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Cardiovascular Hospitalisation Costs and Cost-Effectiveness of Transthyretin Stabilizers in the Slovak Healthcare System

A Markov Model Analysis

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Agenda

01

Heart failure & ATTR-CM (Transthyretin Amyloid Cardiomyopathy)

The economic landscape

02

Clinical evidence

Tafamidis (ATTR-ACT) & acoramidis (ATTRIBUTE-CM)

03

Slovak Markov model

Methods, inputs, results

04

Discussion & policy implications

Value, caveats, the registry call

Heart failure: a growing global economic burden

Inpatient care dominates direct costs

>64 M

people with heart failure
worldwide

ESC HF Review 2025

75–80%

of HF direct costs are inpatient

HFSA economic burden paper

0.5–1.3

HF admissions per patient per
year in Europe

ESC HF Review 2025

***European hospital budgets absorb the bulk of HF cost.
Rare cardiomyopathies with high readmission intensity — such as ATTR-CM
— amplify this pattern.***

ATTR-CM: a progressive, life-limiting cardiomyopathy

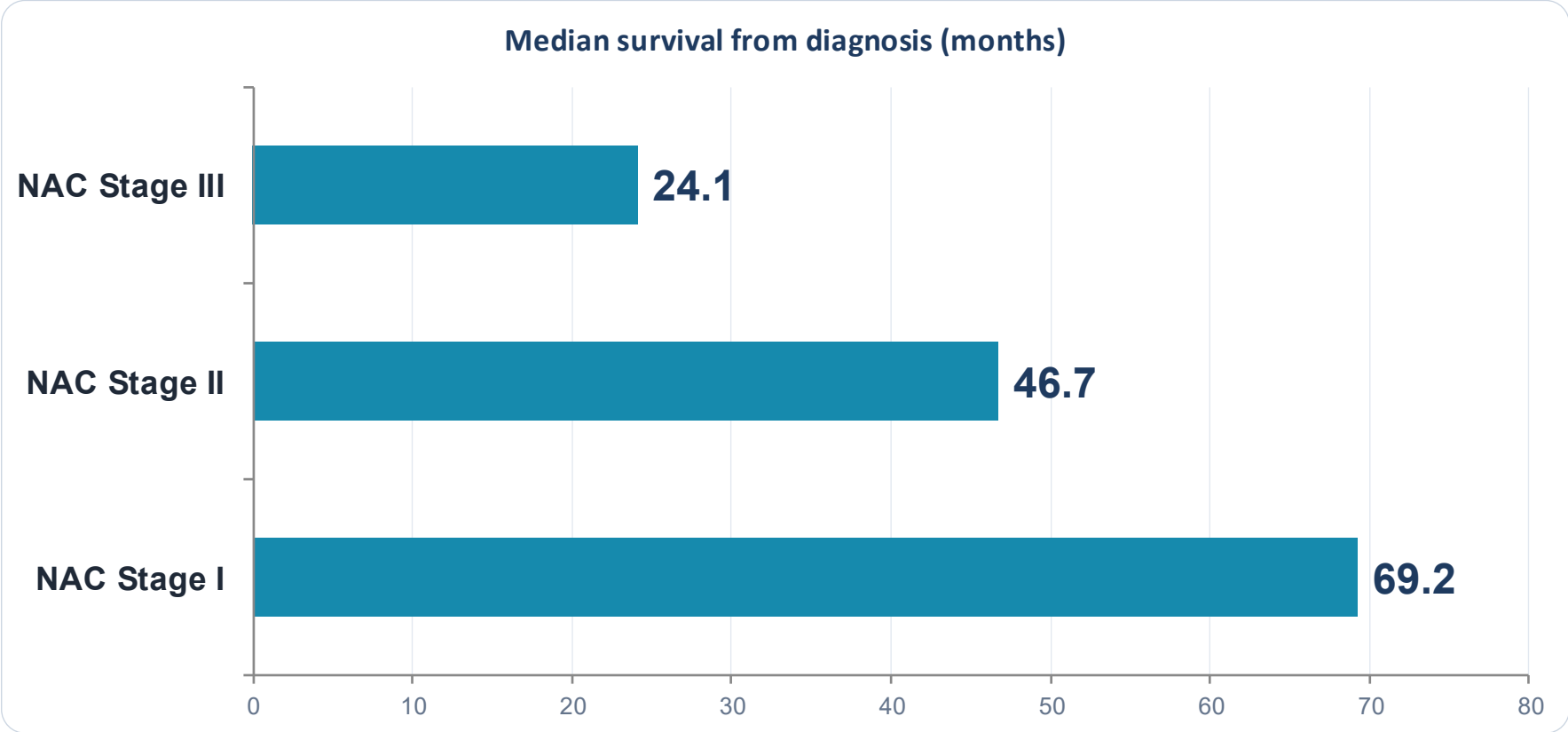
Stage-dependent survival — hospitalisations dominate clinical course

Clinical signature

Heart-failure symptoms, recurrent CV hospitalisations, premature death. NAC staging (Gillmore et al. 2018) stratifies prognosis from diagnosis.

Genotype matters

Median survival 69 mo (non-V122I hereditary), 57 mo (wild-type), 31 mo (V122I) — Lane et al. 2019.



Refs: Gillmore JD, et al. NAC staging, Eur Heart J 2018;39:2799–806. · Lane T, et al. Circulation 2019;140:16–26.

Why ATTR-CM is an economic outlier

Hospitalisations are the dominant direct-cost driver

87 – 93 %

of direct medical costs in ATTR-CM are generated by cardiovascular-related hospitalisations

Delgado D et al. | Gonzalez-Lopez et al.

Compared with general heart failure ($\approx 75\text{--}80\%$ inpatient share), ATTR-CM concentrates an even larger slice of spend into hospital walls.

Consequences for the payer

- Any intervention that reduces admissions has an outsized economic footprint.
- Cost savings are event-driven, not time-driven — they accrue with each avoided admission.
- DRG unit cost becomes the single most influential cost parameter.

International evidence on ATTR-CM resource use

Higher intensity vs. matched HF cohorts across health systems

NORDIC 4-COUNTRY

Denmark · Finland · Norway · Sweden (2008–2018, n=1,831)

Hospitalisations / yr

1.9 vs 1.4

ATTR-CM vs matched HF

Hospitalised days / yr

9.7 vs 7.2

Specialist visits / yr

8.7 vs 4.7

SWEDEN (NYHA)

Cost by severity, healthcare resource use study

Mean cost / patient / yr

€21,200

across NYHA II–IV

Diagnostic delay

3.5 years

prior to ATTR-CM dx

Principal driver

Inpatient

vs other HF types

USA MEDICARE

HF admissions with vs without ATTR-CM

1-yr total cost / pt

\$60,373 vs \$50,247

HF + ATTR-CM vs HF alone

HF readmission (1 yr)

36.8% vs 28.2%

adj HR 1.28 (1.01–1.61)

Outpatient spend

\$20,866 vs \$16,358

primary cost driver

The diagnostic gap is an economic inflection point

Delayed / missed diagnosis drives avoidable admissions

3.5 years

mean diagnostic delay before ATTR-CM diagnosis (Sweden)

17 visits

median hospital contacts in the 3 years before diagnosis (English analysis, n=524)

SPAIN — PROBABILISTIC MARKOV MODEL

Correct vs. missed ATTR-CM diagnosis

Correct diagnosis generates life-years gained and reduces CV hospitalisations for the Spanish NHS.

≈ €2,900

per patient saved at 15-year horizon vs. missed diagnosis

Diagnostic pathways — not just therapy — are a legitimate target for health-economic intervention.

Research question and objectives

“Can the cardiovascular-hospitalisation benefit observed in ATTR-ACT and ATTRIBUTE-CM translate into measurable economic value within the Slovak public-payer setting?”

01

Derive Slovakia-specific per-patient hospitalisation cost offsets for both stabilizers

02

Compute ICERs (€/QALY) from a hospitalisation-only perspective

03

Extend projections to a 6.5-year horizon anchored in NAC staging evidence

04

Identify data gaps that a national ATTR-CM registry should close

Therapeutic breakthrough: two transthyretin stabilizers

Reduced CV morbidity and mortality in pivotal phase-III trials

ATTR-ACT (tafamidis)

CV hosp. rate (events / pt-yr)

0.48 vs 0.70

30-month all-cause mortality

29.5% vs 42.9%

Endpoint

All-cause mortality + CV hospitalisations over 12 mo

Maurer MS, et al. NEJM 2018;379:1007–16.

ATTRibute-CM (acoramidis)

CV hosp. rate (events / pt-yr)

0.22 vs 0.45

30-month all-cause mortality

19.3% vs 25.7%

Endpoint

Hierarchical composite; event curves separated by month 3

Gillmore JD, et al. NEJM 2024;390:132–42. · Judge DP, et al. JACC 2025.

Both trials demonstrate that stabilising TTR translates into fewer CV hospitalisations — the key lever we model for Slovakia.

Slovak health-system context

How HTA and reimbursement frame this analysis

SK-DRG

SK-DRG reimbursement

Diagnosis-Related Group framework anchors unit cost per CV/HF admission in Slovakia.

NIHO

National Institute for Value and Technologies in Healthcare (NIHO)

Institutional framework for HTA appraisal and value-based reimbursement.

ZHL175

Public HTA appraisal — Beyontra (acoramidis), ZHL175

Published dossier provides the reference unit-cost parameter used in this model.

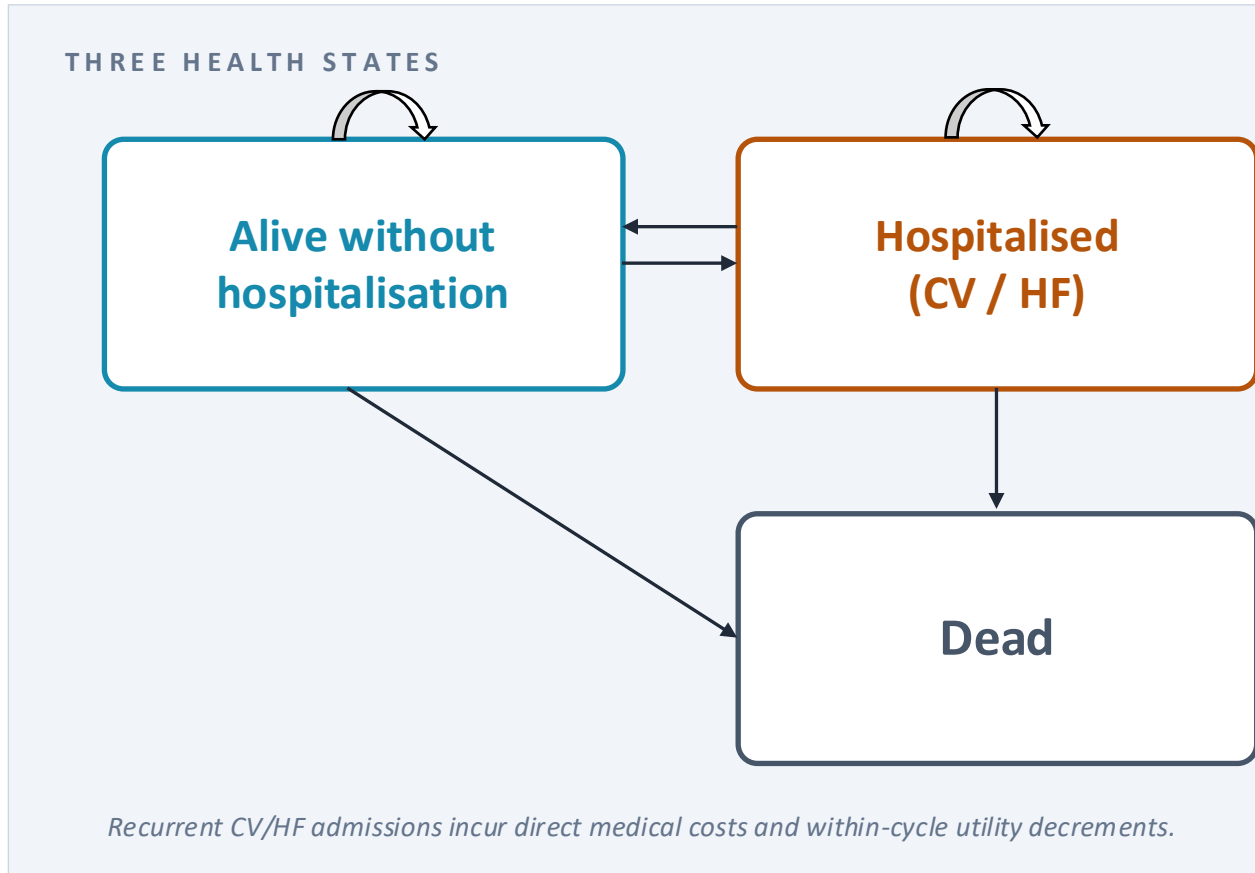
Gap

No national ATTR-CM registry

Absence of Slovak-specific real-world data constrains external validation of any economic model.

Methods — Markov model structure

Cohort-based state-transition model; monthly cycles; half-cycle correction



PERSPECTIVE

Slovak public-payer and hospital · Direct CV / HF hospitalisation costs only.

TIME HORIZONS

12 months (base case) · 6.5 years (scenario — NAC-staging mean survival).

RATE ↔ PROBABILITY

Constant-hazard embedding (ISPOR-SMDM good-practice).

$$p = 1 - e^{-\lambda \Delta t}$$
$$\lambda = -\ln(1 - P) / T$$

VALIDATION

Matrix back-transforms reproduce trial event rates exactly (placebo 0.44 → model 0.443; acoramidis 0.25 → 0.252).

Methods — Input parameters

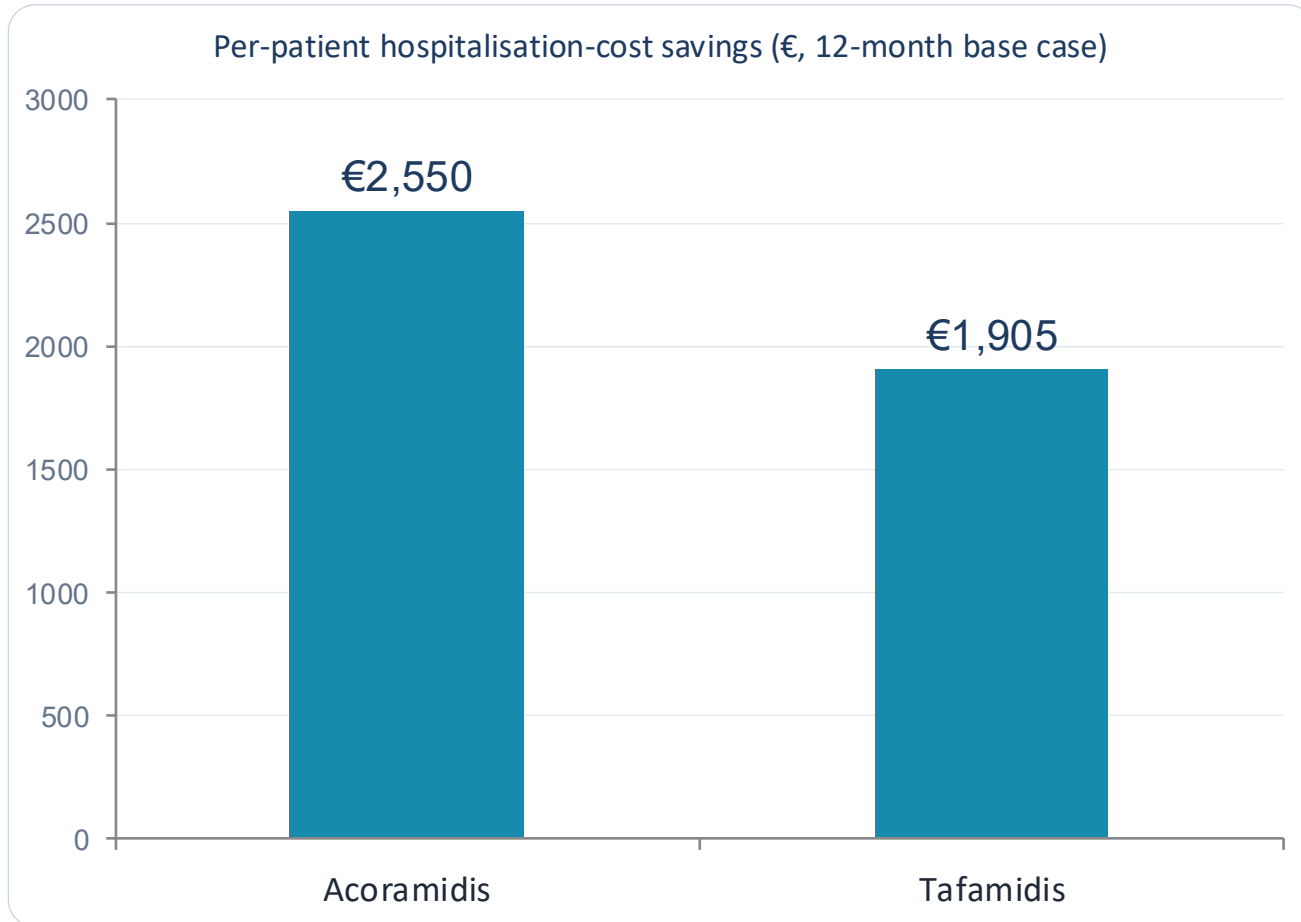
Clinical, economic and utility values (Table 1, Lekársky Obzor 2026)

Parameter	Value	Cycle / unit	Source
P(hospitalised / yr) — standard (ATTRibute-CM arm)	0.443	<i>year</i>	<i>ATTRibute-CM (8)</i>
P(hospitalised / yr) — acoramidis	0.252	<i>year</i>	<i>ATTRibute-CM (8)</i>
P(hospitalised / yr) — standard (ATTR-ACT arm)	0.503	<i>year</i>	<i>ATTR-ACT (7)</i>
P(hospitalised / yr) — tafamidis	0.381	<i>year</i>	<i>ATTR-ACT (7)</i>
P(discharge alive hospitalised)	0.876	<i>year</i>	<i>Trial-derived</i>
Utility — alive, no hospitalisation	0.80	<i>month</i>	<i>Eldhagen 2023 (15)</i>
Utility — hospitalised	0.56	<i>month</i>	<i>Eldhagen 2023 (15)</i>
Unit cost — CV / HF admission (Slovakia)	€3,462.94	<i>per admission</i>	<i>NIHO, ZHL175</i>
Mean survival — extended scenario	6.5 yr	—	<i>NAC staging (1)</i>
Discount rate — costs & effects	5%	<i>per year</i>	<i>MoH SR 422/2011</i>

Base-case event rates are identical to those reported in the pivotal trials (ATTRibute-CM: 0.22 vs 0.45 events/pt-yr; ATTR-ACT: 0.48 vs 0.70 events/pt-yr).

Results — Base case (12 months)

Per-patient hospitalisation-cost savings vs. standard care



ACORAMIDIS

– €2,550

saved per patient in hospitalisation costs

TAFAMIDIS

– €1,905

saved per patient in hospitalisation costs

Results — Extended 6.5-year horizon and cost-effectiveness

Dominant acoramidis profile; highly favourable tafamidis ICER

	Treatment	Standard	Δ (Treatment – Std)	ICER (€ / QALY)
ACORAMIDIS	€9,719	€21,916	– €12,196	– €15,720.22
	4.82 QALY	4.04 QALY	+ 0.78 QALY	
TAFAMIDIS	€16,371	€15,933	+ €437	€450.23
	3.45 QALY	2.48 QALY	+ 0.97 QALY	
	<i>Highly favourable — €/QALY well below Slovak willingness-to-pay thresholds</i>			

***Both agents fall south-east of the origin: QALY-positive.
Acoramidis is also cost-negative (cost-saving, dominant).***

Conclusions — population impact and caveats

POPULATION-LEVEL IMPLICATIONS

≈ **€1.2 million**

cumulative hospitalisation-cost savings projected for a 100-patient acoramidis cohort over 6.5 years.

+ 0.78 QALY / patient (acoramidis)

≈ 9.4 months of perfect health equivalent, ≈ 19.3% improvement in quality-adjusted life expectancy over 6.5 years vs. standard therapy.

IMPORTANT CAVEATS

Hospitalisation-only perspective

Drug acquisition costs, outpatient care, and rehabilitation are outside this scope. A full cost-utility analysis requires acquisition pricing.

Cross-trial heterogeneity

ATTR-ACT and ATTRibute-CM differ in populations, endpoints, and baseline event rates. Any indirect comparison between acoramidis and tafamidis is hypothetical; no head-to-head data exist.

No individual patient data

Matching-adjusted indirect comparison (MAIC) was not possible; residual confounding remains.

No Slovak ATTR-CM registry

Trial-based projections are not validated against Slovak real-world data; DRG unit cost may underestimate complex admissions.

Conclusions - policy implications

Reductions in CV / HF hospitalisations observed with TTR stabilizers translate into tangible Slovak hospital-budget savings.

Acoramidis: dominant profile (saves cost + gains QALYs) under the hospitalisation-only perspective.

Tafamidis: highly favourable ICER (€450.23 / QALY) on the same perspective.

FOUR POLICY ACTIONS FOR SLOVAKIA AND CEE

1 Establish a national ATTR-CM registry

Admissions, length of stay, readmissions, longitudinal HRQoL — validated instruments.

2 Expand specialised ATTR-CM diagnostic & treatment centers

Build a national network of reference centers with cardiac amyloidosis expertise; reduce geographic barriers to specialist access.

3 Improve clinical pathways & primary-care awareness

Deploy red-flag algorithms and referral protocols; raise GP and cardiologist awareness to cut the 3.5-year diagnostic delay — timely diagnosis can extend life expectancy by up to 8 years.

4 Benchmark within European ATTR-CM networks

Collaborate for comparability while preserving Slovak cost structures.

Thank you

Questions & discussion

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Full article: Babela R, Hlavata T, Polak P. Hospitalization-driven economic effects of new treatments for transthyretin amyloid cardiomyopathy in Slovakia. Lekársky Obzor 2026; 75(2): 50–57. · www.lekarskyobzor.sk