
Stroke & TIA @ home: do's & don'ts

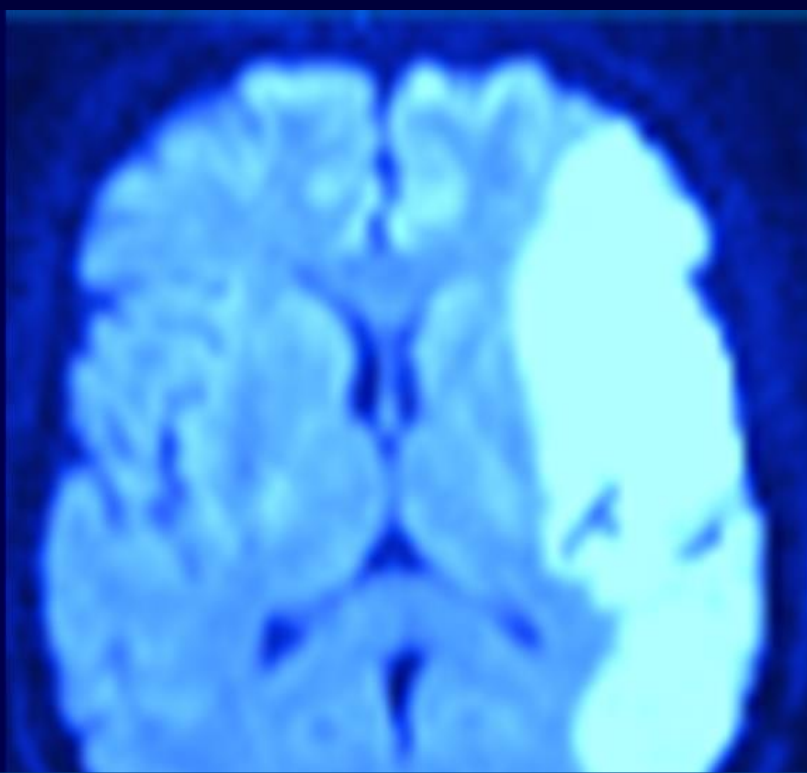
Yves VANDERMEEREN MD, PhD

Neurology Department | Stroke Unit

Vignette clinique 1

- Patient de 87 ans
- Brutalement, troubles de la marche & hémiparésie
- Son fils téléphone au 112
- Réponse du 112 : « *Appelez votre MT* »
- Quelle attitude ?





Acute stroke management

Stroke epidemiology



1 in 4 of us will have a stroke.
**DON'T BE
THE ONE**
World Stroke Day: October 29th



- 1 in 4 of us will have a stroke in our lifetime
- Almost all strokes can be prevented
- The leading (1st) cause of long-term disability
- 2nd leading cause of death



1 in 4

In Belgium, every year
~25.000 stroke victims



Acute stroke management

**RECONNAÎTRE LES SYMPTÔMES D'UN AVC,
C'EST SAUVER DES VIES.**



Observez si la bouche est de travers.

Observez si un bras (ou une jambe) ne bouge plus.

Observez si la personne s'exprime de manière confuse.

Notez l'heure du début des symptômes. Les chances de rétablissement sont plus grandes si le traitement est instauré dans les trois heures.



112
NUMÉRO DE SECOURS

www.ReconnaitreUnAVC.be



Jusqu'à preuve du contraire, toute personne :

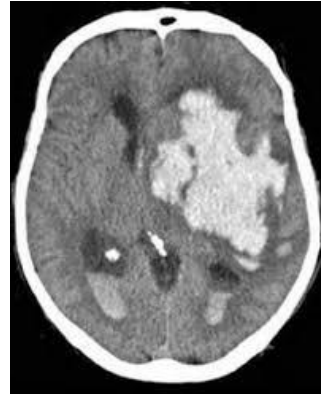
1. qui présente un déficit neurologique brutal est victime d'un AVC
2. victime d'un AVC est candidate à la thrombolyse IV
3. victime d'un AVC est candidate à la thrombectomie

Acute stroke management

Jusqu'à preuve du contraire, toute personne :

1. qui présente un déficit neurologique brutal est victime d'un AVC ...

80%



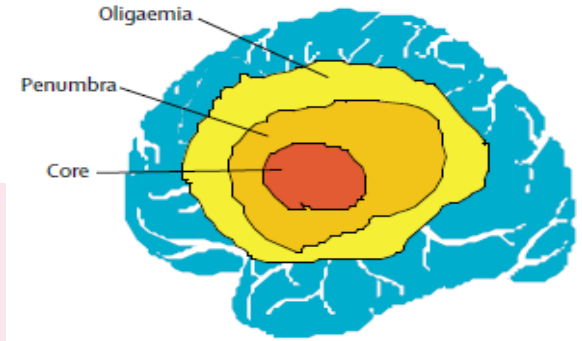
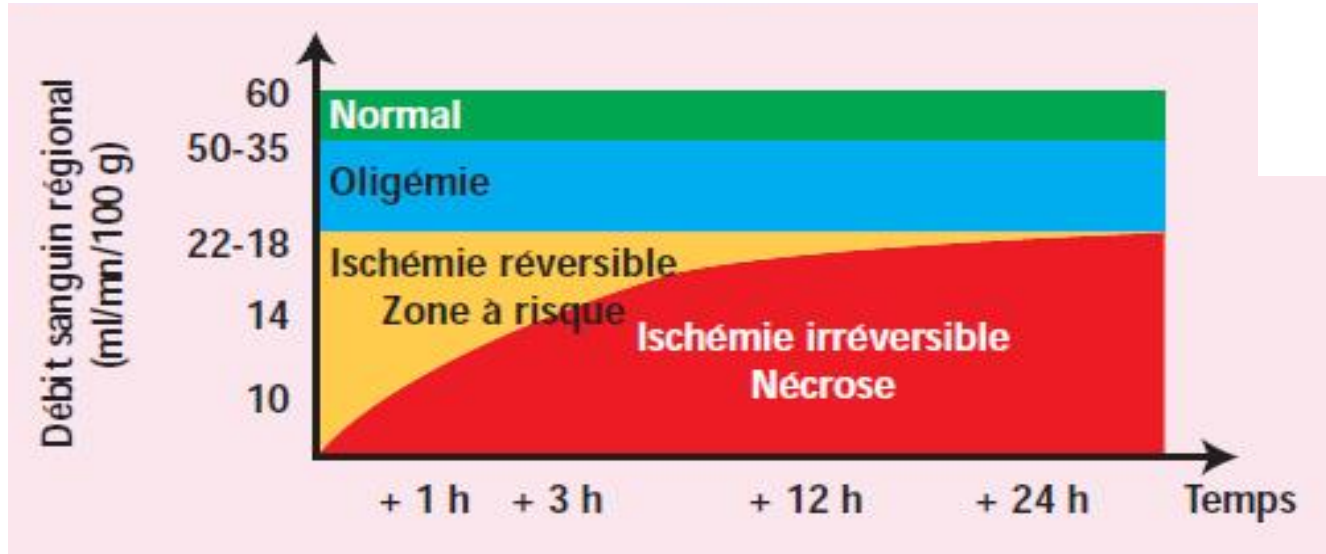
20%

Cliniquement impossible d'exclure formellement un AVC hémorragique

→ Imagerie cérébrale en urgence

Pathophysiology of ischemic stroke

Penumbra model

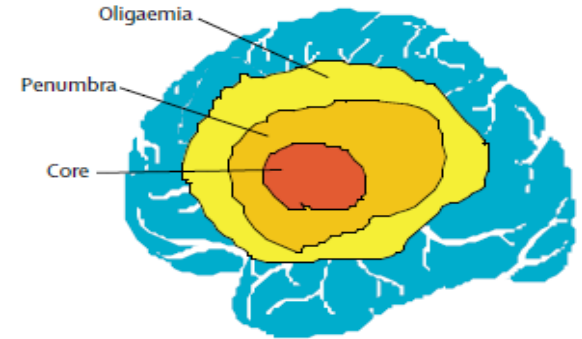


Muir et al., *Lancet Neurology* 2006;5;755-768

Nighoghossian,
Lett Neurol 2006;2;20-24

Pathophysiology of ischemic stroke

The mantra: time is brain



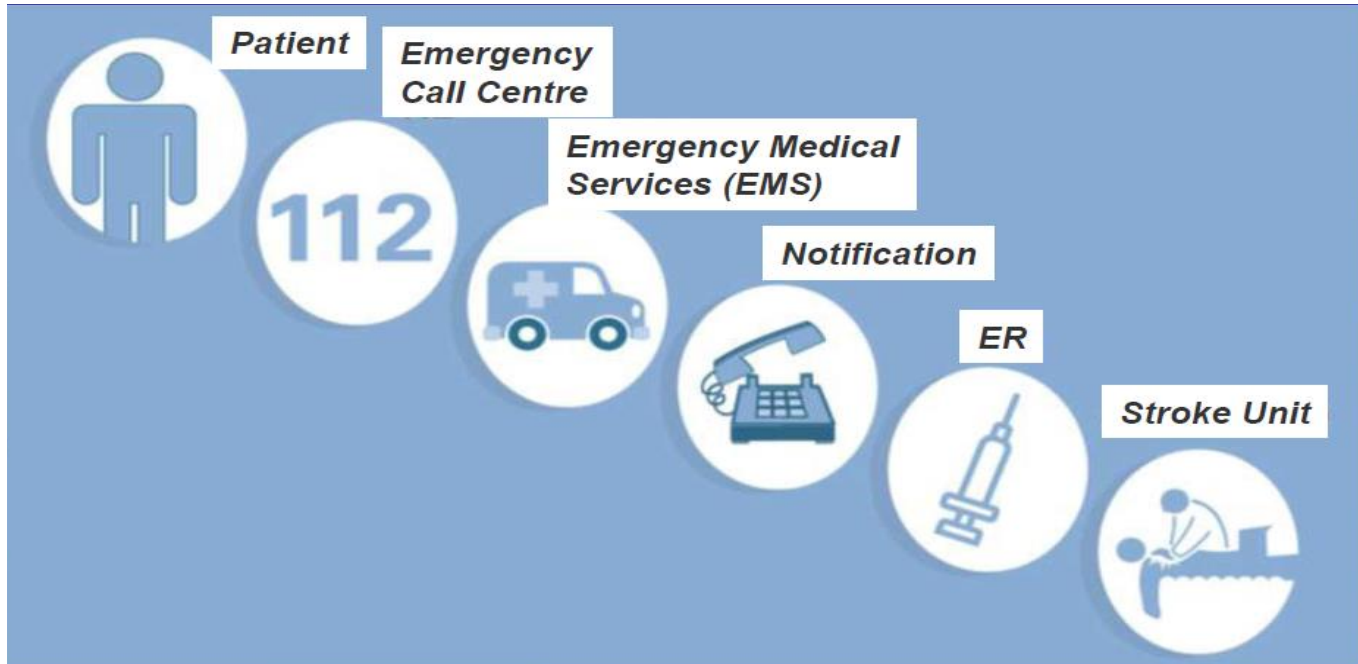
Estimated Pace of Neural Circuitry Loss in Typical Large Vessel, Supratentorial Acute Ischemic Stroke

	Neurons Lost	Synapses Lost	Myelinated Fibers Lost	Accelerated Aging
Per Stroke	1.2 billion	8.3 trillion	7140 km/4470 miles	36 y
Per Hour	120 million	830 billion	714 km/447 miles	3.6 y
Per Minute	1.9 million	14 billion	12 km/7.5 miles	3.1 wk
Per Second	32 000	230 million	200 meters/218 yards	8.7 h

Saver. *Stroke* 2006; 37: 263-266

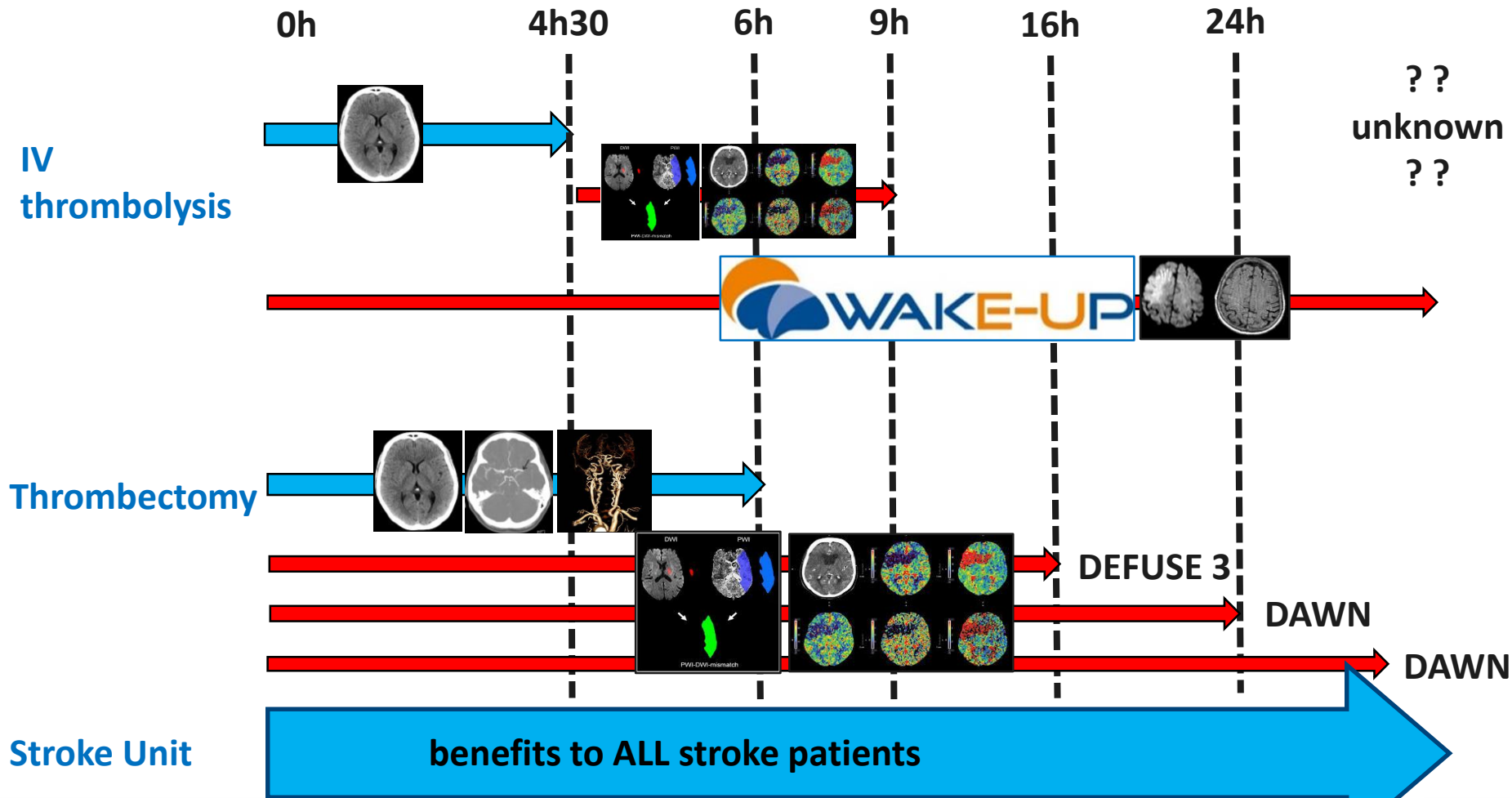
Acute stroke management

Time is brain / chain of survival



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Dptmt of Neurology Helsinki University Central Hospital



Pre-hospital acute stroke management

Referral and Patient Transfer

Guidelines for Management of Ischaemic Stroke and Transient Ischaemic Attack 2008

The European Stroke Organisation (ESO) ESO Writing Committee

Cerebrovasc Dis 2008;25:457–507
DOI: [10.1159/000131083](https://doi.org/10.1159/000131083)

Recommendations 2008

- Transport without delay to the nearest medical centre with a **Stroke Unit** that can provide ultra-early treatment
- Dispatchers & ambulance personnel should be trained to recognise stroke (**FAST**)
- Remote / rural areas → **helicopter** transfer / **tele-medicine** to be considered

ESO-EAN: pre-hospital management of stroke 2018

1. SaO₂ levels <95% → O₂ titrated to maintain normoxia
Routine use of O₂ recommended
not recommended
5. Pre-hospital treatment of **high blood pressure**
in people suspected of acute stroke **not** recommended
6. Pre-hospital administration of **insulin**
in persons with suspected stroke & **hyperglycaemia** **not** recommended
7. Pre-hospital treatment of **elevated body temperature** **no** recommendation (?)
8. All EMS implement a **'code stroke' protocol**: highest priority dispatch,
pre-hospital notification & **rapid transfer** to the closest 'stroke-ready' centre

Kobayashi et al. *Eur J Neurol.* 2018, 25: 425–433

ESO EUROPEAN
STROKE
ORGANISATION

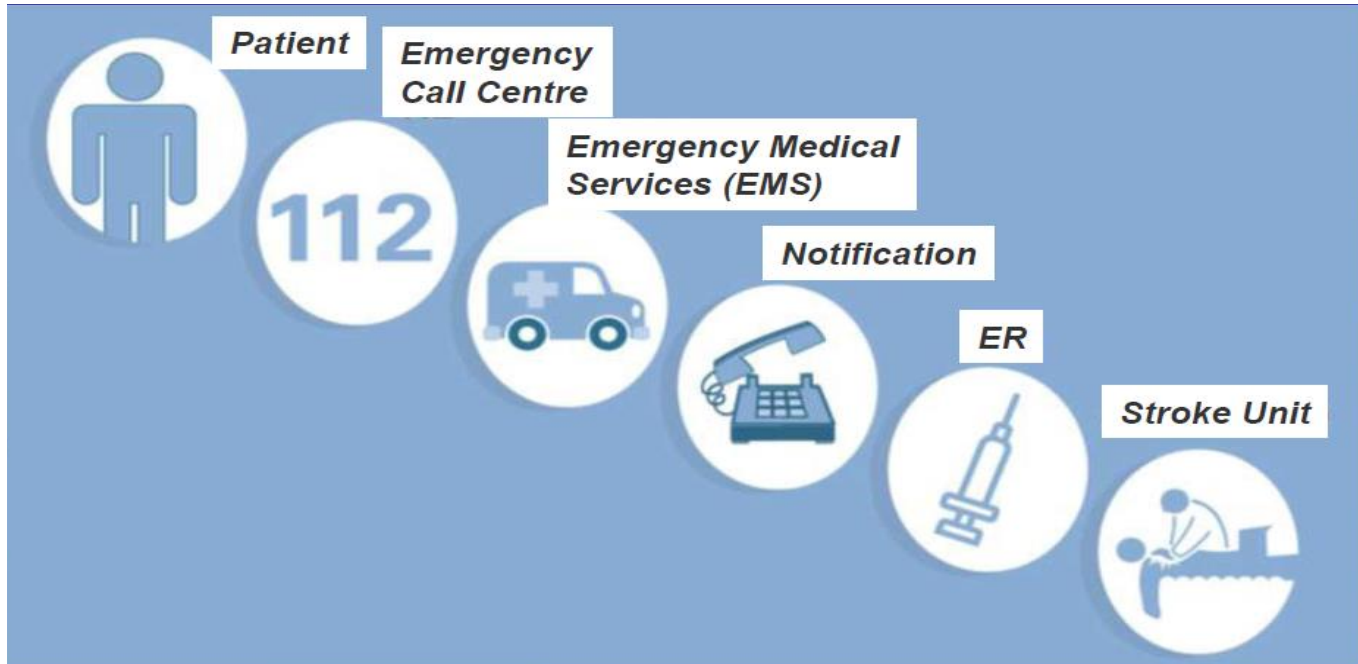
Pre-hospital acute stroke management

@ home, if stroke is the most likely diagnosis :

- | | |
|---|--|
| <ul style="list-style-type: none">■ Aspirin (ASA)■ Low molecular weight heparin | <ul style="list-style-type: none">■ haemorrhagic ? thrombolysis ?■ haemorrhagic ? thrombolysis ? |
| <ul style="list-style-type: none">■ Paracetamol■ Blood pressure lowering drug■ 0° supine position (flat) | <ul style="list-style-type: none">■ if fever (<i>unlikely early</i>)■ impaired cerebral hemodynamic■ impaired cerebral hemodynamic (→ 30°) |
| <ul style="list-style-type: none">■ Oxygen■ Anti-emetic | <ul style="list-style-type: none">■ if severe stroke■ if needed |

Acute stroke management

Time is brain / chain of survival

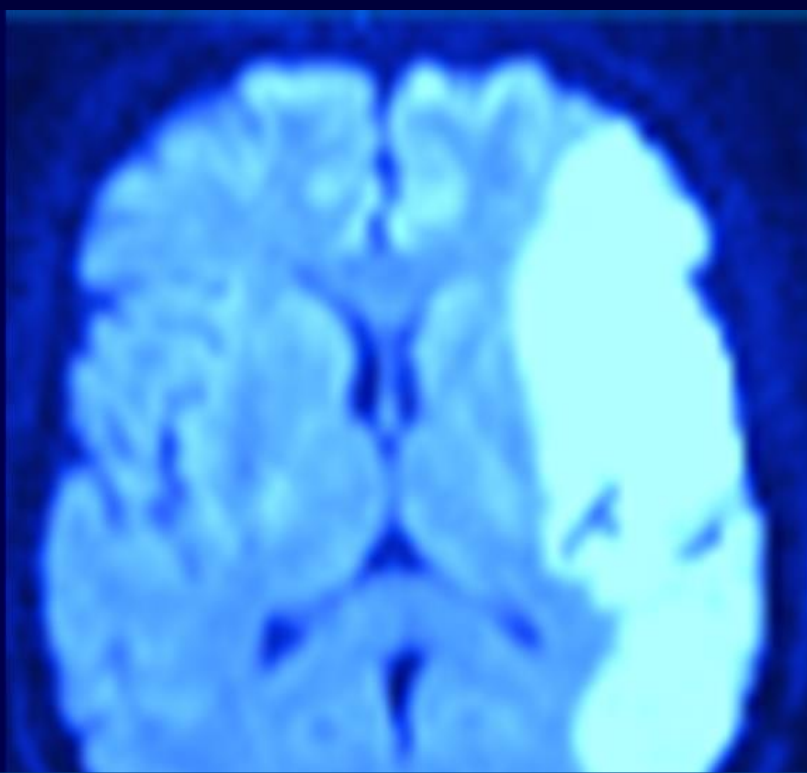


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Vignette clinique 2

- Patiente de 87 ans
- Un épisode transitoire de confusion & d'aphasie ... **il y a 15 j**
- Tout est rentré dans l'ordre depuis.
- Régulièrement suivie par le cardiologue.
- Bilan : biologie, CT-scan cérébral, écho-doppler vaisseaux du cou, consultation neuro.
- **15 j après** → Urgences pour déficit neuro avec perte de connaissance
- ... Nouvel AIT.



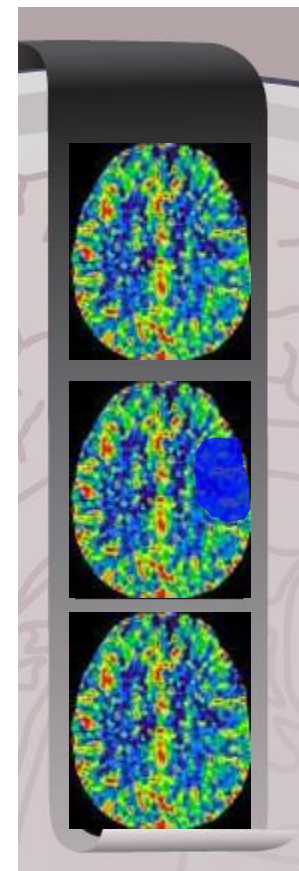
TIA management

Cerebrovascular diseases: definitions

Transient ischemic attack (TIA)

AHA/ASA 2009 tissue-based definition:

- “A transient episode of neurologic dysfunction caused by focal brain, spinal cord or retinal ischemia **without acute infarction**”
- TIA is caused by a clot (*ischemic*)
- **Only ≠ with a stroke: the arterial blockage is transient (temporary)**
- Most TIAs < **5 min**, average **~1 min**
- NB. “spells” in cerebral amyloid angiopathy (CAA)

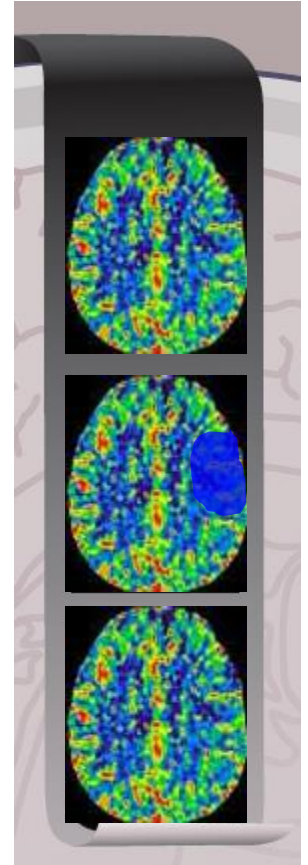


Cerebrovascular diseases: definitions

Transient ischemic attack (TIA)

AHA/ASA 2009 tissue-based definition:

- *“A transient episode of neurologic dysfunction caused by focal brain, spinal cord or retinal ischemia **without acute infarction**”*
- Even if the tissue-based definition is used,
the diagnosis of TIA is still based on clinical grounds



TIA management

In ~25% of stroke patients, a TIA has preceded the stroke

Meta-analysis 14.889 patients, pooled stroke risk:

- @ 2 days **1,4%**
- @ 7 days **2 %**
- @ 30 days **2,8%**
- @ 90 days **3,4%**
- @ 1 year **12%**






Valls et al. *Cerebrovasc Dis.* 2017;
43: 90-98. doi: 10.1159/000452978.

TIA = emergency !

→ Quick management → Hospital: ER – Stroke Unit ?
→ ? TIA clinic ?

- Risk factors ?
- Etiology ?
- → 2^{ary} prevention

European Stroke Organisation (ESO) guidelines on management of transient ischaemic attack

Ana Catarina Fonseca^{1,*} , Áine Merwick^{2,*} , Martin Dennis³,
Julia Ferrari⁴, José M Ferro¹, Peter Kelly⁵ , Avtar Lal⁶,
Angel Ois⁷ , Jean Marc Olivot⁸ and Francisco Purroy⁹ 

European Stroke Journal
2021, Vol. 6(2) CLXIII–CLXXXVI
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DOI: 10.1177/2396987321992905
journals.sagepub.com/home/eso



Table 1. Synoptic table of all recommendations and expert consensus statements.

Topic / PICO Question	Recommendations	Expert consensus statement
<p>Services Organization</p> <p>1.1: In patients suspected of TIA does stroke specialist review of the patient within 24 hours compared to more than 24 hours reduce TIA/stroke recurrence?</p>	<p>In patients with a TIA, we recommend specialist review of the patient within 24 hours after the onset of symptoms compared to assessment more than 24 hours after symptoms onset.</p> <p>Quality of evidence: Low ⊕⊕ Strength of recommendation: Strong for intervention ††</p> <p>In patients with a TIA, we suggest specialist review in a TIA clinic within 24 hours over a conventional outpatient appointment more than 24 hours after the TIA.</p> <p>Quality of evidence: Low ⊕⊕ Strength of recommendation: Weak for intervention †?</p> <p>There is insufficient evidence to provide a recommendation.</p>	
<p>2.1: In patients suspected of TIA does stroke specialist review of the patient in a TIA clinic within 24 hours compared to conventional outpatient appointment more than 24 hours?</p>		
<p>3.1: In patients suspected of high-risk TIA does stroke specialist review of the patient in a TIA clinic within 24 hours compared to hospitalization in a stroke unit reduce stroke recurrence risk?</p>		<p>In patients suspected of high-risk TIA, 9/9 experts suggest that prompt review in a TIA clinic or hospitalization in a stroke unit are reasonable options as settings for evaluation by a stroke specialist, depending on local available resources and the patients' preferences, in the absence of evidence comparing each approach.</p>
<p>Risk prediction tools</p> <p>4.1 In patients suspected of TIA does the use of risk prediction tools by primary care physicians compared to not using risk prediction tools reduce the risk of stroke recurrence, accurately identify high-risk patients, and improve diagnostic accuracy of TIA?</p>	<p>For patients with suspected TIA, we suggest not to use prediction tools alone to identify high risk patients/ make triage and treatment decisions.</p> <p>Quality of evidence: Very low ⊕ Strength of recommendation: Weak against intervention †?</p> <p>There is insufficient evidence to provide a recommendation.</p>	
<p>Imaging</p> <p>5.1: For patients with suspected TIA does the use of MRI (DWI/PWI) or CT Perfusion vs standard CT alone decrease stroke recurrence by accurately identifying an ischaemic mechanism and therefore patients at high stroke risk?</p>		
<p>6.1: In suspected TIA patients is the use of MR angiogram (MRA) compared to CT angiography (CTA) superior for identifying patients with large artery stenosis of 50% or greater and therefore patients with high risk of stroke recurrence?</p>	<p>In TIA patients, we suggest using either MRA or CTA for additional confirmation after ultrasound of large artery stenosis of 50% or greater, to guide further management.</p> <p>Quality of evidence: Very low ⊕ Strength of recommendation: Weak for intervention †?</p> <p>In patients suspected of TIA, if a wait of more than 24 hours to planned imaging is foreseen and a delay is</p>	<p>In suspected TIA patients, to confirm ischaemic pathophysiology of transient neurological symptoms, where it will influence treatment and /or there is diagnostic uncertainty, 8/9 experts suggest to use MR (multimodal) or CT perfusion, if feasible, instead of non-contrast CT.</p>
<p>Secondary prevention</p> <p>7.1: In patients with suspected acute TIA does "de</p>		
<p>novo" antiplatelet usage (prior to imaging) compared to delayed antiplatelet usage reduce stroke recurrence?</p>	<p>judged to increase the risk of further ischaemic events, above the risk of starting antiplatelet medication, we suggest "de novo" antiplatelet monotherapy usage compared to not starting antiplatelet monotherapy.</p> <p>Quality of evidence: Low ⊕⊕ Strength of recommendation: Weak for intervention †?</p> <p>In patients with acute non-cardioembolic high risk TIA (ABCD2 score of 4 or more), we recommend short term dual antiplatelet therapy with aspirin and clopidogrel over monotherapy, subsequently followed by monotherapy.</p> <p>Quality of evidence: High ⊕⊕⊕⊕ Strength of recommendation: Strong for intervention ††</p>	
<p>8.1: In patients with non-cardioembolic acute TIA does dual antiplatelet therapy (DAPT) compared to monotherapy reduce the risk of stroke recurrence?</p>		<p>For patients with acute non-cardioembolic low risk TIA or uncertain TIA diagnosis, 9/9 experts voted against using dual antiplatelet therapy over monotherapy.</p>

Guide

European guidelines

Ana Julia Ang



ESO guidelines | TIA management

Recommendation

In patients with a TIA, we recommend specialist review of the patient within 24 hours after the onset of symptoms compared to assessment more than 24 hours after symptoms onset

Quality of evidence: **Low** ⊕⊕

Strength of recommendation: **Strong for intervention** ↑↑



...

Fonseca et al. *Eur Stroke J.* 2021; 6
doi: 10.1177/2396987321992905.

ESO guidelines | TIA management

Recommendation

In patients with a TIA, we recommend specialist review of the patient within 24 hours after the onset of symptoms compared to assessment more than 24 hours after symptoms onset

Quality of evidence: Low ⊕⊕

Strength of recommendation: Strong for intervention ↑↑

Fonseca et al. *Eur Stroke J.* 2021; 6
doi: 10.1177/2396987321992905.

ESO guidelines | TIA management

Dual antiplatelet therapy (DAPT)

ideally, within the first 24 h



High risk
transient ischaemic
attack (TIA)

A score of 4 or more on the ABCD2 scale,
which estimates the risk of recurrent
stroke after a TIA



- **Day 1 :** clopidogrel 300 mg + ASA 80 mg
- **Days 15-21 :** clopidogrel 75 mg + ASA 80 mg
- **> 21 days :** clopidogrel OR ASA

Prasad et al.
BMJ 2018;363

ABCD²

A	Age \geq 60	1
B	sBP \geq 140 mmHg dBP \geq 90 mmHg	1
C	Clinical : hemiparesis	2
	speech disturbance	1
D	Duration \geq 60 min	2
	10-59 min	1
	< 10 min	0
D	Diabetes	1

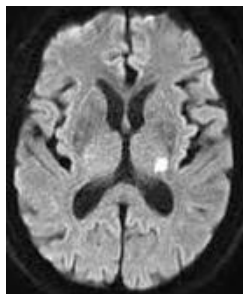
ESO guidelines | TIA management

Recommendation

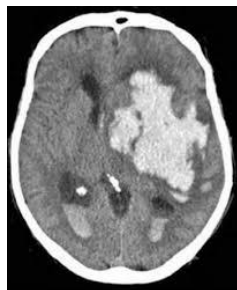
In patients suspected of TIA, if a wait of more than 24 hours to planned imaging is foreseen and a delay is judged to increase the risk of further ischaemic events, above the risk of starting antiplatelet medication, we suggest “de novo” antiplatelet monotherapy usage compared to not starting antiplatelet monotherapy.

Quality of evidence: **Low** ⊕⊕

Strength of recommendation: **Weak for intervention** †?



> ?



Fonseca et al. *Eur Stroke J.* 2021; 6
doi: 10.1177/2396987321992905.

Guidelines | TIA management

Impact of treatments

Treatment	Spared stroke
Antiplatelet drugs	- 25%
Antihypertensive drugs	- 24%
Statins	- 17%
Anticoagulation for AFib	- 62%
Thromboendarterectomy	- 50%

Guidelines | TIA management

1. Manage Blood Pressure
2. Control Cholesterol
3. Reduce Blood Sugar
4. Get Active
5. Eat Better
6. Lose Weight
7. Stop Smoking



AMERICAN HEART ASSOCIATION
JOURNALS

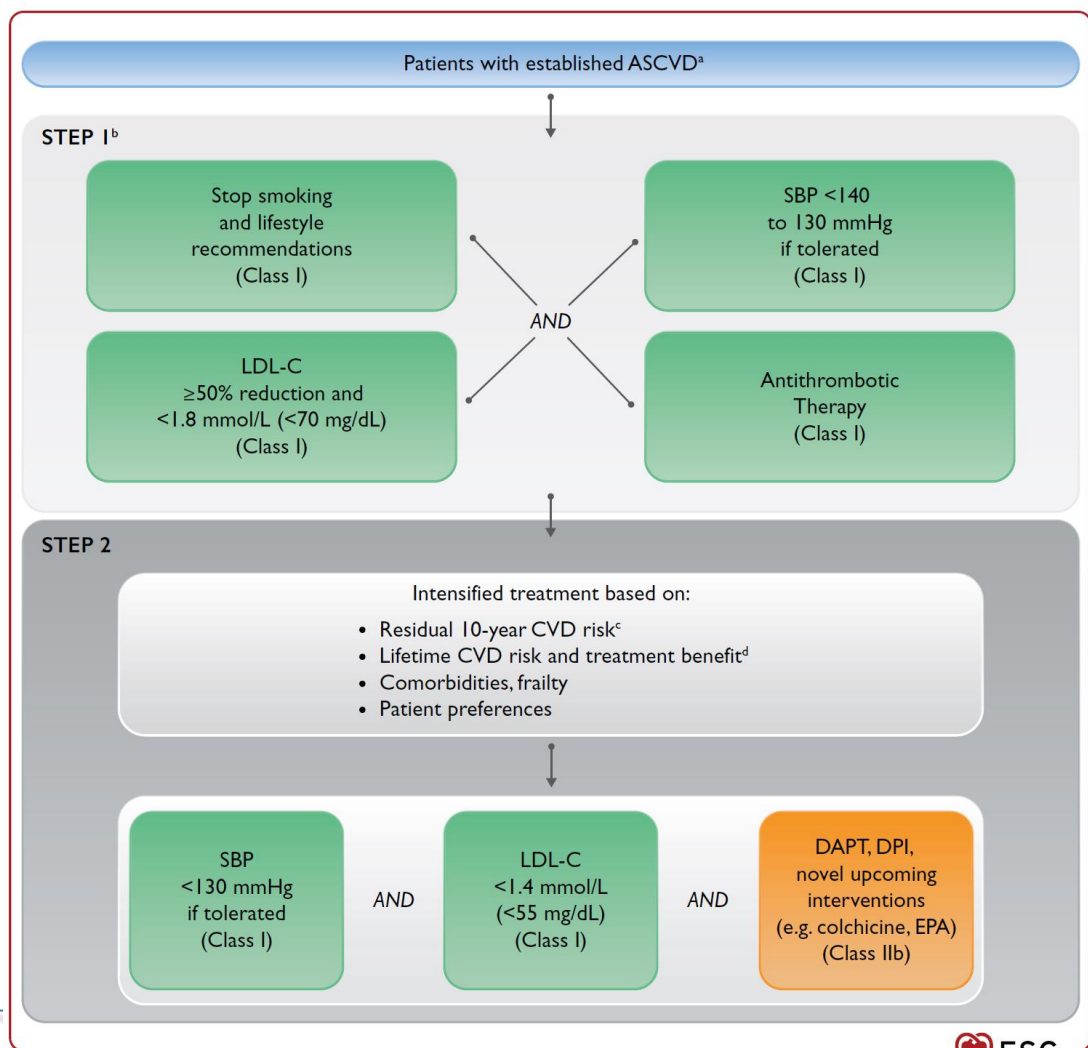
2020 AHA STATISTICAL UPDATE
Heart Disease & Stroke Statistics -
2020 Update
Circulation. 2020;141:e139-
e596.

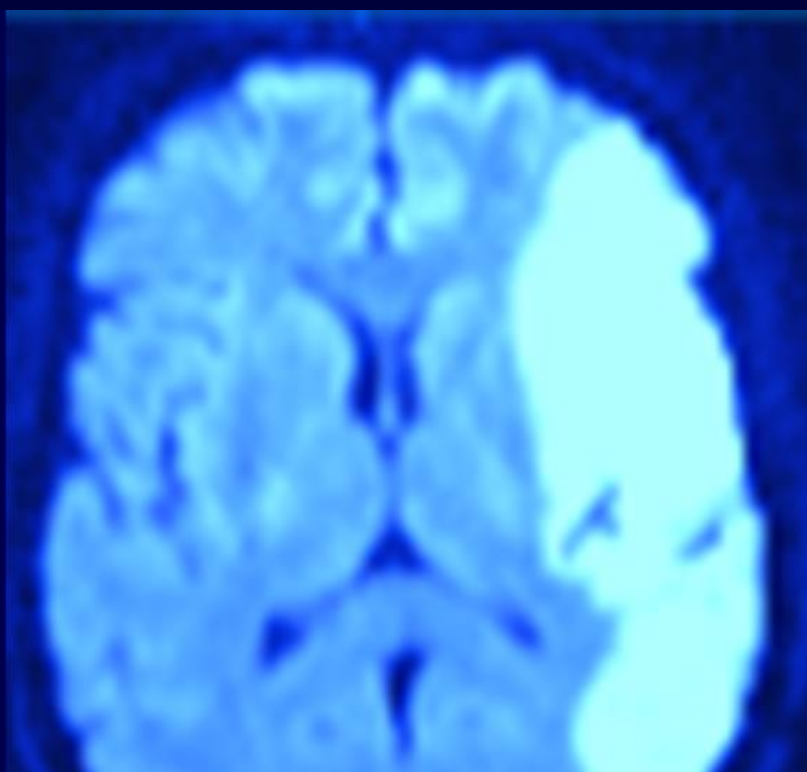
Guidelines | TIA management

2021 ESC Guidelines on cardiovascular disease prevention in clinical practice

Developed by the Task Force for cardiovascular disease prevention in clinical practice with representatives of the European Society of Cardiology and 12 medical societies

With the special contribution of the European Association of Preventive Cardiology (EAPC)





Conclusion

Conclusions

1. Déficit neurologique brusque faisant suspecter un AVC :

- Que faire ? → appeler le **112** / stabiliser le patient (*voies respiratoires*)
→ **Stroke Unit**
- Que **NE PAS** faire ? → **antiagrégant plaquettaire, héparine, anti-hypertenseur**

2. Déficit neurologique d'une durée de qlqs min = AIT :

- Bilan, quel délai ? → **le + court possible ! < 24 h**
→ Stroke Unit / TIA clinic

3. Déficit neurologique d'une durée de qlqs min = AIT ... **il y a 15 jours** :

- Bilan, quel délai ? → **le + court possible !** ... urgence relative ...
→ ? Antiagrégant plaquettaire ?

Merci pour votre attention

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Stroke & TIA @ home: do's & don'ts

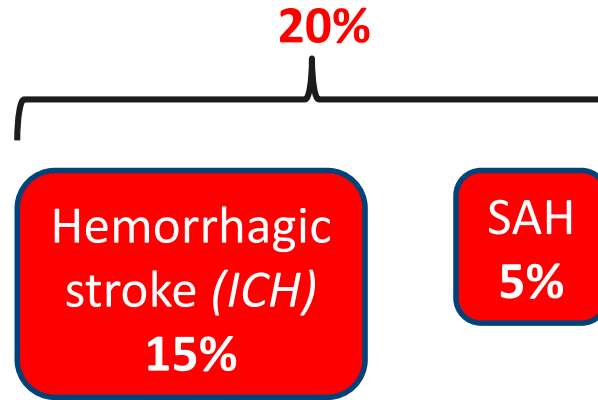
Yves VANDERMEEREN MD, PhD

Neurology Department | Stroke Unit

Cerebrovascular diseases: definitions

Stroke subtypes

Ischemic
stroke
80%



■ CVT
0,5-1%

- ICH: intracerebral hemorrhage
- SAH: subarachnoid hemorrhage
- CVT: cerebral venous thrombosis

Pre-hospital acute stroke management

ESO-EAN Practical guidance for pre-hospital management of stroke 2018

1. Educational campaigns to increase the awareness of immediately calling EMS for people with suspected stroke are recommended.
2. It is recommended that all EMS technicians & paramedics are familiar with a simple pre-hospital stroke scale to identify potential stroke patients.
3. There is insufficient evidence to recommend a pre-hospital stroke scale to predict large vessel occlusion.
4. In patients with **SaO₂ levels <95%** the administration of O₂ titrated to maintain normoxia is recommended.
Routine use of O₂ is not recommended.

Kobayashi et al. *Eur J Neurol.* 2018, 25: 425–433

ESO EUROPEAN
STROKE
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Pre-hospital acute stroke management

ESO-EAN Practical guidance for pre-hospital management of stroke 2018

5. **Pre-hospital treatment of high blood pressure** in people suspected of acute stroke is **not** recommended.
6. Because of safety concerns pre-hospital administration of **insulin** in persons with suspected stroke and hyperglycaemia is **not** recommended.
7. In the absence of clinical studies **no** recommendations can be made on pre-hospital interventions for lowering **elevated body temperature**.
8. It is recommended that all EMS implement a **'code stroke' protocol**, including highest priority dispatch, **pre-hospital notification** & **rapid transfer** to the closest 'stroke-ready' centre.

Pre-hospital acute stroke management

ESO-EAN Practical guidance for pre-hospital management of stroke 2018

9. **No** recommendation on the additional value of **pre-hospital telemedicine** can be made.
10. The routine use of **mobile emergency stroke units** is **not** recommended because there is insufficient evidence that they lead to a better functional outcome.
11. **No** recommendation can be made on the **pre-hospital use of POC laboratory** analysis of blood count & INR.
12. **No** recommendation can be made on the use of currently available **biomarkers** in persons with a suspected stroke.

Pre-hospital acute stroke management

Jusqu'à preuve du contraire, toute personne :

1. avec un déficit neurologique brutal subit un AVC

→ **Pré-notification : appeler la Salle d'Urgence**

- **NOM + PRENOM + date de naissance ?**
- **Médicaments antiagrégants/coagulants ?**
- **Heure de l'AVC ? ...H... ?**
(début des symptômes)
- **Vu bien pour la dernière fois à ...H... ?**
- **Retrouvé symptomatique à ...H... ?**
- **Arrivée estimée à ...H... ?**

TIA management

- In ~25% of stroke patients, a TIA has preceded the stroke
- Meta-analysis 14.889 patients, pooled stroke risk:
 - @ 2 days **1,36% (95% CI 1.15-1.59)**
 - @ 7 days **2,06% (95% CI 1.83-2.33)**
 - @ 30 days **2,78% (95% CI 2.47-3.12)**
 - @ 90 days **3,42% (95% CI 3.14-3.74)**

 - @ 1 year **12%**

Valls et al. *Cerebrovasc Dis.*
2017;43(1-2):90-98. doi:
10.1159/000452978.

TIA = an emergency !

→ quick management → Hospital: ER – Stroke Unit ?
→ ? TIA clinic ?

- Risk factors ?
- Etiology ?
- 2^{ary} prevention

TIA management

Cohorts > 10.000 TIA → stroke risk =

- @ 2 days **3%**
- @ 7 days **5%**

- @ 1 year **12%**

Risk of stroke early after transient ischaemic attack:
a systematic review and meta-analysis

Matthew F Giles, Peter M Rothwell

Lancet Neurol 2007; 6: 1063-72

TIA = an emergency !

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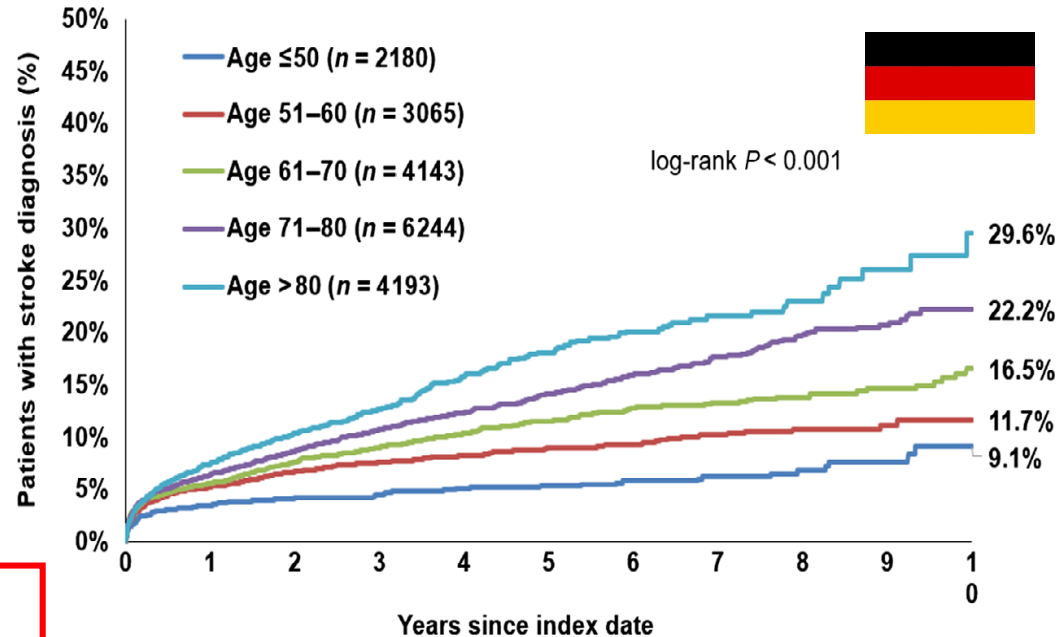
TIA management

- Cohort of 19.824 TIA patients
- **18,3%** diagnosed with stroke within 10 years after TIA index event
- ↔ Age, male sex, hypertension, diabetes mellitus, ischemic heart diseases

TIA = an emergency !

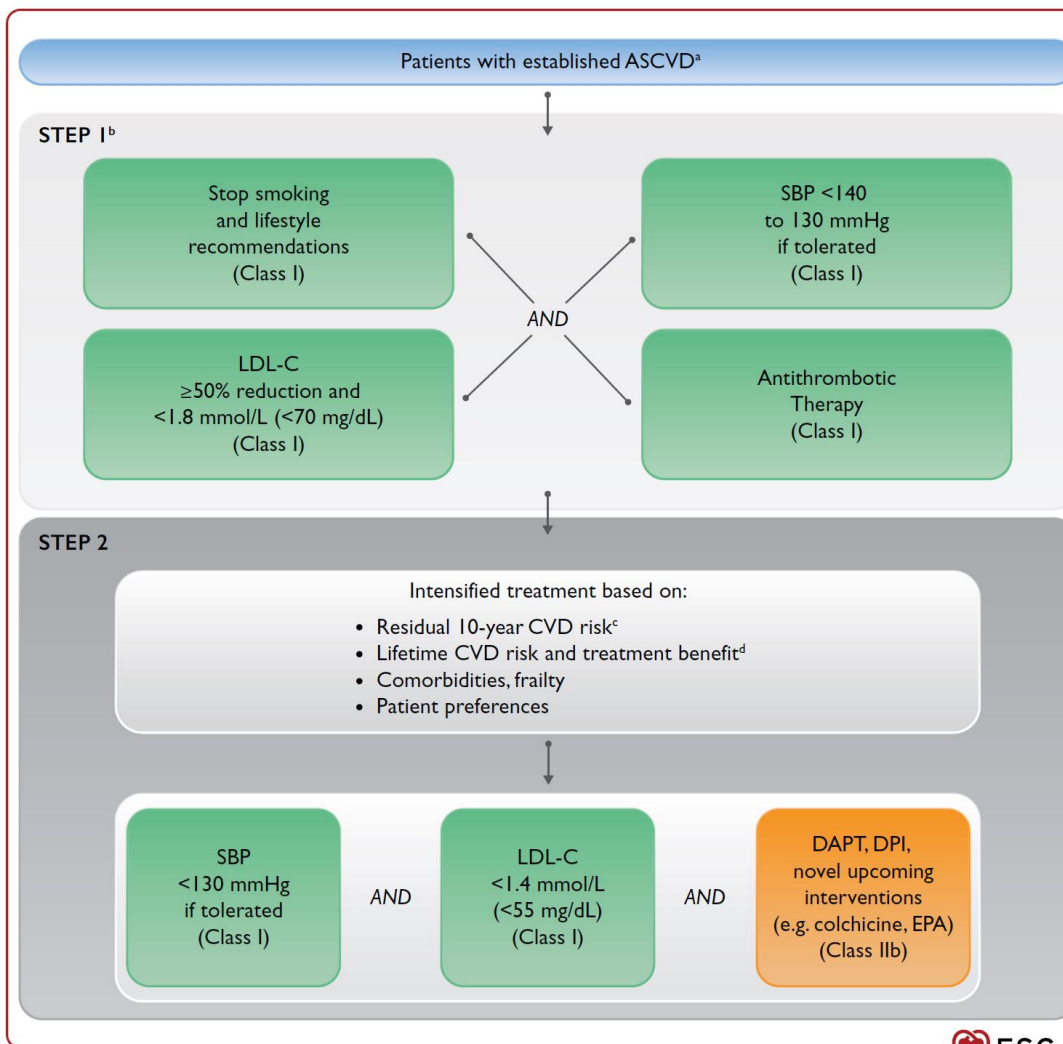
→ quick management → hospital ?

→ ? TIA clinic ?



Jacob et al., *European J of Neurology* 2020, 27: 723–728

Guidelines | TIA management



stroke / TIA : 2^{ary} prevention

RÉSUMÉ DES RECOMMANDATIONS 2019 DE L'ESC/EAS POUR LA PRISE EN CHARGE DES DYSLIPIDÉMIES

ÉVALUATION DES RISQUES	TRÈS HAUT RISQUE	HAUT RISQUE	RISQUE MODÉRÉ	RISQUE FAIBLE
Historique Cardio-vasculaire	MCVAS (clinique/imagerie)			-
Diabète	<ul style="list-style-type: none"> Atteinte d'organe (microalbuminurie, rétinopathie ou neuropathie) Avec ≥ 3 facteurs de risque majeurs ou DT1 depuis > 20 ans 	<ul style="list-style-type: none"> Sans atteinte d'organe Avec ≥ 1 facteur de risque ou Depuis ≥ 10 ans (DT1 ou DT2) 	Patients jeunes <ul style="list-style-type: none"> DT1 < 35 ans DT2 < 50 ans avec durée du diabète < 10 ans sans autres facteurs de risque 	-
Fonction rénale	eGFR < 30 mL/min/1,73m ²	eGFR 30-59 mL/min/1,73m ²	-	-
Facteur hérité	HF & MCVAS ou un autre facteur de risque majeur	HF sans autres facteurs de risque majeur	-	-
Facteurs de risque isolés	-	<ul style="list-style-type: none"> PA > 180/110 mmHg ou CT > 310 mg/dL ou LDL-C > 190 mg/dL 	-	-
SCORE <i>Risque CV fatal sur 10 ans</i>	≥ 10%	≥ 5% et < 10%	≥ 1% et < 5%	< 1%

1 ^{re} CIBLE	TRÈS HAUT RISQUE	HAUT RISQUE	RISQUE MODÉRÉ	RISQUE FAIBLE
LDL-C	< 40 mg/dL ÉVÈNEMENT RECURRENT**	< 55 mg/dL ET ≥ 50% réduction*	< 70 mg/dL ET ≥ 50% réduction*	< 100 mg/dL

2 ^{me} CIBLE	TRÈS HAUT RISQUE	HAUT RISQUE	RISQUE MODÉRÉ	RISQUE FAIBLE
Non-HDL-C	< 85 mg/dL	< 100 mg/dL	< 130 mg/dL	
ou ApoB	< 65 mg/dL	< 80 mg/dL	< 100 mg/dL	

Intervention

1. **Changement de mode de vie ET statine de haute Intensité**
2. **ÉZÉTIMIBE/fibrate (1TG)**
3. **Inhibiteur PCSK9**

Les taux de lipides doivent être réévalués 4 à 6 semaines après le SCA.

1. **Changement de mode de vie**
2. **Statine de haute Intensité**
3. **ÉZÉTIMIBE/fibrate (1TG)**

1. **Changement de mode de vie**
2. **Statine**

Conseils de style de vie

SABEZEN.20.01.0025

* par rapport à la valeur non-traitée. / ** MCVAS avec 2^{me} événement vasculaire < 2 ans. MCVAS: maladie cardiovasculaire athéroscléreuse.

statin


stroke / TIA : 2^{ary} preve ntion

U-Prevent+ CALCULATORS MANUAL ABOUT CONTACT NL EN


U-Prevent: You are in control

Watch a video or jump right in

Start calculator



▶ Clinician video



▶ Patient video

We provide tools for personalized Vascular Medicine. Get more insights by calculating individual cardiovascular risk and the effect of preventive treatment.

SCORE2 and SCORE2-OP are integrated in U-Prevent menu

Dear visitor,

We have updated the U-Prevent site with a new calculator overview that integrates the recently published, [SCORE2](#) and the [SCORE2-OP](#) calculators. Using the updated calculators have been advised by the European Society of Cardiology (ESC) on 30.08.2021. The new guidelines are available via the [European Society of Cardiology website](#) and in the [European Heart Journal](#). Please refer to your local medical guidelines when using these or other guidelines.

Select a calculator

I would like assistance with selecting a calculator

Patient group	10-years cardiovascular risk	Lifetime risk & treatment effect
Previous cardiovascular disease	SMART risk score	SMART-REACH model
Type 2 Diabetes Mellitus	ADVANCE risk score	DIAL model

www.u-prevent.com

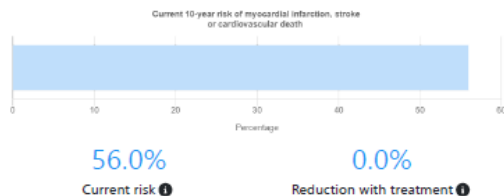
stroke / TIA : 2^{ary} prevention

U-Prevent*

SMART risk score

Personal Risk Profile ⁱ

Gender	Male	Years since first cardiovascular event	5 years	Systolic blood pressure	185 mmHg
Age	70 years	Type(s) of atherosclerotic vascular disease		Creatinin	80 umol/L
Current smoking	+	- Coronary artery disease	-	High Sensitivity CRP	5 mg/L
Antithrombotic treatment	-	- Cerebrovascular disease	+	Total cholesterol	8 mmol/L
		- Peripheral artery disease	-	HDL-cholesterol	2.5 mmol/L
		- Aortic Aneurysm	-	LDL-cholesterol	0.3 mmol/L
		Diabetes mellitus	+		



Future treatment ⁱ

Systolic blood pressure

No treatment target

LDL-cholesterol

No treatment target

Smoking



Antithrombotic treatment



stroke / TIA : 2^{ary} prevention

U-Prevent⁺

SMART risk score

Personal Risk Profile ⓘ

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		Diabetes mellitus	+		

U-Prevent⁺

SMART risk score

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56.0%

Current risk ⓘ

0.0%

Reduction with treatment ⓘ

Future treatment ⓘ

Systolic blood pressure

No treatment target

LDL-cholesterol

No treatment target

Smoking

Antithrombotic treatment



56.0%

Current risk ⓘ

45.7%

Reduction with treatment ⓘ

2

10-years NNT ⓘ

Future treatment ⓘ

Systolic blood pressure

< 120 mmHg

LDL-cholesterol

< 1.0 mmol/L / < 39 mg/dL

Smoking

Antithrombotic treatment