Persistent growth-promoting effects of vosoritide in children with achondroplasia is accompanied by improvement in physical aspects of quality of life

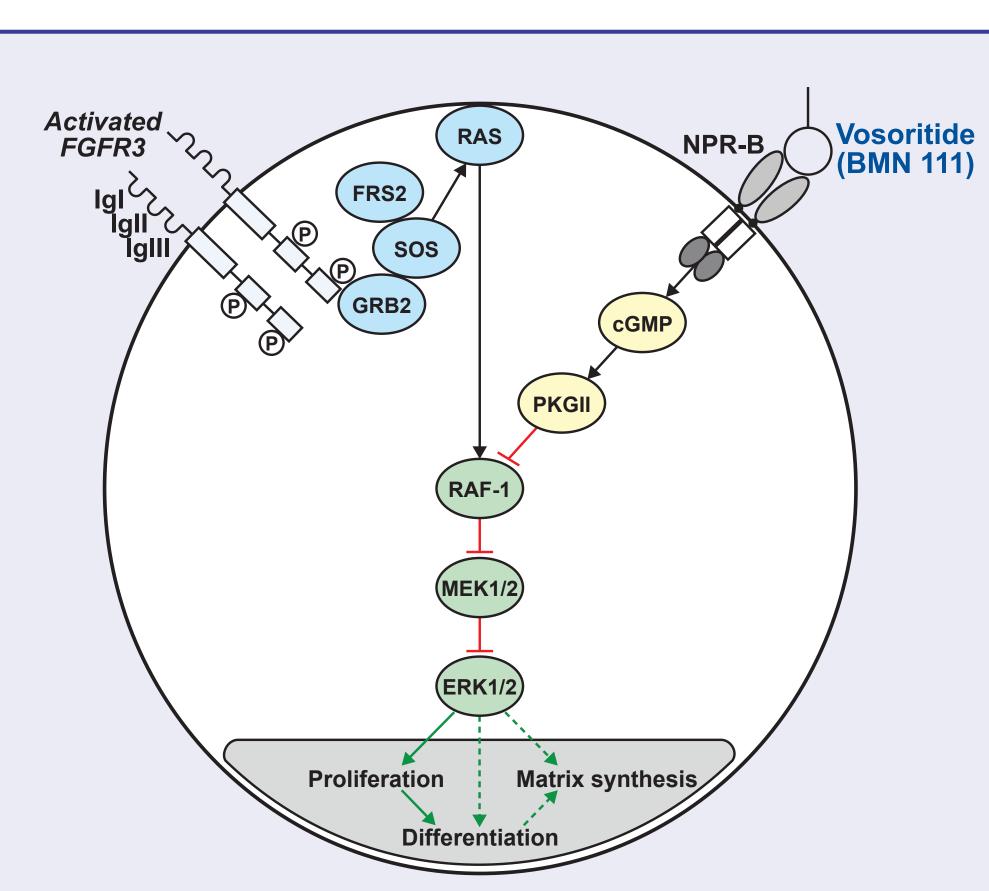
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Background and Objectives

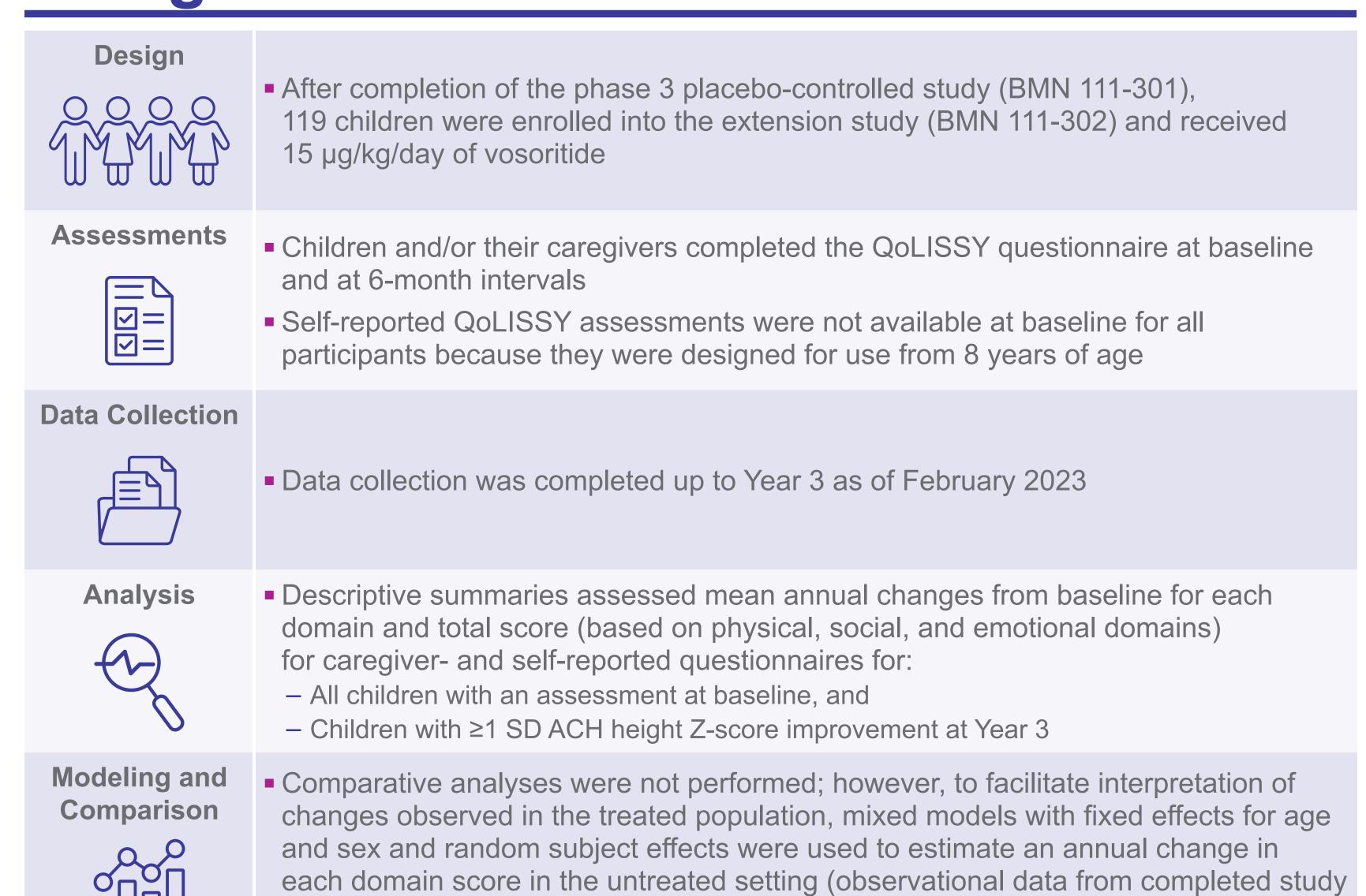
Vosoritide: Targeted therapy for achondroplasia

- Achondroplasia (ACH) is the most common form of disproportionate short stature (1:25,000 live births)^{1,2} and is associated with a high burden of medical complications²⁻⁵ and a reduced quality of life⁶
- ACH is caused by a pathogenic variant in FGFR3 that constitutively activates the downstream inhibitory signaling pathway in chondrocytes, leading to impaired endochondral bone growth and multiple complications^{1,2}



- CNP down-regulates aberrant FGFR3 signaling in chondrocytes by inhibiting the MAPK-ERK pathway^{7,8}
- Vosoritide is based on naturally occurring CNP engineered to resist degradation and increase the half-life⁹
- Vosoritide, a CNP analog, has been shown to increase growth in children with ACH
- An open-label, 52-week phase 2 trial (BMN 111-202) and its extension study (BMN 111-205) in children with ACH aged ≥5 years showed that vosoritide treatment resulted in sustained increases in annualized growth velocity (AGV) for over 7 years^{10,11}
- A phase 3 randomized placebo-controlled trial (BMN 111-301) in children with ACH aged ≥5 years showed a statistically significant improvement in AGV with vosoritide after 52 weeks compared to placebo¹²; AGV improvement was sustained after 3 years of vosoritide treatment in extension study BMN 111-302¹³,¹⁴
- In children with ACH aged 0-5 years, improvement in height Z-score was seen with vosoritide compared to placebo after 52 weeks (111-206)¹⁵
- Vosoritide is approved for use in children with ACH and open epiphyses from birth in the USA, Japan, and Australia, and aged ≥4 months in the EU and ≥6 months in Brazil
- Our objective was to evaluate the impact of vosoritide on health-related quality of life (HRQoL) in children with ACH using Quality of Life in Short Stature Youth (QoLISSY) questionnaires¹⁶

Design and Methods



BMN 111-901 and placebo data from study BMN 111-301)

Untreated Population Treated Population

Results

Participant characteristics

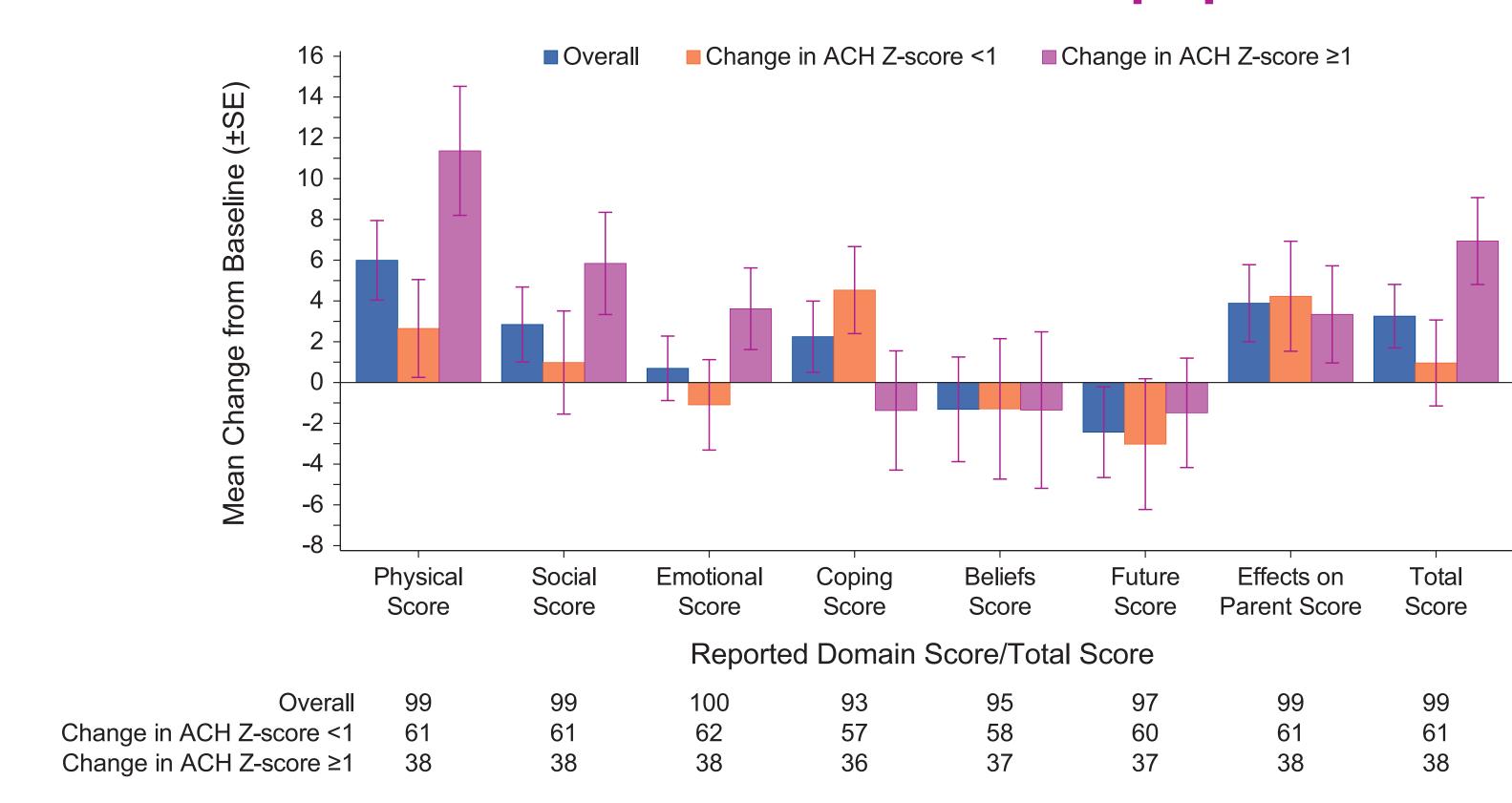
Higher score indicates better QoL —

	(N=150)	(N=119)
Age at first assessment (years)		
Mean (SD)	8.0 (2.4)	9.7 (2.6)
Median (Min, Max)	7.7 (5.0, 13.9)	9.7 (5.6, 16.4)
Age subgroup, n (%)		
≥5 to <8 years	79 (52.7)	40 (33.6)
≥8 to <11 years	51 (34.0)	39 (32.8)
≥11 to <15 years	20 (13.3)	37 (31.1)
≥15 to <18 years	_	3 (2.5)
Sex, n (%)		
Female	72 (48.0)	56 (47.1)
Race, n (%)		
White	118 (78.7)	85 (71.4)
Asian	18 (12.0)	21 (17.6)
Black or African American	7 (4.7)	5 (4.2)
Location, n (%)		
United States	60 (40.0)	53 (44.5)
Spain	34 (22.7)	12 (10.1)
Australia	19 (12.7)	22 (18.5)
United Kingdom	16 (10.7)	13 (10.9)
Germany	11 (7.3)	9 (7.6)
Japan	6 (4.0)	6 (5.0)
Turkey	4 (2.7)	4 (3.4)
Number of assessments		
Median	4	
25th, 75th percentile	3, 5	
Duration between first and last assessments (months)		
Median	13.2	
25th, 75th percentile	9.2, 21.6	
Mean (SD 9.7 (2.6) 5.6 Age (years) 0 5 10	116.4 15 20	N=119
Mean (SD) Baseline (SD) Baseline (SD) Baseline (SD) Baseline (SD) (ST) (ST) (ST) (ST) (ST) (ST) (ST) (ST	■ Caregiver-reported (n=119)	52.9%
64.5 (18.4)		
$= \ddot{\sigma} \ 0 \qquad 20 \qquad 40 \qquad 60 \qquad 80$	100	

Mixed-effect analysis of QoLISSY parameters

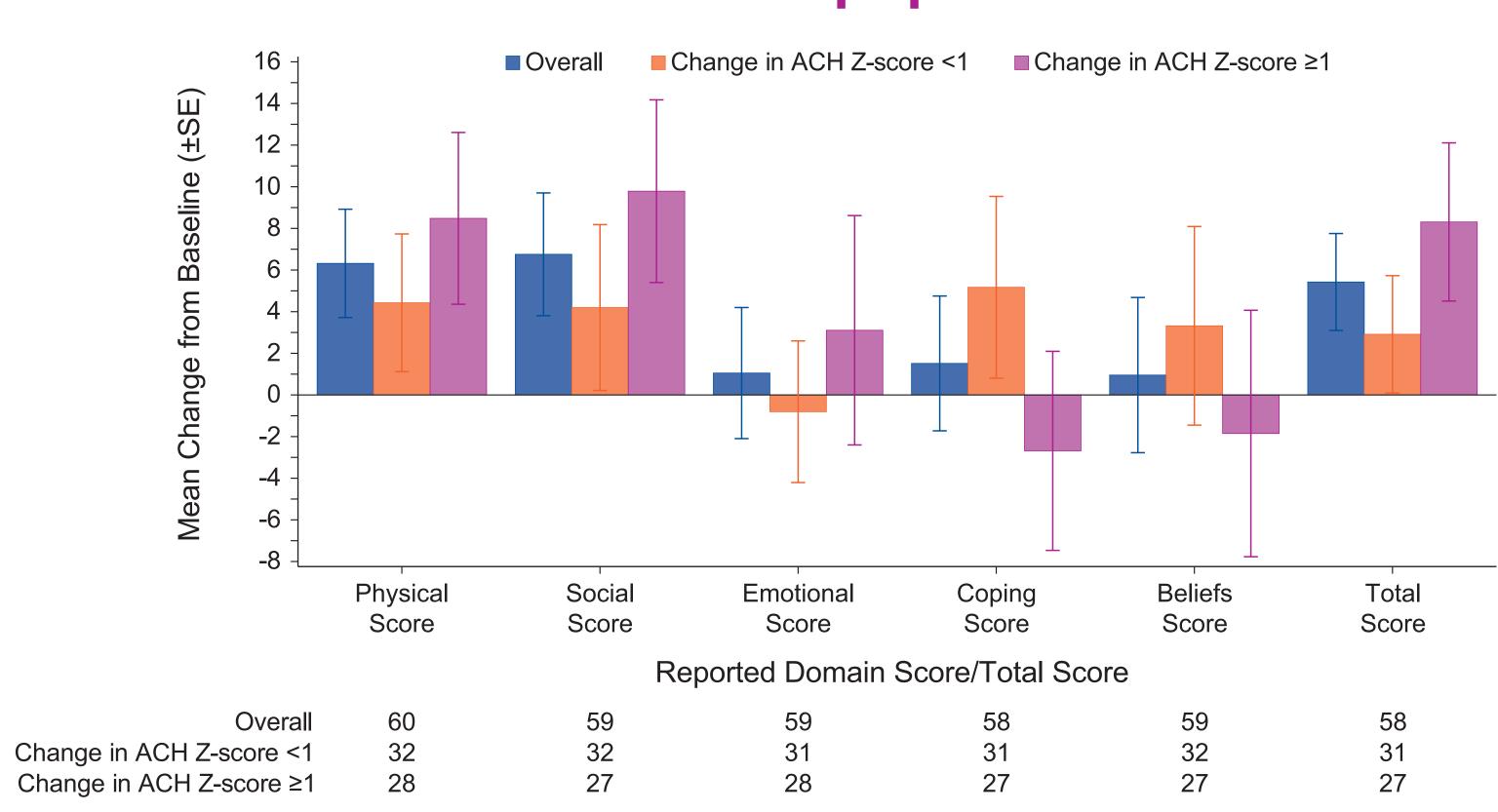
Reported Domain Score/Total Score	Estimated Annual Slope (SE) in the Untreated Population	
Caregiver-reported		
Physical Score	0.16 (0.55)	
Social Score	0.16 (0.50)	
Emotional Score	-1.40 (0.57)	
Coping Score	1.41 (0.48)	
Beliefs Score	-0.70 (0.66)	
Future Score	-1.45 (0.63)	
Effects on Parent Score	1.53 (0.50)	
Total score	-0.27 (0.48)	
Self-reported		
Physical Score	1.45 (0.77)	
Social Score	1.92 (0.77)	
Emotional Score	1.19 (0.70)	
Coping Score	-0.75 (0.93)	
Beliefs Score	1.94 (1.09)	
Total score	1.63 (0.63)	

Mean change from baseline in caregiver-reported QoLISSY scores at Year 3 in the treated population



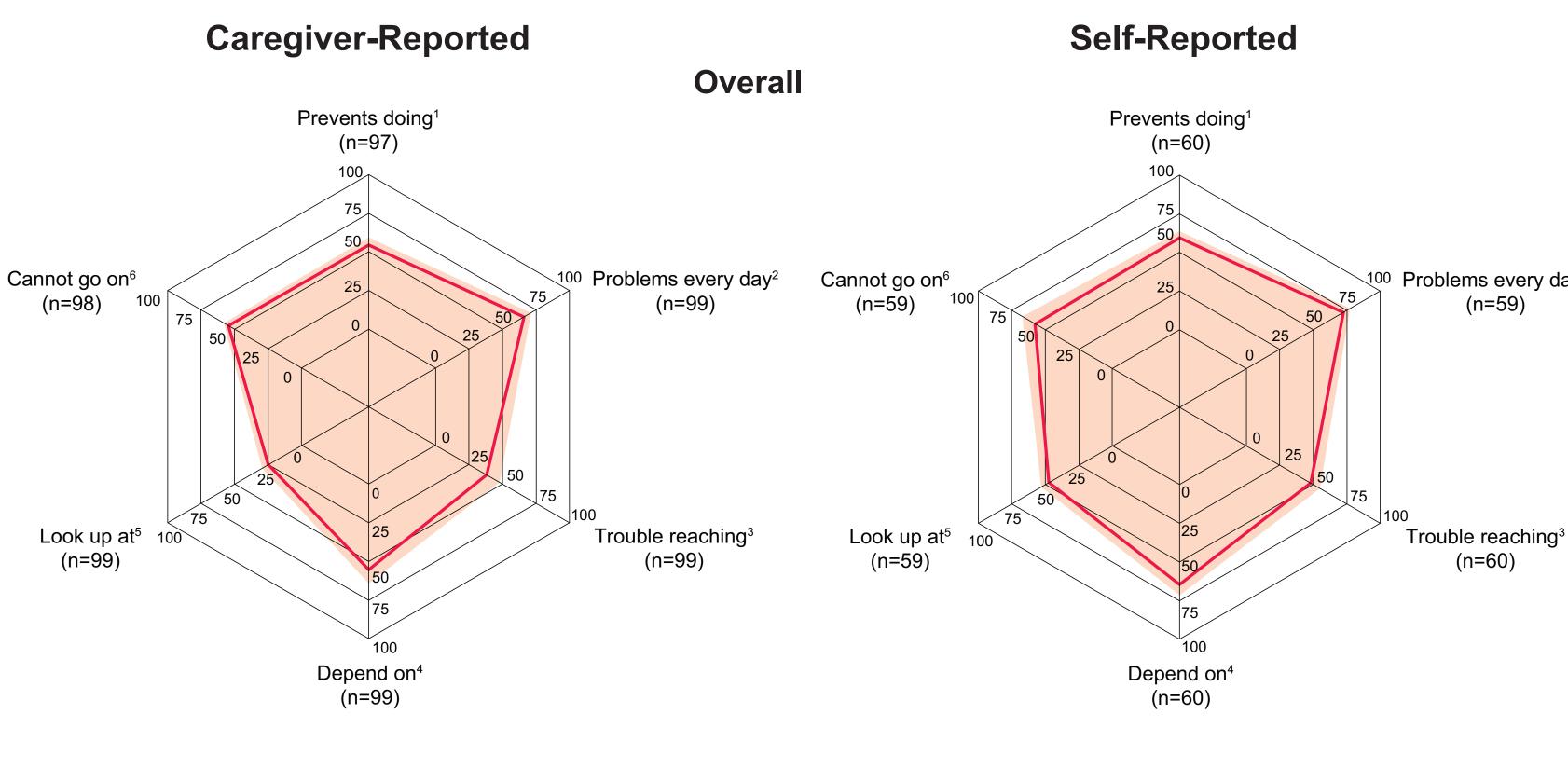
Data cut-off February 25, 2023. A positive change in QoLISSY score is indicative of an improvement in QoL. Z-scores were derived using ACH age/sex-specific reference data (means and SDs) from CLARITY (Hoover-Fong J et al. Orphanet J Rare Dis 2021).

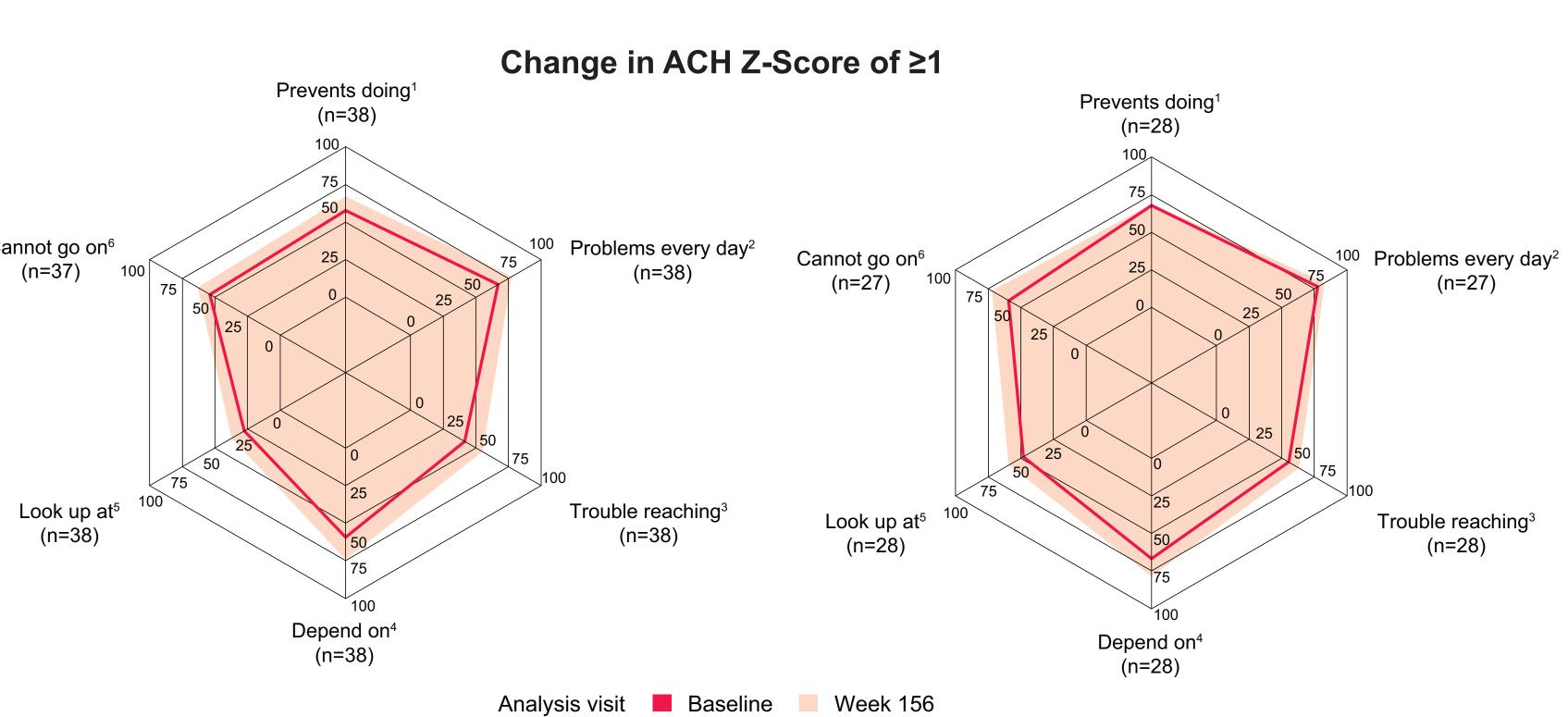
Mean change from baseline in self-reported QoLISSY scores at Year 3 in the treated population



Data cut-off February 25, 2023. A positive change in QoLISSY score is indicative of an improvement in QoL. Z-scores were derived using ACH age/sex-specific reference data (means and SDs) from CLARITY (Hoover-Fong J et al. Orphanet J Rare Dis 2021).

Distribution of mean physical domain scores at baseline and Year 3 in the treated population





¹Prevents doing things; ²Problems every day; ³Trouble reaching things; ⁴Depend on others; ⁵Look up at others when talking; ⁶Cannot go on fairground rides.

Conclusions

- These data suggest that vosoritide improves HRQoL among children with ACH, particularly for the physical domain scores
- There was a more pronounced change in participants with significant improvement in their ACH height Z-score (≥1 SD)

References

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Disclosures

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