

Gruppo di Ricerca Ecografia Toracica nell'Anziano (GRETA)





Simone Scarlata

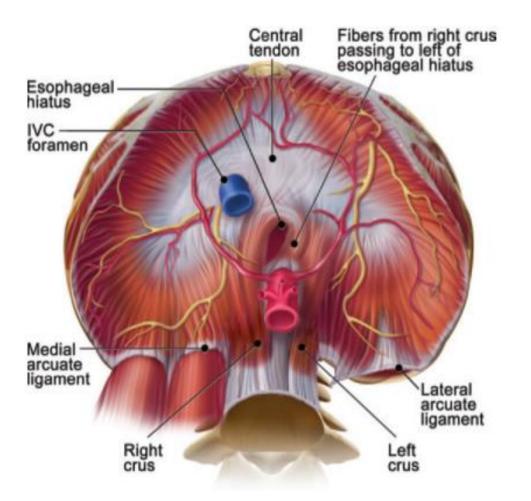
Firenze, <u>13 Dicembre 2023</u> Servizio di Fisiopatologia Respiratoria ed Endoscopia Toracica Dipartimento di Medicina Interna e Geriatria Fondazione Policlinico Universitario Campus Bio Medico di Roma

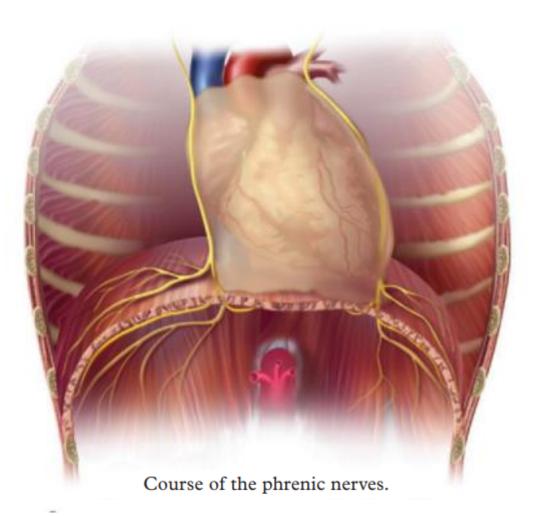




Gruppo di Ricerca Ecografia Toracica nell'Anziano

Imaging of the Diaphragm: Anatomy and Function¹





E51

CAMPUS





Paralysis and Weakness

Eventration

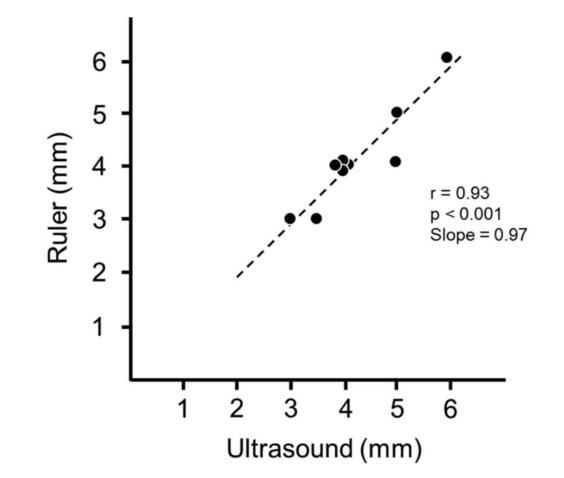
Eventration is a congenital thinning of the diaphragmatic muscle that causes a focal bulge.

Location of the lesion	Disease	Comment
Phrenic	Trauma ³⁰ Compression/infiltration (mediastinal neoplasms) Guillain-Barré ³¹ Infection (Herpes-Zoster virus, pneumonias, ³² Lyme disease, ³³ HIV infection ³⁴) Amyotrophic neuralgia (Parsonage-Turner) ³⁵ Thoracic surgeries ³⁶ Others [malnutrition, ³⁷ diabetes, ³⁸ hypothyroidism, ²³ benign thyroid hypertrophy, ³⁹ porphyria, vasculitis, Charcot-Marie-Toot ⁴¹ disease] Idiopathic ⁴⁶	Guillain-Barré disease is the most frequent cause of acute respiratory muscle paralysis. More than 25% of patients will need non-invasive ventilation. Infection with Herpes-Zoster virus produces diaphragmatic paralysis if it affects the cervical territory and is usually ipsilateral and usually permanent. Amyotrophic neuralgia presents pain and flaccid paralysis of the shoulder muscles. It is associated with both uni and bilateral diaphragmatic involvement. In idiopathic causes the paralysis can be unilateral or bilateral.
Lung	Asthma and chronic obstructive pulmonary disease	The existing pulmonary hyperinflation can deteriorate the diaphragmatic function since the diaphragm does not have an optimal length for its normal functioning. ¹
Neuromuscular junction Muscular	Myasthenia gravis, botulism, ⁴⁰ Lambert–Eaton syndrome ²⁰ Muscular dystrophies, steroid myopathy, ⁴² Pompe disease, ⁴³ myositis, mechanical ventilation ⁴⁵	During an acute myasthenic crisis there may be acute respiratory failure that will require invasive ventilation. In the presence of a diaphragmatic paralysis, acid alfa-glucosidase enzyme levels should be determined to discard late-onset Pompe disease (16.7%) prevalence). ⁴⁴ Mechanical ventilation, both invasive and non-invasive, can produce atrophy of the diaphragm due to disuse.



Ultrasound measurement of diaphragm thickness (zone of apposition)





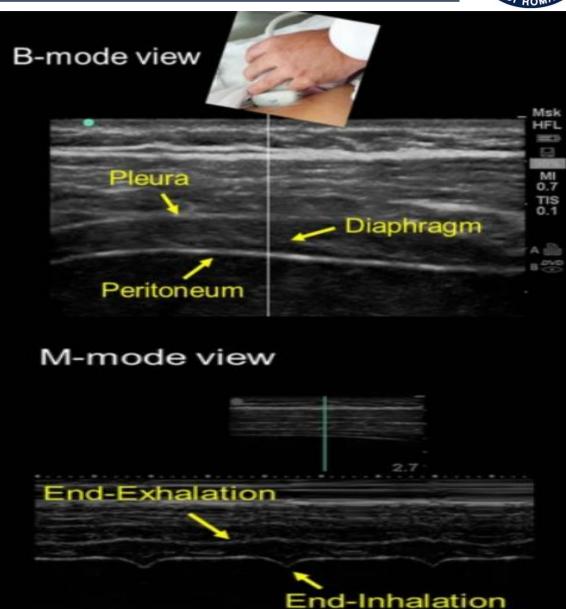
Relationship of diaphragm thickness measured in situ in 10 human cadavers by ultrasound and in vitro by a ruler. Ultrasound measurements of diaphragm thickness in situ are as accurate as measurements in vitro with a ruler.

Reproduced with permission from The American Physiological Society: Wait et al. J Appl Physiol 1989;67(4):1560–1568





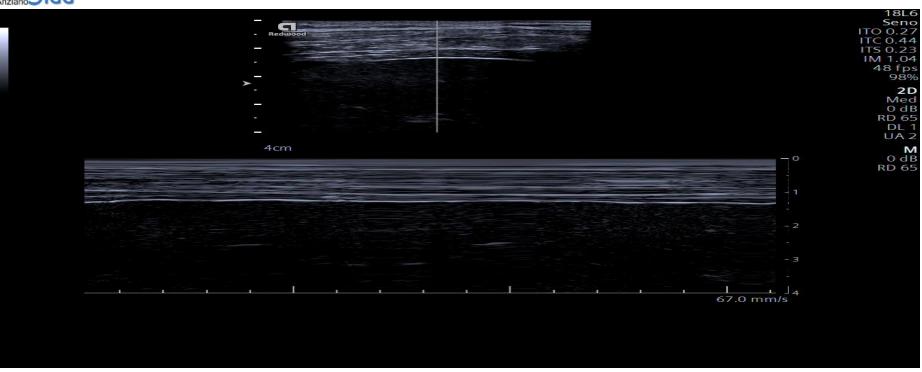
- Operators use linear ultrasound probes to measure diaphragm thickness. These probes use high frequency ultrasound waves (7–18 Hz) to create high resolution images of structures near the body surface .
- To measure diaphragm thickness, operators place the ultrasound probe longitudinally parallel to the long axis of the body, usually between the eighth to tenth intercostal space, at the anterior axillary line or midway between the anterior- and mid-axillary lines.
- The costo-phrenic sinus is identified as the transition between lung and liver (right) or between lung and spleen (left).
- The zone of apposition, where the diaphragm is opposed to the rib cage, is located caudal to the costo-phrenic sinus.
- To identify the diaphragm, subjects are asked to inhale while operators select B-mode imaging.
- The diaphragm is identified as a three-layer structure (two echogenic layers of peritoneum and pleura sandwiching a more hypoechoic layer of the muscle itself) underneath the intercostal muscles that reappear as lung artifact recedes





Ultrasound measurement of diaphragm thickness (zone of apposition)



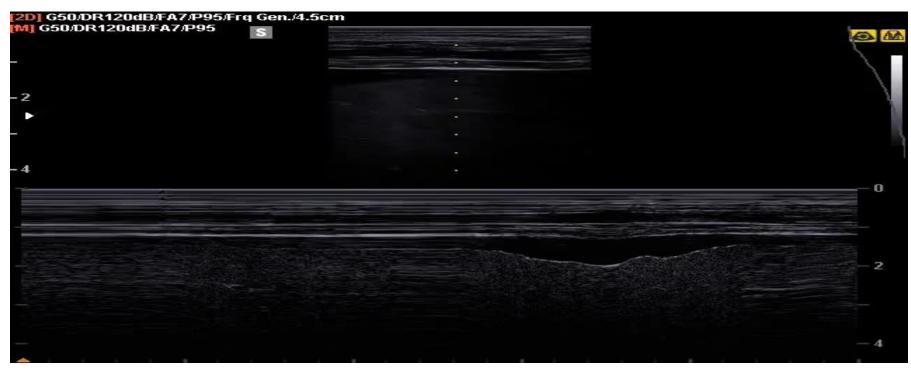


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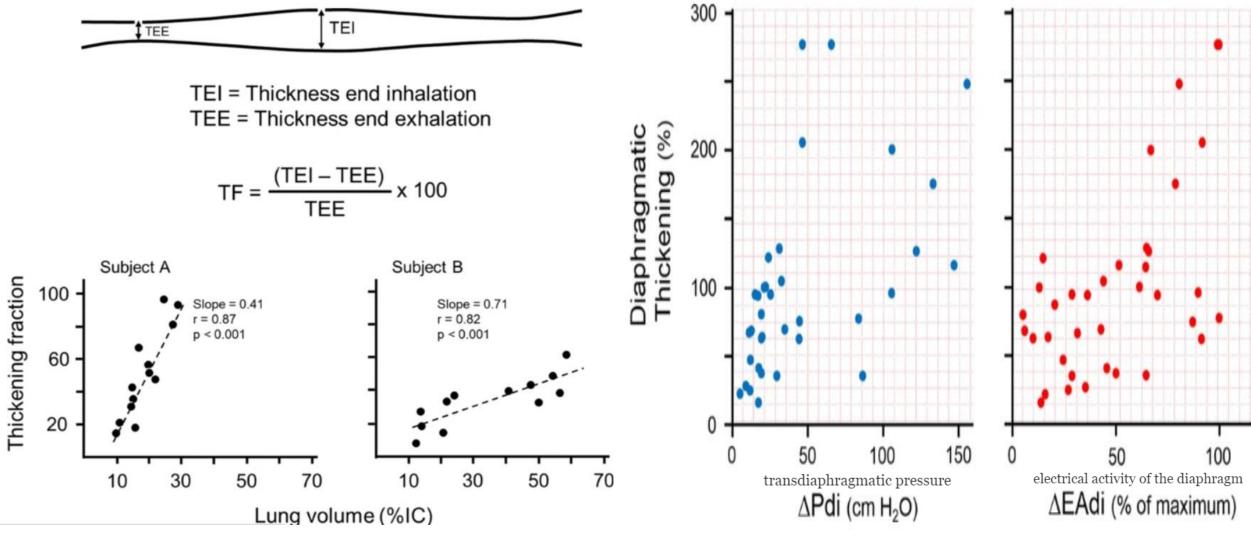




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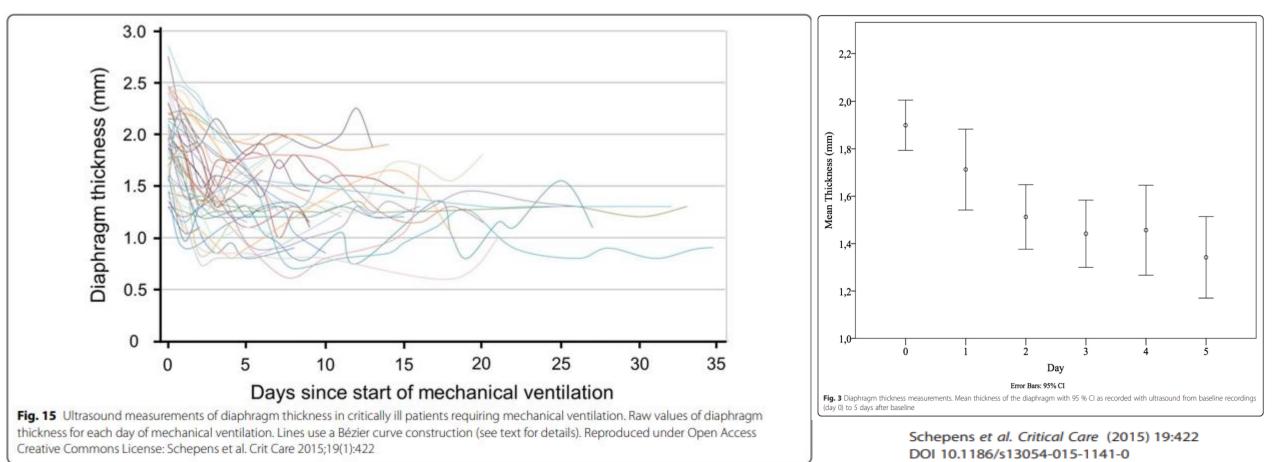


($r^2 = 0.32$ and 0.28, respectively, p < 0.01).





The course of diaphragm atrophy in ventilated patients assessed with ultrasound: a longitudinal cohort study

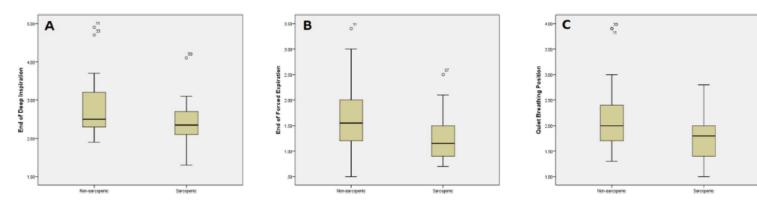






Diaphragmatic muscle thickness in older people with and without sarcopenia

Olgun Deniz^{3,1} · Suheyla Coteli¹ · Nur Betul Karatoprak² · Mehmet Can Pence² · Hacer Dogan Varan¹ · Muhammet Cemal Kizilarslanoglu¹ · Suna Ozhan Oktar² · Berna Goker¹



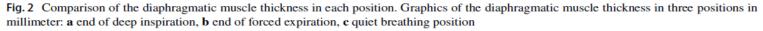


Table 3Ultrasonographicassessment of diaphragmaticthickness, PEF results, andskeletal muscle index of theparticipants

	Sarcopenic $(n=30)$	Non-sarcopenic $(n=30)$	р
End of deep inspiration (mm)	2.3 (1.3-4.1)	2.5 (1.9-4.9)	0.02
End of forced expiration (mm)	1.1 (0.7–2.5)	1.5 (0.5-3.4)	< 0.01
Quiet breathing position (mm)	1.8 (1.0-2.8)	2.0 (1.3-3.9)	0.02
Peak expiratory flow rate (lt/min)	245 (150-500)	310 (220-610)	< 0.01
Skeletal muscle mass index (SMI) (kg/m ²)	7.64 ± 0.83	9.22 ± 1.31	< 0.01

Normally distributed variable was presented as mean \pm standard deviation while skew distributed ones presented as median (min-max)

Bold values denote statistical significance at the p < 0.05 level

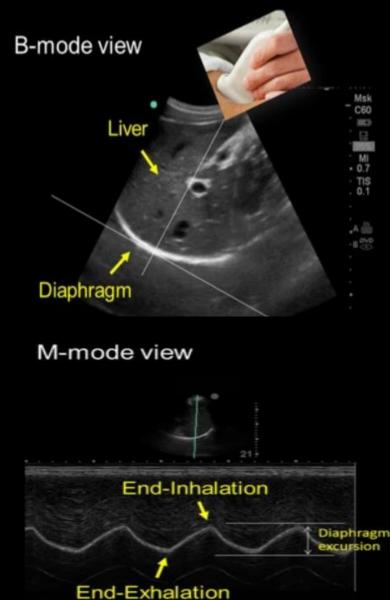
Table 1 Baseline	characteristics	of the	participants
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	Sarcopenic (<i>n</i> =	: 30)	Non- sarcopenic $(n=30)$	р
Age (years)	79.9±5.6		73.3±5.5	< 0.01
BMI (kg/m ²)	23.9 ± 3.6		29.4 ± 4.3	< 0.01
Gender				
Women	16 (53)		19 (63)	0.43
Men	14 (47)		11 (37)	
Diabetes mellitus	8 (26.7)		9 (30.0)	0.39
Hypertension	23 (76.7)		20 (66.7)	0.39
Osteoporosis	10 (34.5)		8 (28.6)	0.6
Coronary artery disease	11 (36.7)		4 (13.3)	0.04
Chronic heart failure	2 (6.7)		0 (0)	0.49
Dementia	2 (6.7)		0 (0)	0.49
Depression	2 (6.7)		2 (6.7)	1.00
Smoking	10 (33.4)		10 (33.4)	1.0
Number of medication	6 (0–15)		4 (0-8)	< 0.01
Katz ADLs	DLs 5(1-6) 6		-6)	< 0.01
Lawton-Brody IADLs	6 (1-8)	8 (5-8)		< 0.01
MNA-SF	10 (5-14)	13 (11-14)		< 0.01
MMSE	23.5 (8-30)	28 (22-30)		< 0.01
GDS	5.5 (0-15)	2 (0-12)		0.05
Gait speed (m/sn)	0.84 (0.17-2)	1.39 (1.03-1.93)		< 0.01
Hand grip strength (kg)	18.6 ± 6.5	24.5	±7.0	< 0.01
Right calf circumference (cm)	32.6±2.9	37.2	±3.2	< 0.01





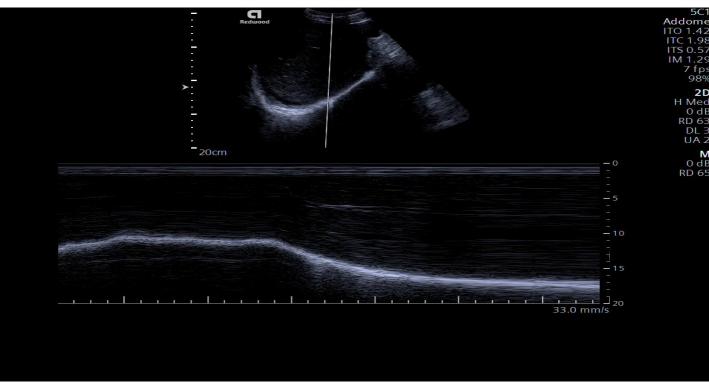
- The cranio-caudal movement of the dome of the diaphragm during quiet breathing and during forceful inspiratory efforts such as sniff maneuvers or maximal inspirations can be monitored using curvilinear ultrasound probes.
- Curvilinear probes use low frequency ultrasound waves (2– 6 Hz) that penetrate deeply in the body giving a wide depth of field.
- On the right, operators position the probe longitudinally in the subcostal area between the mid-clavicular and anterior axillary lines using the liver as acoustic window.
- The probe is directed medially, cephalad and dorsally so that the ultrasound beam reaches the right dome of the diaphragm perpendicularly.
- On the left side, operators use the spleen as an acoustic window. (Less often, operators may use the right or left lateral view (midaxillary lines) or the posterior subcostal view or the subxiphoid view).
- Once a good quality B-mode image is obtained, operators adjust the M-mode interrogation line as to be perpendicular to the movement of the hemidiaphragm. With M-mode ultrasonography, the diaphragm appears as a single thick echogenic line.
- During inhalation the contracting diaphragm moves towards the ultrasound probe.





Ultrasound measurement of diaphragm motion (dome)



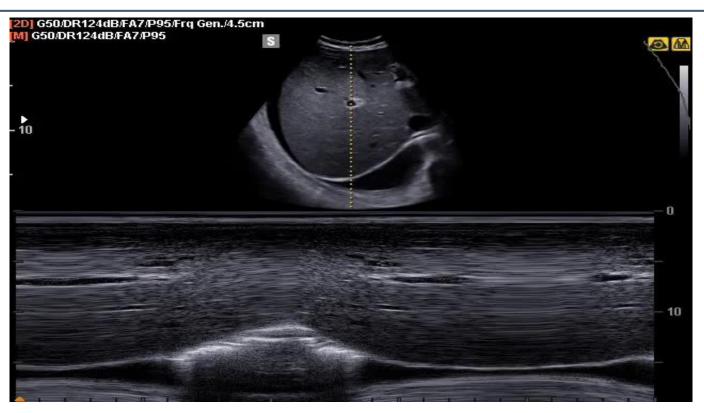


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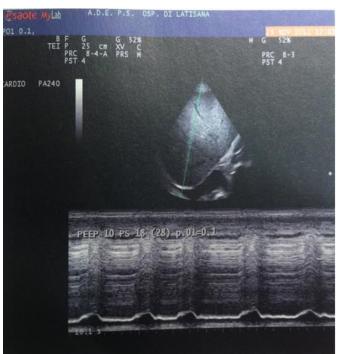




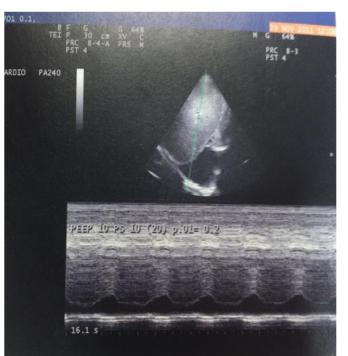
DIAFRAMMA

Lo studio del diaframma in corso di ventilazione meccanica

PEEP 18 cmH2O



PEEP 10 cmH2O



PEEP 6 cmH2O

