

Evidence for Continuing Quality of Life Benefit Following CNM-Au8 ALS Treatment

Preliminary Analyses of the RESCUE-ALS Long Term OLE

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CONCLUSION: In RESCUE-ALS, CNM-Au8 treatment stabilised ALS-Specific QoL decline during the double-blind period, which was maintained for up to 84 weeks post-randomisation

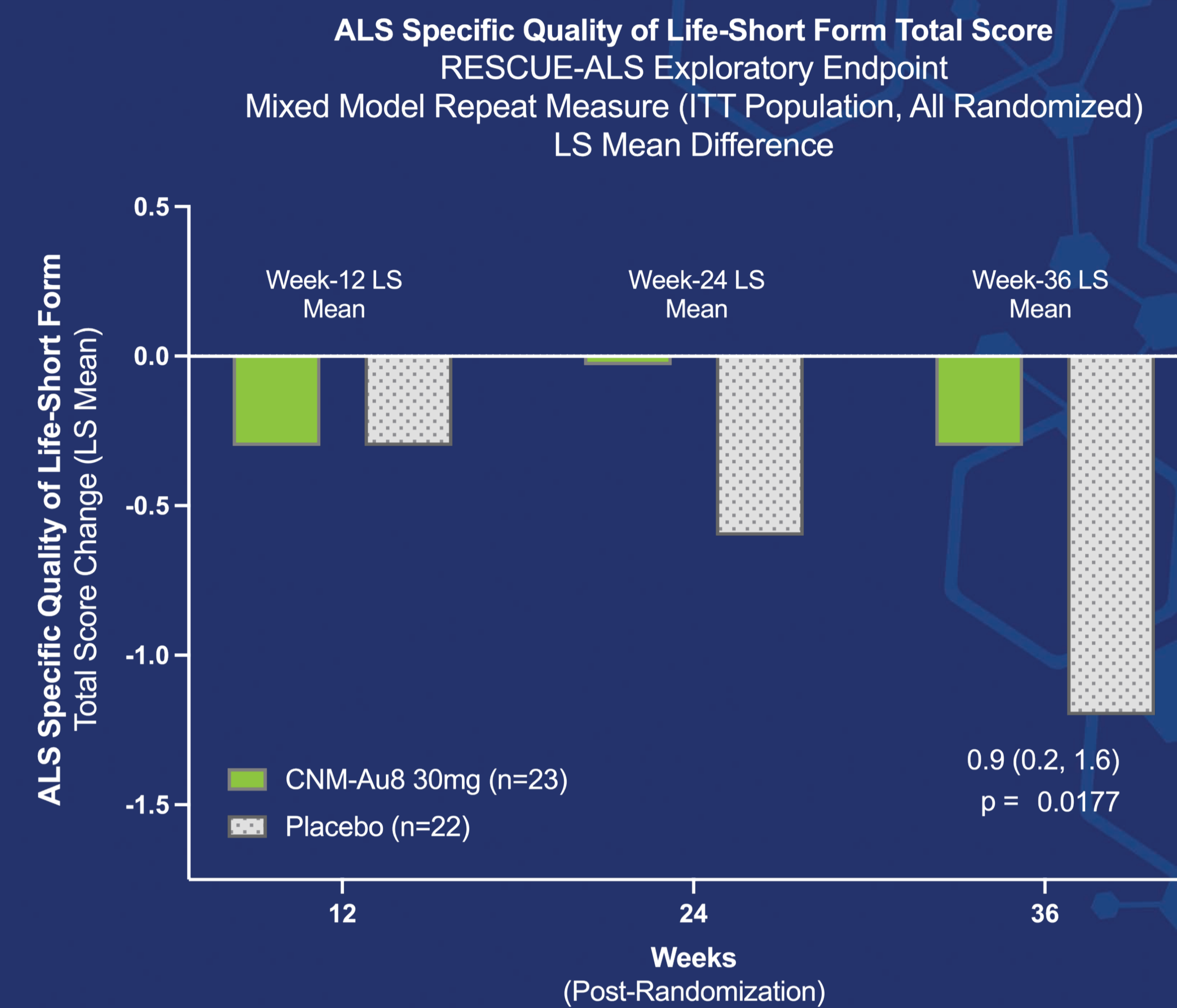
Design Scheme

36-Week Blinded Treatment Period with Long-Term Open-Label Extension



- Early symptomatic ALS (within 2-years onset or 1-year diagnosis)
- Randomized (1:1, CNM-Au8 30 mg or placebo)
- 36-week treatment period with long-term open label extension
- 1st EP: MUNIX(4) summed %change of 4-spinal cord innervated muscles
- 2nd EPs: absolute MUNIX change, % FVC
- Exploratory EPs: disease progression, 6-pt decline in ALSFRS-R, ALSSQOL-SF, & other neurophysiology endpoints

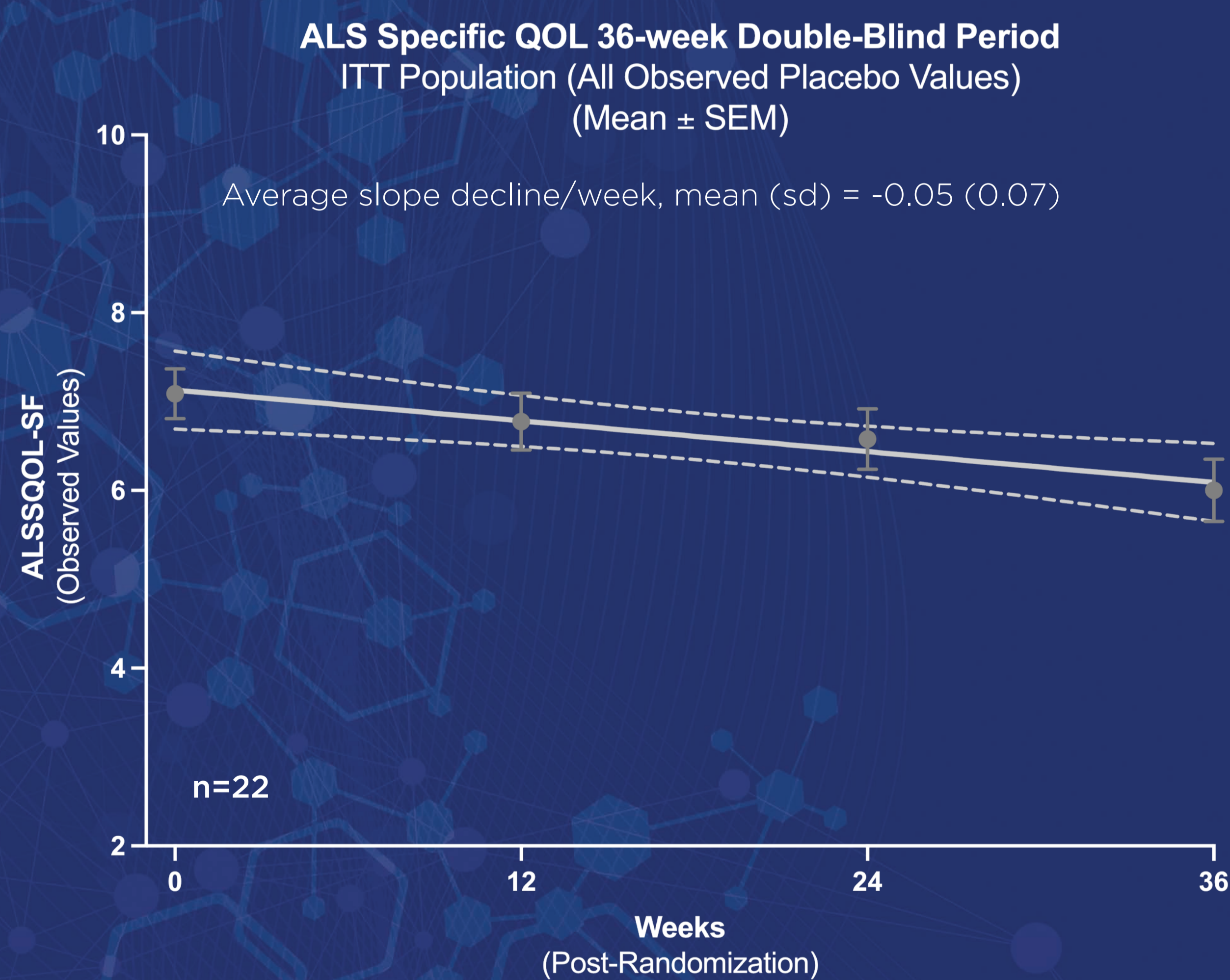
Week 36 Unblinded ALSSQOL Results



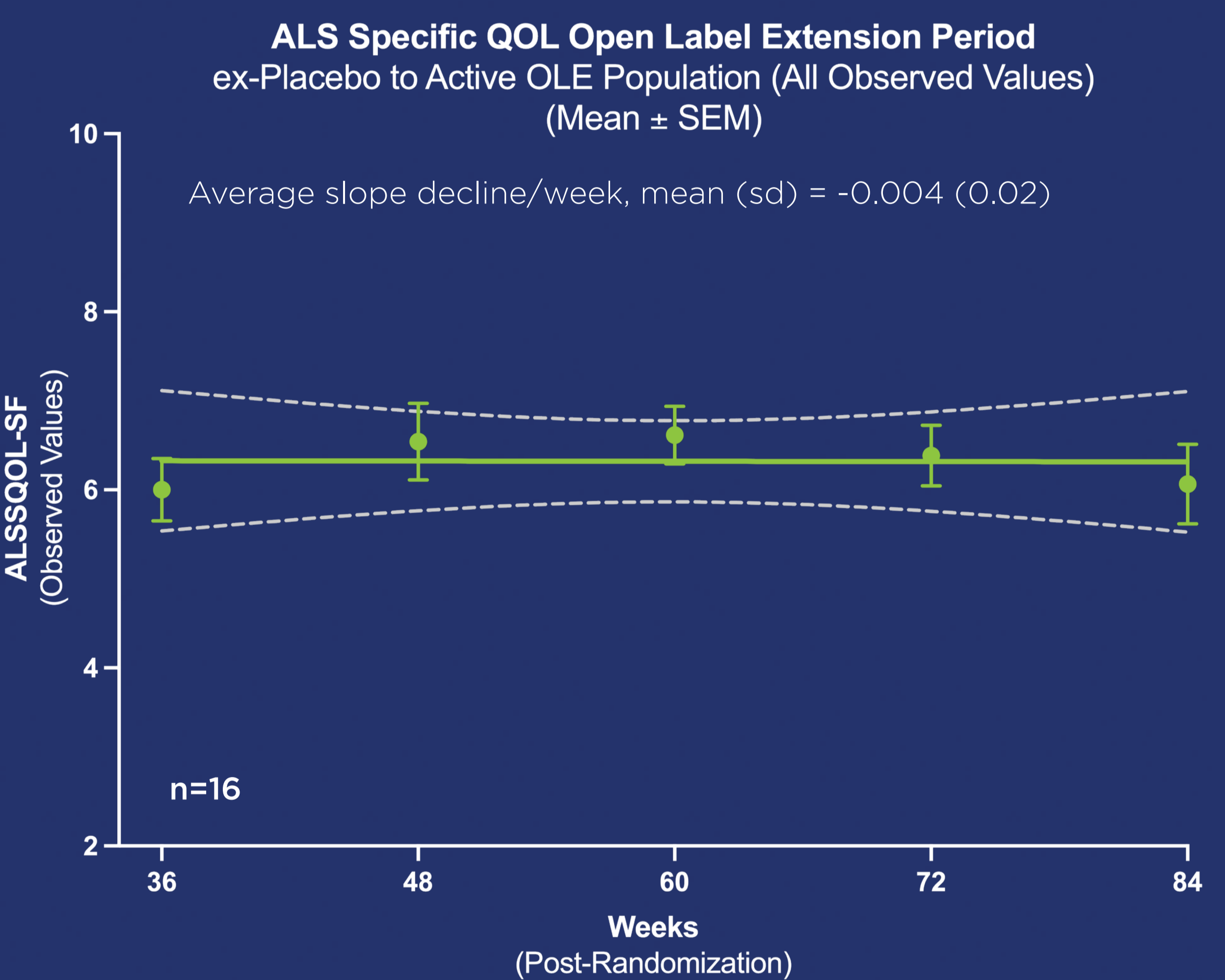
P-value is based on MMRM model with treatment, visit, treatment by visit interaction as fixed effects, and baseline value, and ENCALS score as covariates. An unstructured covariance model was used.

Ex-Placebo OLE CNM-Au8 treated participants stabilized ALS Specific QOL decline during OLE

Original Placebo | 36-Week Randomized Period

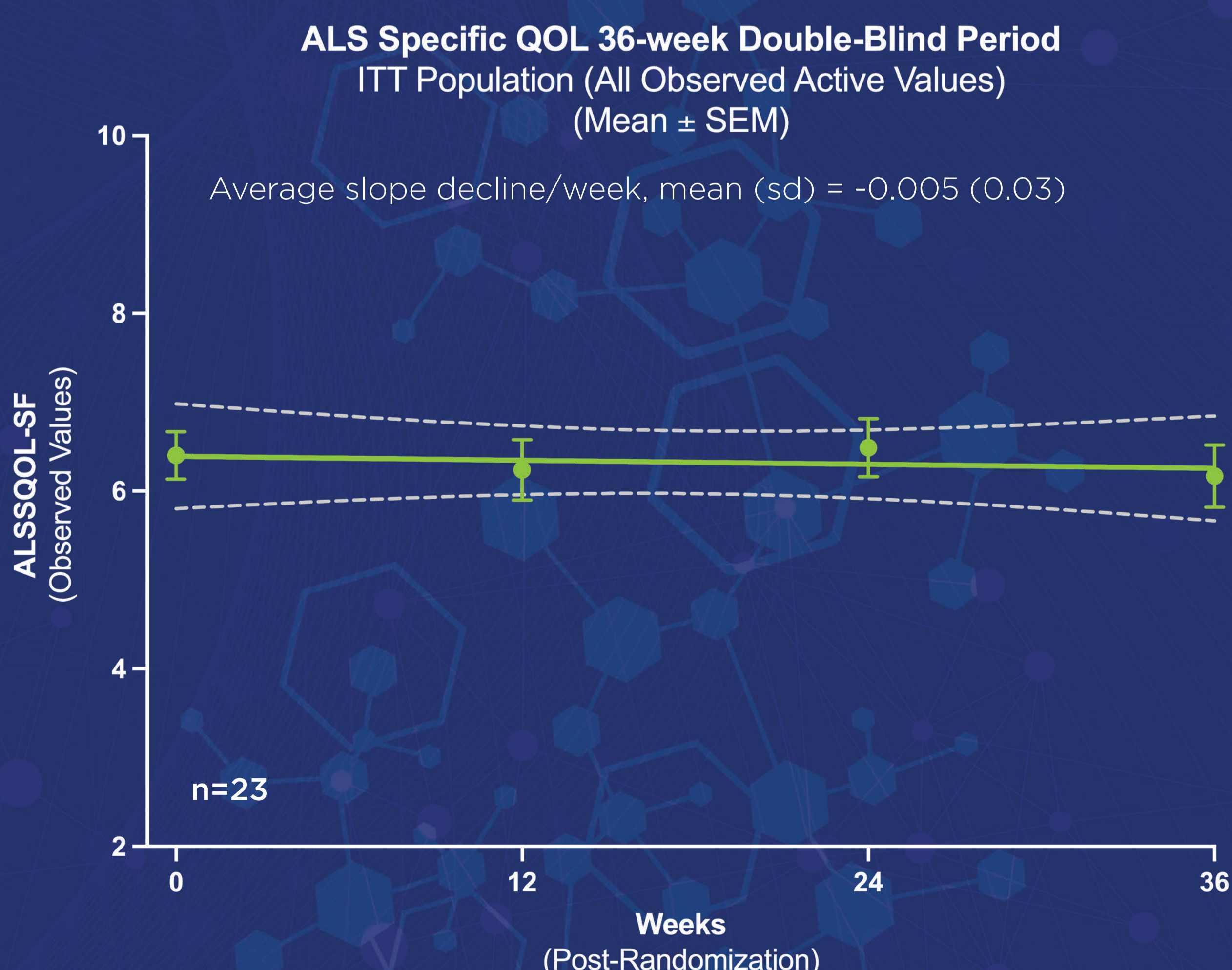


Original Placebo to OLE CNM-Au8 Treatment

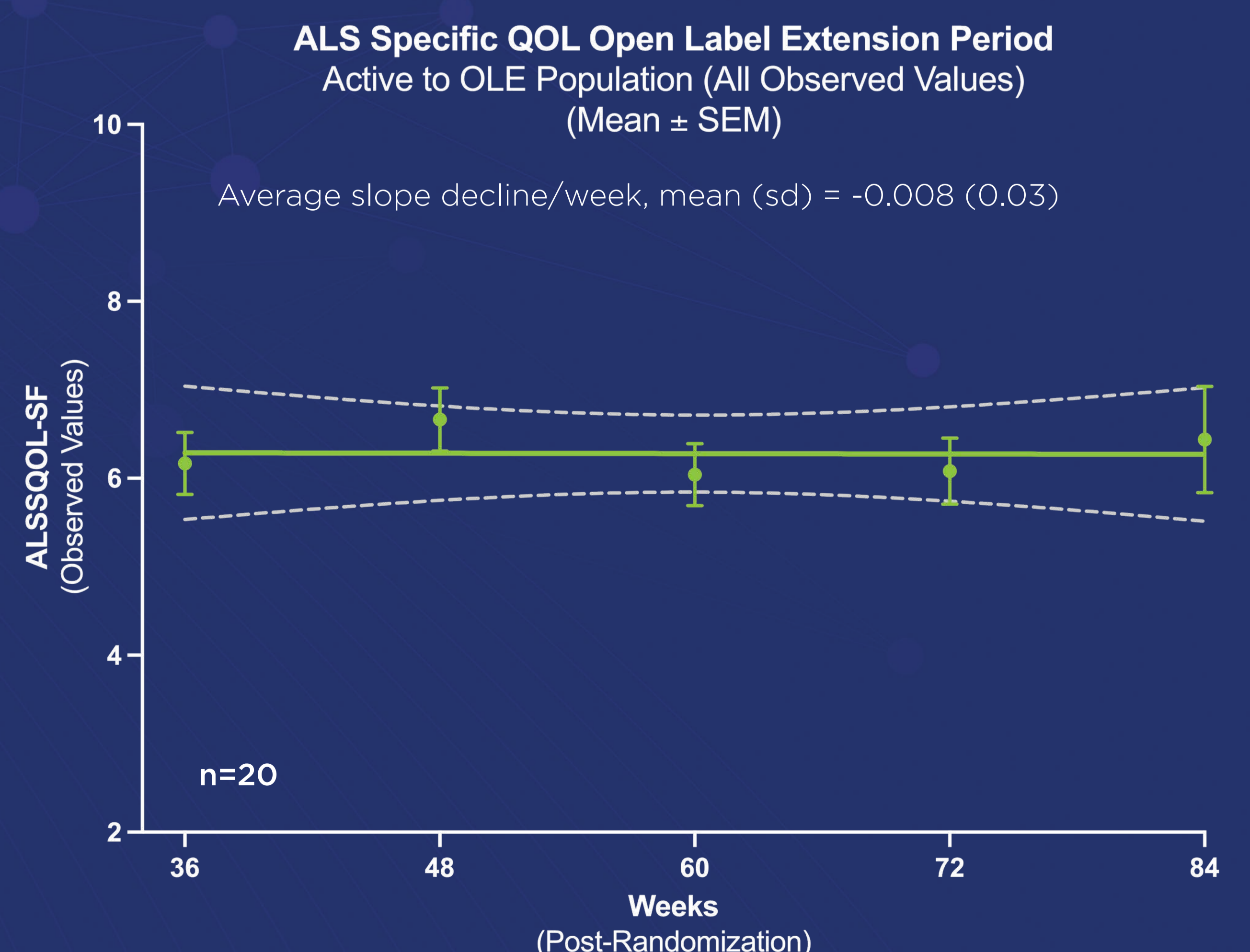


Original CNM-Au8 treated participants maintained ALS Specific QOL values during OLE treatment

Original Active | 36-Week Randomized Period



Original Active to OLE CNM-Au8 Treatment



Notes: Interim data cut as of 15-March-2022. Data include all reported values without imputation for missing data. **Acknowledgements:** We thank the ALS study patients and their families for their support and willingness to engage in clinical research. We thank the site investigators for their research excellence and dedication to patients. We thank FightMND of Australia for substantially funding the RESCUE-ALS trial.