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Background

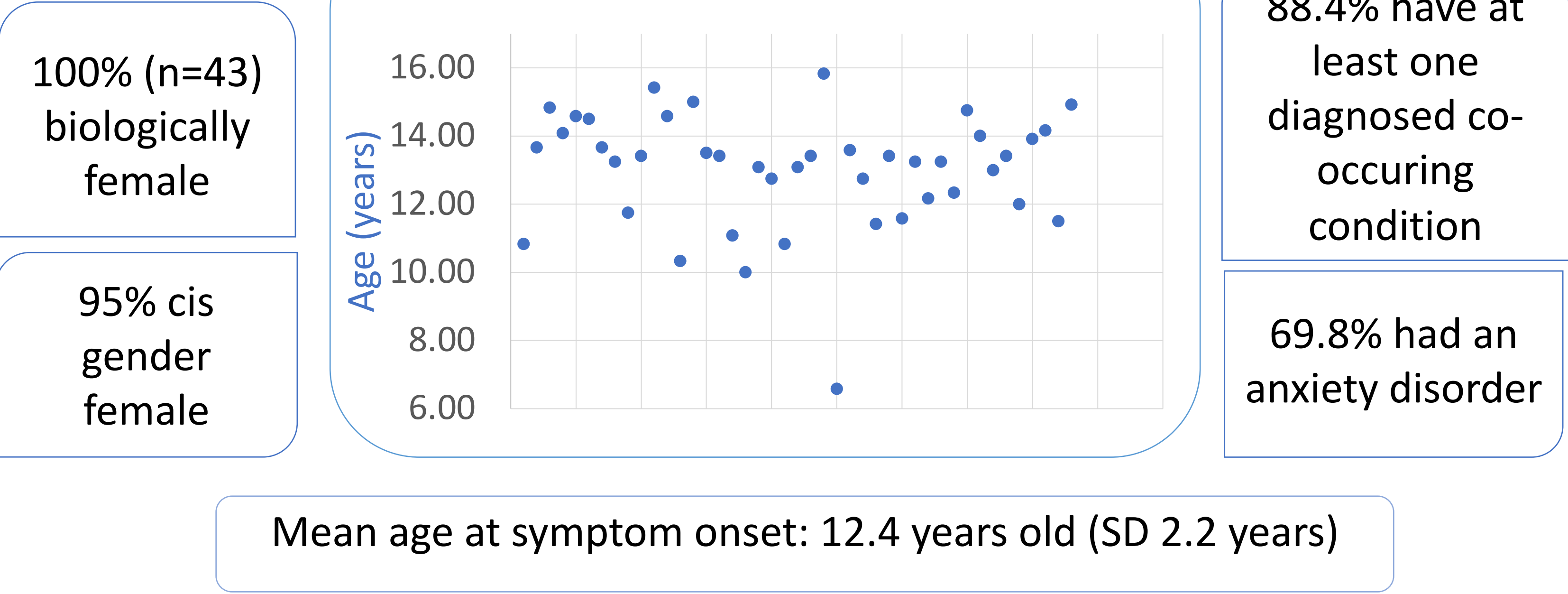
Currently, few studies have investigated the prognosis for children with FTLB:

- One study found 20% of children had remission without active treatment, but 55% relapsed (Martino *et al.*, 2023).
- Another reviewed 11 adolescents with FTLB and showed variable outcomes in anxiety and tics at 1 year follow up (Prato *et al.*, 2023).
- One reported marked improvements in symptoms in the majority of 15 adolescents studied (Howlett *et al.*, 2022). Similarly, a study looking at 28 adolescents found that all had improvements or resolution of FLTB symptoms (Okkels *et al.*, 2023)

Methods

We completed a clinical interview with parents/carers of patients diagnosed with FTLB between 1-4 years after symptom onset (**n=43**). We also administered a structured questionnaire with a collation of qualitative and quantitative data. Inclusion criteria included teenagers with FTLB in the Evelina TANDEM service ( all aged under 18 years at diagnosis). Retrospective Children’s Global Assessment Scale (CGAS) scores were done by the clinicians at diagnosis and repeated at follow-up. CGAS is a measure of general function, wellbeing and impairment.

Demographics:

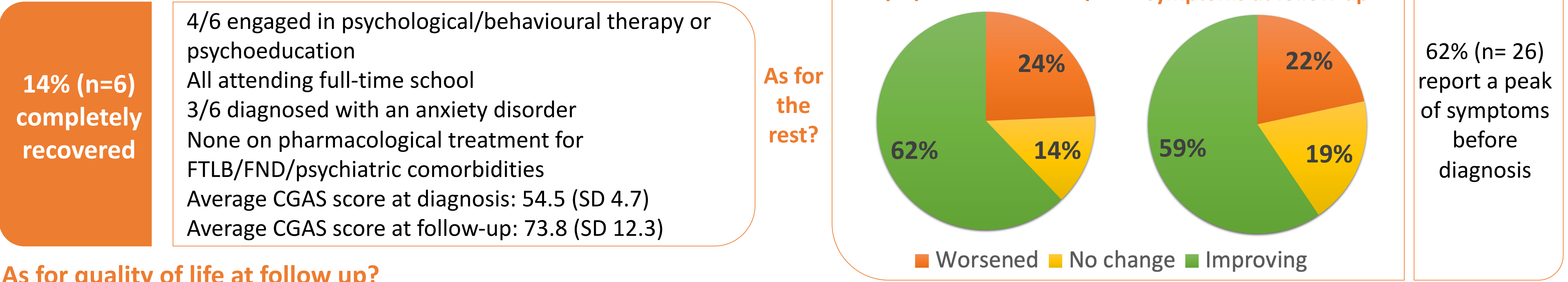


SUMMARY DECILE DESCRIPTIONS FOR CGAS:	
100-91	DOING VERY WELL
90-81	DOING WELL
80-71	DOING ALL RIGHT—MINOR IMPAIRMENT
70-61	SOME PROBLEMS—IN ONE AREA ONLY
60-51	SOME NOTICEABLE PROBLEMS—IN MORE THAN ONE AREA
50-41	OBVIOUS PROBLEMS—MODERATE IMPAIRMENT IN MOST AREAS OR SEVERE IN ONE AREA
40-31	SERIOUS PROBLEMS—MAJOR IMPAIRMENT IN SEVERAL AREAS AND UNABLE TO FUNCTION IN ONE AREA
30-21	SEVERE PROBLEMS—UNABLE TO FUNCTION IN ALMOST ALL SITUATIONS
20-11	VERY SEVERELY IMPAIRED—SO IMPAIRED THAT CONSIDERABLE SUPERVISION IS REQUIRED FOR SAFETY
10-1	EXTREMELY IMPAIRED—SO IMPAIRED THAT CONSTANT SUPERVISION IS REQUIRED FOR SAFETY

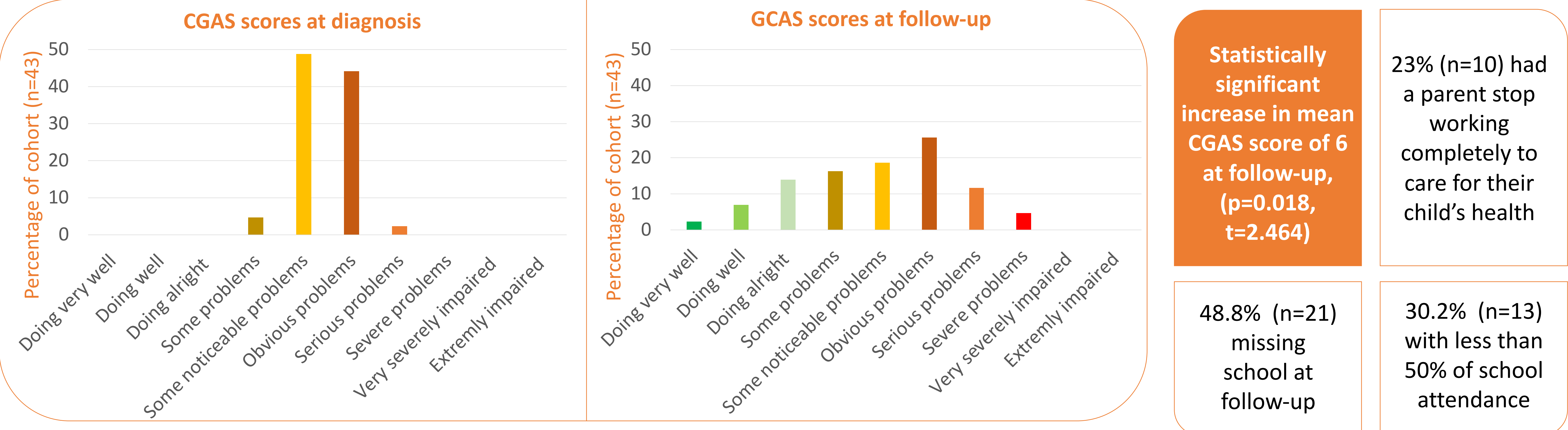
Time of follow-up:

1-2 years after symptoms onset	12%	Average time from symptom onset to diagnosis is 1.2 years (SD=0.8)
2-3 years after symptoms onset	56%	
3 years or more after symptoms onset	33%	

Findings:



As for quality of life at follow up?



Discussion:

- Symptoms overall improve in our cohort however, full resolution within 4 years was only seen in 14 % of our patients.
- Although functional symptoms improved in many, quality of life remained impaired for 76.7% (CGAS <71) and 31 % were not attending full-time school.
- Further analysis is planned to describe the impact of management approaches and of time-to-diagnosis on prognosis.
- We plan to study the emergence of new or different functional symptoms prospectively.

Limitations:

- Our service is not an FND service, therefore support for patients post-diagnosis is limited to psychoeducation and referral on to relevant services. This may have had an impact on the findings.
- The nature of data collection (interview of parents) may have led to bias (response, observer, interviewer). Similarly, retrospective CGAS scores may be subject to recall bias.

References:

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