

L'électrothérapie

**Controverse
CONTRE**

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AUCUN CONFLIT D'INTERÊT

LAROUSSE

**électrothérapie**

nom féminin

Traitement utilisant l'énergie électrique ou l'enregistrement de l'activité électrique musculaire.



TOUS LES AVIS

Par **Véronique D.** le 23 Nov. 2022 :

(5/5) ★★★★★

Produit évalué : **Multisport Pro Filaire sport électrostimulation musculaire**

fonctionne très bien

je suis très satisfaite de ce produit. J'ai repris du muscle et c'est très gratifiant. Je recommande.

L'ÉLECTROSTIMULATION

L'électrostimulation permet de faire travailler la masse musculaire par zone pour des effets de massage de préparation avant effort, de récupération après efforts et de musculation.



2

T'es pas censé défendre
le contre?

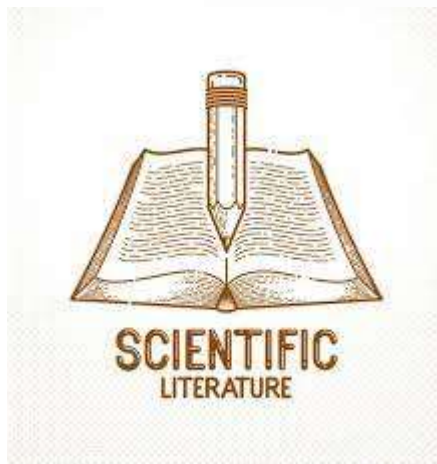


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Appuie toi sur la
LITTÉRATURE SCIENTIFIQUE !



Mon premier argument scientifique

Les muscles des vrais patients de réanimation ne sont répondeurs pas à l'électrostimulation

Pourquoi cela ne fonctionne pas?

We were able to show that at the initiation of NMES, only two thirds of stimulations led to a contractile response in our cohort and that this number declined to one third throughout the first 7 days of stimulation, which we think is due to progressing pathophysiological processes. Established mechanisms that would hypothetically contribute to this effect are channelopathy, decreasing metabolic flexibility, advanced muscle atrophy and edema.

Table 1 Baseline characteristics

n		21
Sex (m/f)		16/76.2% / 5/ 23.8%
Age (years)		53.0 (45.0/70.0)
Weight (kg)		85.0 (75.0/100.0)
Height (m)		1.78 (1.74/1.80)
BMI (kg/m ²)		27.1 (24.2/30.9)
Diagnosis responsible for ICU admission		
	ARDS	8/38.1%
	Sepsis	3/14.3%
	Multiple trauma	7/33.3%
	Neurologic	2/9.5%
	Miscellaneous	1/4.8%
SOFA at ICU admission		13.0 (11.0/15.0)
APACHE II at ICU admission		25.0 (20.0/28.0)
SAPS2 at ICU admission		58.0 (47.0/65.0)
GCS at ICU admission		5.0 (3.0/6.0)
Time until first awakening (days)		17.0 (10.0/25.0)
ICU length of stay (days)		32.0 (21.0/43.0)
Percent of days with RASS > -3 during ICU stay		64.3 (37.5/79.3)
Noradrenalin (µg/kg min)		0.07 (0.05/0.11)
Time requiring noradrenalin (days)		12.0 (9.0/18.0)
Survivors/non-survivors		18/85.7% / 3/ 14.3%
Non-excitabile muscle membrane/excitabile muscle membrane		7/50% / 7/50%
Start of NMES treatment after ICU admission (days)		3.0 (2.0/6.0)

GRUNOW et al. *Critical Care* (2019) 23:309
<https://doi.org/10.1186/s13054-019-2540-4>

Critical Care

RESEARCH

Open Access

Differential contractile response of critically ill patients to neuromuscular electrical stimulation

Julius J. Grunow¹, Moritz Goll¹, Niklas M. Carbon¹, Max E. Liebl², Steffen Weber-Carstens^{1,3} and Tobias Wollersheim^{1,4*}

Check for updates

Patients

a Mechanically ventilated patients ≥ 18 years of age were included on the basis of a Sepsis-related Organ Failure Assessment (SOFA) score ≥ 9 within the first 72 h after ICU admission as well as written informed consent by a legal proxy and excluded if any of the following applied: prior hospital treatment for longer than 7 days, illness prohibiting early mobilisation, pre-existing neuromuscular disease, insulin-dependent diabetes mellitus, Body Mass Index > 35 kg/m², not ambulating before admission or poor prognosis with a high likelihood of death within the next hours.

relative contractile response



C

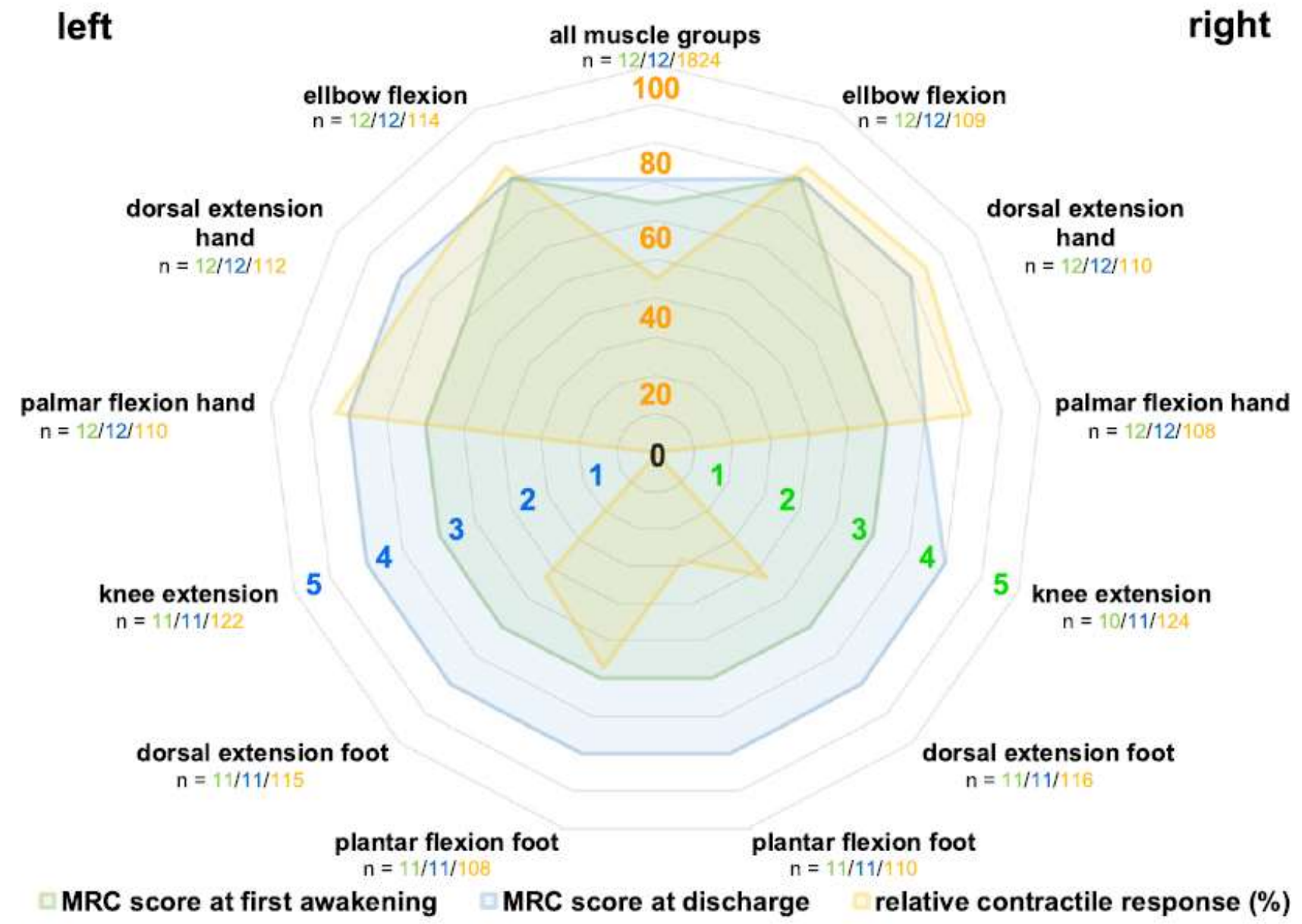
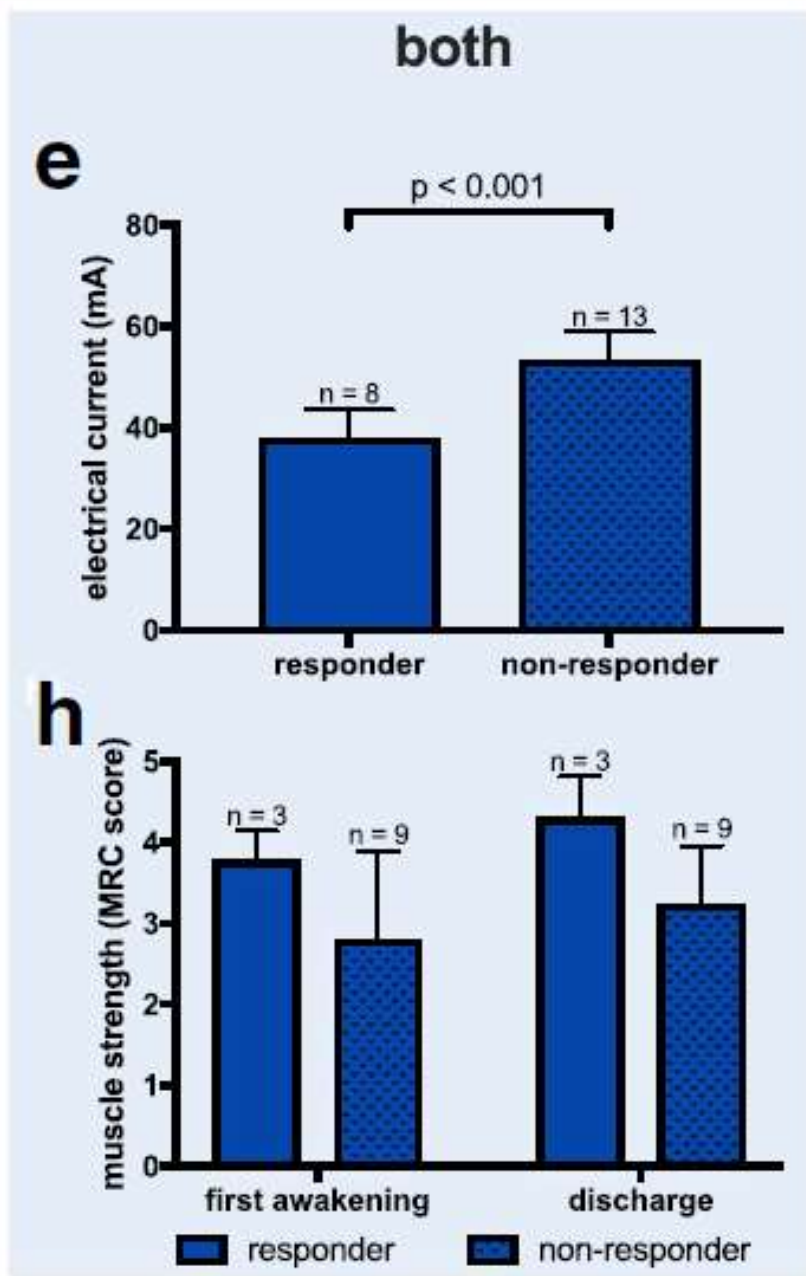
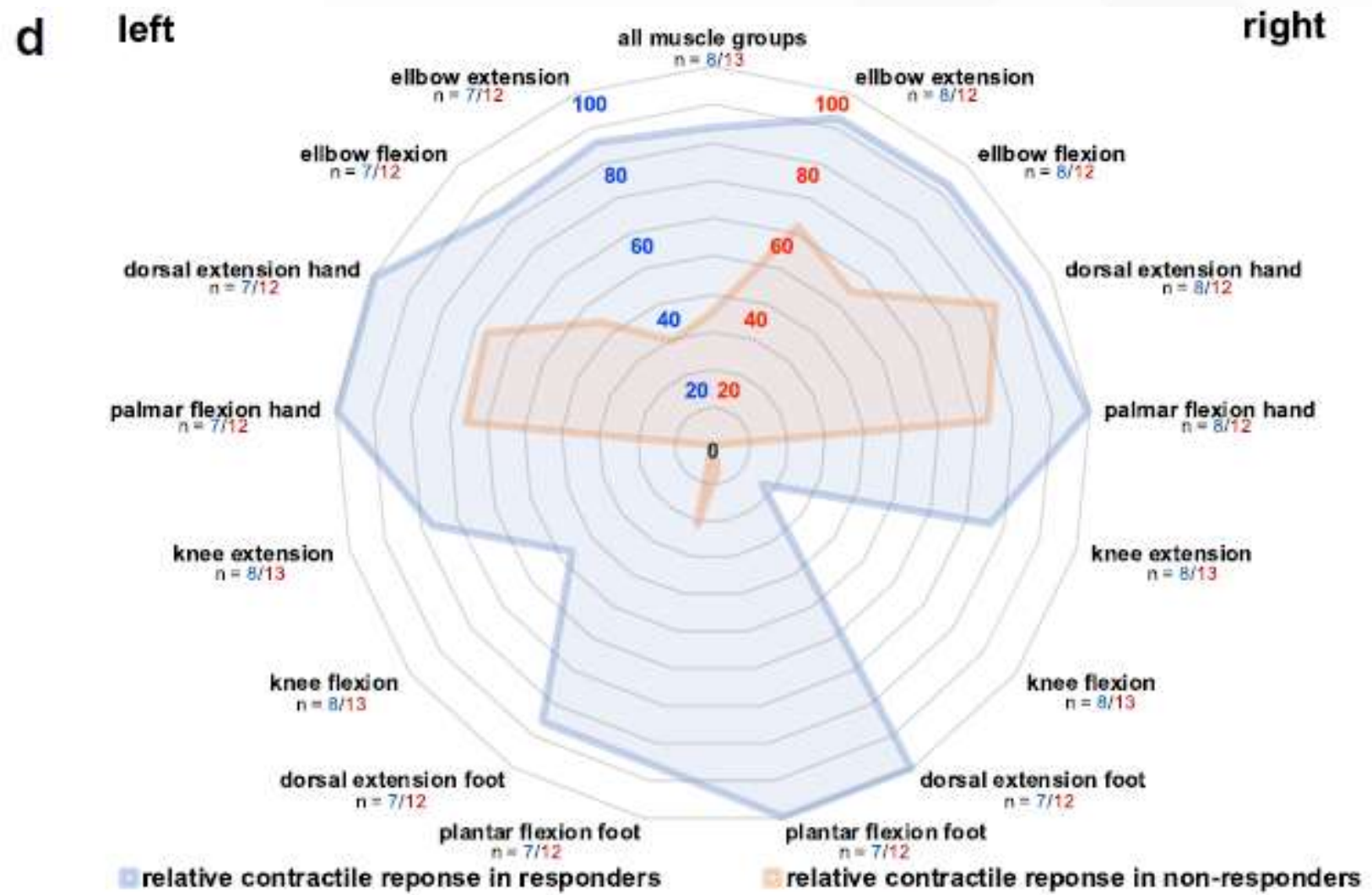


Table 2 Univariate analysis

		Responder	Non-responder	<i>p</i> value
Patients/stimulations (<i>n</i>)		8/702	13/1122	
Diagnosis responsible for ICU admission	ARDS	2/25.0%	6/46.2%	0.118
	Sepsis	0/0.0%	3/23.1%	
	Multiple trauma	4/50.0%	3/23.1%	
	Neurologic	2/25.0%	0/0.0%	
	Miscellaneous	0/0.0%	1/7.7%	
SOFA at ICU admission		12.0 [9.5/13.5]	14 [12.0/16.0]	0.030



Pour que le courant circule, il faut que le patient ne soit

Pas grave (sepsis...)

Pas en surpoids

Sans pacemaker

Non enceinte/allaitante

Non hospitalisé depuis plus de 7 jours avant la réanimation...

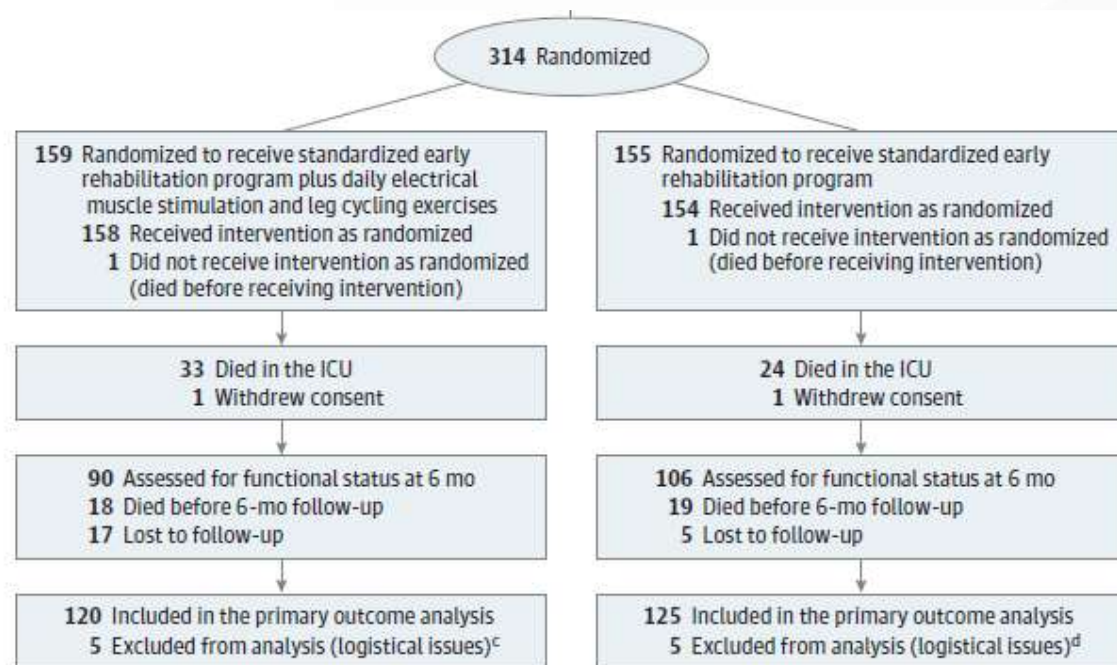
Ne présentant pas une faiblesse musculaire au réveil

Donc pas un patient de réanimation!

Argument n°2

Même dans un grand échantillon

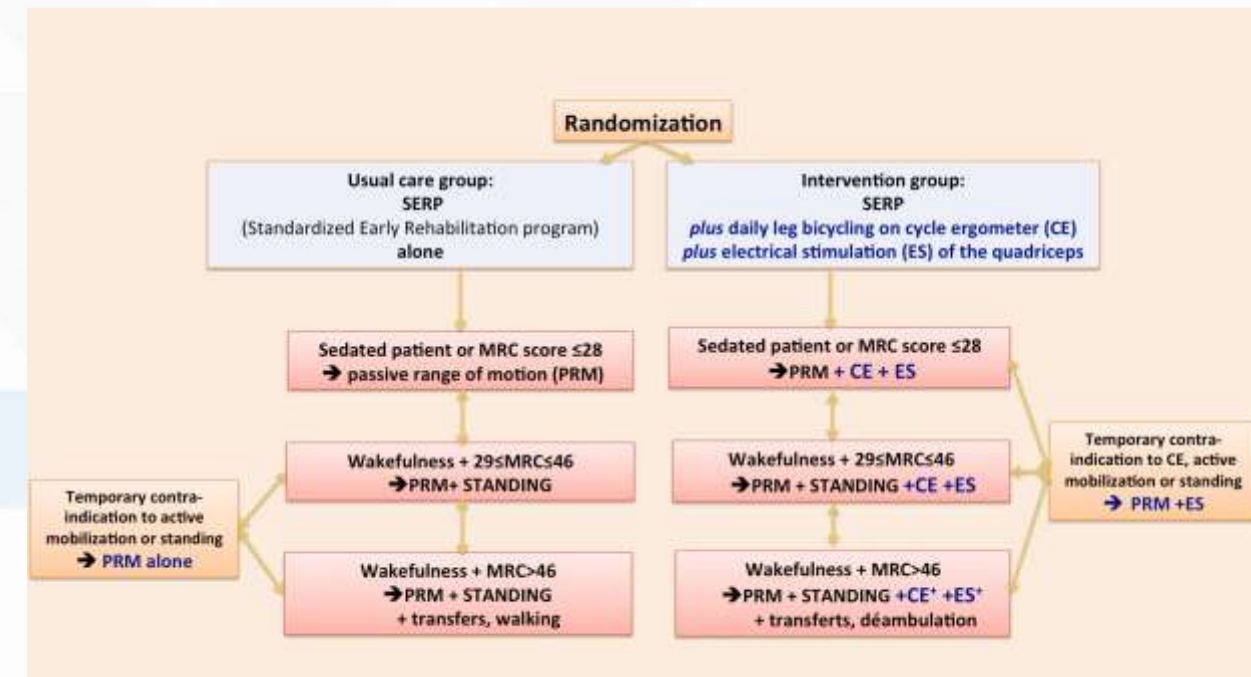
L'électrothérapie ne fonctionne pas



JAMA | Preliminary Communication

Effect of In-Bed Leg Cycling and Electrical Stimulation of the Quadriceps on Global Muscle Strength in Critically Ill Adults A Randomized Clinical Trial

Guillaume Fossat, PT; Florian Baudin, PT; Léa Courtes, PT; Sabine Bobet, PT; Arnaud Dupont, PT; Anne Bretagnol, MD; Dalila Benzekri-Lefèvre, MD; Toufik Kamel, MD; Grégoire Muller, MD; Nicolas Bercault, MD; François Barbier, MD, PhD; Isabelle Runge, MD; Mai-Anh Nay, MD; Marie Skarzynski, MD; Armelle Mathonnet, MD; Thierry Boulain, MD



- Electrical stimulation of the quadriceps following the 54-min “atrophy prevention program”
 - Sedated patient (Ramsay score > 3 and RASS < 0) or in awaking phase and $MRC \leq 28$
 - Patient in the awakening phase and $29 \leq MRC \leq 46$
 - Awake patient and $29 \leq MRC \leq 46$
- Electrical stimulation of the quadriceps following the 20-min “muscle building” program
 - Awake and $47 \leq MRC$

JAMA | Preliminary Communication

Effect of In-Bed Leg Cycling and Electrical Stimulation of the Quadriceps on Global Muscle Strength in Critically Ill Adults
A Randomized Clinical Trial

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	Median (IQR) ^a		Between-Group Difference (95% CI) ^b	P Value
	Intervention Group (n = 158)	Usual Care Group (n = 154)		
Primary Outcome				
Global MRC score at ICU discharge among ICU survivors ^{12,c}	48 (29 to 58)	51 (37 to 58)	MD, -3.0 (-7.0 to 2.8)	.28 ^d
Change from study inclusion to ICU discharge				
Rectus femoris muscle thickness, mm	-1.9 (-8.0 to 0.2)	-2.4 (-7.1 to -0.3)	MD, -0.5 (-1.0 to 2.4)	.17 ^d
Invasive mechanical ventilation, d				
Intention-to-treat population	5 (2 to 10)	5 (2 to 11)	MD, 0 (-2.0 to 1.5)	.69 ^h
ICU survivors				
Frequency of reintubation within 48 h after first extubation, No./total (%) [95% CI]	12/131 (9.2)[5.3 to 15.3]	13/133 (9.8)[5.8 to 16.0]	ARR, 0.7 (-6.7 to 7.9)	>.99 ^g

Argument n°3

Même après la réanimation

L'ES ne fonctionne pas

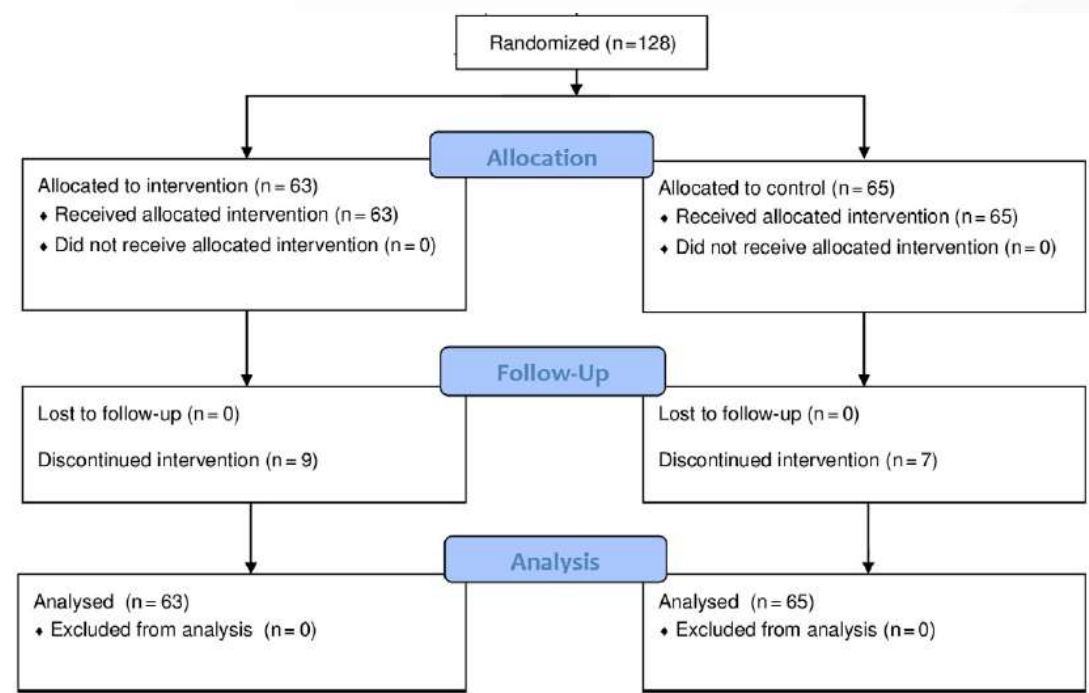


Fig. 1. Flow chart diagram of the patients discharged from the ICU during the study period.

Effect of neuromuscular stimulation and individualized rehabilitation on muscle strength in Intensive Care Unit survivors: A randomized trial

Irini Patsaki, MSc^a, Vasiliki Gerovasili, MD^{b,*}, Georgios Sidiras, PhD^a, Eleftherios Karatzanos, PhD^b, Georgios Mitsiou, MSc^a, Emmanuel Papadopoulos, PhD^a, Anna Christakou, PhD^a, Christina Routsis, MD^a, Anastasia Kotamidou, MD^a, Serafim Nanas, MD^a

2.5. Intervention

NMES implementation has been described in detail [28]. NMES was implemented daily (7 days/week) for 55 min on rectus femoris and peroneus longus of both lower extremities. The stimulator used was Rehab 4 Pro (Cefar Medical AB, Malmö, Sweden) and delivered biphasic, symmetric impulses of 45 Hz, with 400 μ sec pulse duration, 12 s on (0,8 s rise time and 0,8 s fall time) and 6 s off. The intensity levels were aimed to cause visible contractions and be tolerated by the patients. In case of doubt, contraction was confirmed by palpation.

Patients in the control group received sham NMES along with usual care until hospital discharge. Sham NMES was implemented daily (7 days/week) for 55 min on the same muscle groups with the NMES group, but the intensity was set at the minimum value of 1–5 mA, without palpable contractions.

Table 2
The physical rehabilitation program of both groups.

		Intervention group	Control group
Range of motion (ROM)	Upper extremity	Passive and active 10–20 repetitions (resistance when tolerated)	No 10 repetitions (no resistance)
	Lower extremity	Shoulder flexion/abduction Elbow flexion/extension Wrist flexion/extension Hip flexion/abduction/adduction Knee flexion Ankle dorsiflexion/plantar flexion	Shoulder flexion/abduction Elbow flexion/extension Hip flexion Knee flexion Ankle dorsiflexion/plantar flexion
Functional exercises		Bed transfers from side to side, transfers to chair, coming to standing position	Sited position at the edge of the bed
Ambulation		With or without mobility aid	Only if patient able to ambulate with minimal assistance
Balance exercises		From sitting and standing position	No
Frequency		40 min	20 min
Duration		5 days/week until hospital discharge	5 days/week until hospital discharge

2.4. Primary and secondary outcomes

Primary outcome of the study was muscle strength assessed with the MRC score and handgrip dynamometry at hospital discharge.
Secondary outcomes were functional ability assessed with the Functional Independence Measure (FIM) and hospital length of stay.

Table 3
MRC score at ICU discharge and weekly until hospital discharge in patients assigned to the NMES (n = 63) group and the control group (n = 65).

	MRC			ΔMRC%		
	NMES group	Control group	p	NMES group	Control group	p
ICU discharge	50 ± 13 (n = 63)	51 ± 10 (n = 65)	0.5			
1 week post ICU	53 ± 11 (n = 51)	53 ± 9 (n = 56)	0.98	13% ± 22 (n = 51)	7% ± 12 (n = 56)	0.1
2 weeks post ICU	52 ± 10 (n = 28)	51 ± 9 (n = 30)	0.83	34% ± 48 (n = 28)	18% ± 19 (n = 30)	0.1
3 weeks post ICU	51 ± 11 (n = 19)	51 ± 10 (n = 19)	0.95	28% ± 45 (n = 19)	24% ± 26 (n = 19)	0.7
4 weeks post ICU	49 ± 13 (n = 12)	50 ± 9 (n = 14)	0.84	43%59% (n = 12)	59% ± 29 (n = 14)	0.6
5 weeks post ICU	49 ± 14 (n = 9)	53 ± 8 (n = 10)	0.43	57% ± 69 (n = 9)	43% ± 30 (n = 10)	0.55
6 weeks post ICU	48 ± 12 (n = 7)	52 ± 5 (n = 5)	0.46	88% ± 112 (n = 7)	27% ± 17 (n = 5)	0.2
7 weeks post ICU	43 ± 12 (n = 5)	49 ± 4 (n = 4)	0.37	100% ± 120 (n = 5)	27% ± 23 (n = 4)	0.3
8 weeks post ICU	43 ± 14 (n = 4)	48 ± 6 (n = 3)	0.58	108% ± 136 (n = 4)	25%21 (n = 3)	0.3
Hospital discharge	48 ± 21 (n = 63) ^a	50 ± 18 (n = 65) ^a	0.53	3% ± 60 (n = 63)	1% ± 38 (n = 65)	0.85
Hospital discharge (without ITT analysis)	57 ± 7 (n = 54)	57 ± 5 (n = 58)	0.8	19% ± 47 (n = 54)	12% ± 20 (n = 58)	0.3

Values are given as No., mean ± SD (standard deviation); MRC: Medical Research Council score for muscle strength; ICU: Intensive Care Unit; NMES: neuromuscular electrical stimulation.
ΔMRC%: Relative(%) difference in MRC compared to MRC value at ICU discharge. ITT: Intention to treat.
^a Intention to treat analysis: drop outs were assigned a score of 0 for MRC at hospital discharge.

2.4. Primary and secondary outcomes

Primary outcome of the study was muscle strength assessed with the MRC score and handgrip dynamometry at hospital discharge.
Secondary outcomes were functional ability assessed with the Functional Independence Measure (FIM) and hospital length of stay.

5. Conclusion

Table 4
Handgrip dyn

Handgrip d							p
ICU dischar							0,86
1 week pos							0,93
2 weeks po							0,28
3 weeks po							0,96
4 weeks po							0,71
5 weeks po							0,88
6 weeks post ICU	8 ± 9 (n = 8)	9 ± 13 (n = 4)	0.91	22 ± 21	24 ± 29	0,93	
7 weeks post ICU	3 ± 4 (n = 5)	3 ± 3 (n = 4)	0.75	10 ± 11	11 ± 10	0,95	
8 weeks post ICU	5 ± 4 (n = 4)	8 ± 13 (n = 3)	0.66	14 ± 9	32 ± 55	0,54	
Hospital discharge	14 ± 13 (n = 53)	16 ± 11 (n = 58)	0.46	45 ± 29	51 ± 23	0,25	

NMES exercise and personalized physiotherapy in ICU survivors did not result in greater improvement of muscle strength, functional status and length of stay at hospital discharge in comparison to the control group, treated as usual.

Values are given as mean ± SD. Handgrip dynamometry absolute values were transformed to relative values (% predicted), according to the norms provided by Schlusser et al. 2008. SD: standard deviation, ICU: Intensive Care Unit, NMES: neuromuscular electrical stimulation.

^a Two patients were unable to be evaluated due to presence of peripheral intravenous catheter.

ARGUMENT n°4

Même les REVUES SYSTEMATIQUES
montrent cela ne fonctionne pas

Identification

Screening

Eligibility

Included

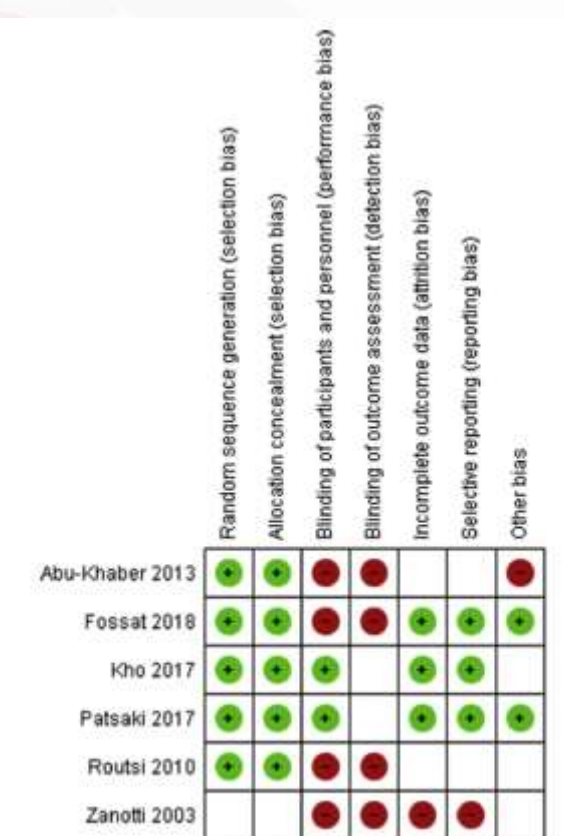
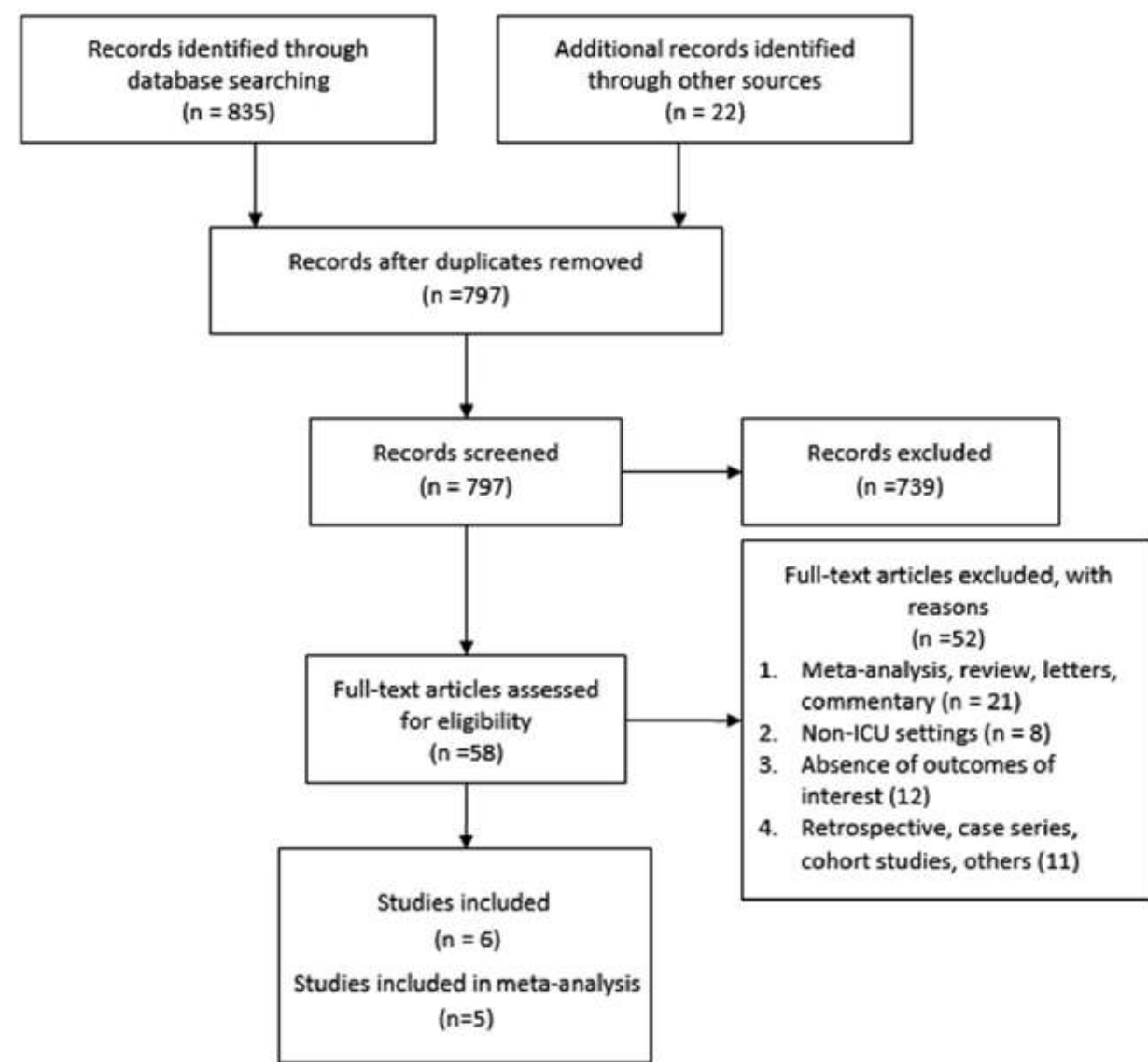


Fig. 2. Risk of bias assessment based on authors' judgement. Blank squares indicate unclear risk of bias.

Fig. 1. Flow diagram for literature search and study selection.

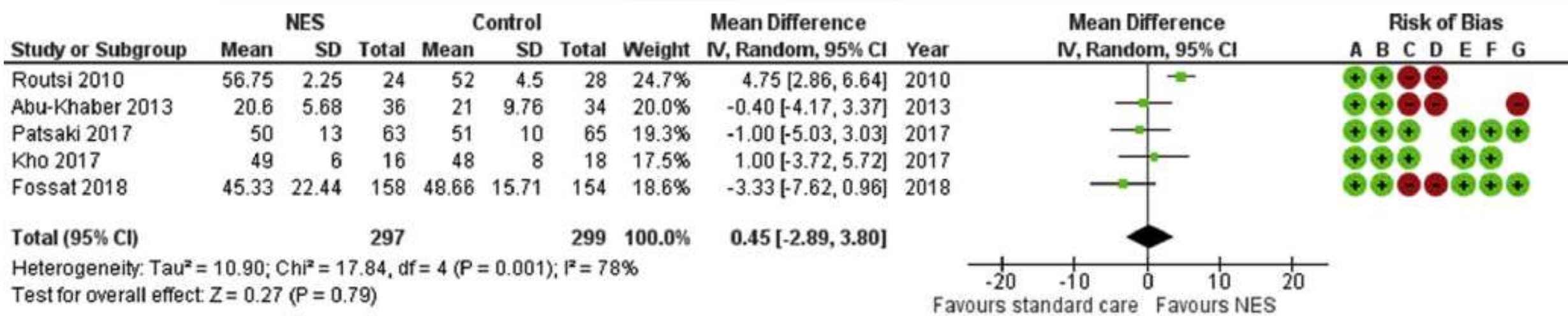


Fig. 3. Forest plots for global muscle strength at ICU discharge measured by modified Medical Research Council (mMRC) grading system.

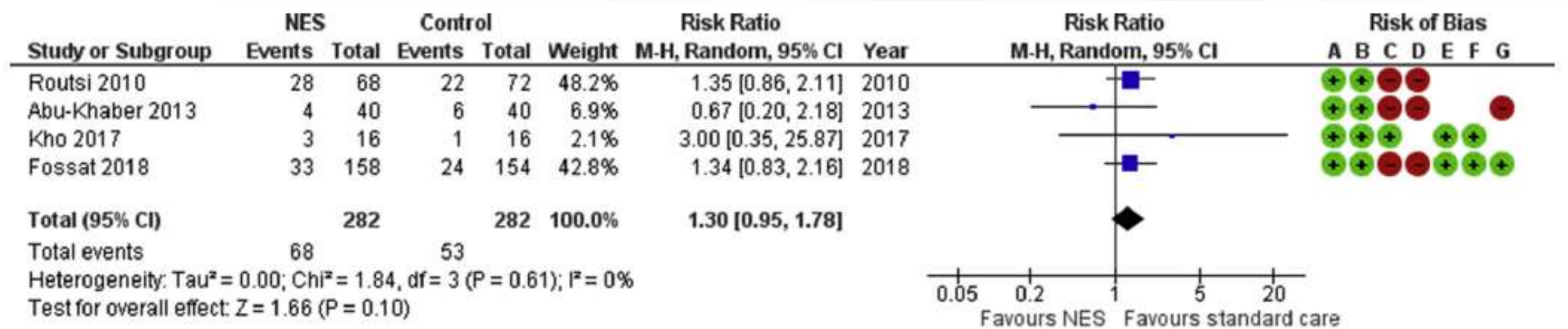


Fig. 4. Forest plot for intensive care unit mortality. NES: neuromuscular electrical stimulation; CI: confidence interval.

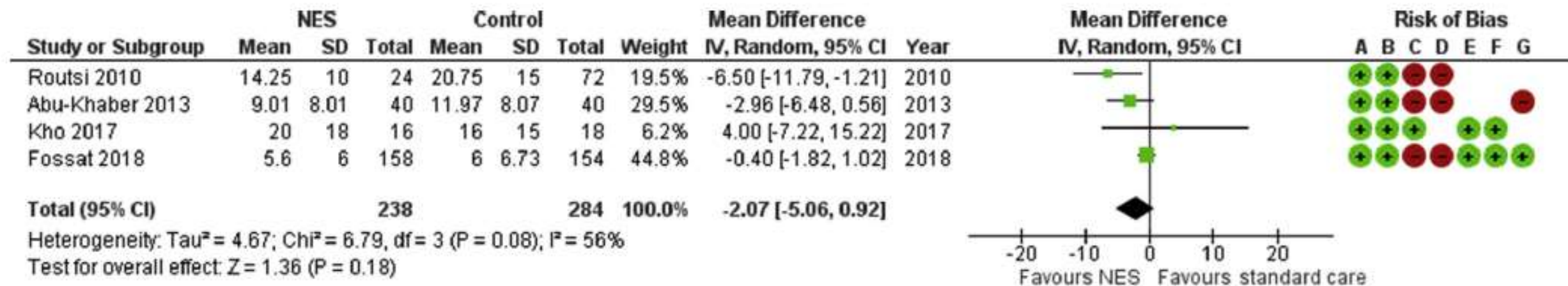


Fig. 5. Forest plot for duration of mechanical ventilation.

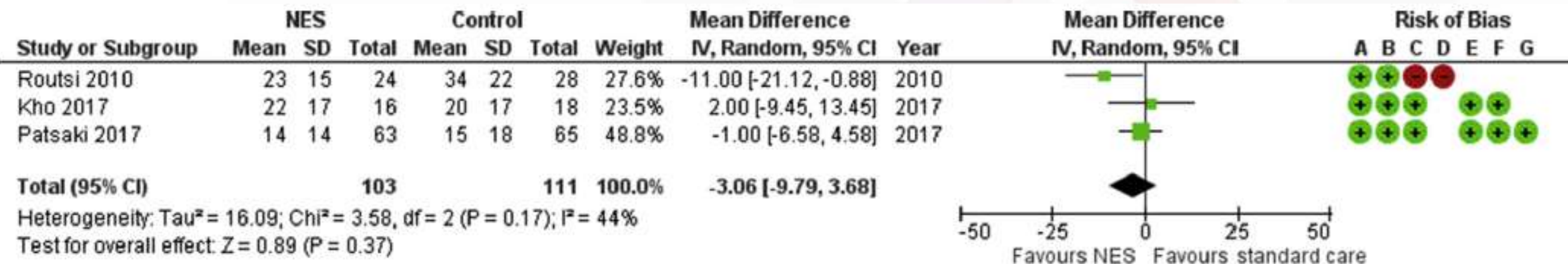
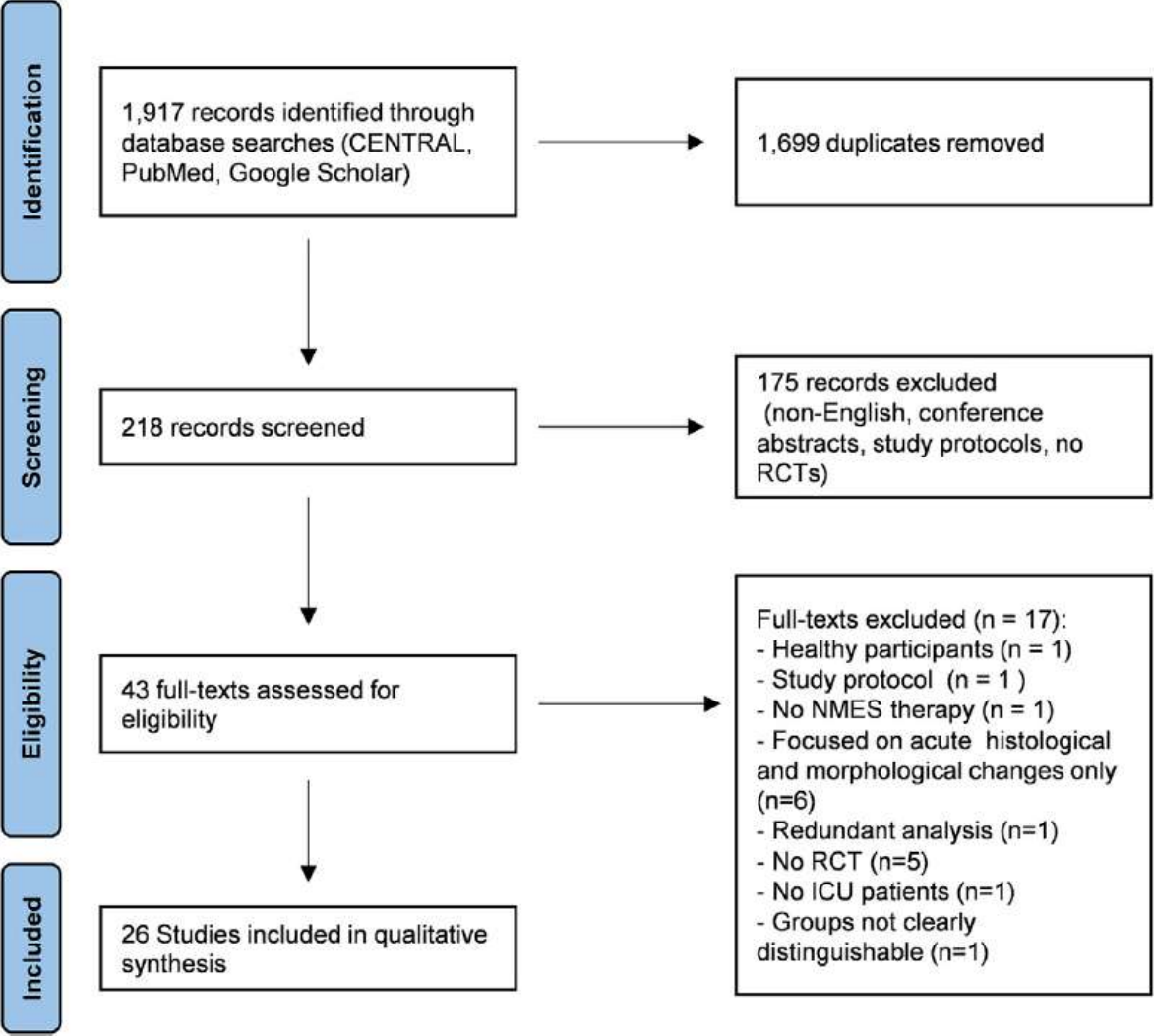


Fig. 6. Forest plot for intensive care unit length of stay.

5. Conclusion

Among critically ill patients, there were no significant differences between NES and the standard-of-care therapy on global muscle strength, ICU mortality, the duration of MV, or ICU length of stay. Further large RCTs are needed to determine which populations may benefit from this modality and to examine its efficacy and safety in adult patients in the ICU.



Therapeutic Potential of Electromyostimulation (EMS) in Critically Ill Patients—A Systematic Review

Maryam Balke^{1,2*}, Marc Teschler^{2,3}, Hendrik Schäfer^{2,3}, Pantea Pape¹, Frank C. Mooren^{2,3} and Boris Schmitz^{2,3}

¹St. Marien Hospital Cologne, Department of Early Rehabilitation, Cologne, Germany, ²Department of Rehabilitation Sciences, Faculty of Health, University of Witten/Herdecke, Witten, Germany, ³DRK Clinic Königsfeld, Center for Medical Rehabilitation, Erkrupetal, Germany

Result Summary

Our investigation indicates large heterogeneity between studies investigating the effects of EMS in critically ill patients on ICU. No pattern in muscular outcomes, post-intervention disease severity (SOFA/APACHE score), functional independence, or ICU length of stay/ventilation with respect to stimulated muscle groups or overall stimulation duration in days was identified. While the effect of sepsis on muscle parameters appears unclear, the effect of sepsis on functional outcomes/independence and duration of mechanical ventilation/ICU length of stay is likely negative and may prevent beneficial responses to EMS. No indications for a modifying effect of age and sex were detected, even though female patients were underrepresented in the analyzed studies.

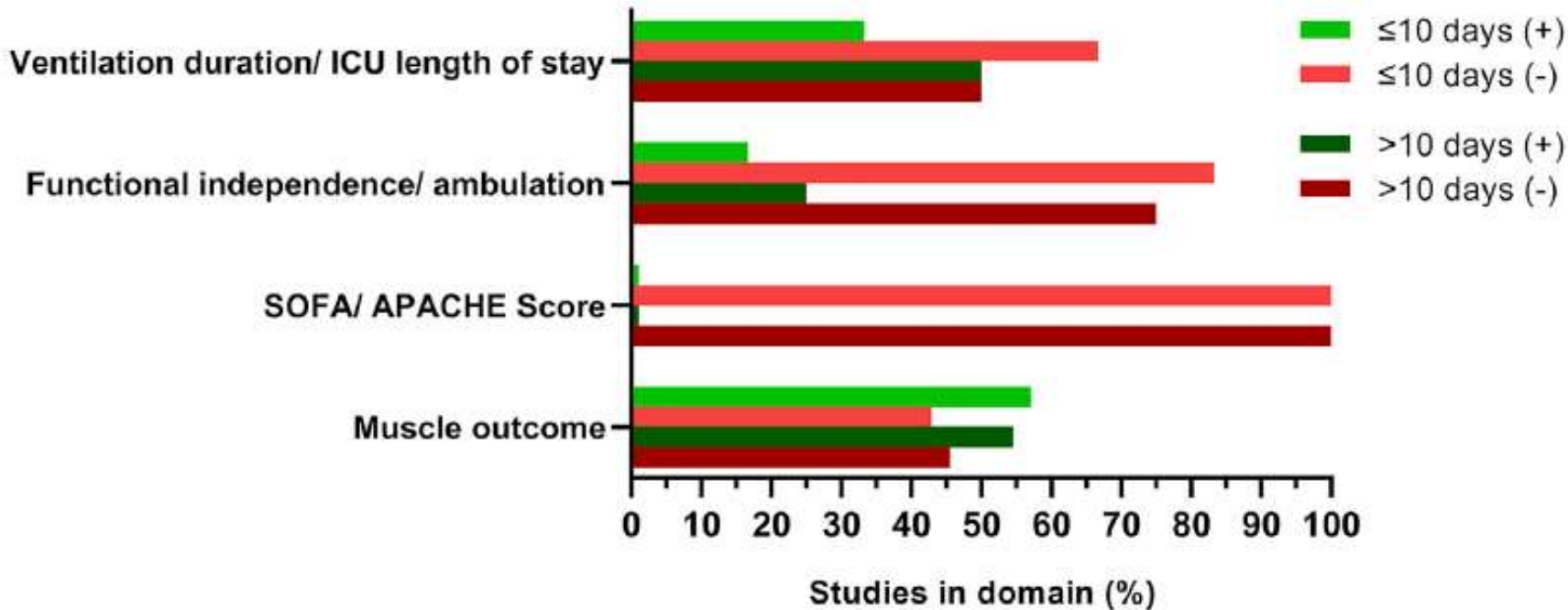


FIGURE 4 | EMS treatment duration did not affect study outcome. Studies were grouped by treatment duration of $\leq/\geq$ 10 days, respective domain investigated, and reported outcome as presented in **Table 1**. Green (+) indicates significant positive effects of EMS intervention, red (-) indicates no EMS effects compared to control. Darker colors represent studies with treatment durations >10 days. Percent refers to number of studies investigating EMS effects on outcome variables in the respective domain. Multiple outcomes per study were considered if applicable. No study reported significant positive effects of EMS on disease severity scores [SOFA (Sepsis Related Organ Failure Assessment) and/or APACHE (Acute Physiology And Chronic Health Evaluation)]. ICU, intensive care unit.

ARGUMENT N°5

Même pour le diaphragme?

Non cela ne fonctionne pas non plus!

Intervention

- 50Hz Bi-directionnal
- 300 ms
- 10s/6s => contraction/rest
- Palpable contraction
- Not synchronized
- Controlled or pressure support
- Once a day / 5d a week
- 20 minutes

Sham

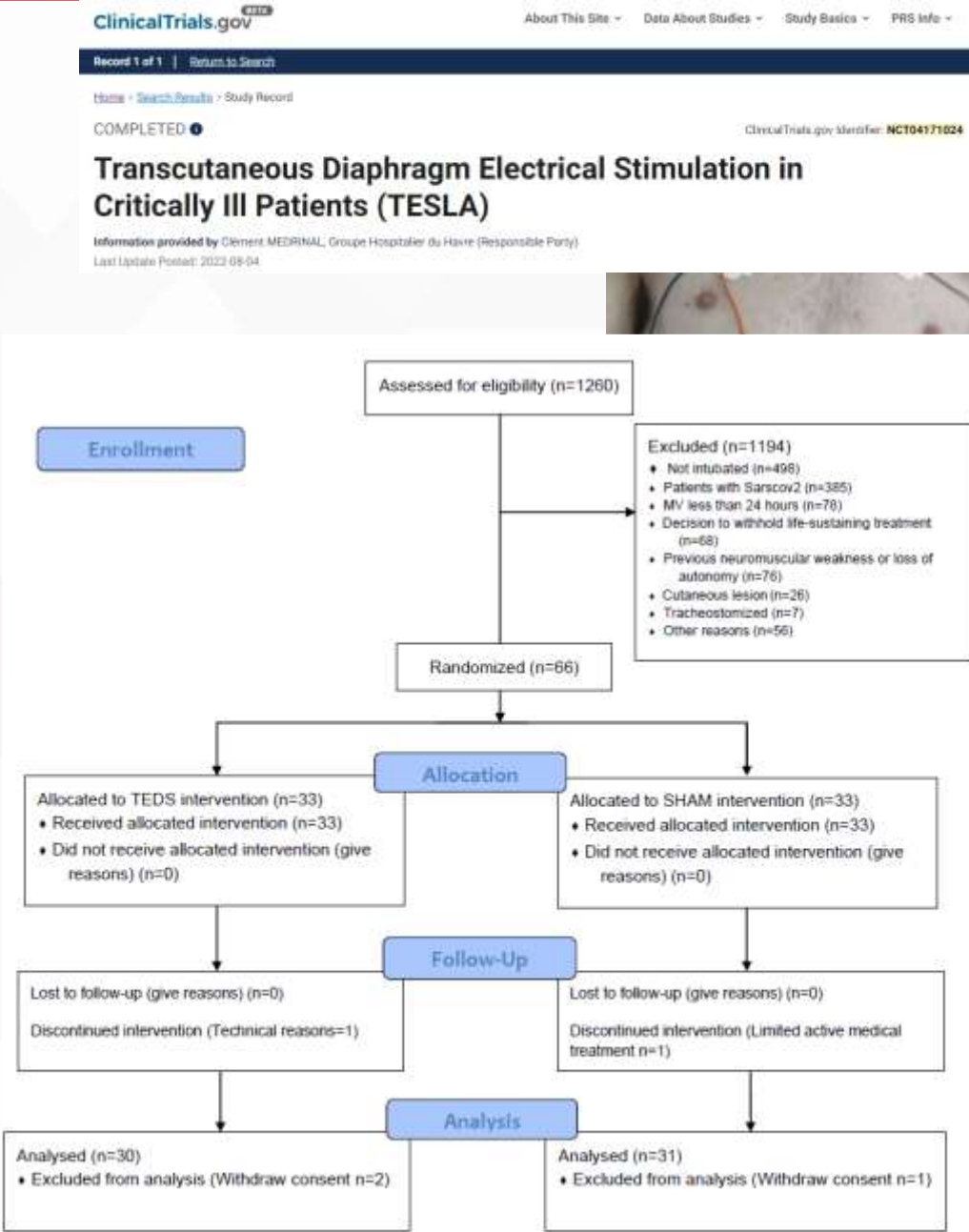
- 2Hz Bi-directionnal
- 300 ms
- No contraction (interferences)
- Not synchronized
- Controlled or pressure support
- Once a day / 5d a week
- 20 minutes

TEDS or SHAM stopped after 1st extubation

Outcomes (Blind)

1. DTF during SBT (PS 10 PEP 0) preceding extubation
< 30% => dysfunction
< 20 % => severe dysfunction

2. MIP start SBT (PS 10 PEP 0)
3. CPEF start SBT
4. End-Tidal Thickness from inclusion to extubation



	TEDS Group	Sham Group	Per Protocol Analysis Adjusted analysis		Intent to treat analysis Adjusted analysis		Cohen’s d
			Estimate (95%CI)	p-value	Estimate (95%CI)	p-value	
Primary Outcome							
DTF > 30% n(%)	16 (67)	13 (54)	1.65 (0.50 to 5.39)	0.411	1.55 (0.47 to 5.10)	0.472	-
DTF > 20% n(%)	22 (92)	20 (83)	2.02 (0.33 to 12.40)	0.449	1.61 (0.28 to 9.25)	0.591	-
Secondary Outcomes							
DTF (%), mean (SD)	47.46 (34.19)	38.67 (23.90)	6.46 (-10.19 to 23.11)	0.438	6.86 (-11.03 to 24.75)	0.442	0.19 (-0.32 to 0.70)
MIP cmH20, mean (SD)	35.57 (11.90)	29.71 (11.15)	5.83 (-1.53 to 13.19)	0.117	2.77 (-4.87 to 10.41)	0.469	0.18 (-0.33 to 0.69)
PEF (L/min), mean (SD)	83.20 (39.57)	75.37 (34.08)	7.80 (-16.59 to 32.19)	0.521	3.58 (-21.78 to 28.94)	0.776	0.07 (-0.44 to 0.82)
Days free of MV at d 28, median (Q1; Q3)	19.5 (4.0; 22.5)	21.0 (13.0; 23.0)	0.68 (0.26 to 1.76)	0.430	0.78 (0.31 to 1.92)	0.583	-
ICU LOS, mean (SD)	14.89 (9.52)	18.21 (12.83)	-4.60 (-10.93 to 1.73)	0.151	-3.40 (-9.43 to 2.63)	0.264	-0.28 (-0.79 to 0.23)

