

Clinical and qEEG Correlates of Tic Severity and Short-Term Outcome in Pediatric Tic Disorders: A Prospective Study



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BACKGROUND

Clinical Problem: Tics impair daily functioning and quality of life. Although many improve during adolescence, ~20% of cases persist/worsen into adulthood.

Pathophysiology: Tic disorders (e.g., Tourette Syndrome) involve corticostriatal-thalamocortical (CSTC) circuit dysfunction, frequently co-occurring with ADHD and OCD.

Quantitative EEG Role: qEEG tracks brain dynamics to identify neurophysiological patterns linked to tic generation and symptom dimensions.

AIM OF THE STUDY: to investigate the relationship between resting-state qEEG parameters, clinical phenotype, and short-term outcome in pediatric tic disorders.

MATERIALS AND METHODS

- 29 patients aged 3-18 years old
- Baseline evaluation of symptom severity using YGTSS, PUTS, CY-BOCS and CGI-S.
- Baseline resting-state EEG recordings and qEEG evaluation: theta/beta ratio (TBR), alpha peak frequency (APF), spectral power, and mean frequency.

Subdivision of patients into two groups:



RESULTS

Baseline Characteristics

- Higher clinical severity in the treated group vs. untreated group (YGTSS: 40.8 vs. 16.4, $p < 0.001$).
- No significant differences between groups in most qEEG parameters at baseline.

Clinical Outcomes and Predictors

- Significant tic improvement over time: patients showed significant tic improvement over time and a high clinical response rate (see charts)
- Baseline clinical severity was the strongest predictor of improvement, showing significant correlations between Δ YGTSS and baseline YGTSS ($\rho = 0.740$, $p < 0.001$), motor tic severity ($\rho = 0.798$, $p < 0.001$) and CGI-S ($\rho = 0.582$, $p = 0.007$)



Reduction in Tic Severity

Global tic severity decreased significantly across the entire cohort ($p = 0.007$).



Clinical Response Rate

Approx. one-third of patients met formal clinical response criteria at follow-up (T1).

qEEG Findings

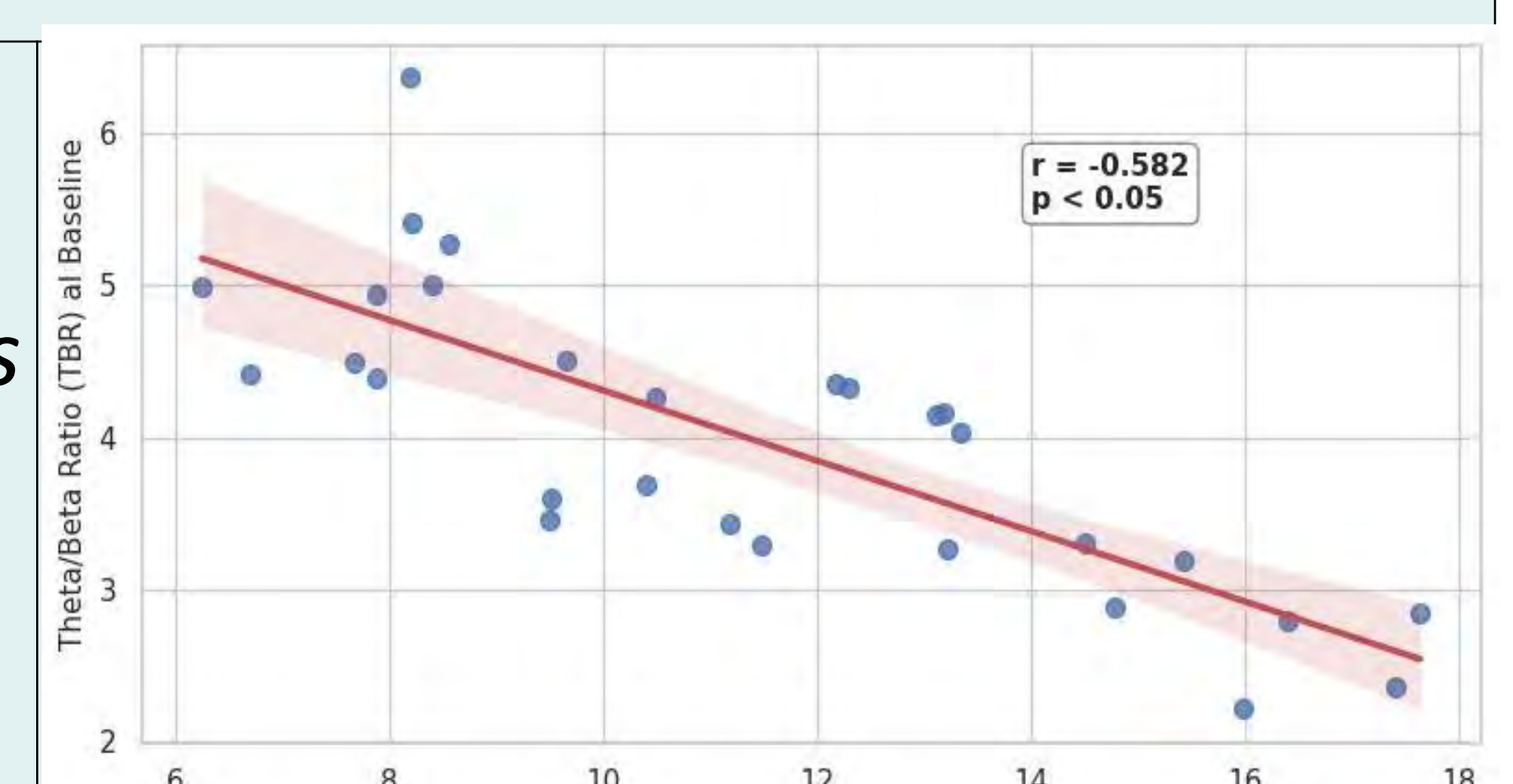
Global tic severity and clinical outcome were not associated with or independently predicted by qEEG parameters.

Specific symptom dimensions showed localized associations:

Premonitory urges showed a trend-level association with beta activity ($r = 0.355$, $p = 0.059$).

Alpha peak frequency (APF) emerged as potentially relevant for prognosis in multivariable models.

Age correlated significantly with qEEG measures, consistent with expected neurodevelopmental patterns.



CONCLUSIONS

Although qEEG markers did not reflect overall tic burden or short-term outcomes, they may capture specific clinical dimensions (e.g., premonitory urges) and developmental processes.