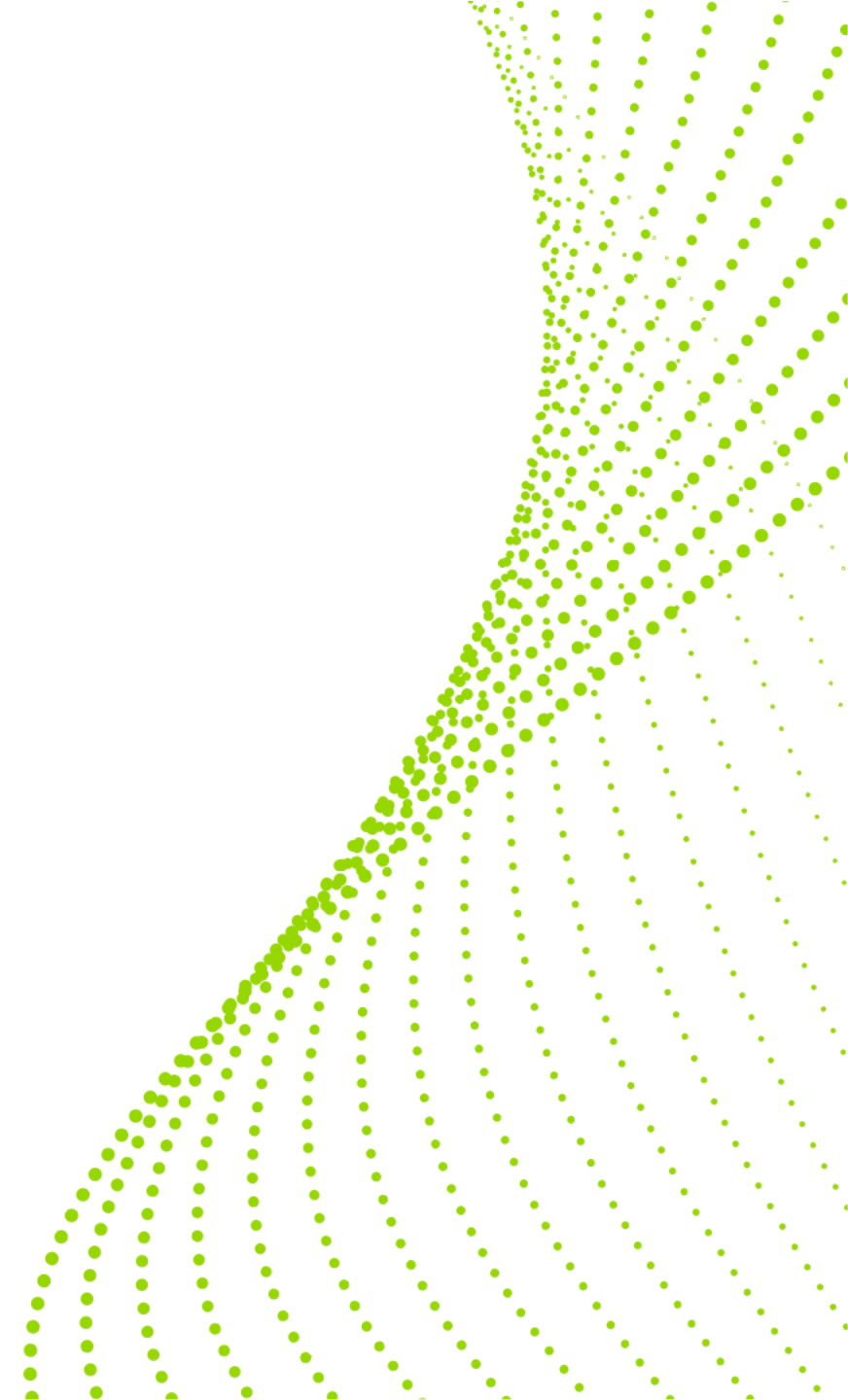




Cytokinetics®

ACACIA-HCM Topline Results

May 5, 2026



ACACIA-HCM: Phase 3 Trial of *Aficamten* in nHCM



Full results to be presented at upcoming medical meeting

Primary Endpoints	Change from Baseline to Week 36 LSM (95% CI)		<i>Aficamten</i> vs Placebo LSM (95% CI)	p-value
	<i>Aficamten</i>	Placebo		
KCCQ-CSS	11.4 (9.6 – 13.2)	8.4 (6.6 – 10.2)	3.0 (0.5 - 5.5)	0.021
pVO₂ (ml/kg/min)	0.64 (0.32 – 0.95)	-0.03 (-0.35 – 0.28)	0.67 (0.22 - 1.1)	0.003

No new safety signals identified

% participants completing planned dosing:
Similar between groups
(88.4% on *aficamten* vs. 90.3% on placebo)

LVEF <50%:
27 (10%) on *aficamten* vs. 2 (1%) on placebo;
2 participants on *aficamten* had a serious adverse event of HF associated with LVEF <50%

Treatment Interruptions
Treatment interruptions due to LVEF <40% occurred in 3% of patients on *aficamten*

Statistically significant (p<0.001) improvements observed in key secondary endpoints:

1. Proportion of participants with improvements in NYHA Functional Class
2. Composite z-score of ventilatory efficiency and pVO₂
3. N-terminal pro-B-type natriuretic peptide (NT-proBNP)

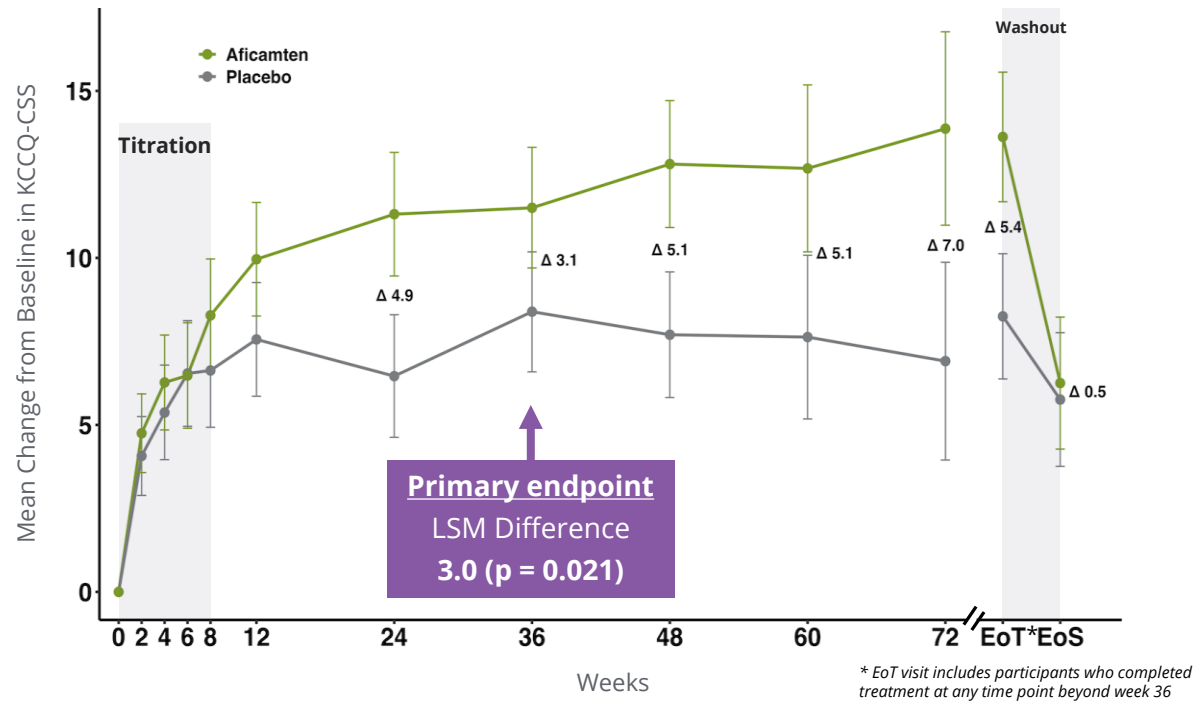
nHCM: non-obstructive hypertrophic cardiomyopathy; pVO₂: peak oxygen uptake; LSM: least squares mean; CI: confidence interval; KCCQ-CSS: Kansas City Cardiomyopathy Questionnaire Clinical Summary Score; NYHA: New York Heart Association; CV: cardiovascular; HF: heart failure; LVEF: left ventricular ejection fraction; AE: adverse event
MYQORZO is only approved in the U.S., China and the EU for the treatment of adults with symptomatic oHCM.

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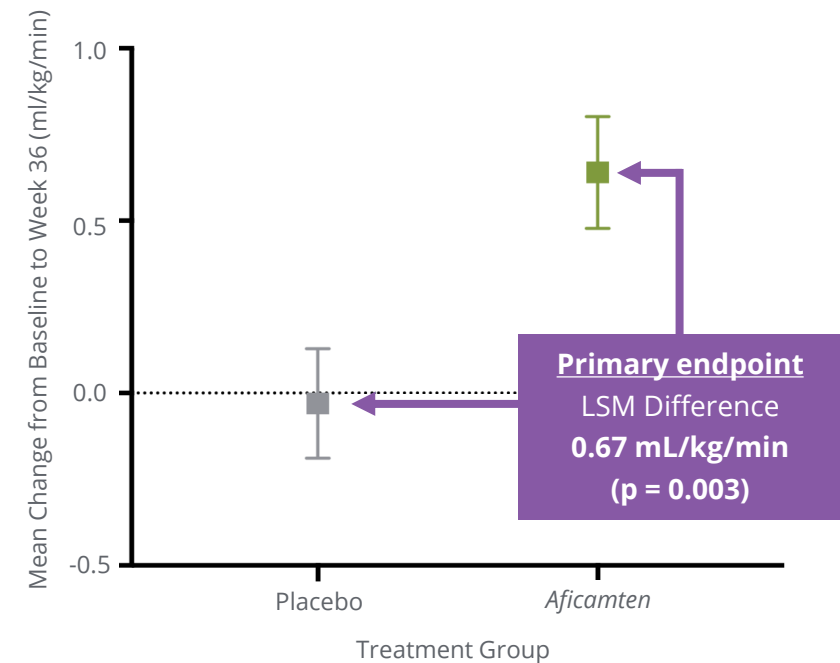


Robust & consistent improvements throughout treatment period

LS Mean Change from Baseline in KCCQ-CSS



Change from Baseline to Week 36 in pVO₂



nHCM: non-obstructive hypertrophic cardiomyopathy; LS: least squares; KCCQ-CSS: Kansas City Cardiomyopathy Questionnaire Clinical Summary Score; LSM: least squares mean; EOT: End of Treatment; EOS: End of Study; pVO₂: peak oxygen uptake
 MYQORZO is only approved in the U.S., China and the EU for the treatment of adults with symptomatic oHCM.