

The STEPCARE-trial criteria for a likely poor outcome

In the STEPCARE trial the prognosis is considered likely poor if criteria A, B and C stated below are all fulfilled.

A. Confounding factors such as severe metabolic derangement and lingering sedation have been ruled out.

B. FOUR-Score Motor: The patient has no response, a stereotypic extensor response or a stereotypic flexor response to bilateral central and peripheral painful stimulation at ≥ 72 hours after randomization (FOUR-M 0-2).

C. At least two of the below mentioned criteria of a poor prognosis are present:

C1. Bilateral absence of pupillary and corneal reflexes at 72h after randomization

C2. Bilaterally absent SSEP N20 potentials

C3. Early status myoclonus within 72h of randomization defined as a continuous and generalized myoclonus persisting for 30 minutes or more.

C4. A highly malignant EEG-pattern without reactivity to sound and painful stimulation on full-montage routine EEG or on simplified continuous EEG more than 24h after randomization and after effects of lingering sedation have been excluded:

i. Suppressed background (amplitude < 10 microV, $> 99\%$ of the recording) with or without superimposed discharges.

ii. Burst-suppression (periods of suppression with amplitude < 10 microV constituting at least 50% of the recording) with or without superimposed discharges.

C5. Neuroimaging: either a CT or MRI with signs of diffuse and extensive hypoxic ischemic injury

C6. Serum-NSE higher than 60 ng/mL at either 48h or 72h after randomization

Prognostication Checklist

Date of prognostication (YY/MM/DD): _____

Time of prognostication (24h clock): _____

Criterion A, confounding factors

Confounding factors such as severe metabolic derangement and lingering sedation have been ruled out

Yes No

When was the last given dose of a sedative agent prior to prognostication

_____ hours

Criterion B, FOUR-Score motor >72 hours after CA

Evaluate the best motoric response to a bilaterally given painful stimulus both centrally and peripherally in patients who are not awake and obeying commands

- 4. Makes sign (thumbs-up, fist or peace sign)
- 3. Localizes painful stimulus
- 2. Flexion response to pain
- 1. Extension response to pain
- 0. No response to pain or generalized status myoclonus

Criteria C (At least 2)

Poor outcome likely

C1. Corneal reflexes*	Bilaterally absent	<input type="checkbox"/>	Present	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
C1. Pupillary reflexes*	Bilaterally absent	<input type="checkbox"/>	Present	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
C2. SSEP N20	Bilaterally absent	<input type="checkbox"/>	Present	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
N20 amplitudes (if available)	Left _____microvolt		Right _____microvolt		Not assessed	<input type="checkbox"/>
C3. Early status myoclonus < 72h	Present	<input type="checkbox"/>	Absent	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
C4. Routine EEG highly malignant and unreactive**	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
C4. Continuous EEG highly malignant and unreactive**	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
C5. CT with diffuse and extensive hypoxic brain injury***	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
C5. MRI with diffuse and extensive hypoxic brain injury***	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
C6. High serial NSE	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>	Not assessed	<input type="checkbox"/>
NSE concentrations (ng/mL)	24h: _____		48h: _____		72h: _____	

*C1: Bilaterally absent pupillary and bilaterally absent corneal reflexes is considered one criterion

**C4: Highly malignant EEG patterns can be diagnosed either with routine or continuous EEG

***C5: Neuroimaging is considered one criterion regardless of whether CT and MRI are pathological

Does this patient fulfil the STEPCARE criteria for a likely poor neurological outcome?

Yes No