 | 20.5[18-25] |
| Right diaphragm thickness* (mm) | 44 | 1.5 [1-2] |
| Right diaphragmatic TF (%) | 30 | 18[11-36.3] |
| Right deep diaphragmatic inspiratory motion(mm) | 74 | 11[7-27] |
| Left diaphragmatic thickness*(mm) | 30 | 1.4 [1-1.7] |
| Left diaphragmatic TF (%) | 15 | 18.5[12.5-46] |
| Left deep diaphragmatic inspiratory motion(mm) | 41 | 8[6-20] |

TF: thickening fraction; BMI, body mass index; FVC, forced vital capacity; IC, inspiratory capacity; FEV1, forced expiratory volume in 1 s ; MIP: maximal inspiratory pressure; PEF: peak expiratory flow; RV: residual volume; LVEF: left ventricular ejection fraction; sPAP: systolic arterial pulmonary pressure; GMW: Gardner Medwin Walton score

*: end expiratory; ** % of theoretical in upright

***: prednisone (0.75 mg/kg/day) was used for all patients expect for one who was treated with deflazacort

Table 2: Correlations between diaphragm parameters and clinical parameters, spirometric and cardiac parameters

| Para meters | Age (y) | BMI (kg/m²) | GMW Score | LVEF (%) | IC (%) | Upright FVC (%) | Supine FVC (%*) | MIP (cmH20) |
|--|---------------------|-----------------------------------|----------------------|-------------------|------------------------|----------------------------|----------------------------|------------------------|
| Right diaph TF (%) | - 0.58 (p 0.001) | 0.18 (p 0.38) | -0.45 (p 0.012) | 0.39 (p 0.03) | 0.56 (p 0.004) | 0.68 (p <0.0001) | 0.56 (p 0.038) | 0.57 (p 0.002) |
| Left diaph TF (%) | -0.76 (p 0.001) | 0.34 (p 0.27) | -0.75 (p0.001) | 0.70 (p 0.004) | 0.67 (p 0.012) | 0.75 (p 0.002) | 0.9 (p 0.037) | 0.79 (p 0.004) |
| Right diaph motion (mm) | -0.5 (p <0.0001) | 0.38 (p 0.002) | -0.53 (p<0.0001) | 0.06 (p 0.6) | 0.62 (p<0.0001) | 0.77 (p <0.0001) | 0.64 (p<0.0001) | 0.63 (p<0.0001) |
| Left diaph motion (mm) | -0.53 (p<0.0001) | 0.41 (p 0.016) | -0.59 (p<0.0001) | 0.025 (p 0.87) | 0.79 (p <0.0001) | 0.85 (p <0.0001) | 0.79 (p<0.0001) | 0.69 (p<0.0001) |

R, Spearman's rho; TF, thickening fraction; diaph, diaphragm; BMI, body mass index; LVEF, left ventricular ejection fraction; IC, inspiratory capacity; FVC, forced vital capacity; MIP, maximal inspiratory pressure; y, years; %*, % of theoretical in upright; GMW: Gardner Medwin Walton score