

Z-proso Meeting 2018

Developmental trajectories of ADHD symptoms

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Developmental subtypes of ADHD symptoms

- **ADHD**

- Affects ~5% of global population but meaningful variation both above and below clinical thresholds
- Traditionally conceptualised as early-onset childhood disorder
 - Age-of-onset <12 required for diagnosis
- In reality substantial variation in age of onset/symptom developmental trajectories in general
- *Question: Can we parse the heterogeneity in ADHD symptom trajectories into meaningful developmental subtypes?*

Inattention

- *Difficulty organising tasks and activities*
- *Easily distracted by extraneous stimuli*
- *Forgetful in daily activities*

Hyperactivity/impulsivity

- *Fidgets, squirms in seat*
- *Often 'on the go'*
- *Difficulty waiting turn*

Example diagnostic indicators from DSM-5

Method

- **Overview**

1. Can developmental trajectories of ADHD symptoms estimated from longitudinal data be summarised in terms of a small number of 'developmental subtypes'?
2. Do these subtypes include a 'late onset' category?
3. Do those in this category differ from those with an early onset in clinically meaningful ways?

Method

- **Participants and Measures**

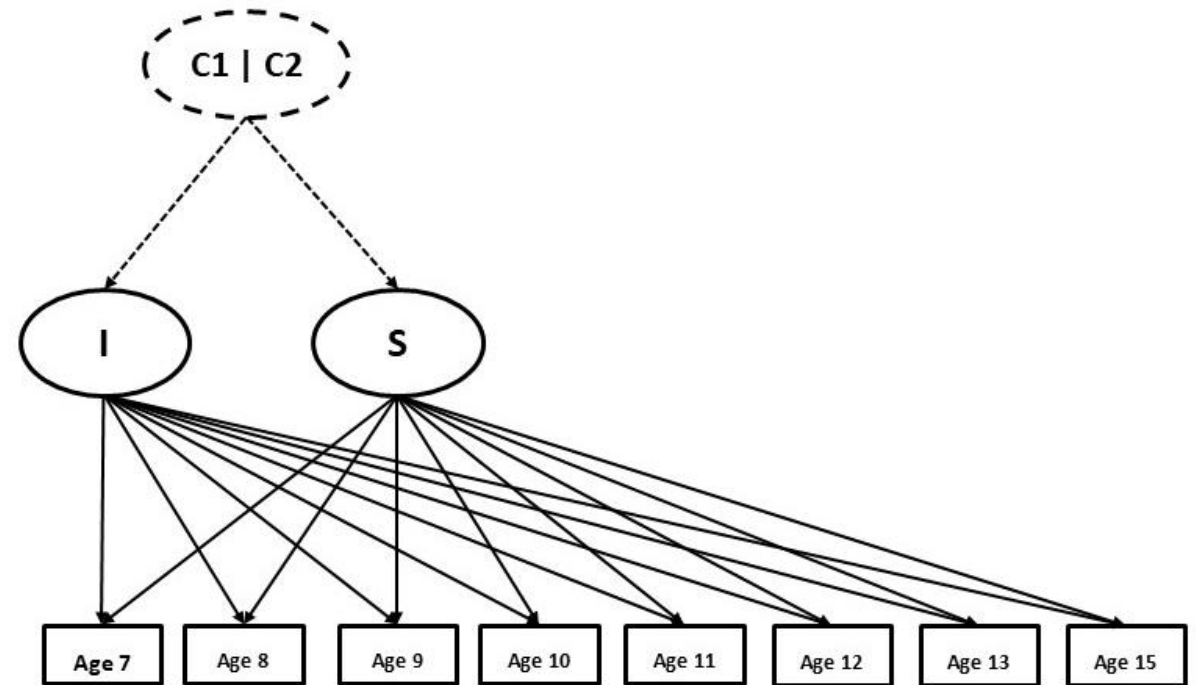
- Data from z-proso study
- n=1572 youth from Zurich, normative sample
- Teacher-reported ADHD symptoms at age 7,8,9,10,11,12,13,15



Method: Identifying developmental subtypes

- **Analysis Method**

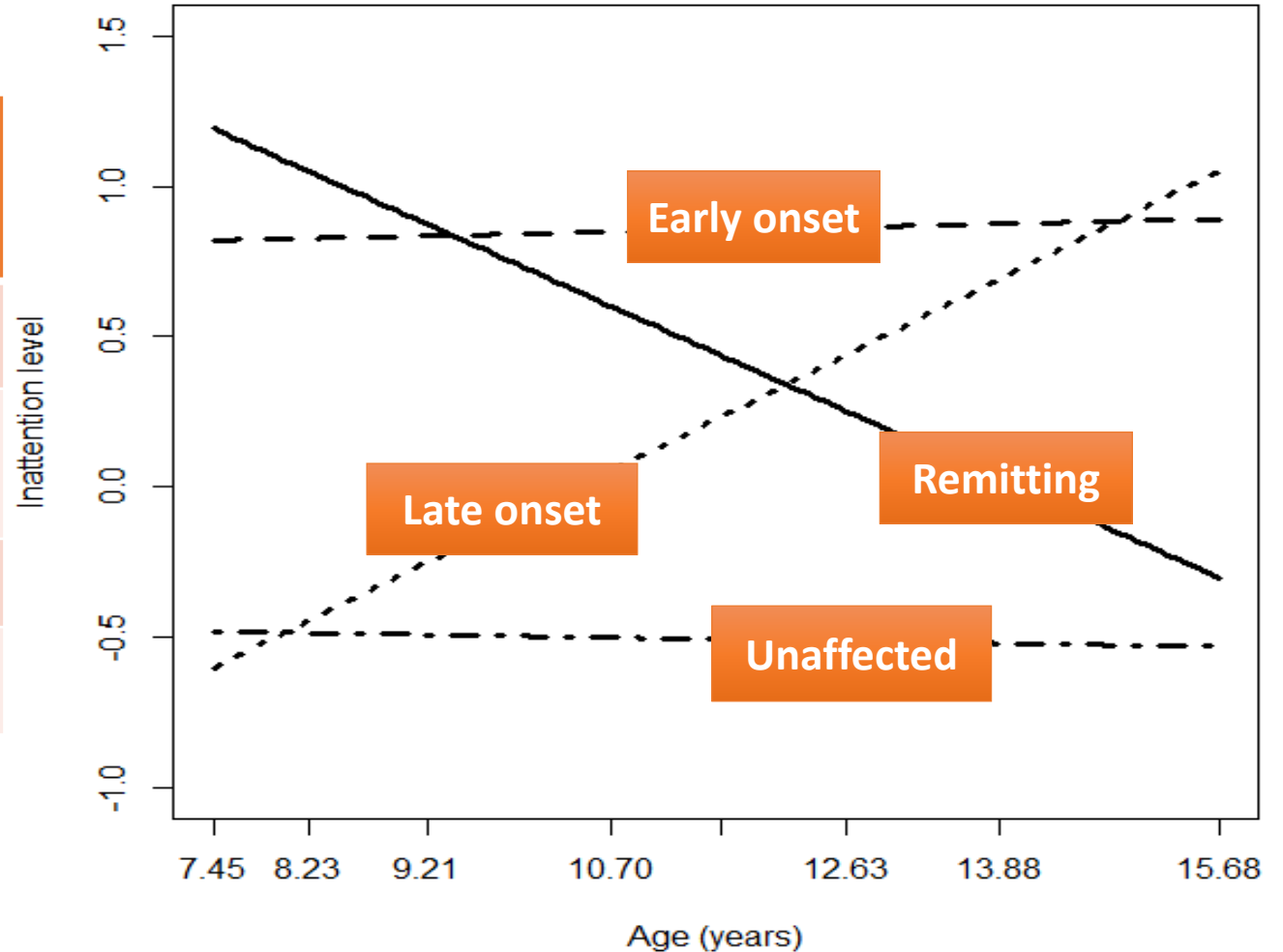
- Inattention and hyperactivity/impulsivity analysed separately.
- Growth mixture modelling (GMM)
 - Form of latent class analysis applied to longitudinal data
 - Identifies classes '**developmental subtypes**' defined by similar developmental trajectories in symptoms over time
 - Number of classes not known a priori but estimated from the data



The basic growth mixture model. I=intercept, S=slope; C1 and C2 are the categories of a latent categorical variable

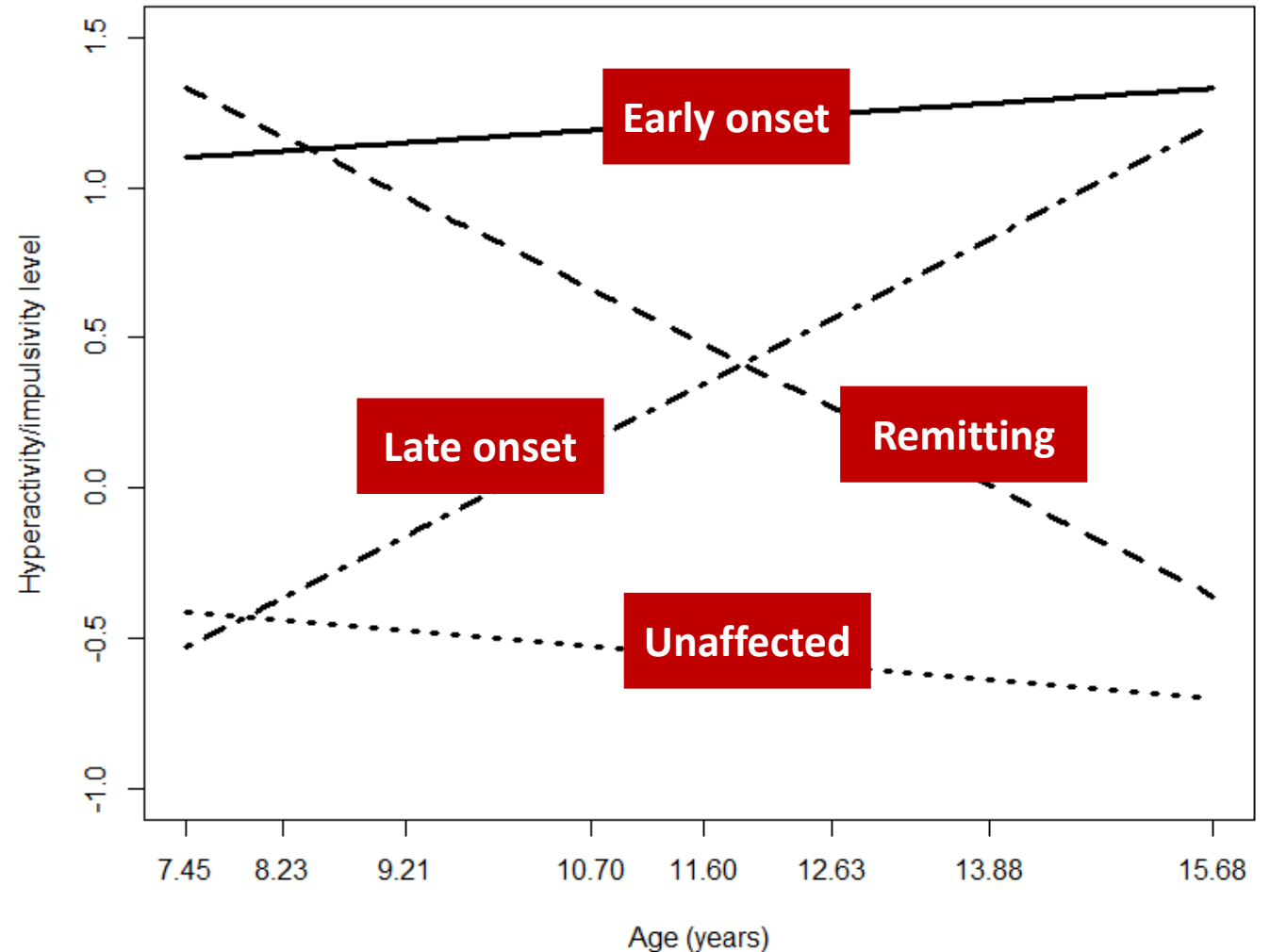
Results: Developmental subtypes of inattention

Inattention subtype	% (n) of Sample
Unaffected	63% (996)
Early onset	20% (311)
Remitting	10% (151)
Late onset	8% (112)



Results: Developmental subtypes of hyperactivity/impulsivity (H/I)

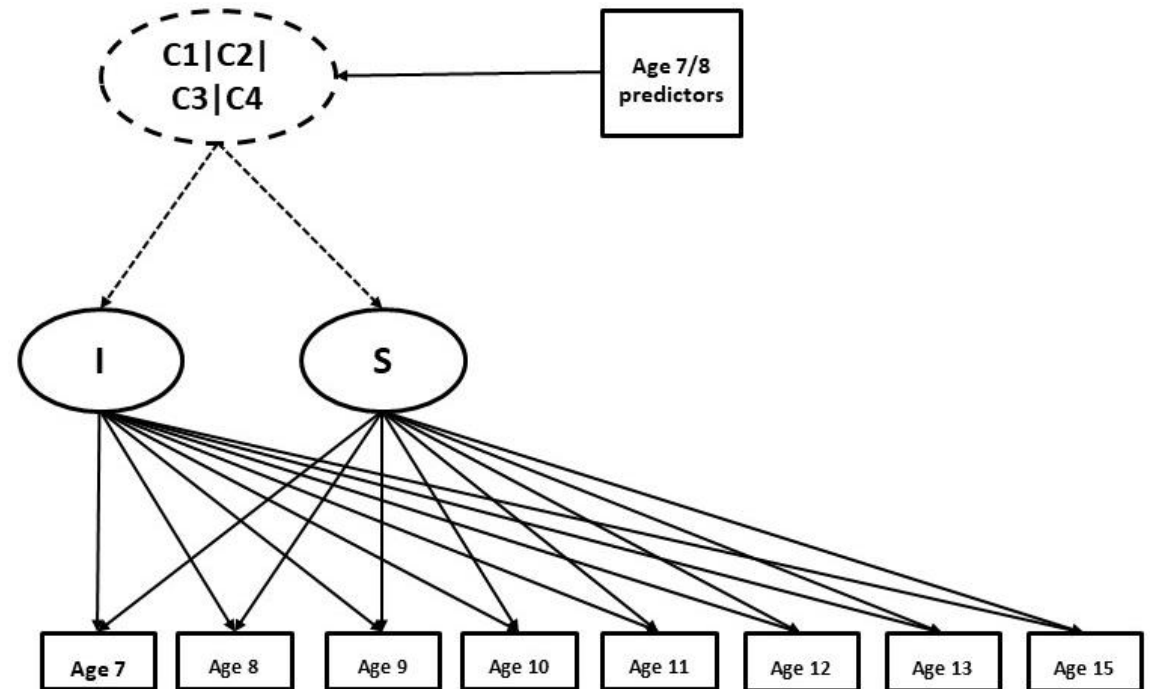
H/I subtype	% (n) of Sample
Unaffected	73% (1144)
Early onset	8% (125)
Remitting	13% (215)
Late onset	5% (86)



Method: Predictors and outcomes of category membership

- **Analysis method**

- Extend GMM to include predictors of class membership (measured at age 7/8)
- Extend the GMM to include outcomes of class membership (measured at age 17)
- Control for gender



Method: the predictors and outcomes

- **Predictors**

- Known childhood ADHD correlates
 - Sensation-seeking (age 7)
 - Risk-taking (age 8)
 - Reactive Aggression (age 7)
 - Anxiety (age 7)

- **Outcomes (age 17)**

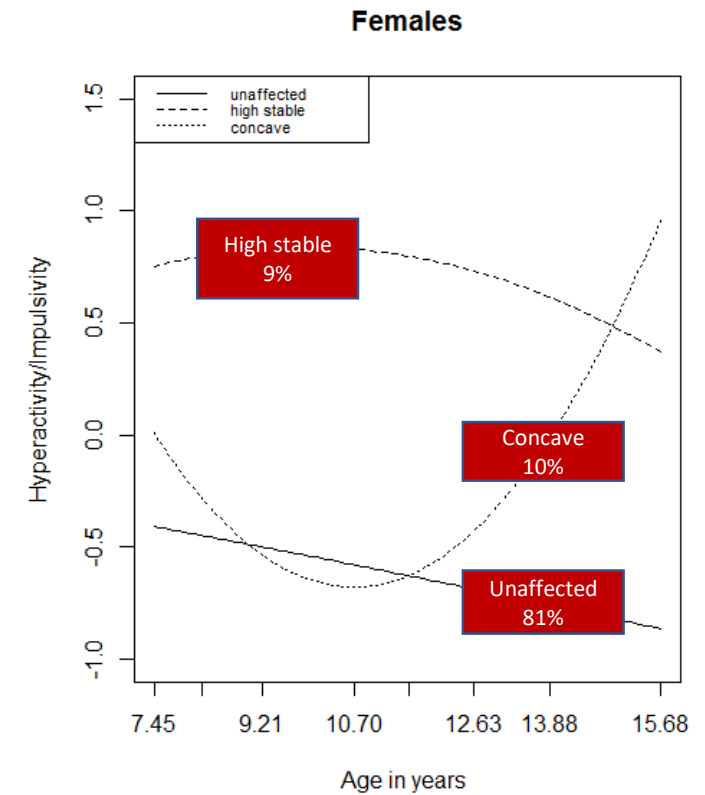
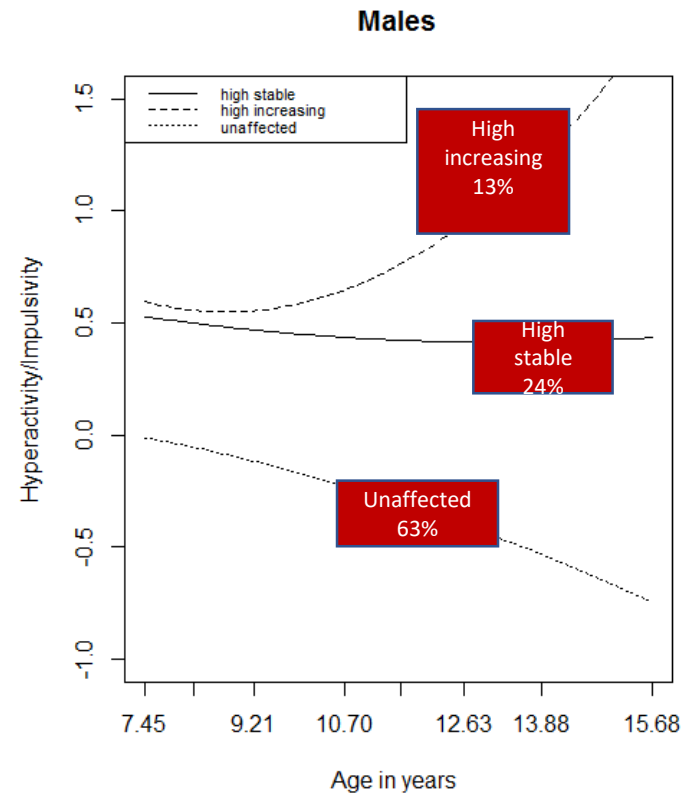
- Substance use
- Delinquency
- Aggression
- Internalising problems
- Violent ideations

Results



- General pattern:
 - Late onset scored higher on ADHD risk factors and outcomes than unaffected
 - Late onset scored lower on ADHD risk factors and outcomes than early onset
- Interpretation:
 - Late onset trajectory seems to show expected hallmarks and sequelae of ADHD but may be considered a milder developmental subtype

And in follow-up work...

- Females more likely to show a later onset
- Late onset category shows a particular increase in symptoms around puberty onset → *critical period for ADHD*

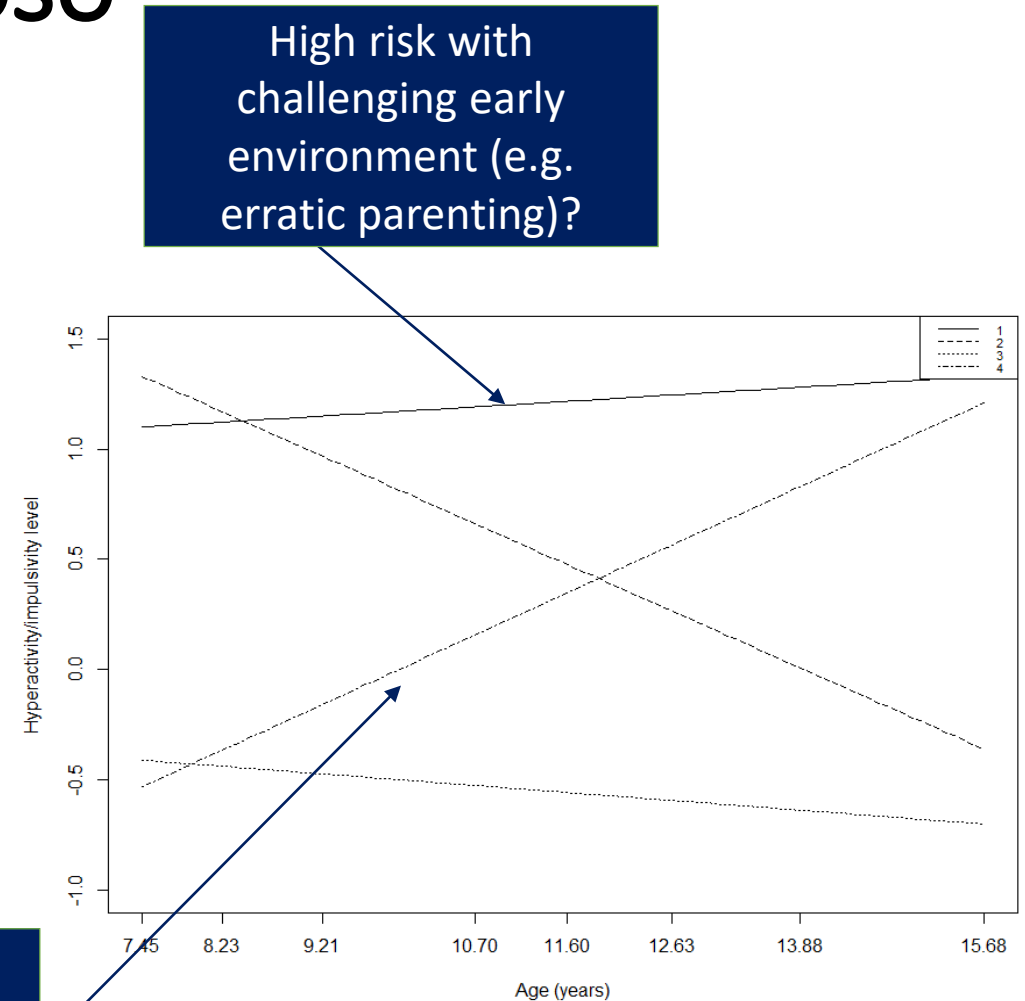


Summary and clinical implications

- Around **5-8%** of individuals show later onsets of ADHD symptoms
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- Scrap onset-before-12 criterion in diagnosis.
 - **Why?** People with later onset who could benefit from intervention may be excluded, especially females
- Those with a later onset show overall less impaired profile
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- Replace age-of-onset restriction with developmental subtypes
 - **Why?** Developmental trajectories may be informative about risks of adverse outcomes and support needs

Future directions in Z-proso

- 1. Follow-up at age 20:
 - Do we see more 'late onsets'?
 - Do late versus early onset show different experience sampling profiles at age 20?
- 2. Do those in the late onset category show a more 'enriched' early environment that delays symptom manifestation relative to early onset?



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References

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Thank You