Patient population

Inclusion criteria

- Ischemic stroke Left middle cerebral artery (MCA) territory
- Non-fluent aphasia
- Enrollment in the study between 3 and 5 days post-stroke
- Right-handed
- Local language as primary language
- Provision of signed informed consent

Exclusion criteria

- History of symptomatic ischemic or hemorrhagic stroke
- Severe comprehension deficits that could compromise the understanding of informed consent or instructions, such as fluent aphasias (e.g. Wernicke aphasia) or global aphasias
- History of epilepsy or EEG-documented epileptic discharges
- Severe chronic renal or liver failure, indicated by aspartate aminotransferase, alanine aminotransferase levels greater than 4 times the normal values, or creatinine levels exceeding 4mg/dL
- Presence of life-threatening diseases
- Uncorrectable auditory or visual deficits that could impair testing
- Preexisting neurodegenerative or psychiatric disease



Combination treatments including Cerebrolysin improve post-stroke complications!

Treat stroke patients with Cerebrolysin as soon as possible

Disorder	Daily dosage	Initiation of treatment	Duration of treatment
Stroke	30 ml	as soon as possible no later than 5 days after stroke	30 days

Cerebrolysin is safe and well tolerated.



Speech therapy combined with Cerebrolysin enhances aphasia recovery after ischemic stroke: ESCAS¹ Randomized Pilot Study

Groups:

Titel:

66 Cerebrolysin group + 66 Placebo group

First cycle starts 3 - 5 days after stroke

¹ESCAS = Efficacy and Saftey of Cerebrolysin in the treatment Aphasia after AIS ²Western Aphasia Battery ³ Speech and Language Therapy

Copyright © 2025 by EVER Neuro Pharma GmbH, Oberburgau 3, 4866 Unterach, Austria. All rights reserved. No part of this brochure may be reproduced in any form or by any electronic or mechanical means, including information storage and retrieval systems, without permission in writing from the publisher. Cerebrolysin is a registered trademark of EVER Neuro Pharma GmbH, 4866 Unterach, Austria

ABBREVIATED PRESCRIBING INFORMATION - Cerebrolysin. Name of the medicinal product: Cerebrolysin - Solution for injection. Qualitative and quantitative composition: One ml contains 215.2 mg of Cerebrolysin concentrate in aqueous solution. List of excipients: Sodium hydroxide and water for injection. Therapeutic indications: For treatment of cerebrovascular disorders. Especially in the following indications: Senile dementia of Alzheimer's type. Vascular dementia. Stroke. Craniocerebral trauma (commotio and contusio). Contraindications: Hypersensitivity to one of the components of the drug, epilepsy, severe renal impairment. Marketing Authorization Holder: EVER Neuro Pharma GmbH, A-4866 Unterach Only available on prescription and in pharmacies. More information about pharmaceutical form, posology and method of administration, special warnings and precautions for use, interaction with other medicinal products and other forms of interaction, fertility, pregnancy and lactation, effects on ability to drive and use machines, undesirable effects, overdose, pharmacodynamics properties, pharmacokinetic properties, preclinical safety data, incompatibilities, shelf life, special precautions for storage, nature and contents of the container and special precautions for disposal is available in the summary of product characteristics. (Reference SPC-CCDS Version 2.0/03.06.2016)

EVER Neuro Pharma GmbH, Oberburgau 3, 4866 Unterach, Austria, www.everpharma.com, www.cerebrolysin.com







Treatment Scheme of ESCAS¹

132 stroke patients in total

Treatment: 1 cycle = 30ml Cerebrolysin + 1 hour SLT³ for 10 days 3 cycles in two-week intervals = d1-14, d29-42, d57-70

Speech therapy combined with **Cerebrolysin®** in enhancing non-fluent aphasia recovery after AIS

Results from the ESCAS' Randomized Pilot Study

Homberg V. et al, et al. Stroke 2025 A prospective, randomized-controlled, multicenter, double blinded-study



Cerebrolysin sets a new treatment standard in post-stroke aphasia

- Aphasia related deficits significantly improve
- Excellent mRS results
- More pronounced response of Cerebrolysin patients
- Better quality of life

CERE/INT/01/2025/2

Cerebrolysin[®]

Reconnecting Neurons Empowering for Life.

Aphasia problems after stroke



Primary objective

ESCAS¹ study is investigating the efficacy and safety of Cerebrolysin and speech therapy versus placebo and speech therapy at day 30, 60 and 90 after baseline.





Aphasia related deficits significantly improve



Excellent mRS results



More pronounced response of **Cerebrolysin patients**



- Patients in Cerebrolysin group shift **from moderate** (9) **to minor** stroke (3)
- Placebo treated patients remain at moderate stroke severity level (9;5)
- Cerebrolysin group improves by 6 points, placebo only 4 points
- According to NINDS criteria: Cerebrolysin treated patients respond to the treatment

Patient benefits

- + Better quality of life
- + Faster reintegration into social- and work-life
- + Less dependence on caregivers