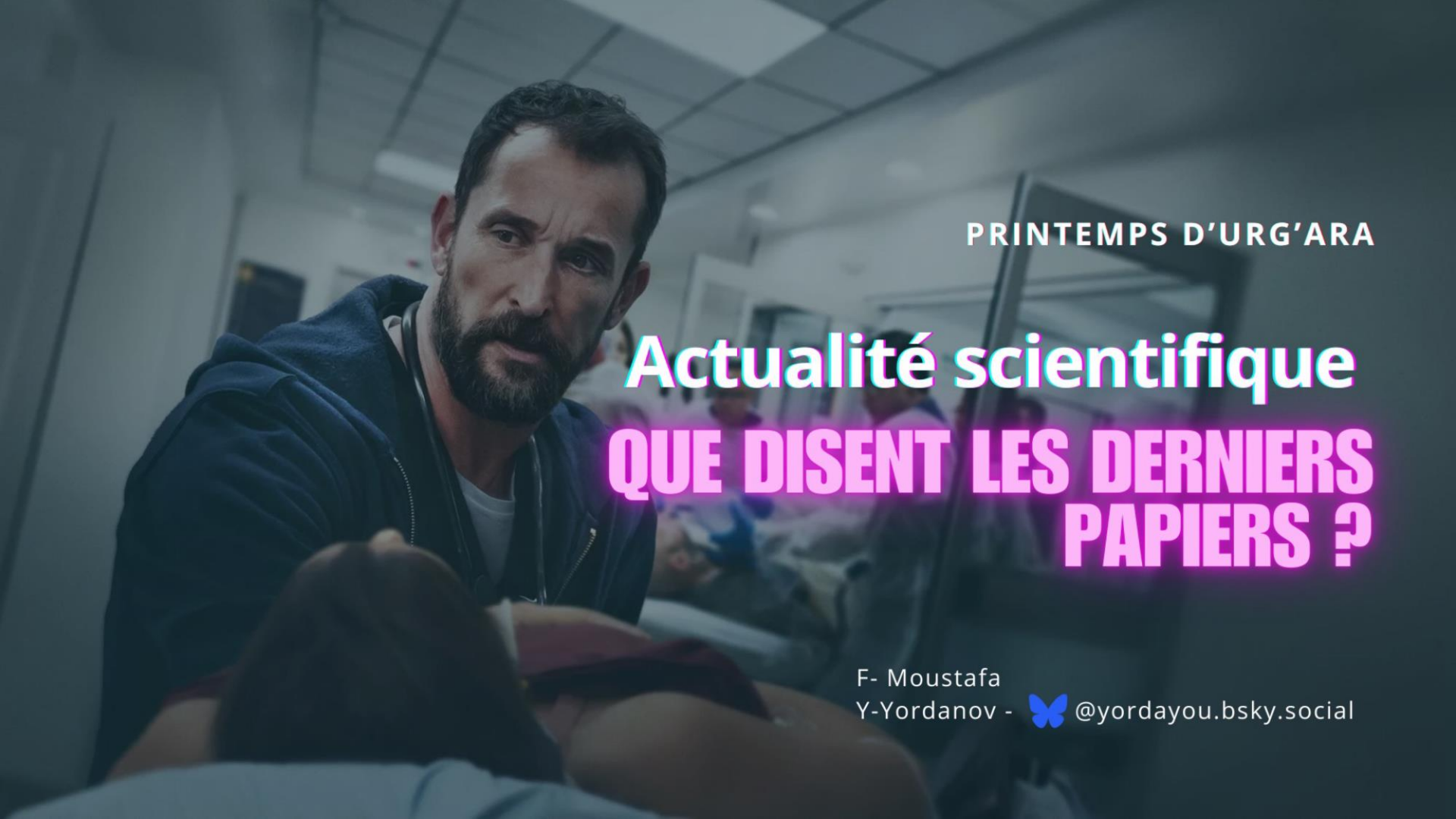




PRINTEMPS D'URG'ARA

Actualité scientifique QUE DISENT LES DERNIERS PAPIERS ?

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Intubation au vidéo-laryngoscope : pour tous ?



État des lieux sur l'usage de la vidéolaryngoscopie par les médecins préhospitaliers en France



Disponibilité et modèle de VL en SMUR									
Disponibilité de la VL	S + D		SMUR		SAUV**	Aucun	NI	N	
Nombre de SAU-SMUR	173		24		17	43	2	259	
Pourcentage	67		9		7	17	1	100	
Vidéolaryngoscope disponible	(*)*		(*)*						
McGrath Mac	86	43	14	7	6				
Airtraq	33	16	7	4	1				41
Glidescope	12	6	0	0	5				17
I View	1	0,5	1	0,5	0				2
NSP	35	17	1	0,5	5				41
Plus d'une réponse	6	3	1	0,5	0	7			

* Le pourcentage est exprimé sur l'ensemble des VL disponibles en SMUR qui est l'objet de l'étude.

**Les modèles disponibles uniquement en intra-hospitalier ne sont donnés qu'à titre informatif.

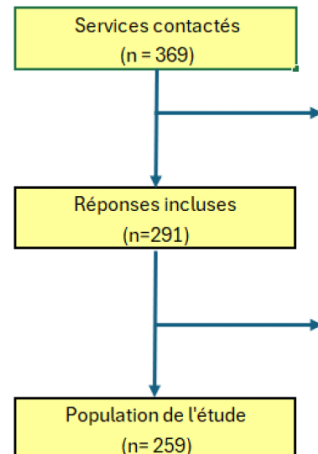
Expérience des médecins urgentistes avec la vidéolaryngoscopie		
Nombre de modèles utilisés en simulation	N (259)	%
Non	54	20,8
Un seul	137	52,9
Plusieurs	68	26,6
Nombre d'intubation en simulation		
0	54	20,8
<5	108	41,6
5 à 20	58	22,3
>20	32	12,3
NI	7	2,7
Intérêt d'une nouvelle formation à la VL		
Indispensable	70	27
Plutôt utile	152	58,8
Inutile	37	14,3

Recommandations selon les urgentistes			
Première intention	n (259)	Deuxième intention	n (258)
LD	157 (61%)	LM	93 (36%)
		VL	39 (15%)
		VM	18 (7%)
		ML	7 (3%)
LM	66 (25%)	VL	32 (12%)
		VM	16 (6%)
		ML	18 (7%)
VL	23 (9%)	VM	18 (7%)
		ML	4 (2%)
VM	12 (5%)	LM	1 (0,5%)
		ML	11 (4%)

LD : Laryngoscopie Direct, LM : Laryngoscopie + Mandrin

VL : Vidéolaryngoscopie, VM = Vidéolaryngoscope + Mandrin

ML : Masque Laryngé



RESEARCH ARTICLE

The use of video laryngoscopy outside the operating room: A systematic review

Emma J. Perkins^{1*}, Jonathan L. Begley^{1,2}, Fiona M. Brewster³, Nathan D. Hanegbi¹, Arun A. Ilancheran¹, David J. Brewster^{2,4}

1 Alfred Health, Melbourne, VIC, Australia, **2** Intensive Care Unit, Cabrini Hospital, Malvern, VIC, Australia, **3** Department of Anaesthesia, Royal Women's Hospital, Parkville, VIC, Australia, **4** Central Clinical School, Monash University, Melbourne, VIC, Australia

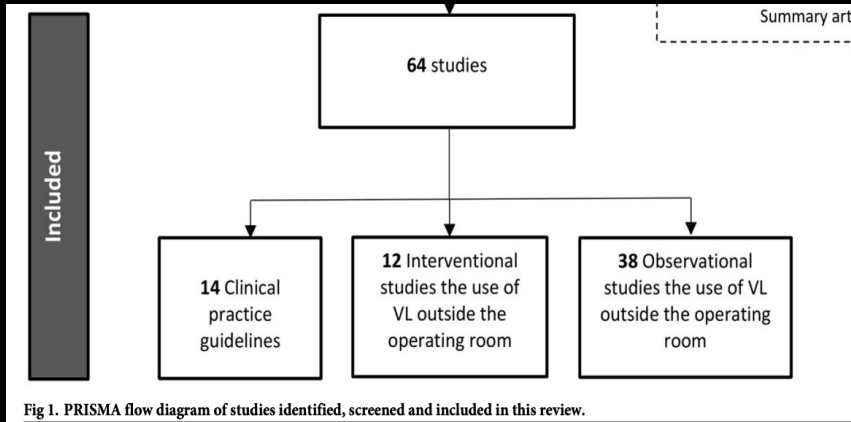


Fig 1. PRISMA flow diagram of studies identified, screened and included in this review.

Conclusion

Ultimately, our results suggest that the use of VL outside the OR has increased in recent years, and particularly through the COVID-19 pandemic. The early use of VL is recommended in most published clinical practice guidelines and is unanimously recommended for management of COVID-19 intubations. It appears that VL is likely to improve glottic view and

decrease incidence of oesophageal intubations; however, it remains unclear as to how this contributes to first-pass success. Within the limitations of our research, we found that VL has yet to show significant improvement in overall intubation success or clinical outcomes such as mortality outside the OR. More directed research is required to further characterise the use of VL outside the OR, the type of blade used and the clinical outcomes associated with its use.

Video Laryngoscopy vs Direct Laryngoscopy for Endotracheal Intubation in the Operating Room

A Cluster Randomized Clinical Trial

Kurt Ruetzler, MD^{1,2}; Sergio Bustamante, MD³; Marc T. Schmidt¹; Federico Almonacid-Cardenas, MD¹; Andra Duncan, MD^{1,3}; Andrew Bauer, MD³; Alparslan Turan, MD^{1,2}; Nikolaos J. Skubas, MD³; Daniel I. Sessler, MD¹; for the Collaborative VLS Trial Group

» Author Affiliations | Article Information

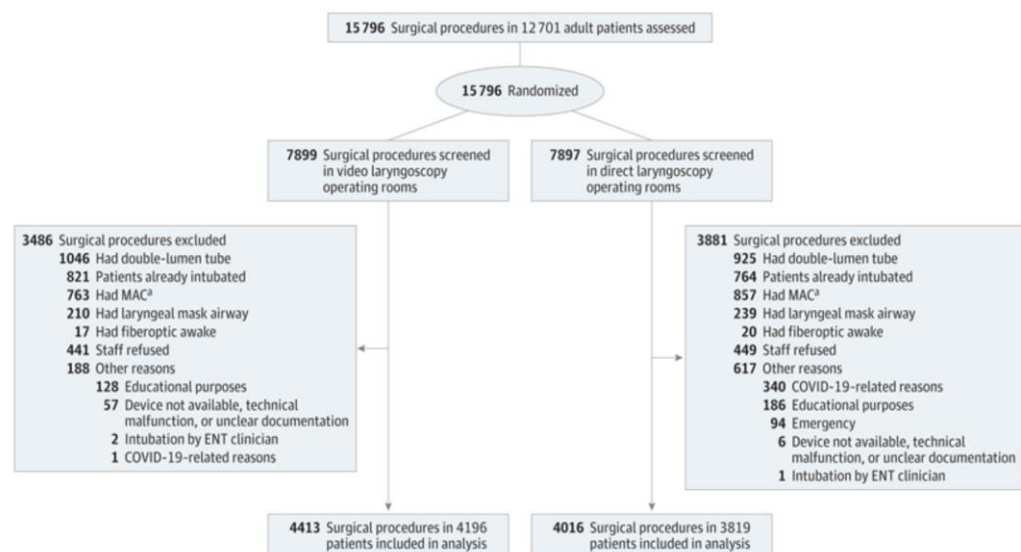
JAMA. 2024;331(15):1279-1286. doi:10.1001/jama.2024.0762

single US academic hospital.

Table 2. Treatment Effect on the Primary and Secondary Outcomes

Outcome	Video laryngoscopy (n = 4413)	Direct laryngoscopy (n = 4016)	Treatment effect estimate (95% CI) ^a	P value ^b
Primary outcome				
Intubation attempts per patient, No. (%)				
1	4336 (98.3)	3710 (92.4)	0.20 (0.14-0.28) ^c	<.001
2	70 (1.6)	277 (6.9)		
3	3 (0.07)	27 (0.67)		
>3	4 (0.09)	2 (0.05)		

Findings In this cluster randomized trial including 8429 surgical procedures in 7736 patients, more than 1 intubation attempt was required in 1.7% of patients randomized to receive video laryngoscopy. More than 1 intubation attempt was required in 7.6% of patients randomized to receive direct laryngoscopy.





Cochrane
Library

Cochrane Database of Systematic Reviews

[590 pages]

Videolaryngoscopy versus direct laryngoscopy for adults undergoing tracheal intubation (Review)

Hansel J, Rogers AM, Lewis SR, Cook TM, Smith AF.

Videolaryngoscopy versus direct laryngoscopy for adults undergoing tracheal intubation.

Cochrane Database of Systematic Reviews 2022, Issue 4. Art. No.: CD011136.

DOI: [10.1002/14651858.CD011136.pub3](https://doi.org/10.1002/14651858.CD011136.pub3).

Analysis 4.1. Comparison 4: VL versus DL (all devices combined), Outcome 1: Failed intubation Analysis 4.1. (Continued)

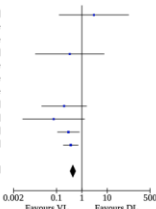
Study or Subgroup	VL		DL		Weight	Risk Ratio M-H, Random, 95% CI	Risk Ratio M-H, Random, 95% CI
	Events	Total	Events	Total			
Abdallah 2019	0	35	0	35		Not estimable	
Abdelgalel 2018	0	80	1	40	0.5%	0.17 [0.01, 4.05]	
Abdelgawad 2015	0	40	0	40		Not estimable	
Acarel 2019	0	30	0	30		Not estimable	
Aggarwal 2019	0	50	0	50		Not estimable	
Agrawal 2020	0	40	0	40		Not estimable	
Ahmad 2015	1	49	3	48	1.0%	0.33 [0.04, 3.03]	
Akbar 2015	0	45	2	45	0.6%	0.20 [0.01, 4.05]	
Al-Ghamdi 2016 (1)	0	65	0	22		Not estimable	
Alkandrovicz 2018	0	20	5	20	0.7%	0.09 [0.01, 1.54]	
Ali 2017	0	40	0	30		Not estimable	
Alhaise 2020	0	50	0	50		Not estimable	
Amor 2013	0	60	0	60		Not estimable	
Anandraj 2021	0	30	0	30		Not estimable	
Ander 2017	0	39	5	39	0.7%	0.09 [0.01, 1.59]	
Andersen 2011	0	50	2	50	0.6%	0.20 [0.01, 4.06]	
Asl 2010	1	18	1	18	0.7%	1.00 [0.07, 14.79]	
Aqil 2016	0	40	0	40		Not estimable	
Aqil 2017	0	70	0	70		Not estimable	
Arici 2014	0	40	0	40		Not estimable	
Arora 2013 (2)	0	54	0	54		Not estimable	
Arslan 2017 (1)	0	80	0	40		Not estimable	
Aziz 2012	6	149	12	147	3.6%	0.49 [0.19, 1.28]	

Jungbauer 2009	1	100	8	100	1.2%	0.14 [0.02, 0.98]
Kaar 2020	0	80	0	40		Not estimable
Kido 2015	0	25	0	25		Not estimable
Kill 2013	0	30	3	30	0.6%	0.14 [0.01, 2.65]
Kim 2013	0	22	0	23		Not estimable
Kim 2018	0	110	0	110		Not estimable
Klein-Bruenger 2017	64	360	52	120	7.3%	0.41 [0.30, 0.55]
Kornicke 2014 (1)	10	83	10	30	4.5%	0.36 [0.17, 0.78]
Koh 2010	1	25	4	25	1.1%	0.25 [0.03, 2.08]
Komatsu 2010	1	50	0	50	0.5%	3.00 [0.13, 71.92]
Kucukozman 2020	0	30	0	30		Not estimable
Lascarrrou 2017	3	186	2	185	1.5%	1.49 [0.25, 8.83]
Lee 2009	0	41	0	44		Not estimable
Lee 2012 (1)	3	75	1	25	1.0%	1.00 [0.11, 9.18]
Lim 2005	0	30	0	30		Not estimable
Lin 2012	2	85	3	85	1.5%	0.67 [0.11, 3.89]
Liu 2016	2	90	1	90	0.9%	2.00 [0.18, 21.67]
Liu 2019	0	179	10	181	0.7%	0.05 [0.00, 0.82]
Massens 2012	0	40	0	40		Not estimable
Macke 2020	0	76	3	76	0.6%	0.14 [0.01, 2.72]
Maharaj 2006	0	30	0	30		Not estimable
Maharaj 2007	0	20	1	20	0.6%	0.33 [0.01, 7.72]
Maharaj 2008	0	20	4	20	0.7%	0.11 [0.01, 1.94]
Malik 2008 (1)	3	90	2	30	1.6%	0.50 [0.09, 2.85]
Malik 2009a	0	30	0	30		Not estimable
Malik 2009b	1	50	4	25	1.1%	0.13 [0.01, 1.06]
Mathew 2018	0	33	0	33		Not estimable

Analysis 4.1. (Continued)

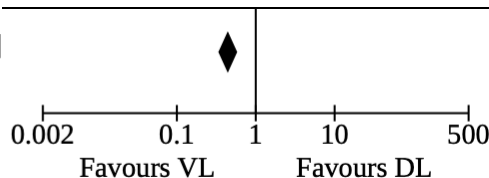
Walker 2009	1	60	0	60	0.5%	3.00 [0.12, 72.20]
Wallace 2015	0	52	0	53		Not estimable
Wasson 2013	0	30	0	30		Not estimable
Wu 2019	0	23	1	23	0.6%	0.33 [0.01, 7.78]
Wu 2012	0	50	0	109		Not estimable
Xue 2007	0	30	0	27		Not estimable
Yao 2015	0	48	0	48		Not estimable
Yao 2018	1	22	5	22	1.2%	0.20 [0.01, 1.58]
Yusef 2012	0	30	6	30	0.7%	0.08 [0.00, 1.31]
Yusuf 2016	6	90	7	31	3.4%	0.30 [0.11, 0.81]
Zhao 2014	9	74	25	75	4.9%	0.36 [0.18, 0.73]

Total (95% CI) 8800 7428 100.0% 0.44 [0.35, 0.56]
 Total events: 221 375
 Heterogeneity: $\tau^2 = 0.19$; $\chi^2 = 97.56$, $df = 76$ ($P = 0.05$); $I^2 = 22\%$
 Test for overall effect: $Z = 6.60$ ($P < 0.00001$)
 Test for subgroup differences: Not applicable



Analysis 4.1. Comparison 4: VL versus DL (all devices combined), Outcome 1: Failed intubation

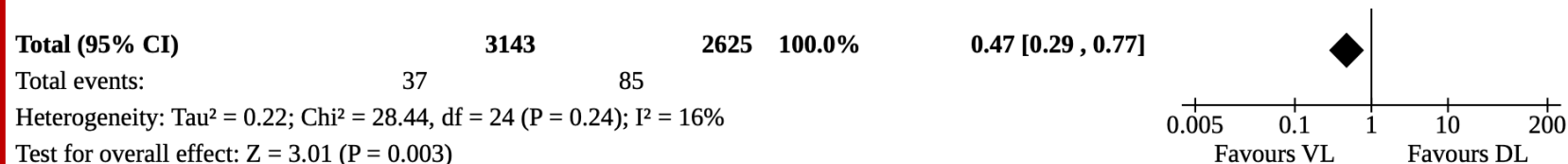
Total (95% CI)	8800	7428	100.0%	0.44 [0.35, 0.56]
Total events:	221	375		
Heterogeneity: $\tau^2 = 0.19$; $\chi^2 = 97.56$, $df = 76$ ($P = 0.05$); $I^2 = 22\%$				
Test for overall effect: $Z = 6.60$ ($P < 0.00001$)				



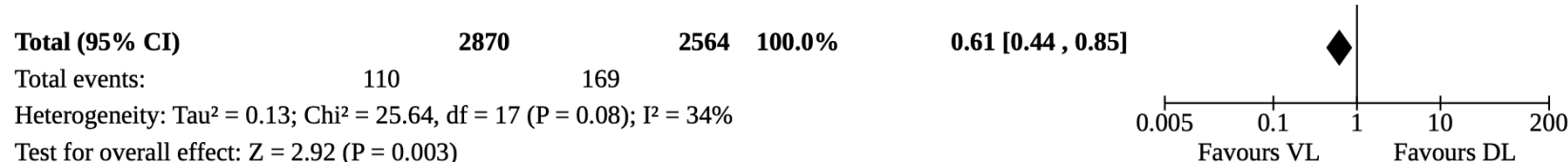
Enomoto 2008	0	99	11	104	0.7%	0.05 [0.00, 0.76]
Erden 2010	1	17	0	16	0.6%	2.83 [0.12, 64.89]
Erdin 2018	13	388	22	388	5.0%	0.59 [0.30, 1.16]
Erturk 2015	0	40	0	40		Not estimable
Ferrando 2011	1	30	0	30	0.5%	3.00 [0.13, 70.83]
Fosale 2018b	0	24	7	25	0.7%	0.07 [0.00, 1.15]
Frulich 2011	12	30	0	30	0.7%	25.00 [1.55, 403.99]
Gao 2018	6	81	8	82	3.4%	0.76 [0.28, 2.09]
Gunes 2020	0	90	0	90		Not estimable
Gupta 2013	0	60	0	60		Not estimable
Hirabayashi 2009	0	264	2	256	0.6%	0.19 [0.01, 4.02]
Hosaili 2017	0	30	0	30		Not estimable
Hosic 2016	1	100	1	40	0.7%	0.40 [0.03, 6.24]
Hu 2017 (4)	0	100	0	96		Not estimable
Ilyas 2014	5	64	0	64	0.7%	11.00 [0.62, 194.90]
Inal 2016	0	50	0	50		Not estimable
Jafra 2018	0	100	0	100		Not estimable
Jungbauer 2009	1	100	8	100	1.2%	0.13 [0.02, 0.98]
Kaar 2020	0	80	0	40		Not estimable

Shah 2016 (5)	0	30	1	30	0.5%	0.33 [0.01, 7.87]
Shinazaki 2018	0	20	0	20		Not estimable
Shippey 2013	0	24	1	25	0.6%	0.35 [0.01, 8.12]
Shukla 2017	0	40	2	40	0.6%	0.20 [0.01, 4.04]
Siddiqui 2009	0	20	0	20		Not estimable
Silveberg 2015	5	57	16	60	3.7%	0.33 [0.13, 0.84]
Sun 2005	0	100	1	100	0.5%	0.33 [0.01, 8.09]
Takenaka 2011	0	35	5	34	0.7%	0.09 [0.01, 1.54]
Taylor 2013	0	44	18	44	0.7%	0.03 [0.00, 0.43]
Tempe 2016	1	40	1	20	0.7%	0.50 [0.03, 7.59]
Tesh 2010	0	300	0	100		Not estimable
Tolson 2012	0	20	0	20		Not estimable
Tosh 2018	0	65	0	65		Not estimable
Tsan 2020	0	69	0	69		Not estimable
Turkstra 2009	0	24	0	24		Not estimable
Varsha 2019	0	35	0	35		Not estimable
Vijayakumar 2016	0	45	0	45		Not estimable
Walker 2009	1	60	0	60	0.5%	3.00 [0.12, 72.20]
Wallace 2015	0	52	0	53		Not estimable

Analysis 4.4. Comparison 4: VL versus DL (all devices combined), Outcome 4: Oesophageal intubation



Analysis 4.2. Comparison 4: VL versus DL (all devices combined), Outcome 2: Hypoxia



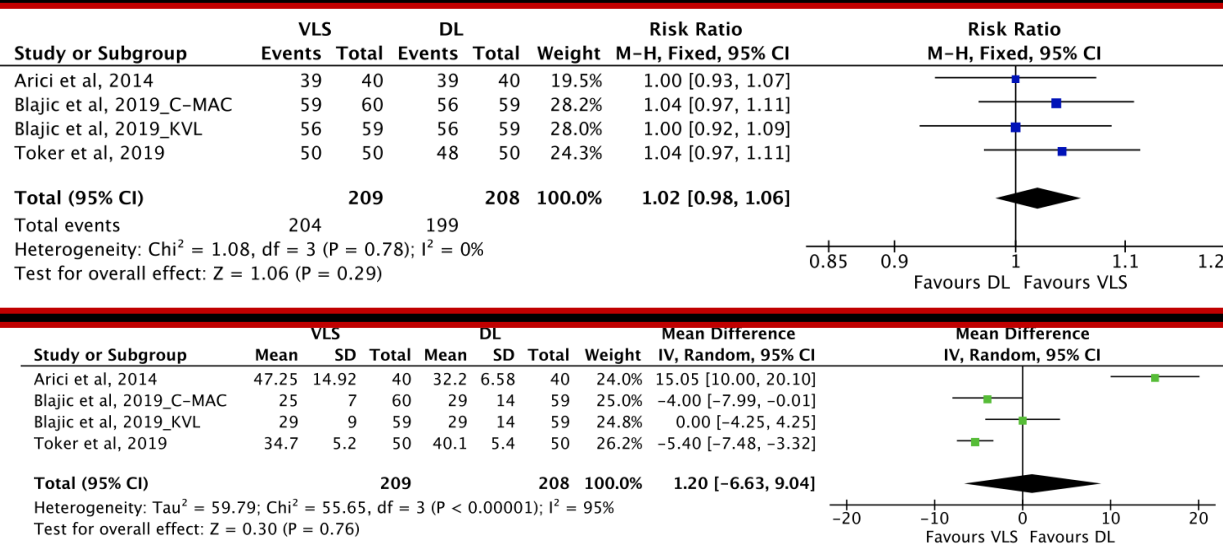
Y a-t-il des
différences?





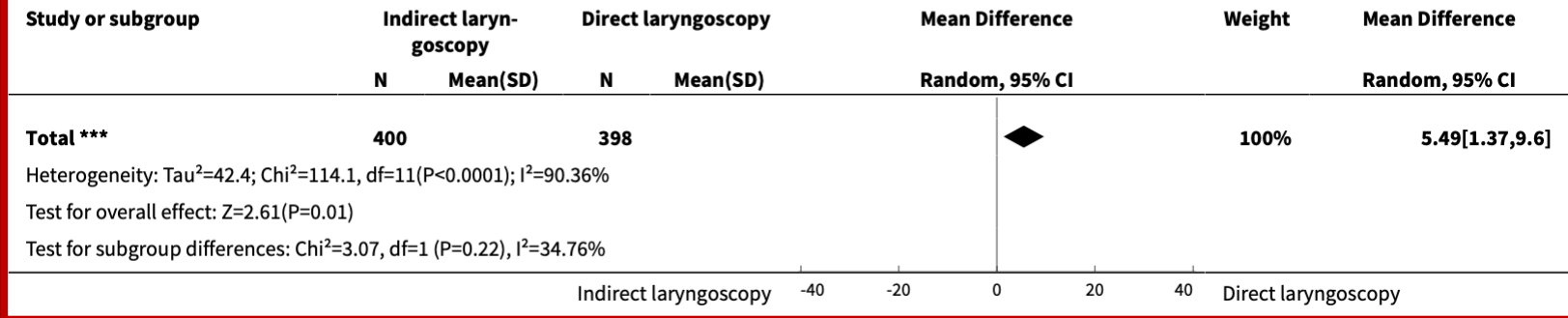
REVIEW ARTICLE/BRIEF REVIEW

Comparison of videolaryngoscopy and direct laryngoscopy for tracheal intubation in obstetrics: a mixed-methods systematic review and meta-analysis





Analysis 1.1. Comparison 1 Indirect videolaryngoscope vs conventional laryngoscope for intubation of children, Outcome 1 Unsuccessful or more than 2 intubation attempts.



ORIGINAL ARTICLE

Video versus Direct Laryngoscopy for Tracheal Intubation of Critically Ill Adults

M.E. Prekker, B.E. Driver, S.A. Trent, D. Resnick-Ault, K.P. Seitz, D.W. Russell,

AUGUST 3, 2023

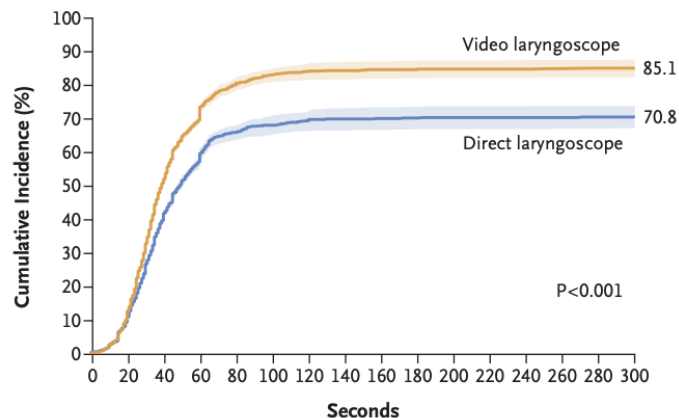


Figure 1. Cumulative Incidence of Successful Intubation on the First Attempt.

étude multicentrique, randomisée sur un ratio 1/1 qui pour des raisons évidentes n'est pas menée en double aveugle

population concernée est Nord-Américaine, répartie sur 17 sites dans 11 hôpitaux (7 services d'urgences et 10 services de réanimation)

Subgroup	Video Laryngoscope no. of events/total no. (%)	Direct Laryngoscope no. of events/total no. (%)	Absolute Risk Difference (95% CI) percentage points
Overall	600/705 (85.1)	504/712 (70.8)	
Location in hospital			
Emergency department	425/495 (85.9)	352/493 (71.4)	
Intensive care unit	175/210 (83.3)	152/219 (69.4)	
Body-mass index			
<30	402/468 (85.9)	343/483 (71.0)	
≥30	179/217 (82.5)	155/216 (71.8)	
Traumatic injury			
Yes	151/171 (88.3)	114/167 (68.3)	
No	449/534 (84.1)	390/545 (71.6)	
Anticipated difficulty of intubation			
Easy	206/232 (88.8)	172/223 (77.1)	
Moderate	266/317 (83.9)	235/331 (71.0)	
Difficult	51/67 (76.1)	30/62 (48.4)	
Not reported	77/89 (86.5)	67/96 (69.8)	
No. of operator's previous intubations			
<25	128/160 (80.0)	83/154 (53.9)	
25–100	379/441 (85.9)	330/448 (73.7)	
>100	93/104 (89.4)	91/109 (83.5)	
Proportion of previous intubations performed with a video laryngoscope			
<0.25	39/44 (88.6)	27/34 (79.4)	
0.25–0.75	335/398 (84.2)	303/429 (70.6)	
>0.75	226/262 (86.3)	174/248 (70.2)	

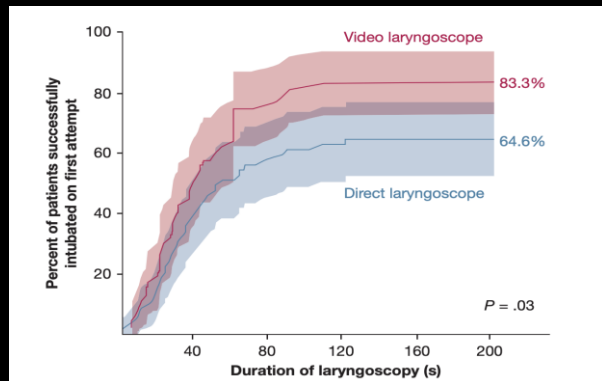
-50 -40 -30 -20 -10 0 10 20 30 40 50
 Direct Laryngoscope Better Video Laryngoscope Better

Video vs Direct Laryngoscopy for Tracheal Intubation After Cardiac Arrest

A Secondary Analysis of the Direct vs Video Laryngoscope Trial



Sur 1417 patients, 113 avec arrêt cardiaque



Results: Among adults undergoing tracheal intubation after experiencing cardiac arrest, use of video laryngoscopy increased the incidence of successful intubation on the first attempt by 18.7% and shortened the mean duration of laryngoscopy by 50.0 seconds compared with use of direct laryngoscopy.

We conducted a secondary analysis of the DEVICE trial. The DEVICE trial was a randomized controlled study that enrolled adults who were critically ill undergoing intubation at 17 sites in the Pragmatic Critical Care Research Group, including 7 EDs and 10 ICUs.⁹ All

TABLE 2] Outcomes

Outcome	Video Laryngoscope Group	Direct Laryngoscope Group	Absolute Risk Difference or Mean Difference (95% CI)	P Value
Overall				
No. of patients	48	65	NA	NA
Successful intubation on the first attempt	40 (83.3%)	42 (64.6%)	18.7 (1.2-36.2)	.03
Duration of laryngoscopy, s	48.0 (37.3)	98.0 (122.4)	-50.0 (-86.8 to -13.3)	.004
Grade I view	34 (70.8%)	30 (46.2%)	24.7 (5.2-44.2)	.01
Death				
By 1 h	11 (22.9%)	24 (36.9%)	-14.0 (-32.5 to 4.5)	.11
By ICU discharge	33 (68.8%)	45 (69.2%)	-0.5 (-18.2 to 17.3)	.96
By 28 d	35 (72.9%)	46 (70.8%)	2.1 (-16.4 to 20.7)	.80
Among those who did not receive sedation				
No. of patients	24	28	NA	NA
Successful intubation on the first attempt	21 (87.5%)	16 (57.1%)	30.4 (7.8-53.0)	.02
Duration of laryngoscopy, s	44.4 (40.6)	111.1 (115.4)	-66.7 (-15.4 to -118.0)	.006
Death				
By 1 h	8 (33.3%)	20 (71.4%)	-38.1 (-63.3 to -12.9)	.006
By ICU discharge	18 (75.0%)	27 (96.4%)	-21.4 (-40.1 to -2.8)	.02
By 28 d	20 (83.3%)	27 (96.4%)	-13.1 (-29.5 to 3.3)	.11

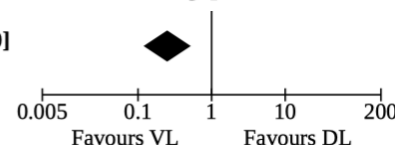
Intubation au vidéo-laryngoscope **S** : pour tous ?





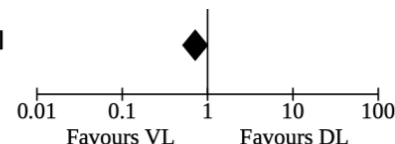
Analysis 3.2. Comparison 3: Channelled VL versus DL, Outcome 2: Hypoxaemia

Total (95% CI) 985 981 100.0% 0.25 [0.12, 0.50]
Total events: 8 41
Heterogeneity: $\text{Tau}^2 = 0.00$; $\text{Chi}^2 = 4.01$, $\text{df} = 6$ ($P = 0.68$); $I^2 = 0\%$
Test for overall effect: $Z = 3.94$ ($P < 0.0001$)



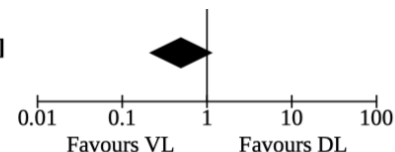
Analysis 1.2. Comparison 1: Macintosh-style VL versus DL, Outcome 2: Hypoxaemia

Total (95% CI) 1070 1057 100.0% 0.72 [0.52, 0.99]
Total events: 81 112
Heterogeneity: $\text{Tau}^2 = 0.05$; $\text{Chi}^2 = 8.11$, $\text{df} = 6$ ($P = 0.23$); $I^2 = 26\%$
Test for overall effect: $Z = 2.02$ ($P = 0.04$)



Analysis 2.2. Comparison 2: Hyperangulated VL versus DL, Outcome 2: Hypoxaemia

Total (95% CI) 850 841 100.0% 0.49 [0.22, 1.11]
Total events: 21 37
Heterogeneity: $\text{Tau}^2 = 0.38$; $\text{Chi}^2 = 8.24$, $\text{df} = 5$ ($P = 0.14$); $I^2 = 39\%$
Test for overall effect: $Z = 1.70$ ($P = 0.09$)





Analysis 4.8. Comparison 4: VL versus DL (all devices combined), Outcome 8: Subgroup analysis of failed intubation: intubator experience

Study or Subgroup	VL		DL		Weight	Risk Ratio	Risk Ratio
	Events	Total	Events	Total		M-H, Random, 95% CI	M-H, Random, 95% CI
4.8.1 Expert							
Subtotal (95% CI)		5987		4952	71.8%	0.41 [0.33 , 0.50]	
Total events:	145		223				
Heterogeneity: Tau² = 0.00; Chi² = 42.74, df = 47 (P = 0.65); I² = 0%							
4.8.2 Non-expert							
Subtotal (95% CI)		1099		1057	28.2%	0.62 [0.32 , 1.18]	
Total events:	66		110				
Heterogeneity: Tau² = 0.58; Chi² = 32.21, df = 13 (P = 0.002); I² = 60%							
Test for overall effect: Z = 1.46 (P = 0.14)							

Video Versus Direct Laryngoscopy in Novice Intubators: A Systematic Review and Meta-Analysis

Published 09/25/2022

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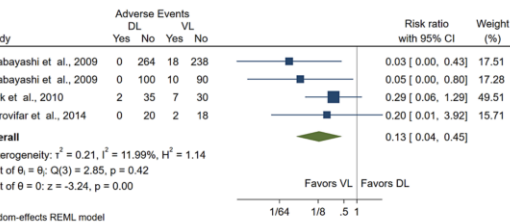
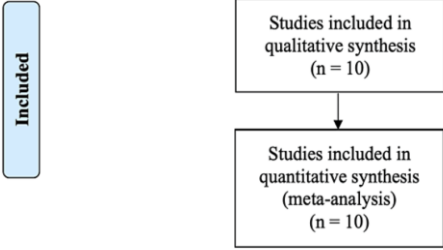


FIGURE 6: Forest plot of pooled risk ratios for the rate of esophageal intubation in novices using VL versus DL.

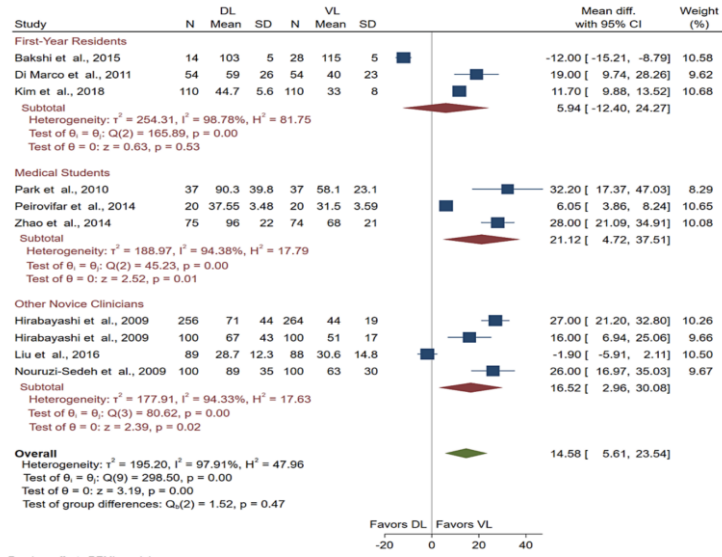


FIGURE 4: Forest plot of the difference in the mean time to intubate (seconds) for novices using VL or DL.

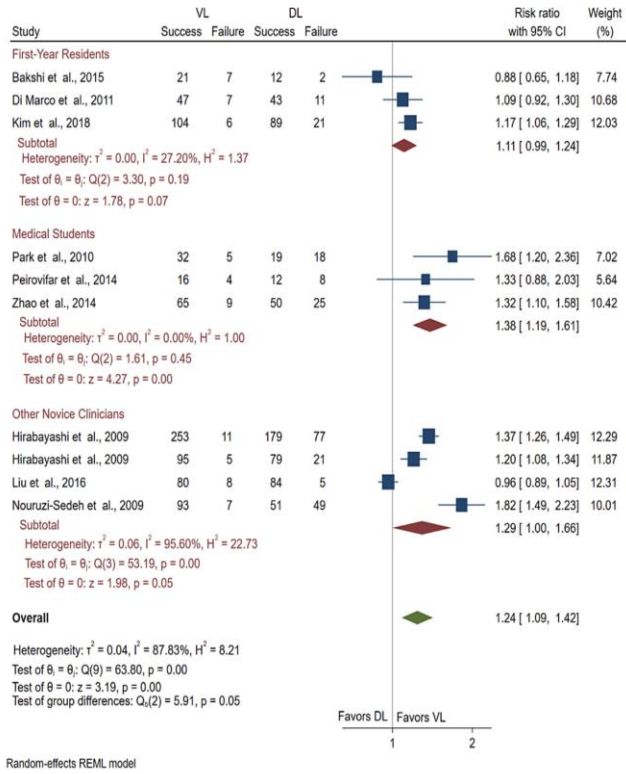


FIGURE 2: Forest plot of pooled risk ratios for the success of novices using VL or DL.

Effets secondaires

Review of Videolaryngoscopy Pharyngeal Wall Injuries

Devon Greer, MD; Kathryn E. Marshall, PhD; Scott Bevans, MD; Aurora Standlee, MD;
Patricia McAdams, MD; Wayne Harsha, MD

13147 intubations au DL vs. 1713 au VL

Lésions oro-pharyngées : 0,015% vs. 0,23%



TABLE III.
Site of Injury and Type of Repair.

Patient Number	Injury Location	Repair
1	Right tonsillar pillar	Primary
2	Right soft palate	Primary
3	Right palatopharyngeal fold	Electrocautery
4	Right palatopharyngeal arch	Primary
5	Right palatopharyngeal arch	Electrocautery
6	Soft palate	None
7	Base of tongue	NR
8	Palatoglossal arch	None
9	Right soft palate	None
10	Right palatoglossal arch	NR
11	Tonsillar pillar perforation	Primary
12	Right palatopharyngeal wall perforation	None
13	Right retromolar trigone	None
14	Soft palate	NR
15	Right soft palate	Primary
16	Right anterior tonsillar pillar	Primary
17	Right soft palate	None
18	Right soft palate	Primary
19	Right soft palate	Primary
20	Right soft palate	Primary
21	Right tonsillar pillar	Palatoplasty
22	Right soft palate	Primary
23	Right tonsillar pillar	Primary



CONCLUSION



RECOMMANDATIONS FORMALISEES D'EXPERTS

De la Société Française d'Anesthésie et Réanimation (SFAR)

ET

De la Société Française de Médecine d'Urgence (SFMU)

Intubation en urgence d'un adulte hors bloc opératoire et hors unité des soins critiques

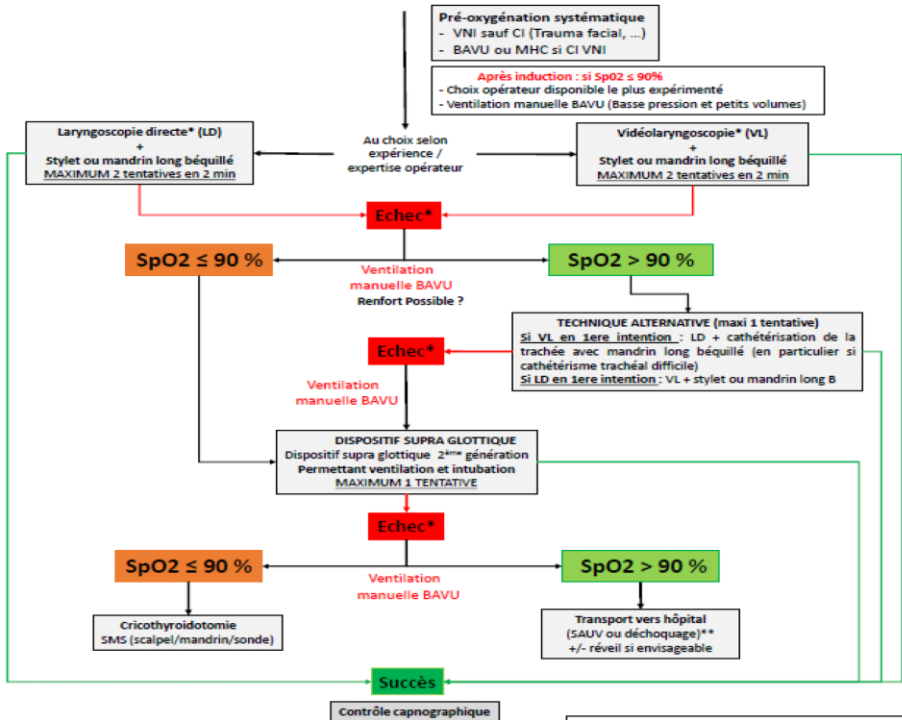
Emergency intubation of an adult outside the operating room and intensive care unit

Extra



Intra

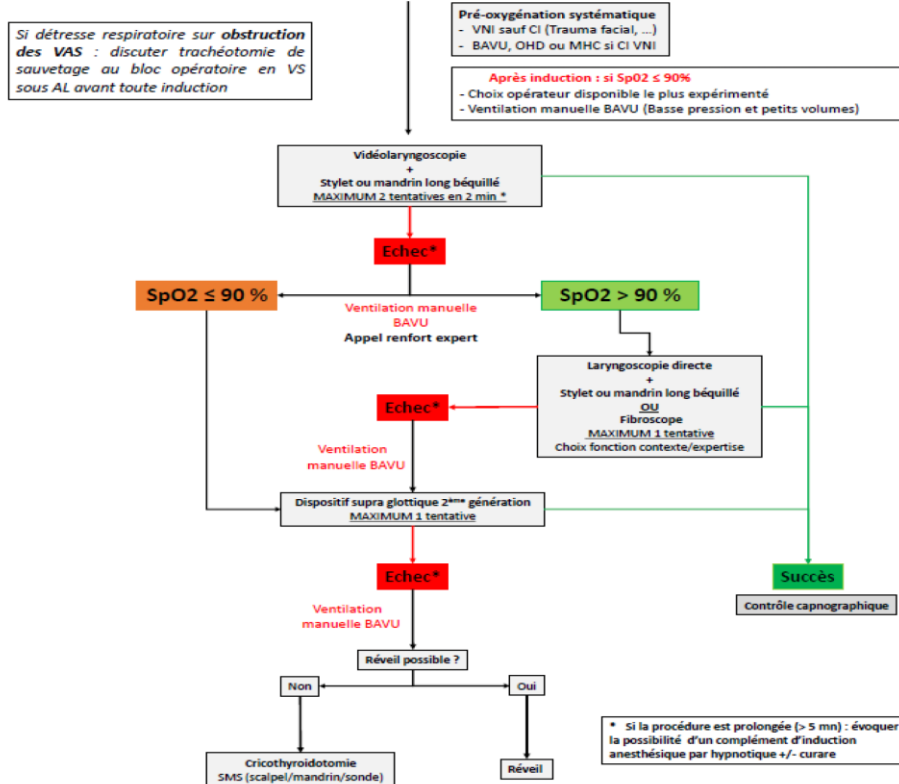
Intubation en urgence extrahospitalière



* Si la procédure est prolongée (> 5 mn) : évoquer la possibilité d'un complément d'induction anesthésique par hypnotique +/- curare

** Après information du service receveur

Intubation en urgence intrahospitalière



* Si la procédure est prolongée (> 5 mn) : évoquer la possibilité d'un complément d'induction anesthésique par hypnotique +/- curare

VL = expérience de 50
Video = expérience de 15

R2.4.1 – Si la procédure d'intubation en urgence repose sur la laryngoscopie directe, il est probablement recommandé que l'expérience minimale de l'opérateur positionné en 1ère ligne sur cette procédure soit d'au moins 50 laryngoscopies directes réussies sur des patients afin de réduire la morbi-mortalité.

GRADE 2 (Accord Fort)

R2.4.2 – Si la procédure d'intubation en urgence repose sur la vidéolaryngoscopie, les experts suggèrent que l'expérience minimale de l'opérateur positionné en 1ère ligne sur cette procédure soit d'au moins 15 vidéo-laryngoscopies réussies sur des patients afin de réduire la morbi-mortalité.

AVIS D'EXPERTS (Accord Fort)

De nombreuses études montrent que le taux de succès à la 1ère tentative est plus élevé avec un vidéolaryngoscope qu'en laryngoscopie directe chez les médecins en formation intubant en dehors du bloc opératoire [11-13]. Ces travaux suggèrent ainsi que l'acquisition de la compétence d'intuber en vidéolaryngoscopie est plus rapide à acquérir que celle d'intuber en laryngoscopie directe, avec un nombre d'intubations réussies probablement moindre pour estimer que l'opérateur est compétent.

Expérience des médecins urgentistes avec la vidéolaryngoscopie

Nombre de modèles utilisés en simulation	N (259)	%
Non	54	20,8
Un seul	137	52,9
Plusieurs	68	26,6
Nombre d'intubation en simulation		
0	54	20,8
<5	108	41,6
5 à 20	58	22,3
>20	32	12,3
NI	7	2,7
Intérêt d'une nouvelle formation à la VL		
Indispensable	70	27
Plutôt utile	152	58,8
Inutile	37	14,3

*Tu vas plus vite en
NRI,
place de l'Actilyse et
de la Métalyse*





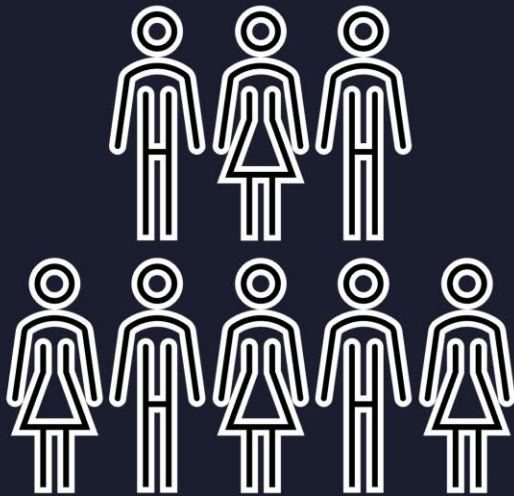
Rappels

Généralités

*1 AVC toutes les 4
minutes en France !*

- AVC ischémique =
infarctus cérébral =
80%
- AVC hémorragique =
rupture d'un vaisseau
= 20%





→ Age moyen = 73 ans MAIS 25% ont moins de 65 ans !!!

→ Plus d'une personne sur 10 après 85 ans



- INCIDENCE EN FRANCE =
130 000 cas par an
- > 110 000 hospitalisations/an
- Vieillissement de la population
 - 1ère cause de handicap
- = PROBLÈME DE SANTÉ
PUBLIQUE +++

Mortalité

- 2ème cause de décès dans le monde (OMS 2016)
- 3ème cause en France pour l'homme et 1ère cause chez la femme (HAS 2019)
- Environ 20% des patients décèdent dans le mois suivant l'AVC, 40 à 50% des survivants décèdent dans les 5 ans suivants



Dans le cadre d'une télé-expertise ou télé-médecine (TéléAVC)

Dès la sortie de l'imagerie avec confirmation d'un AVC ischémique et dans l'attente de la décision thérapeutique du médecin neurovasculaire

IDE

- Reprendre les constantes (PA, pouls, SpO2, température +/- glycémie capillaire)
- Mise en place scope
- Peser ou récupérer le poids du patient
- S'assurer de l'absence de globe vésical (bladder scan)

Médecin

NIHSS de sortie d'imagerie
Rechercher les CI à une TIV
Vérifier les traitements habituels du patient

Stratégie de reperfusion indiquée par le neurovasculaire

Si nécessité de **transfert secondaire urgent (TM)** => **alerter immédiate Centre15**

TIV Indiquée

oui

Obtention prudente d'une tension < 185/110 mmHg avant toute TIV
Au besoin protocole Nicardipine (Loxen®) et/ou Urapidil (Eupressyl®)
Mesures associées : pose de SAD, antalgiques

Présence d'anti-coagulant

non

Sans attendre les résultats du bilan de coagulation

oui

Bilan de coagulation compatible avec TIV?
Résultat rendu **en urgence**

oui

Tenecteplase (Metalyse®) selon fiche médicament AVC
0,25 mg/kg, 5mg = 1000 UI
ou
Alteplase (Actilyse®) selon fiche médicament AVC
0,9 mg/kg avec 10 % de la dose totale en bolus
et 90 % de la dose restante en 1 heure

Surveillance post TIV selon référentiel

Thrombectomie mécanique indiquée

oui

Relancer centre 15 pour transfert **URGENT**

non

Si transfert et hospitalisation en UNV indiqués
Appel C15 pour transfert non urgent

TIV : Thrombolyse Intra-Veineuse / TM : Thrombectomie Mécanique / NRI : Neuro-Radiologie Interventionnelle

4H30



6h00



Et la littérature alors?



Tenecteplase versus alteplase for acute stroke within 4.5 h of onset (ATTEST-2): a randomised, parallel group, open-label trial

Lancet Neurol 2024; 23: 1087–96

Keith W Muir, Gary A Ford, Ian Ford, Joanna M Wardlaw, Alex McConnachie, Nicola Greenlaw, Grant Mair, Nikola Sprigg, Christopher I Price, Mary Joan MacLeod, Sofia Dima, Marius Venter, Liqun Zhang, Eoin O'Brien, Ranjan Sanyal, John Reid, Laszlo K Sztrika, Syed Haider, William N Whiteley, James Kennedy, Richard Perry, Sekaran Lakshmanan, Annie Chakrabarti, Ahmad Hassan, Richard Marigold, Senthil Raghunathan, Don Sims, Mohit Bhandari, Ivan Wiggam, Khalid Rashed, Chris Douglass, on behalf of the ATTEST-2 Investigators*

Pkoi:
La ténecteplase présente des avantages potentiels par rapport à l'altéplase: administrée en un seul bolus et pourrait être plus efficace.

L'essai ATTEST-2 a évalué si la ténecteplase était non inférieure ou supérieure à l'altéplase lorsqu'elle est administrée dans un délai de 4,5 heures après le début des symptômes.

	Tenecteplase (n=885)	Alteplase (n=892)	Tenecteplase vs alteplase	p value
Primary outcome				
Modified Rankin Scale score distribution at day 90	..	..	1.07 (0.90 to 1.27)	<0.0001 (non-inferiority); 0.43 (superiority)
Secondary outcomes				
Excellent neurological recovery (modified Rankin Scale score 0–1) at day 90	369/832 (44%)	352/831 (42%)	Absolute difference 2.03 (–2.71 to 6.77); OR 1.05 (0.85 to 1.30)	0.0018 (non-inferiority); 0.40 (superiority)*; 0.66 for the OR
Independent neurological recovery (modified Rankin Scale score 0–2) at day 90	565/832 (68%)	536/831 (65%)	Absolute difference 3.41 (–1.14 to 7.95); OR 1.15 (0.92 to 1.45)	0.14*; 0.23 for the OR

ATTEST-2 corroborates the non-inferiority of tenecteplase (0.25 mg/kg) compared with alteplase (0.9 mg/kg) in acute ischaemic stroke. Taken together with all available trial evidence, tenecteplase (0.25 mg/kg) is associated with a better likelihood of excellent recovery compared with alteplase (superiority of tenecteplase for excellent recovery: mRS 0–1 vs 2–6, pooled OR 1.15 [95% CI 1.03–1.28], p=0.013; appendix 2

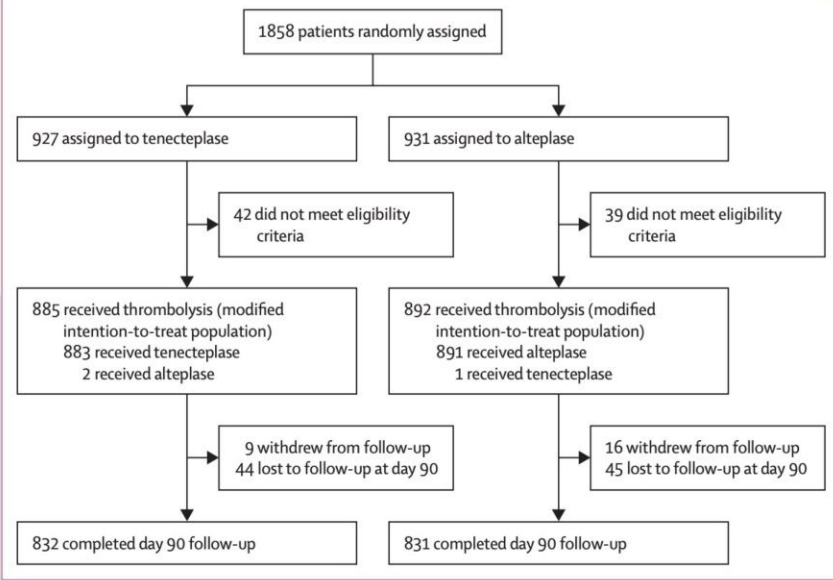


Figure 1: Patient flowchart
Safety was analysed in participants according to the treatment received (ie, 884 received tenecteplase and 893 received alteplase).

	Tenecteplase (n=884)	Alteplase (n=893)	Tenecteplase vs alteplase	p value
Mortality	68 (8%)	75 (8%)	HR 0.96 (0.69–1.33)	0.80
Symptomatic intracerebral haemorrhage—SITS-MOST criteria	20 (2%)	15 (2%)	1.37 (0.69–2.70)	0.37
Symptomatic intracerebral haemorrhage—ECASS-3 criteria	29 (3%)	21 (2%)	1.44 (0.81–2.56)	0.21
Parenchymal haematoma type 2	37 (4%)	26 (3%)	1.48 (0.89–2.48)	0.14
Any intracerebral haemorrhage	94 (11%)	78 (9%)	1.26 (0.91–1.74)	0.16
Significant extracranial haemorrhage	13 (1%)	6 (1%)	2.39 (0.89–6.39)	0.083
Neurological deterioration >3 NIHSS points by day 1 or day 5	81 (9%)	84 (9%)	0.98 (0.71–1.35)	0.90
Angioedema	6 (1%)	6 (1%)	1.03 (0.33–3.20)	0.96

Data are n (%) and OR (95% CI), unless otherwise stated. All analyses are adjusted for the variables used in the randomisation minimisation (age group, stroke severity, and onset to randomisation time) and sex. HR=hazard ratio. NIHSS=National Institutes of Health Stroke Scale score. OR=odds ratio. SITS-MOST=Safe Implementation of Stroke Thrombolysis, Monitoring Study. ECASS-3=third European Cooperative Acute Stroke Study.

Table 3: Safety outcomes



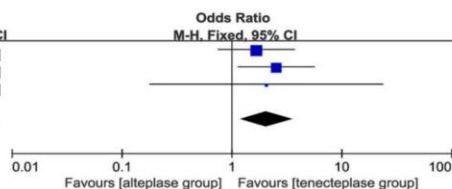
Review Article

Efficacy and safety outcomes of Tenecteplase versus Alteplase for thrombolysis of acute ischemic stroke: A meta-analysis of 9 randomized controlled trials



A: Early vessel recanalization

Study or Subgroup	tenecteplase group		alteplase group		Weight	Odds Ratio	
	Events	Total	Events	Total		M-H, Fixed, 95% CI	
ATTEST,2015	28	47	23	49	51.0%	1.67 [0.74, 3.74]	
EXTEND-IA TNK,2018	22	101	10	101	43.8%	2.53 [1.13, 5.67]	
TASTE-A,2022	34	35	33	35	5.3%	2.06 [0.18, 23.83]	
Total (95% CI)		183		185	100.0%	2.07 [1.19, 3.59]	
Total events	84		66				
Heterogeneity: $\chi^2 = 0.52$, $df = 2$ ($P = 0.77$); $I^2 = 0\%$							
Test for overall effect: $Z = 2.57$ ($P = 0.01$)							



9 études

In this analysis of nine RCTs in patients with AIS, we found that TNK and alteplase had similar rates of efficacy outcomes and safety outcomes. Our study supports the conclusion that TNK is comparable to alteplase when applied within a 4.5-h occurrence of AIS. TNK 0.25 mg/kg significantly increased the possibility of early vessel recanalization and excellent recovery relative to TNK 0.9 mg/kg, which suggested that TNK might be a better choice for patients with major artery disease with a large volume of salvageable tissue.

B: Excellent recovery (90 days mRS 0-1)

Study or Subgroup	tenecteplase group		alteplase group		Weight	Odds Ratio	
	Events	Total	Events	Total		M-H, Fixed, 95% CI	
ACT,2022	296	802	266	765	44.4%	1.10 [0.89, 1.35]	
ATTEST,2015	13	47	10	49	1.8%	1.49 [0.58, 3.83]	
EXTEND-IA TNK,2018	49	101	41	101	5.5%	1.38 [0.79, 2.41]	
Haley,2010	15	31	13	31	1.7%	1.30 [0.48, 3.54]	
TASTE-A,2022	24	55	22	49	3.4%	0.95 [0.44, 2.06]	
TRACE,2021	35	57	35	59	3.4%	1.09 [0.52, 2.30]	
TRACE-2,2023	439	705	405	696	39.8%	1.19 [0.96, 1.47]	
Total (95% CI)		1798		1750	100.0%	1.15 [1.01, 1.32]	
Total events	871		792				
Heterogeneity: $\chi^2 = 1.28$, $df = 6$ ($P = 0.97$); $I^2 = 0\%$							
Test for overall effect: $Z = 2.06$ ($P = 0.04$)							

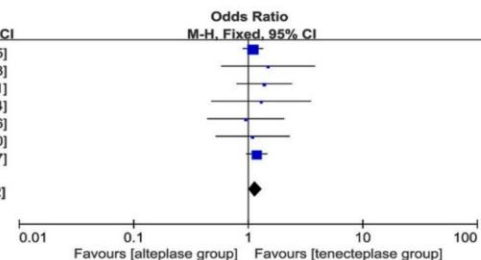
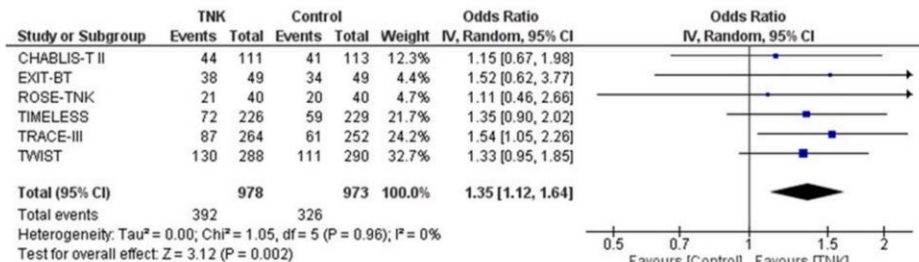


Fig. 2. Forest plots for the meta-analysis of the efficacy outcomes between alteplase(0.9 mg/kg) and TNK(0.25 mg/kg) groups. ENI: early neurological improvement.



Efficacy and safety of tenecteplase administration in extended time window for acute ischemic stroke: An updated meta-analysis of randomized controlled trials

A) Excellent functional outcome

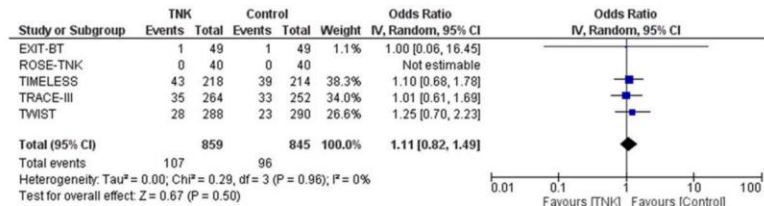


TNK (0.25 mg/kg) vs 0. après 4h30

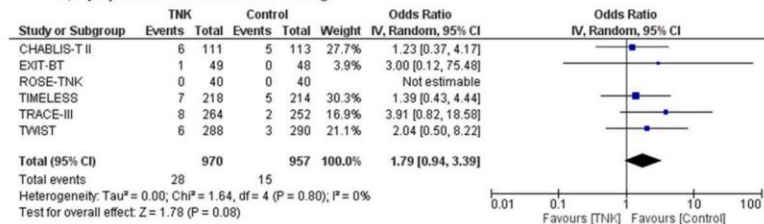
Conclusion: TNK administration in an extended time window for AIS patients leads to favorable neurological outcomes with a good safety profile.

6 études RCTs

A) All-cause death



B) Symptomatic intracranial hemorrhage



C) Any intracranial hemorrhage

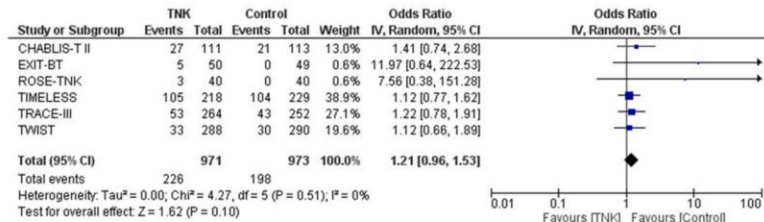


Fig. 4. Forest plots for (A) All-cause death, (B) Symptomatic intracranial hemorrhage, and (C) Any intracranial hemorrhage.

CLINICAL AND POPULATION SCIENCES

Outcomes of Bridging Intravenous Thrombolysis Versus Endovascular Therapy Alone in Late-Window Acute Ischemic Stroke

Table 1. Characteristics of Acute Ischemic Stroke Patients in the 6- to 24-Hour Time Window, According to Treatment Modality (Direct Endovascular Therapy [EVT] Versus Bridging Intravenous Thrombolysis [Bridging IVT+EVT]) (Table view)

	Direct EVT, n=2200	Bridging IVT+EVT, n=549	P value
Age, y, mean (SD)	70 (14.3)	70 (14.7)	0.52
Female, n (%)	1189 (54)	311 (57)	0.27
Prestroke mRS, median (IQR)	0 (0–2)	0 (0–1)	<0.01
Atrial fibrillation, n (%)*	733 (36)	162 (31)	0.04
Diabetes, n (%)†	552 (25)	123 (23)	0.19
Hypertension, n (%)‡	1607 (73)	394 (72)	0.53
Baseline NIHSS, median (IQR)	16 (10–20)	16 (12–20)	0.05
ASPECTS, median (IQR)	8 (7–9)	8 (7–9)	0.22
Occlusion site, n (%)			0.08
Cervical carotid	167 (8)	54 (10)	
Carotid terminus	391 (18)	112 (20)	
M1-segment MCA	1241 (56)	298 (54)	
M2-segment MCA	401 (18)	85 (15)	
Baseline imaging modality, n (%)			<0.01
NCCT+CTA	773 (35)	187 (34)	
NCCT+CTA+CTP	1097 (50)	185 (34)	
MRI	330 (15)	177 (32)	
Transferred from non-EVT stroke center, n (%)	1212 (55)	425 (77)	<0.01
Witnessed stroke onset, n (%)§	213 (12)	117 (25)	<0.01
TLKW pre-EVT brain imaging, min, median (IQR)	666 (482–864)	498 (382–701)	<0.01
TLKW-CSC arrival, min, median (IQR)	725 (544–911)	560 (432–791)	<0.01
TLKW-groin puncture, min, median (IQR)¶	724 (544–912)	560 (432–791)	<0.01
EVT procedure time, min, median (IQR)#	43 (27–72)	44 (26–67)	0.13
General anesthesia, n (%)**	556 (27)	162 (31)	0.06
Balloon-guide catheter, n (%)††	1080 (55)	234 (47)	0.002

2749 patients : 549 Thrombolyse avant thrombectomie

- AVC par occlusion d'un gros vaisseau
- Score NIHSS ≥ 6
- 20 centres répartis dans 10 pays
- dans le cadre de l'étude multicentrique rétrospective CLEAR (CT for Late Endovascular Reperfusion)
- entre janvier 2014 et mai 2022.

Table 2. Inverse Probability of Treatment Weighting (IPTW) Analysis of Clinical and Safety Outcomes, According to Treatment Modality (Direct EVT Versus Bridging IVT+EVT) in Patients in the 6- to 24-Hour Window (Table view)

	Direct EVT	Bridging IVT+EVT	OR (95% CI)	P value
90-d mRS, median (IQR)	3 (2–5)	3 (2–5)	1.03 (0.89–1.19)*	0.72
Functional independence (mRS, 0–2, 3 mo), n (%)	764 (38)	215 (41)	1.09 (0.87–1.38)	0.44
Good outcome (mRS, 0–2 or return to baseline mRS), n (%)	887 (44)	234 (45)	1.06 (0.88–1.28)	0.52
Successful reperfusion (eTICI score, 2b–3), n (%)†	1873 (86)	458 (84)	1.19 (0.81–1.75)	0.39
slCH, n (%)‡	134 (6)	28 (5)	0.75 (0.38–1.48)	0.40
Mortality, n (%)§	471 (23)	126 (24)	1.14 (0.89–1.46)	0.31

In conclusion, in this large, multicenter, retrospective study of patients with acute ischemic stroke due to occlusion of the internal carotid artery or proximal middle cerebral artery, we found no difference with regards to clinical outcomes or safety between patients who underwent EVT 6 to 24 hours after TLKW with versus without prior IVT.

Endovascular Thrombectomy Outcomes with and without Intravenous Thrombolysis for Large Ischemic Cores Identified with CT or MRI

Radiology 2023; 309(1):e230440

Imad Derraz, MD • Solène Moulin, MD • Benjamin Gory, MD • Maéva Kyheng, BST • Caroline Arquizan, MD • Vincent Costalat, MD • Bertrand Lapergue, MD • for the ETIS Investigators

1408 patients (mean age, 68.3 years ± 15.4)

654 (46.4%) were treated with IVT prior to EVT

January 2015 and January 2022

Table 2: Outcomes according to IVT Use in Unadjusted and IPTW-adjusted Analyses in the Overall Study Sample

Outcome	Study Group*		Unadjusted Analysis		IPTW-adjusted Analysis	
	No IVT (n = 754)	IVT (n = 654)	Odds Ratio†	P Value	Odds Ratio†	P Value
Primary outcome						
Favorable outcome (mRS score 0–3)	267 (35.4)	291 (44.5)	1.46 (1.17, 1.82)	<.001	1.24 (1.05, 1.46)	.01
Secondary outcomes						
Functional independence (mRS score 0–2)	167 (22.1)	214 (32.7)	1.71 (1.34, 2.18)	<.001	1.47 (1.22, 1.77)	<.001
Early neurologic improvement	246 (32.6)	251 (38.4)	1.29 (1.03, 1.60)	.02	1.26 (1.07, 1.49)	.005
Successful reperfusion (mTICI score 2b–3)	590 (78.2)	552 (84.4)	1.50 (1.13, 1.98)	.005	1.43 (1.14, 1.79)	.002
Complete reperfusion (mTICI score 3)	259 (34.4)	240 (36.7)	1.11 (0.88, 1.38)	.38	1.18 (0.99, 1.39)	.06
Safety outcomes						
90-day mortality	287 (38.1)	211 (32.3)	0.78 (0.61, 0.98)	.04	0.88 (0.75, 1.04)	.13
Hemorrhagic complications						
Any ICH	434 (57.6)	384 (58.7)	1.05 (0.83, 1.33)	.66	1.18 (0.95, 1.47)	.12
sICH	109 (14.5)	103 (15.7)	1.11 (0.81, 1.52)	.53	1.16 (0.85, 1.60)	.34
PH	154 (20.4)	153 (23.4)	1.19 (0.89, 1.59)	.23	1.17 (0.88, 1.56)	.28

Note.—Descriptive parameters and odds ratios were calculated after handling missing values for variables included in the propensity score and each outcome using a multiple imputation procedure. The propensity score was calculated with all parameters in Table 1. ICH = intracerebral hemorrhage, IPTW = inverse probability of treatment weighting, IVT = intravenous thrombolysis, mRS = modified Rankin Scale, mTICI = modified Thrombolysis in Cerebral Ischemia, PH = parenchymal hemorrhage, sICH = symptomatic ICH.

* Data are numbers of patients, with percentages in parentheses.

† Data in parentheses are 95% CIs.

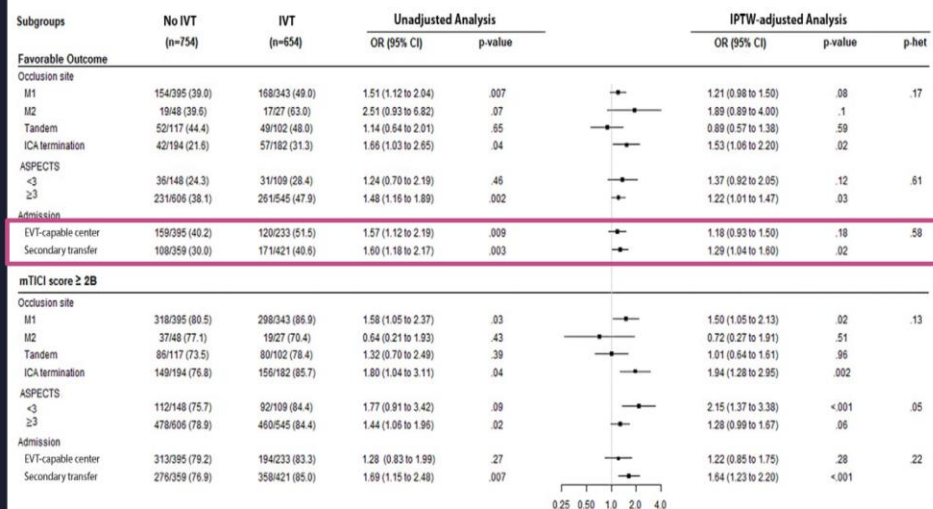


Figure 3: Comparisons of favorable outcome (modified Rankin Scale score 0–3) and successful reperfusion rates according to intravenous thrombolysis (IVT) use in key subgroups before and after inverse probability of treatment weighting (IPTW) adjustment in the overall study population. Odds ratios (ORs) were calculated after handling missing values for variables included in the propensity score and each outcome using a multiple imputation procedure. The propensity score was calculated with all parameters in Table 1. ASPECTS = Alberta Stroke Program Early CT Score, EVT = endovascular thrombectomy, ICA = internal carotid artery, M1 = middle cerebral artery M1 segment, M2 = middle cerebral artery M2 segment, mTICI = modified Thrombolysis in Cerebral Ischemia, p-het = P value for heterogeneity.

Time to Treatment With Intravenous Thrombolysis Before Thrombectomy and Functional Outcomes in Acute Ischemic Stroke

A Meta-Analysis

JAMA. 2024;331(9):764-777. doi:10.1001/jama.2024.0589

6 RCT includes

2 hours 20 minutes = *Bénéfice*

Table 3. Interaction of Symptom Onset to Expected Administration of Intravenous Thrombolysis (IVT) With the Association of Treatment Allocation and Outcomes

	Estimated association for time from symptom onset to expected treatment with IVT + thrombectomy, adjusted OR (95% CI) ^a				Ratio of adjusted OR (95% CI) per 1-h delay	P value for interaction	Direction of the association with increasing time from symptom onset to expected administration of IVT
	At 1 h	At 2 h	At 3 h	At 4 h			
Primary outcome							
Shift in modified Rankin Scale score, adjusted common OR (95% CI)	1.49 (1.13 to 1.96)	1.25 (1.04 to 1.49)	1.04 (0.88 to 1.23)	0.87 (0.68 to 1.13)	0.84 (0.72 to 0.97)	.02	The benefit associated with IVT + thrombectomy lessened
Predictions based on the primary outcome							
Predicted ARD for participants with better outcomes (≥1 point lower on modified Rankin Scale), % (95% CI) ^{b,c}	12 (3 to 21)	7 (1 to 12)	1 (−4 to 6)	−4 (−13 to 4)			
Predicted NNT (95% CI) so that 1 additional patient has a better outcome (≥1 point lower on modified Rankin Scale) ^{b,c,d}	9 (5 to 30)	16 (9 to 99)	91 (16 to −23)	−22 (27 to −7)			
Predicted ARD for participants with a modified Rankin Scale score of 0, 1, or 2, % (95% CI) ^b	9 (3 to 16)	5 (1 to 9)	1 (−3 to 5)	−3 (−8 to 3)			
Predicted NNT (95% CI) so that 1 additional patient has a modified Rankin Scale score of 0, 1, or 2	11 (7 to 35)	20 (11 to 106)	107 (22 to −35)	−34 (40 to −11)			
Secondary outcomes							
Modified Rankin Scale score of 0, 1, or 2 ^f	1.54 (1.10 to 2.17)	1.26 (1.02 to 1.56)	1.03 (0.84 to 1.27)	0.84 (0.61 to 1.17)	0.82 (0.68 to 0.98)	.03	The benefit associated with IVT + thrombectomy lessened
Early recanalization	0.95 (0.36 to 2.54)	1.82 (0.89 to 3.73)	3.48 (1.44 to 8.42)	6.66 (1.76 to 25.15)	1.91 (1.05 to 3.48)	.03	The benefit associated with IVT + thrombectomy increased
Expanded Thrombolysis in Cerebral Infarction score of 2b, 2c, or 3 at the end of the intervention ^g	2.06 (1.29 to 3.29)	1.77 (1.26 to 2.47)	1.52 (1.10 to 2.08)	1.30 (0.85 to 1.99)	0.86 (0.66 to 1.11)	.26	No statistically significant interaction
Intracranial hemorrhage							
Any	1.10 (0.76 to 1.58)	1.17 (0.93 to 1.47)	1.25 (1.02 to 1.52)	1.33 (0.98 to 1.80)	1.07 (0.89 to 1.28)	.49	No statistically significant interaction
Symptomatic	1.46 (0.68 to 3.11)	1.42 (0.83 to 2.40)	1.37 (0.88 to 2.15)	1.33 (0.74 to 2.39)	0.97 (0.70 to 1.35)	.86	No statistically significant interaction

Abbreviations: ARD, absolute risk difference; NNT, number needed to treat; OR, odds ratio.

^a The associations were derived from the primary analysis model. Instead of an adjusted OR, the first row contains adjusted common ORs (as indicated in column 1). Under predictions based on the primary outcome, 2 of the rows contain NNTs and 2 of the rows contain ARDs (as indicated in column 1). The bottom half of the Table (under secondary outcomes) contains adjusted ORs.

^b Positive values represent patients having a better outcome (≥1 point lower on the modified Rankin Scale indicates less disability) if treated with IVT plus thrombectomy. Negative values represent patients having a poorer outcome (≥1 point higher on the modified Rankin Scale indicates more disability).

^c The ARDs and NNTs were estimated using simulation processes, marginal probabilities at the mean, and generalized ORs (eMethods 4 in Supplement 1), therefore, the results and cutoff points may differ slightly compared with the model presented in Figure 1.

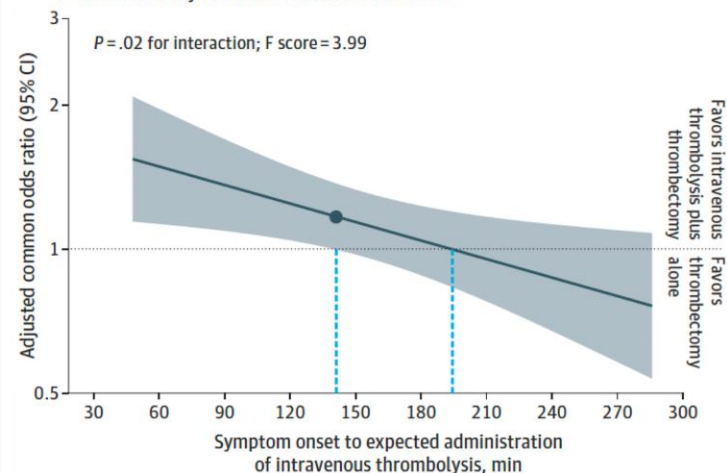
^d The NNTs are expressed so that 1 patient has at least 1 point lower on the modified Rankin Scale (less disability) if treated with IVT plus thrombectomy. Negative values represent the number needed to harm so that 1 patient has at least 1 point higher on the modified Rankin Scale (more disability) if treated with IVT plus thrombectomy.

^e The ARDs were derived from predicted probabilities of a modified Rankin Scale score of 0, 1, or 2 derived directly from the ordinal model (primary analysis) using marginal probabilities at the mean (eMethods 4 in Supplement 1).

^f Scores of 0, 1, and 2 indicate functional independence.

^g A score of 2b indicates a good recanalization result; 2c, near-perfect reperfusion; and 3, complete or 100% reperfusion.

B Modeled association of intravenous thrombolysis plus thrombectomy vs thrombectomy alone and functional outcomes



CONCLUSIONS AND RELEVANCE In patients presenting at thrombectomy-capable stroke centers, the benefit associated with IVT plus thrombectomy vs thrombectomy alone was time dependent and statistically significant only if the time from symptom onset to expected administration of IVT was short.

Prehospital stroke management and mobile stroke units

Curr Opin Neurol 2023, 36:140–146

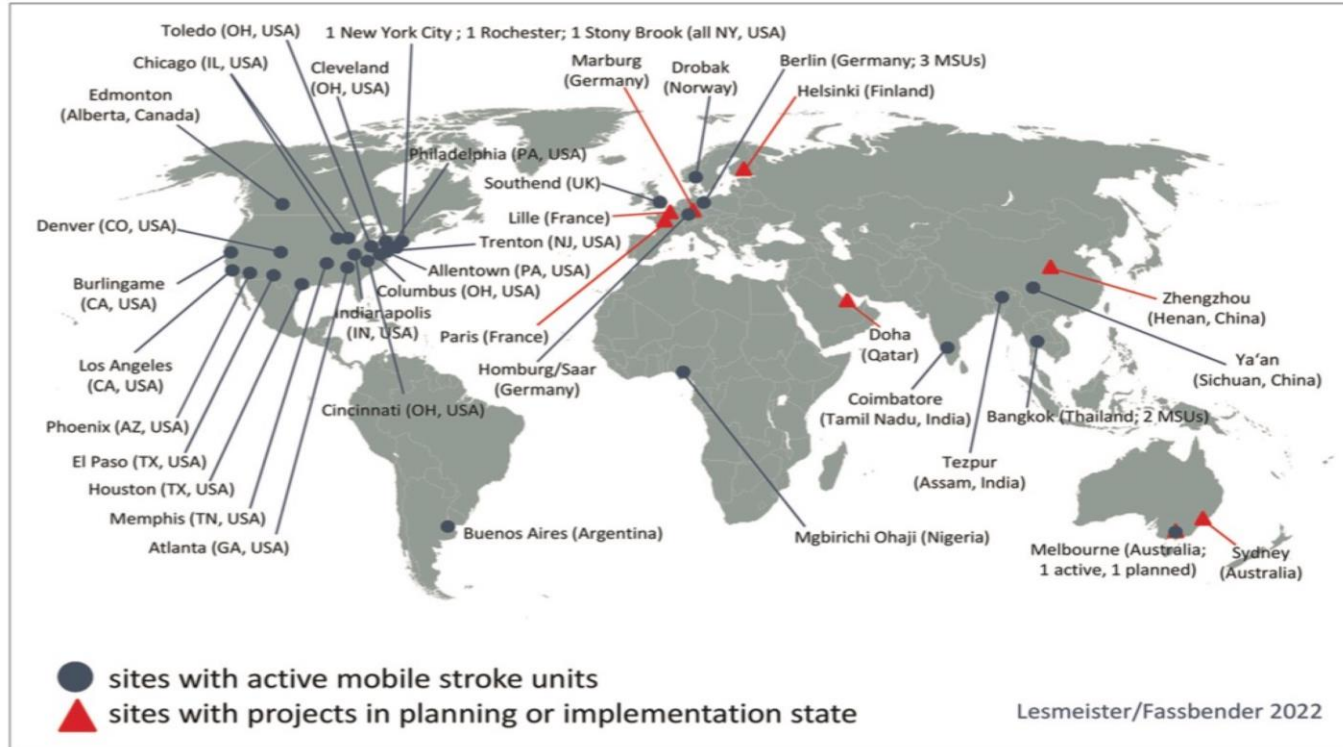


FIGURE 2. World map of Mobile Stroke Unit projects [28**].

A background image showing a business meeting. On the left, a man in a dark suit and striped tie is visible from the chest down, holding a white coffee cup. In the center, a woman in a dark blazer is gesturing with her hand while talking. On the right, another person is partially visible, also holding a coffee cup. In the foreground, a tablet displays a chart with two large circles. The overall scene is dimly lit, suggesting an indoor office environment.

En conclusion

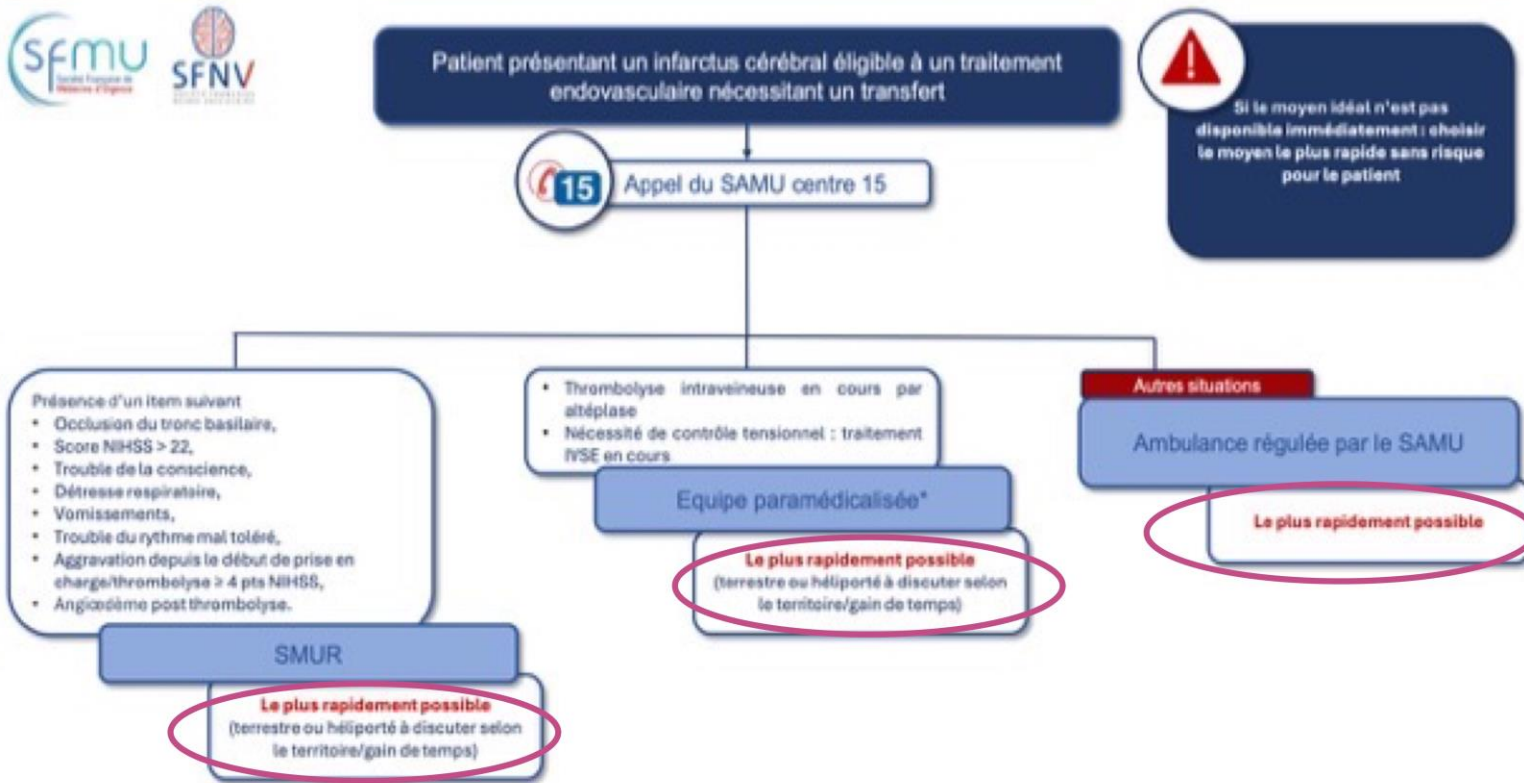


Figure 1. Algorithme de prise en charge d'un patient présentant un accident vasculaire ischémique éligible à un traitement endovasculaire nécessitant un transfert vers un centre de neuroradiologie interventionnelle.

* Equipe paramédicalisée : SMUR ou ambulance paramédicalisée

2 Thrombolyse IV

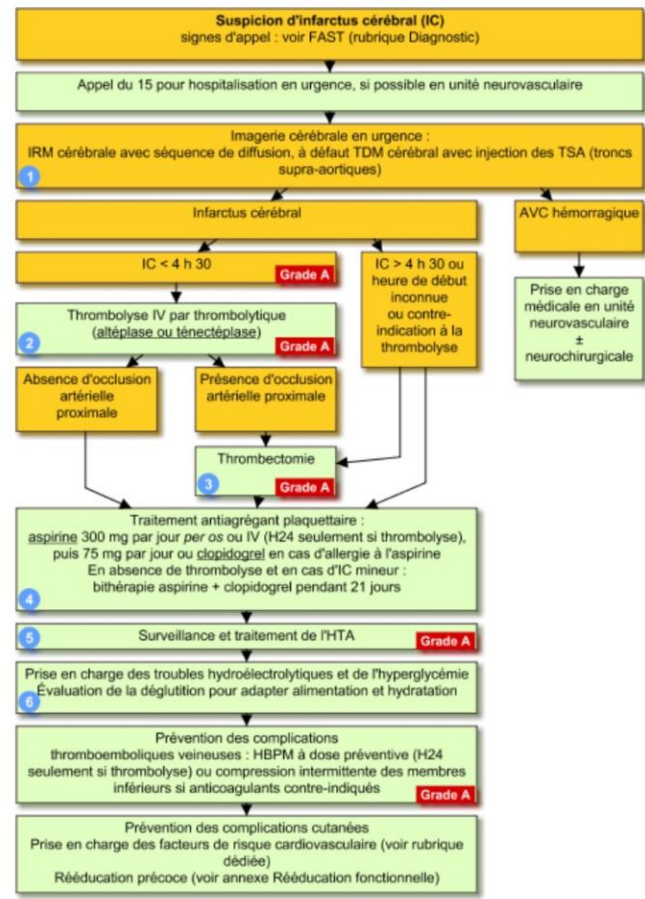
Elle est recommandée pour les IC datant < 4h30, en l'absence de contre-indication (hémorragie cérébrale). L'altéplase et la ténecteplase sont à administrer par un personnel spécialisé. **Grade A**

3 Thrombectomie

Elle est recommandée jusqu'à 6 heures après le début des symptômes si l'occlusion des artères cérébrales est proximale (carotide, cérébrale moyenne, tronc basilaire), en complément de la thrombolyse IV, ou en cas de contre-indication à la thrombolyse IV. **Grade A** Lorsque l'heure de début de l'IC est inconnue ou que le déficit neurologique est constaté au réveil, elle peut être discutée en cas de normalité de la séquence FLAIR de l'IRM, en centre spécialisé.

2. La TM est réalisée en complément de la thrombolyse intraveineuse (TIV) lorsqu'elle est indiquée (4h30) ou d'emblée en cas de contre-indications à la TIV. (Grade A, Niveau 1a)

3. La décision de réaliser une TM ne doit pas retarder la réalisation de la TIV. De même, la réalisation de la TIV ne doit pas retarder la TM (Grade A, Niveau 1a).



← URGENCES



Le bon usage des anticoagulant

Etude TRiP CAST



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Conflits d'intérêt

Speaker: Pr MOUSTAFA Farès

Member of a scientific committee

☒ YES ☐ NO

Bayer Healthcare, Boehringer Engelheim

Speaking or writing in exchange for remuneration

☒ YES ☐ NO

Bayer Healthcare, Sanofi-Aventis, Boehringer Engelheim, Aspen, Roche, Viartis

**Travel expenses and/or registration to congresses
or other events covered**

☒ YES ☐ NO

Sanofi-Aventis, Bayer Health Care, Daiichy Sankio, Boehringer Engelheim, Mundipharma

Leader of research of clinical study

☐ YES ☒ NO

If so: ...



Traumatologie Urgences et MTEV

Thromboprophylaxie



Traumatologie des membres inférieurs aux urgences

- 49% (médiane; [28-72%]) de l'activité des services d'urgence (2004)*
- Une majorité traitée en ambulatoire par une immobilisation orthopédique et décharge*.
- Responsable de complications thrombotiques (4,3% à 40%)**
 - TVP asymptomatique*
 - TVP symptomatique* (0 à 5,5%***)
 - Embolie pulmonaire*
- MVTE = cause majeure de morbidité et mortalité* en Europe
 - Embolie pulmonaire (12% des décès)****

*Société Française de Médecine d'Urgence. Le Gall et al Traumatologie des membres inférieurs : prévention de la maladie veineuse thromboembolique. Urgences 2012; 52:1-16.

** Testroote M, Stigter W, de Visser DC, Janzing H. Low-molecular-weight heparin for prevention of venous thromboembolism in patients with lower-leg immobilization. Cochrane Database Syst Rev 2013; 4: CD006681.

*** Van Adrichem Ra et al. Below-knee cast immobilization and the risk of venous thrombosis: results from a large population based case-control study. J Thromb Haemost 2014; 12: 1461-9.

****Cohen A.T., Agnelli G., Anderson F.A. et al. Venous thromboembolism (VTE) in Europe. The number of venous thromboembolism events associated morbidity and mortality. Thromb Haemost 2007; 98: 756-64.

- Facteurs de risque de thrombose veineuse: triade de Virchow (stase veineuse, lésions de la paroi veineuse et hypercoagulabilité)

Hypercoagulabilité

Acquise ou congénitale :

- Patient : **âge, obésité, antécédent de MTEV**, etc.
- Comorbidités
- Thrombophilies

THROMBOSE

Stase veineuse

Immobilisation, absence d'appui, alitement, chirurgie

Altération endothéliale

Traumatisme (fractures, lésions des tissus mous), inflammation chronique

Traumatologie

C.-M. Samama et al. Prévention de la maladie thromboembolique veineuse postopératoire. Actualisation 2011. Annales Françaises d'Anesthésie et de Réanimation 30 (2011) 947-951

	TVP TOTALES	ETE SYMPTO	Niveau de risque
PTH	50-60%	3-5%	Elevé
PTG	50-60%	2-3%	Elevé
Fracture hanche	50-60%	4-6%	Elevé
Polytrauma Grave	50-70%	-	Elevé
Fracture plateau tibial, fémur	30-40%	1%	Elevé
• Ligamentoplastie • Rotule • Fracture Tibia, cheville • tendon Achille • plâtre	10-20%	1%	Modéré
• Trauma mb inf sans fracture • Arthroscopie simple • Meniscectomie • Chirurgie du pied • Ablation de matériel	0-5%		Faible

Revue des guidelines

Table 1 Overview of guidelines on prevention of thromboembolism

Intervention	Prophylaxis	ACCP	AAOS	Australia/ New Zealand	Nice	ICS	France	Brazil	South Africa	Japan	Germany	SIGN
Total hip arthroplasty	Heparin	x	x	x	x	x	x	x	x	x	x	x
	Fondaparinux	x	x	x	x	x	x				x	
	VKA	x	x			x		x	x		x	x
	Aspirin		x					x				x
Total knee arthroplasty	Heparin	x	x	x	x	x	x	x	x	x	x	x
	Fondaparinux	x	x	x	x	x	x				x	
	VKA	x	x			x		x	x		x	x
	Aspirin		x									x
Hip fracture surgery	Heparin	x		x	x	x	x				x	x
	Fondaparinux	x		x	x	x	x				x	
	VKA	x				x					x	
	Aspirin											x
Knee arthroscopy	None	x				x	x	x				x
	Heparin	RF				RF	RF					
Immobilization of lower extremities	None	x				x					x	
	Heparin					RF	x					
	Fondaparinux										x	
	VKA										x	

VKA, vitamin K antagonist; RF, risk factors present.

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Evaluation du risque

Immobilisation avec plâtre

Facteur de risque ?

Below-knee cast immobilization and the risk of venous thrombosis: results from a large population-based case-control study

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*Department of Clinical Epidemiology, Leiden University Medical Center; †Department of Orthopedic Surgery, Leiden University Medical Center; ‡Department of Trauma Surgery, Leiden University Medical Center; and §Department of Thrombosis and Hemostasis, Leiden University Medical Center, Leiden, the Netherlands

Table 2 Treatment type and indication for below-knee cast immobilization, and the risk of venous thrombosis within 1 year

Treatment	Patients*	Controls	OR _{adj} (95% CI)	OR _{adj} (95% CI)
None	4191	6073	1 (Reference)	1 (Reference)
Below-knee cast	134	23	8.5 (5.4–13.2)	8.3 (5.3–12.9)
Operative	41	11	5.4 (2.7–10.4)	4.9 (2.5–9.6)
Conservative	93	12	11.4 (6.2–20.7)	11.4 (6.2–20.9)
Traumatic	86	10	12.6 (6.5–24.3)	12.7 (6.6–24.6)
Non-traumatic	5	1	7.6 (0.9–65.5)	7.6 (0.9–66.4)

CI, confidence interval; OR_{adj}, adjusted odds ratio (adjusted for sex and age); OR_{adj}, adjusted odds ratio (adjusted for sex, age, BMI, and regular exercise). *For two patients, no information on indication for cast immobilization was available.

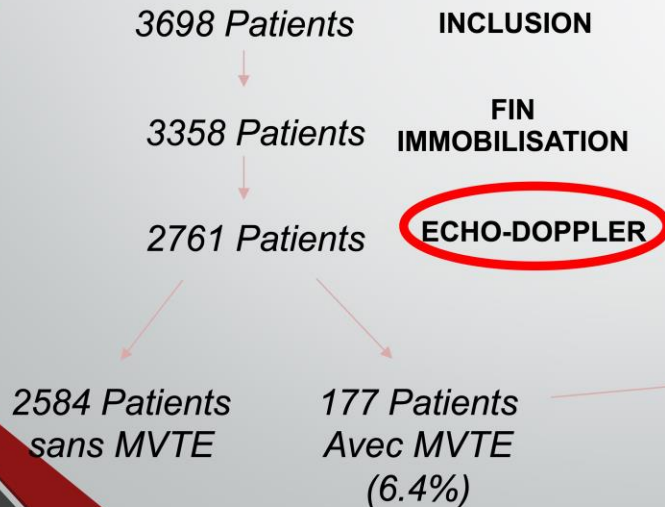
Data based from a large based case control study (MEGA Study)

Plâtre sous le genou = augmentation du risque de MTEV (ORajusté = 8,5)

Facteurs de risques

Patients avec traumatisme du MI avec immobilisation

- Objectif : analyse des facteurs de risque de survenue de thrombose
- Étude prospective, multicentrique
- Patients pris en charge pour traumatisme du membre inférieur traité de façon orthopédique/plâtre/attelle
- Pratiques des urgences français en terme de prévention des MVTE



Original Contribution

Incidence and risk factors for venous thromboembolism in patients with nonsurgical isolated lower limb injuries☆☆

Bruno Riou MD, PhD^{a,*}, Christophe Rothmann MD^b, Nathalie Lecoules MD^c, Eric Bouvat MD^d, Jean-Luc Bosson MD^e, Philippe Ravaud MD, PhD^f, Charles Marc Samama MD, PhD^g, Moussa Hamadouche MD^h

Table 3 Multivariate analysis of risk factors for VTE (n = 2755 patients)

Variables	Odds ratio (95% confidence interval)	P
No weight bearing	4.11 (1.72-9.86)	.0015
Age >50 y	3.14 (2.27-4.33)	<.0001
Rigid immobilization	2.70 (1.66-4.38)	<.0001
Severe injury	1.88 (1.34-2.62)	.0002

See text for definitions.

La décharge, l'âge, l'immobilisation et la sévérité du traumatisme = facteurs de risque de MVTE

L-TRIP(CAST) score

- Médecine personnalisée
- Balance bénéfice-risque

RESEARCH ARTICLE

Venous Thrombosis Risk after Cast Immobilization of the Lower Extremity: Derivation and Validation of a Clinical Prediction Score, L-TRIP(cast), in Three Population-Based Case-Control Studies

Banne Nemeth^{1,2}, Raymond A. van Adrichem^{1,2}, Astrid van Hylckama Vlieg¹, Paolo Bucciarelli³, Ida Martinelli³, Trevor Baglin⁴, Frits R. Rosendaal^{1,5}, Saskia le Cessie^{1,6}, Suzanne C. Cannegieter^{1*}

November 10, 2015

4446 cas. Vs 6118 sans VTE

**Score > ou = 7 points ,
haut risque
→ Thromboprophylaxie**

Table 4. L-TRIP(cast) score based on the clinical risk prediction model.

Environmental Predictor Variable	Point Value
Age ≥ 35 and < 55 y	2
Age ≥ 55 y	3
Male sex	1
Current use of oral contraceptives	4
Cancer within the past 5 y	3
Pregnancy or puerperium	3
BMI ≥ 25 and < 35 kg/m ²	1
BMI ≥ 35 kg/m ²	2
Pneumonia	3
Family history of VTE (first-degree relative)	2
Comorbidity (rheumatoid arthritis, chronic kidney disease, COPD, multiple sclerosis)	1
Hospital admission within the past 3 mo	2
Bedridden within the past 3 mo	2
Surgery within the past 3 mo	2
Superficial vein thrombosis	3
Plaster cast: complete leg	5
Plaster cast: circular knee cast (ankle free)	2
Plaster cast: foot	2
Plaster cast: lower leg	4

This L-TRIP(cast) score was derived from the regression coefficients (betas) of the clinical prediction model: 0.20 < beta ≤ 0.75, 1 point; 0.75 < beta ≤ 1.25, 2 points; 1.25 < beta ≤ 1.75, 3 points; 1.75 < beta ≤ 2.25, 4 points; beta > 2.25, 5 points

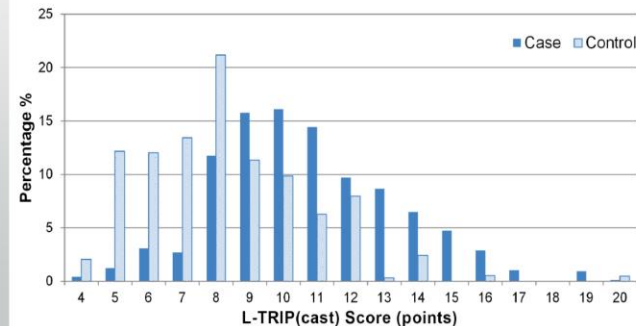


Fig 3. Distribution of individual L-TRIP(cast) scores in the plaster cast subgroup derived from the MEGA study.



Clinical risk assessment model to predict venous thromboembolism risk after immobilization for lower-limb trauma

Banne Nemeth^{a,b,1,*}, Delphine Douillet^{c,1}, Saskia le Cessie^{a,d}, Andrea Penaloza^a, Thomas Moumneh^e, Pierre-Marie Roy^e, Suzanne Cannegieter^{a,d}

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B. Nemeth et al. / EClinicalMedicine 20 (2020) 100270

Step 1: Risk scores

TIP score

(Delphi Method)

- Developed in 2017
- Validated in MEGA study (Douillet et al. PLoS One 14(6): e0217748)

L-TRIP(cast) score

(Case-control data)

- Developed in 2015 in the MEGA study and validated in two other case-control studies. (PLoS Med 12(11): e1001899)

Step 2: Development phase, using data from the MEGA study

- Compare AUC of TIP and L-TRIP(cast)
- Merge both scores into one single score (The TRIP(cast) score) by expert opinion
- Calculate AUC of TRIP(cast) score in MEGA

Step 3: Validation of the TRIP(cast) score in the POT-CAST trial

TRIP(CAST) score

The NEW ENGLAND JOURNAL of MEDICINE

ESTABLISHED IN 1812 FEBRUARY 9, 2017 VOL. 376 NO. 6

Thromboprophylaxis after Knee Arthroscopy and Lower-Leg Casting

Raymond A. van Adrichem, M.D., Banne Nemeth, M.D., Ale Algra, M.D., Ph.D., Saskia le Cessie, Ph.D., Frits R. Rosendaal, M.D., Ph.D., Inger B. Schipper, M.D., Ph.D., Rob G.H.M. Heijmans, M.D., Ph.D., and Suzanne C. Cannegieter, M.D., Ph.D., for the POT-ICAST and POT-CAST Group^a



Using a cutoff score of 7 points in the POTCAST trial (incidence 1.6%):

Se : 76.1%,
Sp : 51.2%,
VPP : 2.5%,
VPN: 99.2%,

Table 1

TRIP(cast) score.^a

	Points
Trauma^b	
High-risk trauma	3
Fibula and/or tibia shaft fracture	
Tibial plateau fracture	
Achilles tendon rupture	
Intermediate-risk trauma	2
Bi or tri-malleolar ankle fracture	
Patellar fracture	
Ankle dislocation, Lisfranc injury	
Severe knee sprain (with edema/haemarthrosis)	
Severe ankle sprain (grade 3)	
Low-risk trauma	1
Single malleolar ankle fracture	
Patellar dislocation	
(Meta)Tarsal bone(s) or forefoot fracture	
Non-severe knee sprain or ankle sprain (grade 1 or 2)	
Significant muscle injury	
Immobilization^c	
Upper-leg cast	3
Lower-leg cast	2
Foot cast (ankle free) or any semi-rigid without plantar support	1
Other cast or bracing with plantar support	0
Patient characteristics^d	
Age <35 years	0
Age ≥35 and <55 years	1
Age ≥55 and <75 years	2
Age ≥75 years	3
Male sex	1
Body Mass Index BMI ≥25 and <35 kg/m ²	1
Body Mass Index BMI ≥35 kg/m ²	2
Family history of VTE (first-degree relative)	2
Personal history of VTE or known major thrombophilia	4
Current use of oral contraceptives or Estrogenic hormone therapy	4
Cancer diagnosis within the past 5 years	3
Pregnancy or puerperium	3
Immobilization (other) within the past 3 months ^e	2
Hospital admission, bedridden or flight > 6 h, Lower limb paralysis	
Surgery within the past 3 months	2
Comorbidity	1
Heart failure, rheumatoid arthritis, chronic kidney disease, COPD, IBD	
Chronic venous insufficiency (varicose veins)	1

^a Thrombosis Risk Prediction in patients with cast immobilization score TRIP (cast) score is the sum of the Trauma, Immobilization and Patient components.

^b Trauma: Choose one, (the most severe trauma).

^c Immobilization: Choose one.

^d Patient: multiple points can be scored.

^e Other immobility next to cast immobilization.

Targeted prophylactic anticoagulation based on the TRiP(cast) score in patients with lower limb immobilisation: a multicentre, stepped wedge, randomised implementation trial

Delphine Douillet, Andrea Penaloza, Damien Viglino, Jean-Jacques Banihachemi, Anmar Abboodi, Mathilde Helderlé, Emmanuel Montassier, Frédéric Balen, Christian Brice, Saïd Laribi, Thibault Ducheno, Philippe Vives, Louis Soulat, Nicolas Marjanovic, Thomas Moumneh, Dominique Savary, Jérémie Riou, Pierre-Marie Roy



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2024



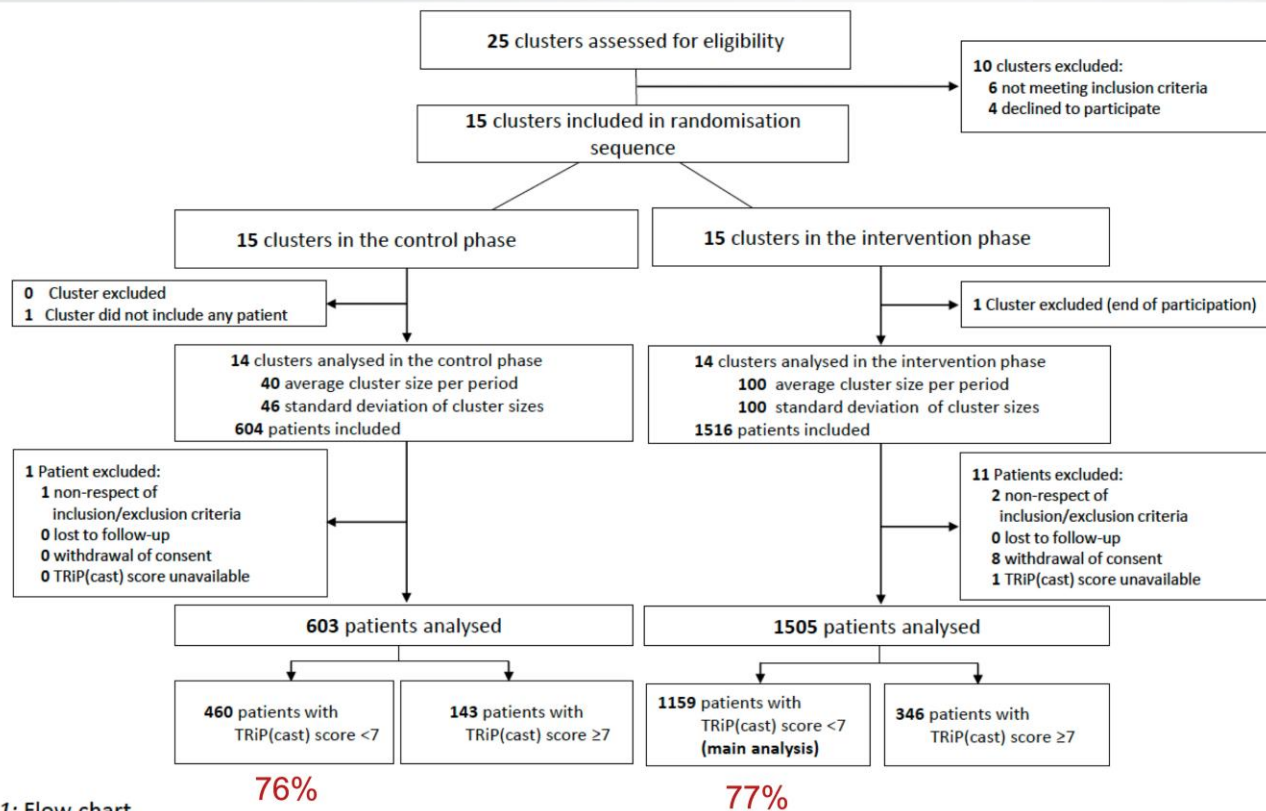
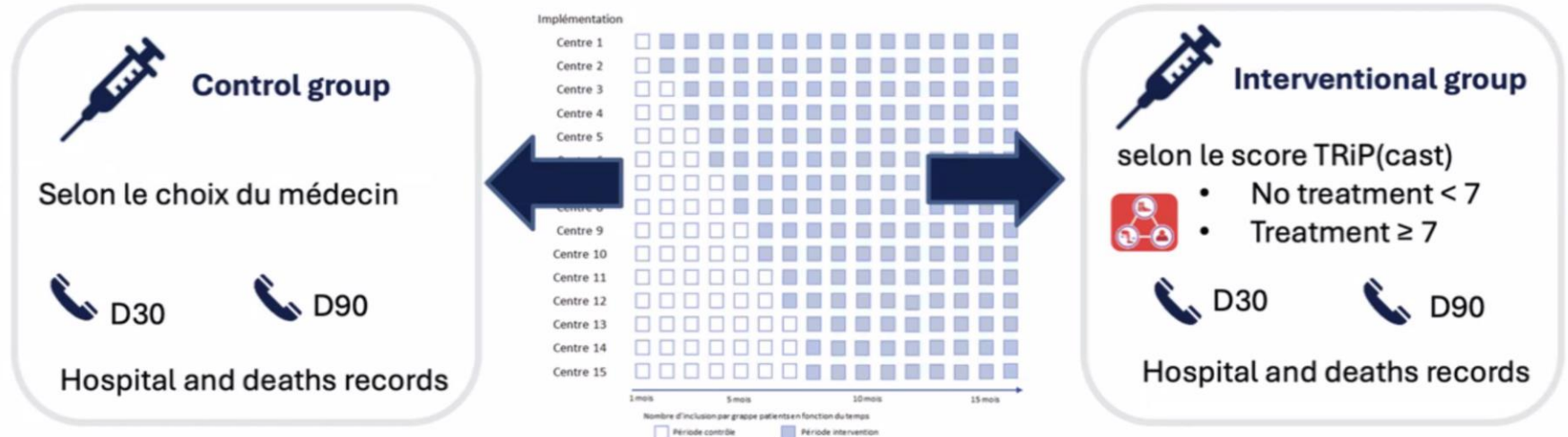


Figure 1: Flow chart

Essai randomisé stepped-wedge 15 centres en France et en Belgique

2120 patients



Comité d'adjudication

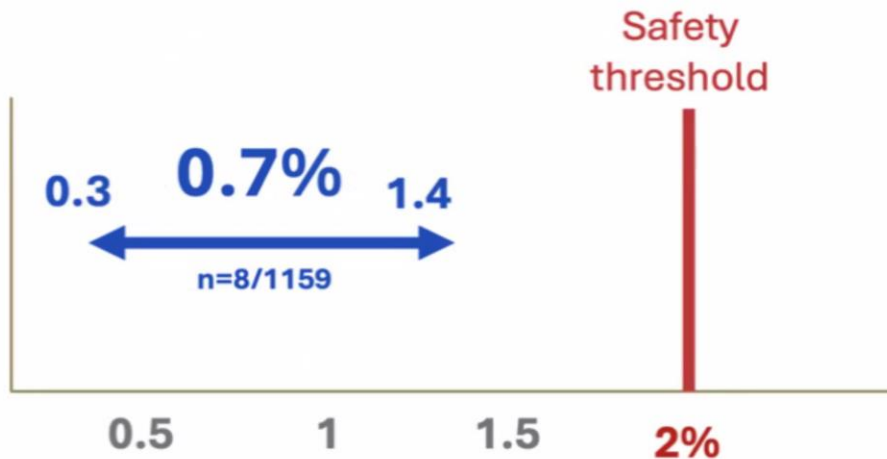


Taux d'ETEV symptomatique à 3 mois



As randomized

1159 patients à faible risque pendant la période d'intervention



Le critère de jugement principal:

le taux cumulé de MTEV symptomatique à 3 mois pendant la phase d'intervention chez les patients ayant un score TRiP(cast) < 7

Per-Protocol

0.8% (95%CI 0.3 to 1.5)

n=8/1048

Table 3: Primary and Secondary Outcomes

Outcome	As-Randomised Population			Per-Protocol Population
	Intervention phase (N=1505)	Control phase (N=603)	Difference (CI)	Intervention phase (N=1382)
	<i>no. (% 90-day probability)</i>		<i>percentage points</i>	
Primary outcome: symptomatic VTE in low-risk* patients	8/1159 (0.7; 95%CI 0.3 to 1.4)	-	-	8/1048 (0.8; 95%CI 0.3 to 1.5)
Secondary outcomes				
Anticoagulant prescription rate, no. of patients (%)	368 (24.5)	304 (50.4)	- 26.0 (-30.2 to -21.7)	-
VTE rate at 3 months, no. of patients (%)	17 (1.1)*	6 (1.0)	0.1 (-0.8 to 1.1)	-
VTE rate at 3 months in low-risk patients	8/1159 (0.7)	2/460 (0.4)	0.3 (-0.2 to 0.5)	
VTE rate at 3 months in high-risk patients	9/346 (2.6)	4/143 (2.8)	- 0.02 (-0.3 to 0.3)	
Fatal PE	0 (0)	(0)	-	-
Unexplained sudden death	0 (0)	1 (0.2)	-	-
Symptomatic PE	2 (0.1)	0 (0)	-	-
Symptomatic proximal DVT	2 (0.1)	1 (0.2)	-	-
Symptomatic distal DVT	11 (0.7)	4 (0.7)	-	-
Major bleeding	1 (0.1)	0 (0)	-	-
Clinically relevant non-major bleeding	1 (0.1)	0 (0)	-	-

Abbreviations: CI= confidence interval; DVT=deep-vein thrombosis; PE=pulmonary embolism; VTE=venous thrombo-embolism

* Low-risk patients identified by a TRIP(cast) score <7, high-risk patients identified by a score ≥7.

*Limited information on symptomatic DVT as defined by the Adjudication Committee was available for 2 patients (no ultrasound results).

Abbreviations: CI= confidence interval; DVT=deep-vein thrombosis; PE=pulmonary embolism; VTE=venous thrombo-embolism

Taux d'anticoagulation préventive



Control group

50.4%

N=304/603



Interventional group

24.5%

N=368/1505



Absolute difference
**- 26% (95%CI -30.2 to -
21.7) (p<0.001)**

Recommandations sur la gestion de la prévention antithrombotique dans les structures d'urgence

• Champ 1. Traumatisme des membres inférieurs nécessitant une immobilisation

Question 1 : Chez les patients ayant un traumatisme d'un membre inférieur, non opérés, nécessitant une immobilisation et pris en charge en ambulatoire, l'anticoagulation préventive permet-elle de réduire le taux d'événements thromboemboliques veineux et la morbi-mortalité associée ?

Borne ≥ 7 anticoagulation

R1.1.1. Chez les patients pris en charge aux urgences ayant un traumatisme isolé d'un membre inférieur nécessitant une immobilisation orthopédique par orthèse amovible ou immobilisation rigide et ne nécessitant *a priori* pas de prise en charge chirurgicale : il n'est probablement pas recommandé de prescrire systématiquement un traitement anticoagulant préventif.

GRADE 2 – (Accord fort)

R1.1.2. Il est recommandé de réaliser une évaluation du risque thromboembolique veineux en prenant en compte les facteurs de risque thromboembolique liés au patient (âge, antécédents personnels de thrombose...), le type de traumatisme (rupture du tendon d'Achille, fracture...) et le type d'immobilisation (immobilisation rigide, absence d'appui...) afin de juger de l'indication éventuelle d'une thromboprophylaxie.

GRADE 1+ (Accord fort)

R1.1.3. Il est probablement recommandé de se baser sur un score évaluant le risque thromboembolique.

GRADE 2 + (Accord fort)

R1.1.4. Les experts suggèrent de se baser sur le score TRiP(cast) pour évaluer le risque thromboembolique.

AVIS D'EXPERTS (Accord fort)

Take home message

Primum Non Nocere



We know that patients with lower limb fractures are particularly vulnerable to thromboembolism and we know that the vast majority of these patients present initially to the Emergency Department. Initiation of prophylactic therapy may or may not be appropriate at this time but we do have a responsibility to ensure that the risk of harm from the treatment we provide – the plaster cast – is minimised.

- **Recommandations Françaises modifiées en 2024**
- **Etude CASTING, Lancet 2024**
- **TRIP CAST score > ou= à 7 → bénéfice de la Thromboprophylaxie**
- **Prise en charge personnalisée**





Fractures de l'enfant



C'est cassé !

Immobilisation of torus fractures of the wrist in children (FORCE): a randomised controlled equivalence trial

Lancet 2023
Perry et al

Rationnel

Fracture en motte de beurre, de l'extrémité inférieure du radius

Plus commune

Prise en charge souvent maximaliste (immobilisation + suivi)

Méthode

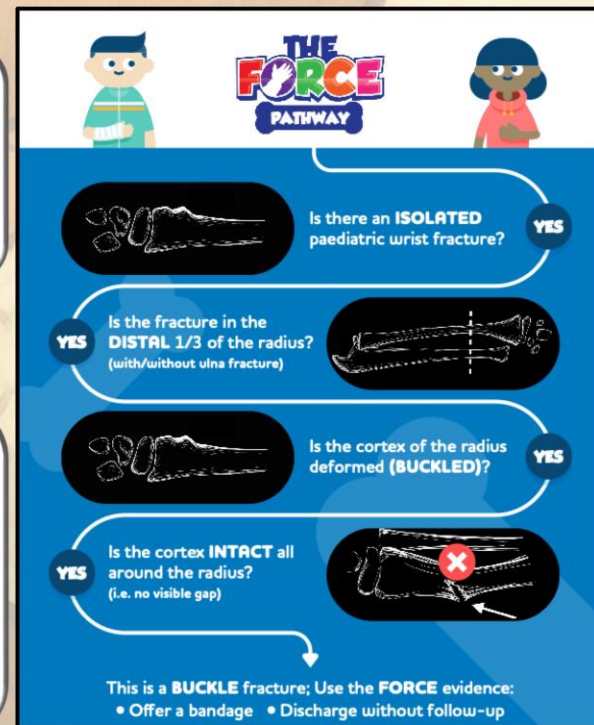
Essai contrôlé randomisé, d'**EQUIVALENCE**

Enfants de 5 à 15 ans, Fracture isolée ou avec fracture ulnaire

CJP : Wong-Baker scale, douleur à J 3, recueillie par sms/web

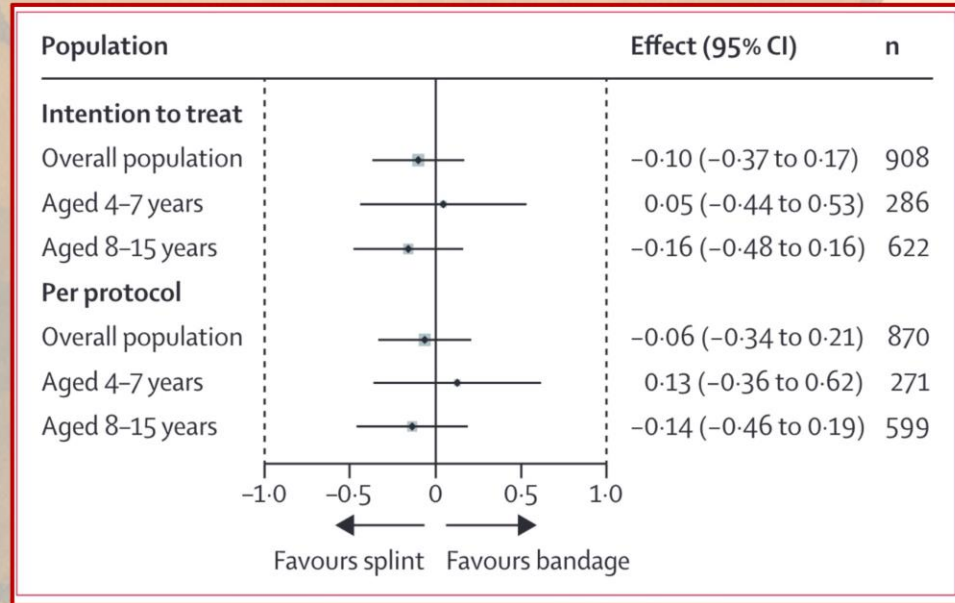
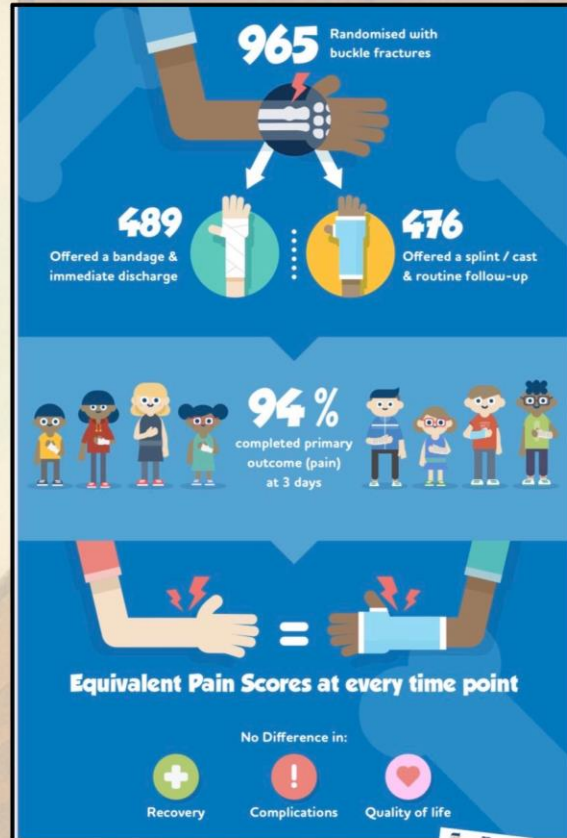
Immobilisation rigide et suivi

Proposer un bandage, et RAD

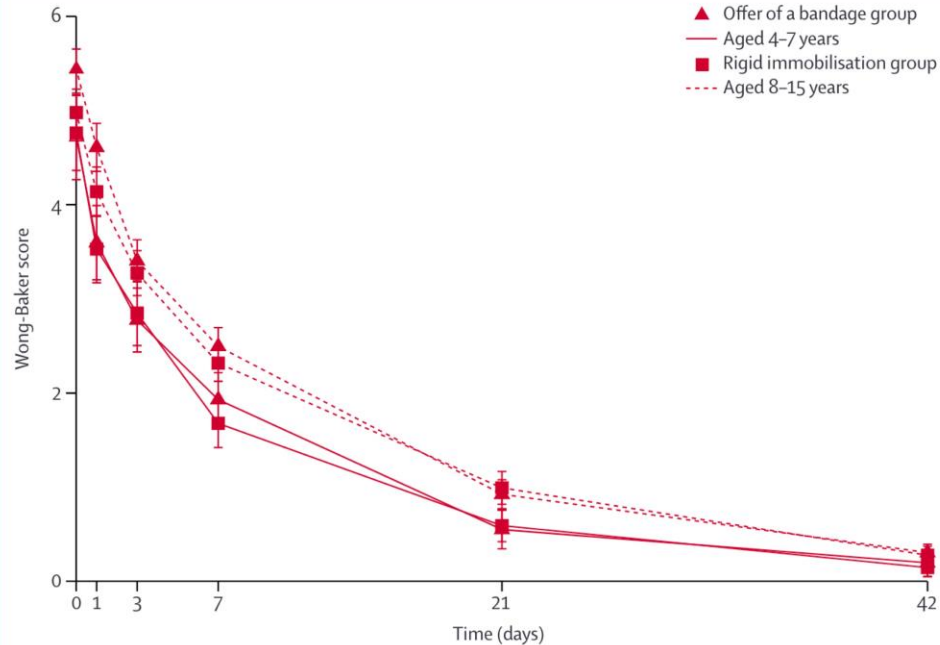


Immobilisation of torus fractures of the wrist in children (FORCE): a randomised controlled equivalence trial

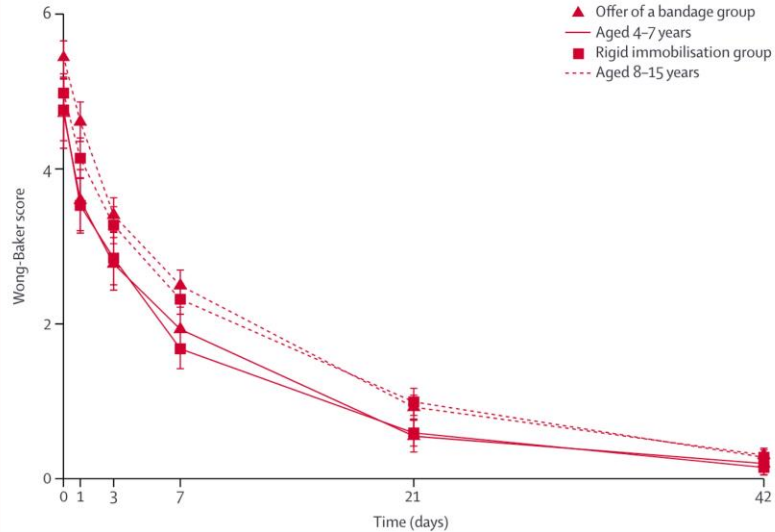
Lancet 2023
Perry et al



Immobilisation of torus fractures of the wrist in children (FORCE): a randomised controlled equivalence trial



Immobilisation of torus fractures of the wrist in children (FORCE): a randomised controlled equivalence trial



	Offer of bandage group (n=489)	Rigid immobilisation group (n=476)	Effect size (95% CI)*	p value
Satisfaction				
Day 1	2 (1, 2), 406	1 (1, 2), 380	..	<0.0001
Day 42	1 (1, 2), 433	1 (1, 2), 425	..	0.12
Use of any analgesia within the previous 24h				
Day 1	337/408 (83%)	297/382 (78%)	OR 0.53 (0.28 to 0.98)	0.04
Day 3	264/465 (57%)	227/442 (51%)	OR 0.60 (0.36 to 0.99)	0.05
Day 7	116/459 (25%)	100/439 (23%)	OR 0.70 (0.40 to 1.22)	0.21
School absence				
Participants who missed school	112/430 (26%)	93/425 (22%)	OR 0.79 (0.57 to 1.08)	0.14
Number of days of school missed	1.5 (1-2); n=112	1.5 (1-2); n=93	..	0.37
Any complication				
Alternative fracture: greenstick	1 (0.2%)	1 (0.2%)	..	..
Alternative fracture: complete but remains undisplaced	3 (0.6%)	2 (0.4%)	..	..
Other	1 (0.2%)	0	..	..



Co-ordinated by KADOORIE



965 Randomised with buckle fractures

489

Offered a bandage & immediate discharge



476

Offered a splint / cast & routine follow-up

94 %

completed primary outcome (pain) at 3 days



Equivalent Pain Scores at every time point

No Difference in:



Recovery



Complications



Quality of life

Use the force pathway

www.FORCEstudy.org



YES

Is the fracture in the **DISTAL 1/3** of the radius? (with/without ulna fracture)



YES

Is there an **ISOLATED** paediatric wrist fracture?

YES

Is the cortex **INTACT** all around the radius? (i.e. no visible gap)



YES

Is the cortex of the radius deformed (**BUCKLED**)?

This is a **BUCKLE** fracture; Use the **FORCE** evidence:
• Offer a bandage • Discharge without follow-up

To see the published evidence and patient materials, go to

www.FORCEstudy.org



The FORCarm fracture Recovery in Children Evaluation Study

Torus Fractures: Evidence from the FORCE study

Torus fractures (also called buckle fractures) of the wrist are the most common type of broken bone in children. They are also the mildest form of broken bone, in which the bone *crushes* (or *buckles*), instead of breaking.



Treatment

Despite being so common, there is no 'standard' or 'agreed' treatment. Traditionally doctors have used plaster casts and arranged outpatient follow-up. However, medical research suggested that wearing a bandage, or even having no treatment, might be similarly good.



About the FORCE Study

The FORCE study looked into whether or not a bandage (which was optional to wear) and no further follow-up resulted in the same levels of pain and speed of recovery as a hard splint and usual follow-up.

A total of 965 children aged 4–15 years from 23 emergency departments in the UK took part in the study. Children were fairly divided between the bandage and hard splint groups in a process called randomisation. Prior to the study, families told us that managing pain after injury was the most important issue to them. We asked children and their families to tell us about pain, recovery using the arm, quality of life, complications encountered and school absences. We also looked at the financial costs to families and the NHS.



What did the trial find?

The two treatments resulted in the same outcomes. The majority of those offered a bandage chose to wear it immediately. There was no difference at all in the levels of pain between those treated with a hard splint and usual outpatient follow-up and those offered a bandage and discharge (i.e. no further follow up) from hospital the same day. Similarly, there was no difference in the recovery using the arm, quality of life, complications encountered or school absences. There was a very slight increase in pain killer use in the bandage group at day 1, but not at any other time point. Overall, the cost of the offer of a bandage was slightly lower for families and the NHS.

In conclusion, the findings of this study support offering a bandage to be used at the discretion of families to treat children with a torus fracture of the wrist.



Scientific Article

Scientific Article

The FORCE study has been published in the *Lancet*.

[Go to the Lancet >](#)

Trial Report

Full trial report

The full FORCE study report is published in the *NHR* journals library.

[Go to report >](#)

Pathway

FORCE pathway

A pathway to help clinicians implement the FORCE study results.

[Download pathway >](#)



Patient Info Leaflet

Oxford Trauma on Twitter...



C'est cassé ?

Ultrasonography or Radiography for Suspected Pediatric Distal Forearm Fractures

NEJM 2023
Snelling et al

Rationnel

La fracture en motte de beurre, de l'extrémité inférieure du radius est toujours la **Plus** commune
La radio est l'examen de référence mais expose à un rayonnement et reste inaccessible à 2/3 de la population
Echographie ?

Méthode

Essai contrôlé randomisé, multicentrique, ouvert et de non-inf
Mené en Australie – Su pédiatriques ou mixtes
Enfants de 5 à 15 ans, traumatisme isolée ou non déformé de l'avant bras

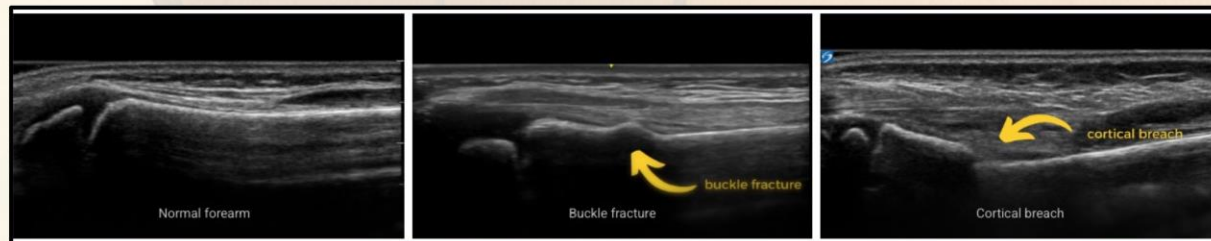
CJP : PROMIS, à 4 semaines

CJS: diagnostics , suivi des complications, de la douleur, de la satisfaction et des temps de prise en charge.



Ultrasonography or Radiography for Suspected Pediatric Distal Forearm Fractures

NEJM 2023
Snelling et al



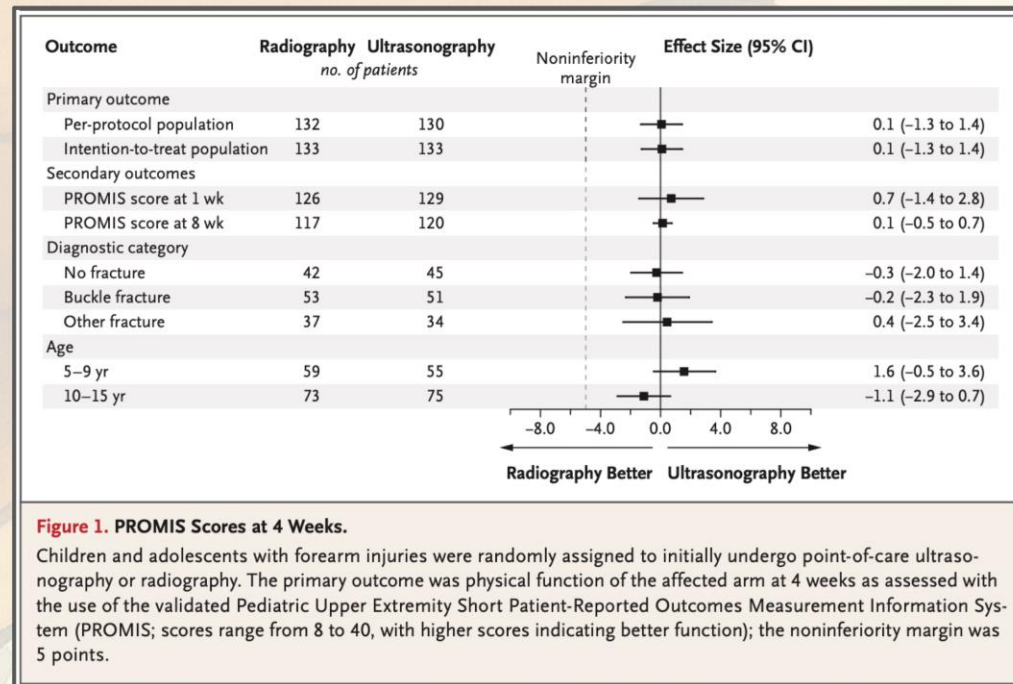
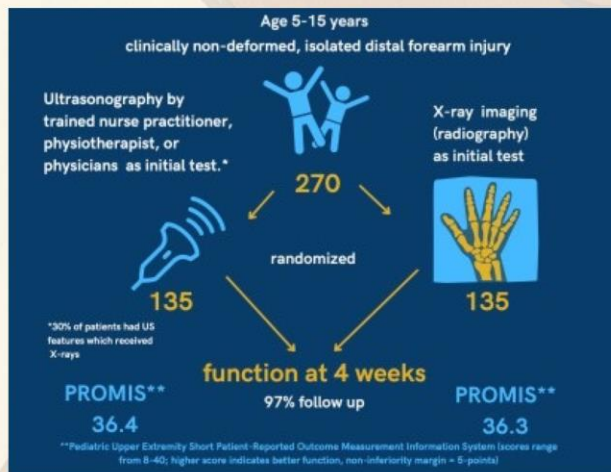
CJP: Fonction physique du bras à 4 semaines, mesurée par le score PROMIS (8 à 40, plus élevé = meilleure fonction), via questionnaire en ligne chez les enfants de 5 à 15 ans.

Sources: Don't Forget the bubbles, SKGEM

Ultrasonography or Radiography for Suspected Pediatric Distal Forearm Fractures

NEJM 2023
Snelling et al

Résultats



Ultrasonography or Radiography for Suspected Pediatric Distal Forearm Fractures

NEJM 2023
Snelling et al

Résultats

- ↑ **satisfaction du patient/parent**
- ↓ douleur à 1, 4 et 8 semaines
- ↓ **durée du séjour aux urgences**
- ↓ durée du traitement
- ↓ **jours d'absence scolaire**

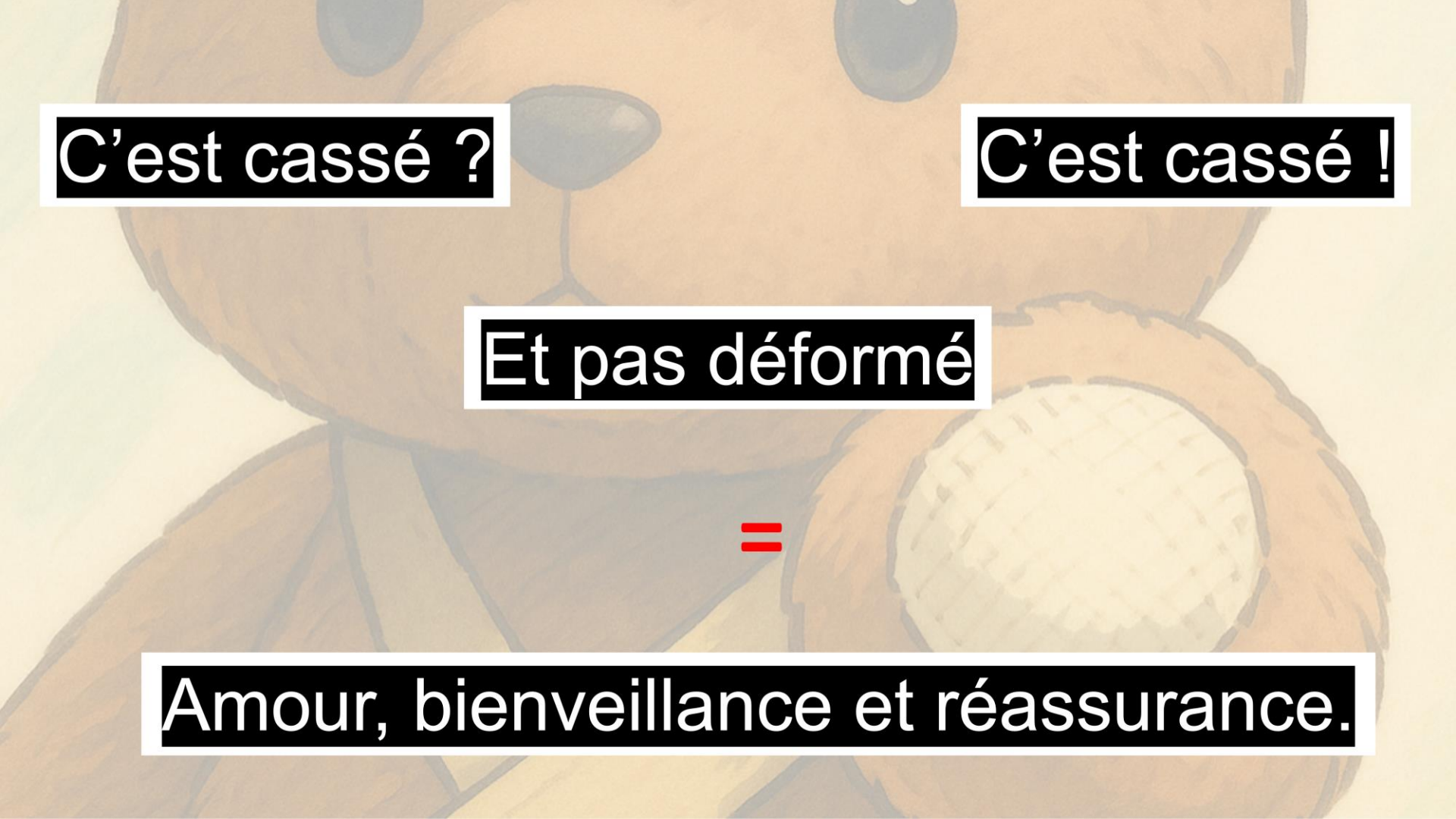
Conclusion

Des auteurs:

Chez les enfants et adolescents présentant un traumatisme distal de l'avant-bras, l'utilisation de l'échographie comme méthode d'imagerie initiale s'est révélée non inférieure à la radiographie en ce qui concerne la fonction physique du bras à 4 semaines.

La mienne:

L'échographie peut être une méthode appropriée pour diagnostiquer les fractures en cas de **traumatismes distaux de l'avant-bras sans déformation clinique**, notamment lorsque l'accès à la radiographie est difficile.



C'est cassé ?

C'est cassé !

Et pas déformé

=

Amour, bienveillance et réassurance.

IA et triage :
gadget, aide ou
danger ?





IA et triage : gadget, aide ou danger ?

Rationnel



Triage = étape critique, souvent sous pression.



Problèmes : subjectivité, sous/sur-triage, surcharge.



IA permettrait de :

- Réduire les biais
- Identifier les patients critiques
- Optimiser les ressources

IA et triage : gadget, aide ou danger ?

Rationnel



L'IA analyse des données cliniques structurées et non structurées.



Exemples : motif de consultation, paramètres vitaux, mais aussi texte libre du dossier.



Outils : NLP, modèles prédictifs, LLM (grands modèles de langage).

AI-driven triage in emergency departments: A review of benefits, challenges, and future directions

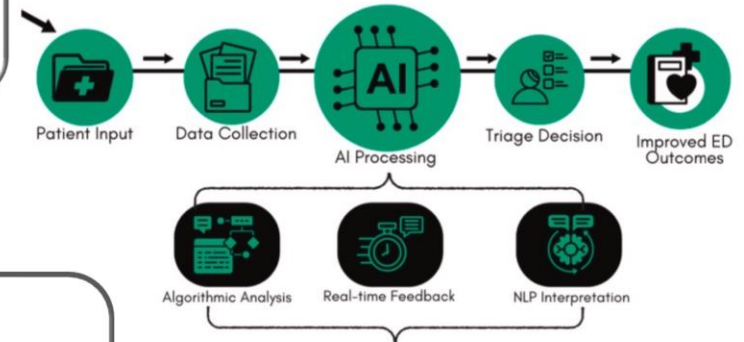
Int J Med Inform
2024
Da'Costa et al

Rationnel

Objectif : Faire le point sur les **bénéfices, défis, composants techniques et perspectives** des systèmes de tri basé sur de l'IA

Méthode

- Revue narrative de la littérature (2015–2024) sur l'IA appliquée au tri en MU
- Sources : PubMed, Scopus, IEEE Xplore, Google Scholar



AI-driven triage in emergency departments: A review of benefits, challenges, and future directions

Int J Med Inform
2024
Da'Costa et al

Résultats

Composants clés des systèmes IA de triage

🇫🇷 Collecte et traitement des données
(antécédents, paramètres vitaux, données démographiques)

🧠 Modèles algorithmiques : arbres de décision, réseaux de neurones, régression

🕒 Analyse en temps réel : priorisation dynamique selon les ressources disponibles

📄 NLP : analyse des observations cliniques et données non structurées



1. Amélioration de la priorisation des patients et réduction des délais de soins
2. Décisions moins subjectives
3. Allègement de la charge cognitive des soignants
4. Optimisation des ressources (filières, personnel etc.)



AI-driven triage in emergency departments: A review of benefits, challenges, and future directions

Int J Med Inform
2024
Da'Costa et al

MAIS

- 🔍 Qualité et diversité des données
- ⚖️ Biais algorithmiques (ex. sous-évaluation de patients selon genre/ethnie)
- 🔒 Enjeux éthiques et de confidentialité des données
- ❓ Acceptabilité par les soignants (manque de transparence, "boîte noire")

Et après

- 🎯 Amélioration continue des algorithmes avec données diversifiées
- 🕒 Intégration avec dispositifs connectés (wearables)
- 🎓 Formation des soignants à l'usage de l'IA
- 📖 Développement de cadres éthiques (transparence, supervision humaine)

Use of a Large Language Model to Assess Clinical Acuity of Adults in the Emergency Department

Jama Open 2024
Williams et al

Rationnel

Objectif : Évaluer si un grand modèle de langage (LLM, ici GPT-4) peut correctement classer l'acuité des patients

Méthode

- Analyse de 251 401 passages aux urgences adultes (UCSF, 2012–2023)
- Constitution de 10 000 paires de patients avec niveaux ESI différents (5 niveaux)
- Le LLM devait désigner, à partir du texte seul, le patient le plus grave dans chaque paire
- Comparaison avec un médecin humain sur un sous-échantillon de 500 paires

Use of a Large Language Model to Assess Clinical Acuity of Adults in the Emergency Department

Jama Open 2024
Williams et al

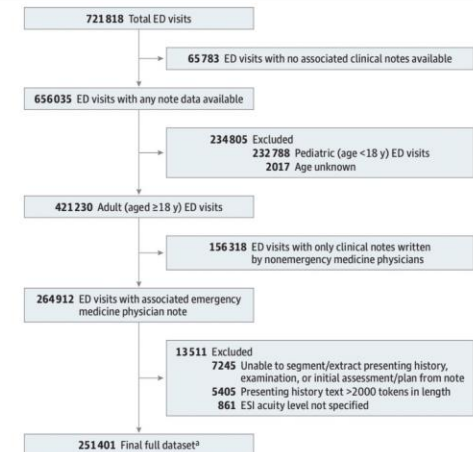
Résultats

GPT-4 : 89 % de précision globale (vs 84 % pour GPT-3.5)

Comparable à un médecin (88 % LLM vs 86 % humain sur 500 paires)

Performance maximale (100 %) sur les cas très contrastés (ex : ESI 1 vs ESI 5)

Figure 1. Flowchart of Included Emergency Department (ED) Visits



Use of a Large Language Model to Assess Clinical Acuity of Adults in the Emergency Department

Jama Open 2024
Williams et al



Interprétation

LLM capable de **stratifier**

**Comparable au
raisonnement clinique
humain** sur des tâches de
catégorisation

Mais

Utilisation de l'histoire clinique
seule

Données post-tri

Performances moindres pour
les distinctions fines tri 4 vs 5

Impact of Artificial Intelligence–Based Triage Decision Support on Emergency Department Care

NEJM AI 2025
Taylor et al

Rationnel

Objectif : Évaluer l'impact de la mise en place d'un outil de tri, se basant sur l'IA, sur la performance du tri et la fluidité des parcours patients.

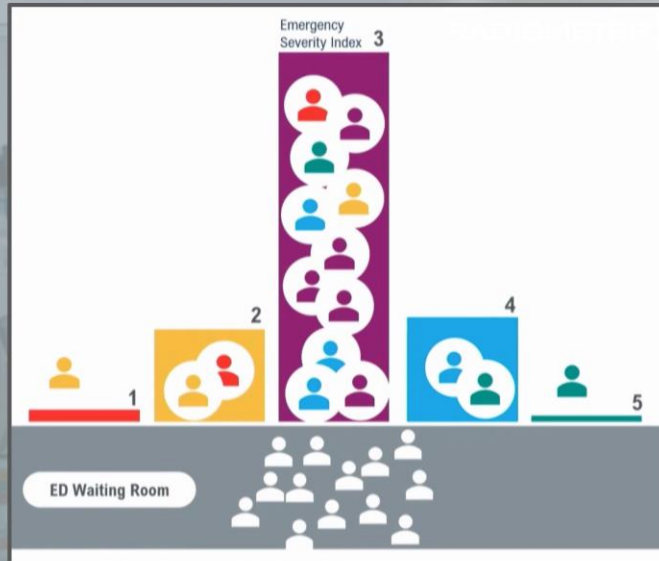
Méthode

- Etude Avant/Après, dans 3 SU, sur un an (180 jours avant et après)
- Algo/outil TriageGO: intégré au dossier patient, utilisant ; âge, sexe, paramètres vitaux, mode d'arrivée, antécédents, motif de recours
- **CJP:** sensibilité/spécificité du besoin en soins critiques, chirurgie urgente et hospi.
- **CJP « flux »:** temps avant prise en charge, avant décision et avant sortie.

Impact of Artificial Intelligence–Based Triage Decision Support on Emergency Department Care

NEJM AI 2025
Taylor et al

Résultats



Impact of Artificial Intelligence–Based Triage Decision Support on Emergency Department Care

NEJM AI 2025
Taylor et al

Résultats

- Sensibilité et Spécificité améliorée: 78,8 % à 83,1 % (soins critiques)
- Impact sur les flux patients (médianes) :
 - Temps arrivée → prise en charge : -33 %
 - Temps arrivée → décision d'orientation : -4,2 %
 - Temps arrivée → sortie des urgences : -6,1 %

Mais

- Taux d'adhésion médian : **70 %**, avec forte variabilité inter-individuelle

Lien homme/machine

IDE + IA : surpassaient IA seule

À l'inverse, IDE qui ne suivaient jamais l'IA : moins bonne performance que le modèle seul.

IA et triage : gadget, aide ou danger ?

Conclusion

-  Amélioration des patients les plus graves
-  Optimisation des flux

Mais quel est le bon critère?

-  Acceptabilité variable
-  Outils d'aide et non remplaçants

