

Genomic contributions to infant and toddler vocabulary scores: Implications for association with health-, cognition-, and behaviour-related outcomes

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Key Points

Question: What is the genetic architecture underlying vocabulary acquisition during language development, and does it affect links with later-life outcomes?

Findings: At least two genetic components contribute to vocabulary size, predominantly distinguishing infant expressive from toddler receptive vocabulary. Matching patterns of genetic overlap were found with later-life outcomes: Larger infant expressive but smaller toddler receptive vocabulary size was correlated with higher ADHD risk and/or childhood maltreatment exposure (a behavioural proxy). Consistently, later-life cognition was associated with toddler vocabulary scores only, irrespective of power.

Meaning: The genetic architecture underlying vocabulary acquisition is dynamic, shaping polygenic associations with later-life behaviour and cognition.

Abstract

Importance: The number of words a child produces (expressive vocabulary) and understands (receptive vocabulary) change rapidly during infancy and toddlerhood, partially due to genetic factors. However, the genetic architecture underlying vocabulary development and association patterns with later-life outcomes, have not yet been fully characterised.

Objective: To (i) study the multivariate genetic architecture underlying vocabulary size during infancy and toddlerhood, and (ii) describe polygenic association patterns with childhood behavioural and health measures, as well as adult cognition-related outcomes.

Design: Meta-genome-wide association study (meta-GWAS) of expressive and receptive vocabulary (age: 15-38 months) performed within the Early Genetics and Life Course Epidemiology (EAGLE) Consortium. Structural equation modelling techniques were applied to study multivariate genetic architectures.

Setting: Children of European descent across seven independent population-based cohorts.

Participants: 37,913 observations from 17,298 individuals.

Main Outcome and Measure: Meta-analyses were performed for early-phase expressive vocabulary (15-18 months), late-phase expressive vocabulary (24-38 months), late-phase receptive vocabulary (24-38 months), and combinations thereof. Vocabulary size was assessed by parent report using standardised psychological instruments.

Results: Common genetic variation explained a modest proportion of phenotypic variation across all vocabulary measures (Single-Nucleotide Polymorphism heritability: 0.08(SE=0.01) to 0.24(SE=0.03)). Genetic correlation (r_g) analyses showed that late-phase expressive vocabulary largely shared genetic influences with both early-phase expressive ($r_g=0.69$ (SE=0.14)) and late-phase receptive vocabulary ($r_g=0.67$ (SE=0.16)). However, the latter two measures were genetically unrelated ($r_g=0.07$ (SE=0.10)),

suggesting different underlying genetic factors. Consistently, we observed differences in polygenic association patterns: Larger early-phase expressive vocabulary size was genetically correlated with increased ADHD risk ($r_g=0.23(SE=0.08)$) and childhood maltreatment exposure ($r_g=0.19(SE=0.07)$), a behavioural proxy. In contrast, larger late-phase receptive vocabulary size was genetically correlated with lower childhood maltreatment exposure ($r_g=-0.33(SE=0.08)$). Finally, toddler, but not infant, vocabulary size was linked to cognitive skills (e.g. late-phase expressive vocabulary and intelligence: $r_g=0.32(SE=0.08)$), despite comparable power.

Conclusions and Relevance:

There are at least two distinct genetic components contributing to vocabulary development during infancy and toddlerhood that shape polygenic association patterns with later-life cognition and ADHD-related traits. Our findings suggest differences in biological mechanisms during a phase where children “learn to speak” (infancy) compared to a phase where children mastered some fluency and “speak to learn” (toddlerhood).

Introduction

Language development in infants and toddlers is often assessed with measures of expressive and receptive vocabulary^{1,2}. These constructs relate to language production and understanding, respectively, and can be relatively easily (albeit indirectly) measured through parental reports. The first spoken words, representing one of the milestones in language development, typically emerge between the ages of 10 to 15 months². Receptive vocabulary development, usually, precedes expressive vocabulary development, emerging at six to nine months of age³. Consequently, the number of words children understand is often larger than the number of words they produce, and exceeds the latter at least four-fold based on parent-reported measures at 16 months of age⁴. Once children reach an expressive vocabulary size of ~50 words at an age of 12 to 18 months, there is often a period of rapid vocabulary growth around 16 to 22 months of age⁵, resulting in a vocabulary size of 100 to 600 words at ~24 months⁴. During the early stages of language learning in infancy (≤ 18 months of age) children typically produce words in isolation², followed by a period of two-word combinations and more complex grammatical structures^{4,6}.

Individual differences in early-life vocabulary development can, partially, be explained by genetic factors⁷⁻¹⁰. Twin heritability (twin- h^2) estimates for expressive vocabulary range between 10% and 25% (24-36 months)⁷⁻⁹, reflecting phenotypic variation due to all possible genetic influences. These findings are corroborated by genetic research investigating Single-Nucleotide Polymorphisms (SNPs), with SNP- h^2 estimates of 13% and 14% (15-30 months)⁷. For receptive vocabulary at 14 months of age, a twin- h^2 estimate of 28% has been reported¹¹. Evidence for SNP- h^2 at a similar age was poor¹², based on a single-cohort study, but was present at 38 months of age (12% SNP- h^2)¹².

Population-based studies of English-speaking children showed that the genetic architecture of language development spanning infancy to early childhood is complex, with evidence for both stability and change in underlying genetic contributions^{7,8,12,13}. At the genome-wide level, genetic correlations (r_g) for measures of expressive vocabulary between 15 and 38 months of age ranged from 0.48 to 0.74^{7,8,12},

suggesting moderate-to-strong genetic stability. At the individual SNP-level, a previous meta-genome-wide association study (meta-GWAS, N=8,889) identified a GWAS signal at rs7642482 on chromosome 3p12.3, near the *ROBO2* gene⁷ that was associated with expressive vocabulary in infants (age: 15-18 months) but attenuated in toddlers (age: 24-30 months)⁷. These changes might be due to age-specific genetic mechanisms, highlighting the need for more powerful studies to identify and characterise genetic association.

Genetic influences underlying early-life vocabulary are shared with many later childhood abilities. In UK twins, for example, early expressive language skills (24-48 months) were moderately genetically correlated ($r_g=0.36$) with childhood reading abilities¹³. Similarly, in a UK population-based genomic study, receptive vocabulary scores (38 months) showed moderate-to-strong genetic correlations ($r_g=0.58-0.92$) with mid-childhood reading skills¹⁴. Genetic influences underlying early-life vocabulary may also be shared with other childhood behavioural and health measures, including childhood-onset neurodevelopmental conditions as Attention-Deficit/Hyperactivity Disorder (ADHD) and Autism Spectrum Disorder (ASD). For example, children with ADHD often experience difficulties with mastering language and literacy skills¹⁵⁻¹⁷ and poor language skills at the age of three years were found to be predictive of inattention and hyperactive symptoms two years later in life¹⁸. More specifically, there is evidence for genetic overlap between higher ADHD risk and lower mid-childhood/early-adolescent language- and literacy-related abilities, primarily implicating reading performance¹⁹⁻²², underscoring the need to study also genetic links with early-life language measures. For children diagnosed with ASD, the phenotypic spectrum is wider, including children with little or no spontaneous spoken language by the time they reach school age²³ as well as individuals with, comparably, few problems in the language domain²⁴. Both, children with ADHD and ASD are at increased risk for childhood maltreatment compared to typically developing children^{25,26}, consistent with moderate genetic trait correlations (ADHD: $r_g=0.56$, ASD: $r_g=0.41$)²⁷. Such genetic relationships may capture gene-environment correlation, as children's behaviour (partially influenced by genetic factors) elicits

responses from parents and others²⁷. In addition, we assess evidence for genetic links between children's vocabulary and head circumference measures, a characteristic that may indicate abnormal development²⁸ and is highly correlated with MRI brain volume^{29,30}.

In this study, we aim to elucidate the polygenic architecture of vocabulary acquisition. We investigate developmental changes in genetic contributions at the single-variant and trait covariance level by performing genome-wide association meta-analyses of expressive and receptive vocabulary size at different developmental stages during infancy and toddlerhood, including an early, single-word phase (15-18 months) and a late (24-38 months) phase during which children start using two-word combinations and more complex grammatical structures. Furthermore, we characterise polygenic links with childhood behavioural and health measures, as well as adult cognition-related outcomes, fitting structural models.

Methods

Phenotype selection and study design

Cohorts with quantitative vocabulary scores assessed during the first three years of life and genome-wide genotypes were invited to participate in this study, embedded within the Early Genetics and Life Course Epidemiology (EAGLE) consortium³¹ (<https://www.eagle-consortium.org/working-groups/behaviour-and-cognition/early-language/>). Expressive vocabulary scores were assessed between 15 and 38 months of age and analysed across two developmental stages to allow for age-specific genetic influences: an early phase (15-18 months) and a late phase (24-38 months). Scores for receptive vocabulary were included for the late phase only (24-38 months), due to low measurement availability, low reliability and little evidence for SNP- h^2 during the early phase¹² (eMethods 1).

Up to seven population-based cohorts (eMethods 2) participated in the meta-analyses, of which two had longitudinal vocabulary assessments (Figure 1, eTable 1). Vocabulary scores were ascertained by

parental report using age-specific word lists that were adapted from the MacArthur Communicative Development Inventory (CDI)^{9,32–36} or the Language Development Survey (LDS)³⁷ (eTable 1, eMethods 3). Ethical approval was obtained by the local research ethics committee for each participating study, and all parents and/or legal guardians provided written informed consent (eMethods 2).

Genotyping and imputation

Genotyping within each cohort was conducted using high-density SNP arrays and quality control followed standard procedures³⁸ (eTable 2). In total, between 440,476 and 608,517 high-quality autosomal genotyped markers were imputed against a Haplotype Reference Consortium (HRC) r1.1 panel³⁹ (eTable 2).

Single variant association analyses and meta-analyses

Within each cohort, vocabulary scores were adjusted for potential covariates and rank-transformed to achieve normality and to allow for comparisons of genetic association effects across different psychological instruments (eMethods 4). SNP-vocabulary associations were then estimated within each cohort using linear regression of rank-transformed residuals on posterior genotype probability, except for the Longitudinal Study of Australian Children (LSAC) where we analysed best-guess genotypes, assuming an additive genetic model (eTable 2, eMethods 4). Prior to meta-analysis, GWAS summary statistics underwent extensive quality control using the EasyQC R package⁴⁰ (v9.2) (eTable 2, eMethods 4).

As part of analysis stage I, single-trait meta-analyses were performed for early-phase expressive vocabulary, late-phase expressive vocabulary and late-phase receptive vocabulary using either METAL⁴¹ and/or multi-trait analysis of genome-wide association (MTAG)⁴² software (Figure 1, eMethods 4). As part of analysis stage II, multi-trait meta-analyses were performed with MTAG⁴² combining genetically correlated vocabulary traits to increase statistical power (Figure 1).

The number of independent vocabulary measures analysed in this study was 2.38, as estimated with a Matrix Spectral Decomposition (matSpD)⁴³ based on bivariate genetic correlations (see below) across the three single-trait vocabulary meta-analyses (stage I). This resulted in a multiple-testing-adjusted genome-wide association significance threshold of $P < 2.10 \times 10^{-8}$ ($5 \times 10^{-8} / 2.38$).

FUMA analyses

SNP-vocabulary associations passing the unadjusted genome-wide significance threshold ($P < 5 \times 10^{-8}$) were identified and annotated using Functional Mapping and Annotation of genetic associations⁴⁴ software (FUMA v1.3.6). In addition, gene-based genome-wide, gene-set and gene-property analyses were conducted with Multi-marker Analysis of GenoMic Annotation (MAGMA, v1.08) within FUMA⁴⁴ (v1.3.6a) (eMethods 5).

SNP-heritability and genetic relationship analyses

Using GWAS summary statistics, SNP- h^2 and bivariate genetic correlations (r_g) were estimated for early-life vocabulary (meta-analysis stages I and II) and/or later-life traits using High-Definition Likelihood⁴⁵ (HDL, eMethods 6, see below).

Given evidence of HDL-SNP- h^2 ($P < 0.05$), HDL- r_g analyses (eMethods 6) were performed to assess genetic overlap (i) across single-trait vocabulary measures (stage I) and (ii) across early-life vocabulary measures and later-life health-, cognition-, and behaviour-related outcomes (eMethods 6): reading performance in the general population (8-22 years, N=13,027; GWAS summary statistics were created as described in eMethods 7, eTable 3), intelligence⁴⁶ (5-98 years, N=279,930), educational attainment⁴⁷ (>30 years, N=766,345), infant head circumference⁴⁸ (6-30 months, N=10,768), childhood head circumference⁴⁹ (6-9 years, N=10,600), childhood aggressive behaviour⁵⁰ (1.5-18 years, N=151,741), childhood internalising

symptoms⁵¹ (3-18 years, N=64,641), childhood maltreatment exposure⁵² (<18 years, N=150,290), ADHD⁵³ (N=53,293; N_{cases}=19,099) and ASD⁵⁴ (N=46,350; N_{cases}=18,381). The multiple-testing-adjusted threshold for HDL- r_g analyses was defined at 5.32×10^{-3} , reflecting a correction for 9.39 independent traits considered (0.05/9.39), estimated using matSpD^{43,55} and a bivariate genetic correlation matrix (eFigure 1).

Polygenic prediction of late-phase expressive vocabulary was carried out using polygenic scoring⁵⁶ (eMethods 8).

Structural equation models

We modelled the multivariate genetic architecture between early-life vocabulary measures and genetically associated later-life outcomes (as identified with HDL- r_g analyses, $P < 5.32 \times 10^{-3}$) using genomic structural equation modelling (genomic SEM)⁵⁷ and genome-wide summary statistics (eMethods 9). Follow-up analyses of phenotypic, genetic and residual correlations of mid-childhood and early adolescent ADHD symptoms with infant and toddler vocabulary measures were performed using individual-level data from the Avon Longitudinal Study of Parents And Children (ALSPAC) cohort and genetic-relationship-matrix structural equation modelling⁵⁸ (GRM-SEM) (eMethods 10, eTable 4).

Results

Single-trait and multi-trait meta-GWAS

Single-trait genome-wide association analyses based on children of European descent from seven independent cohorts were performed for early-phase expressive vocabulary (15-18 months, N=8,799), late-phase expressive vocabulary (24-38 months, N=16,615) and late-phase receptive vocabulary (24-38 months, N=6,291) (stage I, Figure 1). There was little evidence for novel SNP association signals with vocabulary size at the multiple-testing-adjusted genome-wide significance level ($P < 2.10 \times 10^{-8}$, eFigure 2a-

c). For early-phase expressive vocabulary, a single GWAS signal passed the unadjusted genome-wide significance threshold (rs9854781, $P < 5 \times 10^{-8}$), consistent with a known locus (rs764282, LD- $r^2 = 0.78$) identified through a previous meta-GWAS studying overlapping samples⁷. Genome-wide gene-based, gene-set and gene-property analyses with MAGMA⁵⁹ did not provide evidence for gene-based association passing the multiple-testing-adjusted significance thresholds (eFigure 3, eTable 5).

All vocabulary measures were modestly heritable (Figure 2a, eTable 6), with SNP- h^2 estimates of 0.24(SE=0.02), 0.08(SE=0.01), and 0.20(SE=0.04) for early-phase expressive vocabulary, late-phase expressive vocabulary, and late-phase receptive vocabulary, respectively. Given limited data availability, polygenic prediction (out of the meta-analysis samples) was carried out for late-phase expressive vocabulary only ($\beta = 0.04$ (SE=0.04), $P = 0.35$, $R^2 = 0.14\%$), though power was low (≤ 0.11) due to a combination of low SNP- h^2 and low target sample size ($N = 639$). In contrast, genetic correlations between early- and late-phase expressive vocabulary ($r_g = 0.69$ (SE=0.14)), as well as between late-phase expressive and receptive vocabulary ($r_g = 0.67$ (SE=0.16)) were moderate (Figure 2b), suggesting some stability in genetic factors during development. Genetic influences underlying early-phase expressive vocabulary were, however, largely independent of genetic influences related to late-phase receptive vocabulary ($r_g = 0.07$ (SE=0.10)). Given comparable power to detect $r_g = 0.70$ with late-phase receptive vocabulary for both expressive vocabulary measures (power: early-phase=0.83, late-phase=0.71), these findings suggest developmental genetic heterogeneity.

To maximise power for single-variant discovery, genetically correlated vocabulary measures were combined as part of two multi-trait meta-analyses using MTAG (stage II, Figure 1, eTable 7). However, we neither identified further SNP-vocabulary associations (eFigure 2d-e) nor increased evidence for SNP- h^2 (Z-scores, Figure 2a, eTable 6).

Genetic relationships with later-life outcomes

We investigated genetic links between vocabulary traits (stage I) and multiple heritable health-, cognition-, and behaviour-related traits, especially during childhood (see eTable 8 for SNP- h^2) by estimating HDL genetic correlations⁴⁵ (multiple-testing-adjusted threshold: $P < 5.32 \times 10^{-3}$). Consistent with the identified heterogeneous genetic architecture (see above), polygenic association patterns were highly divergent for early-phase and late-phase vocabulary (Figure 3a). Larger early-phase expressive vocabulary size was genetically correlated with increased ADHD risk ($r_g = 0.23$ (SE=0.08)), exposure to childhood maltreatment ($r_g = 0.19$ (SE=0.07)) and, at the nominal level ($P < 0.05$), childhood aggressive behaviour ($r_g = 0.42$ (SE=0.16)). The direction of the association effect reversed, however, in toddlerhood, such that lower late-phase receptive vocabulary size was genetically related to increased childhood maltreatment exposure ($r_g = -0.33$ (SE=0.08)). In addition, genetic links with cognition-related later-life outcomes became detectable in toddlerhood only (Figure 3a), despite comparable power estimates across infant and toddler measures (eTable 9). Both, larger late-phase expressive and receptive vocabulary size were genetically correlated with higher intelligence across the lifespan (late-phase expressive vocabulary: $r_g = 0.32$ (SE=0.08); late-phase receptive vocabulary: $r_g = 0.36$ (SE=0.12)) and higher educational attainment (late-phase expressive vocabulary: $r_g = 0.26$ (SE=0.05); late-phase receptive vocabulary: $r_g = 0.37$ (SE=0.06)).

To integrate genetic covariance patterns into a structural model, we adopted a genomic SEM⁵⁷ approach that was informed by exploratory factor analyses (eTable 10). Specifically, we studied all three early-life vocabulary measures and genetically correlated later-life traits (HDL- r_g - $P < 5.32 \times 10^{-3}$: intelligence, educational attainment, exposure to childhood maltreatment and ADHD). The best-fitting model was a correlated 3-factor model (Figure 3b, eTable 11), implicating two uncorrelated genomic dimensionalities. For comparison with estimates presented above, we report, here, unstandardised factor loadings, representing the direction and strength of association between a trait and model-implied latent genetic factor: The first genetic factor ($F1_g$) reflected shared genetic influences between early-phase

($\lambda=0.25$ (SE=0.09)) and late-phase expressive vocabulary ($\lambda=0.31$ (SE=0.10)). This factor was modestly correlated ($r_g=0.25$ (SE=0.09)) with a second genetic factor ($F2_g$) accounting for the majority of genetic variance in educational attainment ($\lambda=0.33$ (SE=0.01)) and intelligence ($\lambda=0.32$ (SE=0.01)). The third genetic factor ($F3_g$), which was, largely, independent of $F1_g$ ($r_g=-0.09$ (SE=0.09)), captured the entirety of genetic ADHD liability ($\lambda=0.48$ (SE=0.04)) and, to a lesser extent, genetic influences contributing to childhood maltreatment exposure ($\lambda=0.13$ (SE=0.01)). Notably, early-phase expressive ($\lambda=0.16$ (SE=0.12)) and late-phase receptive ($\lambda=-0.19$ (SE=0.04)) vocabulary were associated with this factor with an opposite direction of effect. While, largely, consistent with reported SNP- h^2 estimates and genetic relationships above (Figure 2, 3a), the identified model structure lacked genetic overlap between late-phase expressive and receptive vocabulary (eTable 12), underlining the limitations of the model.

To confirm the change in genetic association pattern with direct genotyping data, we studied children of the ALSPAC cohort with genetic, vocabulary and ADHD symptom information and modelled jointly the phenotypic (r_p), genetic (r_g) and residual (r_e) correlations fitting a saturated (Cholesky) structural model using GRM-SEM⁵⁸ (eMethods 10). Genetic correlations in ALSPAC (eFigure 4) had the same direction of effect as meta-analytic HDL-derived estimates (Figure 3a), underlining the robustness of our findings. However, 95%-confidence intervals (CIs) were wide and analyses have, thus, an exploratory character only. Increased ADHD symptoms were phenotypically very modestly correlated with smaller vocabulary size ($r_p=-0.06$ (SE=0.02)), irrespective of developmental stage (eFigure 4). Thus, during infancy only, genetic correlations ($r_g=0.51$ (SE=0.26)) may have been masked by opposite residual correlations ($r_e=-0.18$ (SE=0.06)).

Discussion

Here, we present findings from genome-wide association meta-analyses of expressive and receptive vocabulary size across infancy and toddlerhood. Genomic covariance analyses revealed heterogeneity within the genetic architecture underlying vocabulary size across different developmental stages that matched distinct polygenic association patterns with cognitive and behaviour-related traits during later life.

Bivariate genetic correlation patterns and structural models suggested two independent genetic factors contributing to early-life vocabulary size, confirming the heterogeneous multivariate genetic architecture observed in previous reports^{7,8,12,13}. Both factors may reflect aetiological differences in vocabulary acquisition stages, given limited genetic overlap between infant expressive and toddler receptive vocabulary size. Genetic influences contributing to utterances in infancy, approximated by early-phase expressive vocabulary size (15-18 months), may capture the first stages of language learning related to emerging speech, where words are usually produced in isolation². During this phase of “learning to speak”, children acquire phonological skills to identify phonemes and sequences from speech and store them for future production⁶⁰, but also develop oral motor⁶¹ and speech motor skills⁶². Despite sufficient power, there was little evidence for genetic overlap between early-phase vocabulary scores and later-life cognition. Such an association only emerged for late-phase vocabulary scores during toddlerhood (24-38 months), in line with previous work¹² and suggested specificity. The genetic overlap with cognition may reflect the onset of a phase of “speaking to learn”, where toddlers have mastered some language fluency and start to use word combinations and more complex grammatical structures^{4,6}. Together, our findings highlight rapid changes in the genetic architecture of vocabulary acquisition across a period of only two years, even when assessed with similar psychological instruments. Shared genetic influences across both developmental phases underline the dynamic character of this process, illustrated by the genetic overlap of late-phase expressive vocabulary with both, early-phase expressive and late-phase receptive vocabulary.

The heterogeneity in genetic components contributing to vocabulary acquisition was further reflected by distinct polygenic association patterns with later-life behaviour and related proxies. Larger infant expressive but lower toddler receptive vocabulary size was genetically correlated with increased ADHD risk and/or childhood maltreatment exposure. Consistently, younger age at first walking, another early developmental milestone, has been linked to higher polygenic ADHD risk⁶³. During a developmental phase of “learning to speak”, potentially involving motor skills that shape children’s learning environment and, in turn, behaviour and language learning⁶⁴, children with a higher genetic predisposition for ADHD may show larger rather than smaller vocabulary size. In contrast, the direction of genetic overlap between ADHD risk and toddler receptive vocabulary is consistent with the known inverse polygenic association pattern of ADHD risk with child/adolescent verbal and cognitive abilities²². Genetic links implicating ADHD and childhood maltreatment exposure were similar, reflecting the genetic trait overlap²⁷.

This work builds on a previous GWAS effort⁷ by increasing the number of studied children by ~50% and adopting a multivariate analysis approach to maximise statistical power while extending the studied phenotypic vocabulary spectrum. However, the power to detect single variant contributions of small effect (e.g. 0.1%) remained low (eMethods 4), especially for receptive vocabulary. Given limited data availability, the study focussed also exclusively on European children and languages. Furthermore, to harmonise vocabulary measures across different developmental stages and instruments, all measures had to be rank-transformed. Finally, although structural models built from genetic summary statistics had an acceptable model fit, the model could not capture all aspects of the underlying multivariate genetic architecture, limiting its interpretation. Future studies may increase the sample size further and boost study power through multivariate analysis of vocabulary with genetically related aspects of language, such as grammatical abilities^{8,9}, preferably in genetically diverse populations.

In summary, there are at least two distinct genetic factors contributing to vocabulary size during infancy and toddlerhood that match distinct polygenic association patterns with later-life outcomes. Our

findings highlight the importance of studying genetic influences underlying early-life vocabulary acquisition to unravel the aetiological processes that shape future behaviour, health and cognition.

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OAA is a consultant to HealthLytix. All other authors declare no conflict of interest.

Data sharing statement

Derived single-trait (stage I) and multi-trait (stage II) vocabulary summary statistics will be made available upon publication of the manuscript via a data repository.

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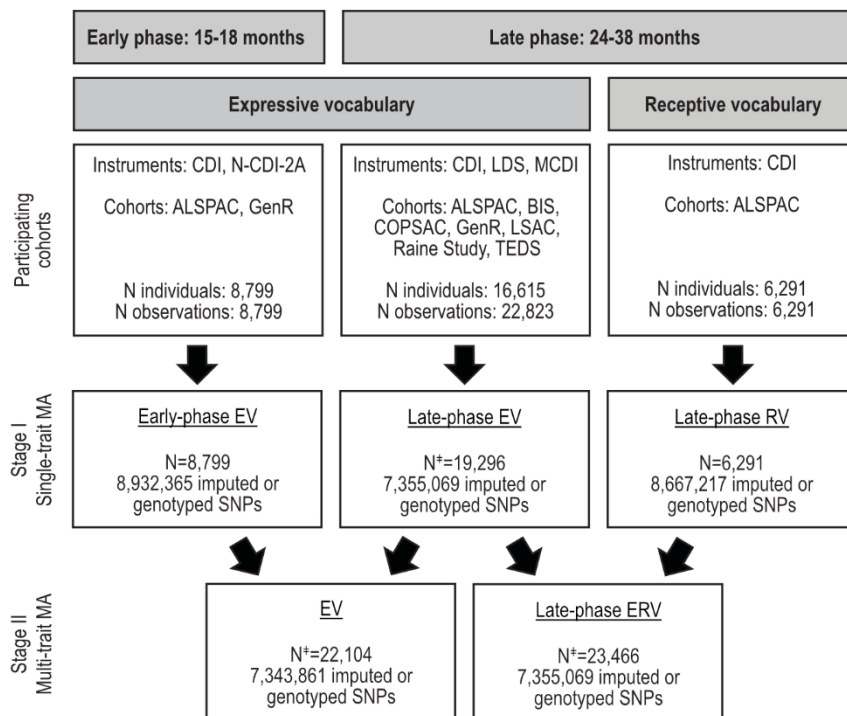


Figure 1: Meta-analysis study design

Vocabulary scores were assessed between 15-38 months of age and divided into an early phase (15-18 months) and late phase (24-38 months) of language acquisition allowing for age-specific genetic influences. Scores for receptive vocabulary were included in the late-phase only. In stage I, three single-trait meta-analyses were conducted: early-phase expressive vocabulary, late-phase expressive vocabulary and late-phase receptive vocabulary. In stage II, multi-trait genome-wide analyses were performed across early-phase and late-phase expressive vocabulary, as well as across late-phase expressive and receptive vocabulary to increase statistical power.

† Estimated sample size based on the increase in mean χ^2 statistic using multi-trait analysis of genome-wide association.

Abbreviations: ALSPAC, Avon Longitudinal Study of Parents and Children; BIS, Barwon Infant Study; CDI, Communicative Development Inventory; COPSAC, Copenhagen Prospective Studies on Asthma in Childhood; EV, expressive vocabulary; ERV, expressive and receptive vocabulary; GenR, Generation R Study; LDS, Language Development Survey; LSAC, Longitudinal Study of Australian Children; MA, meta-analysis; RV, receptive vocabulary; TEDS, Twins Early Development Study

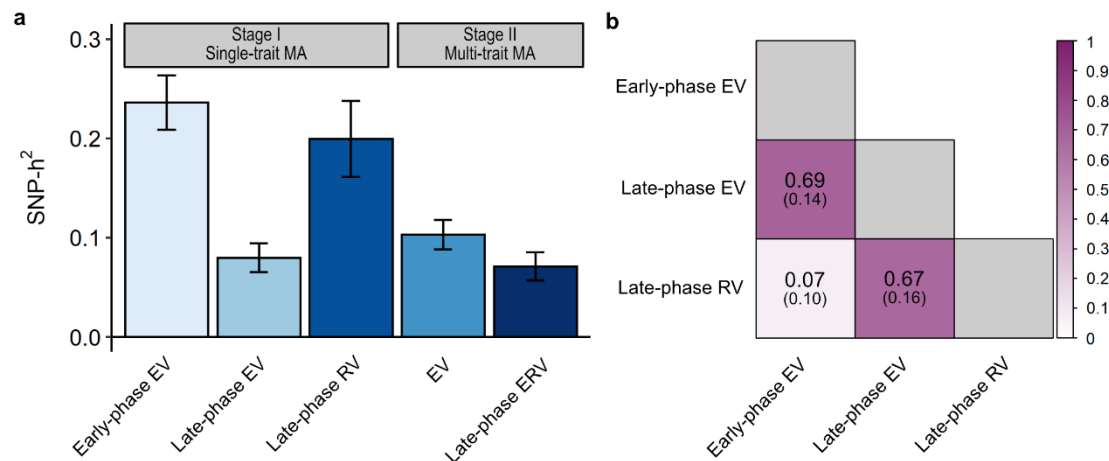


Figure 2: SNP-heritability and genetic correlations of vocabulary traits

(a) SNP-heritability estimates for single- and multi-trait vocabulary summary statistics were estimated with High-Definition Likelihood⁴⁵ software. Error bars represent standard errors. **(b)** Genetic correlations (r_g) among single-trait vocabulary summary statistics were estimated with High-Definition Likelihood⁴⁵ software. Corresponding standard errors are shown in brackets.

Abbreviations: EV, expressive vocabulary; ERV, expressive and receptive vocabulary; MA, meta-analyses; RV, receptive vocabulary; $SNP-h^2$, Single-Nucleotide Polymorphism heritability

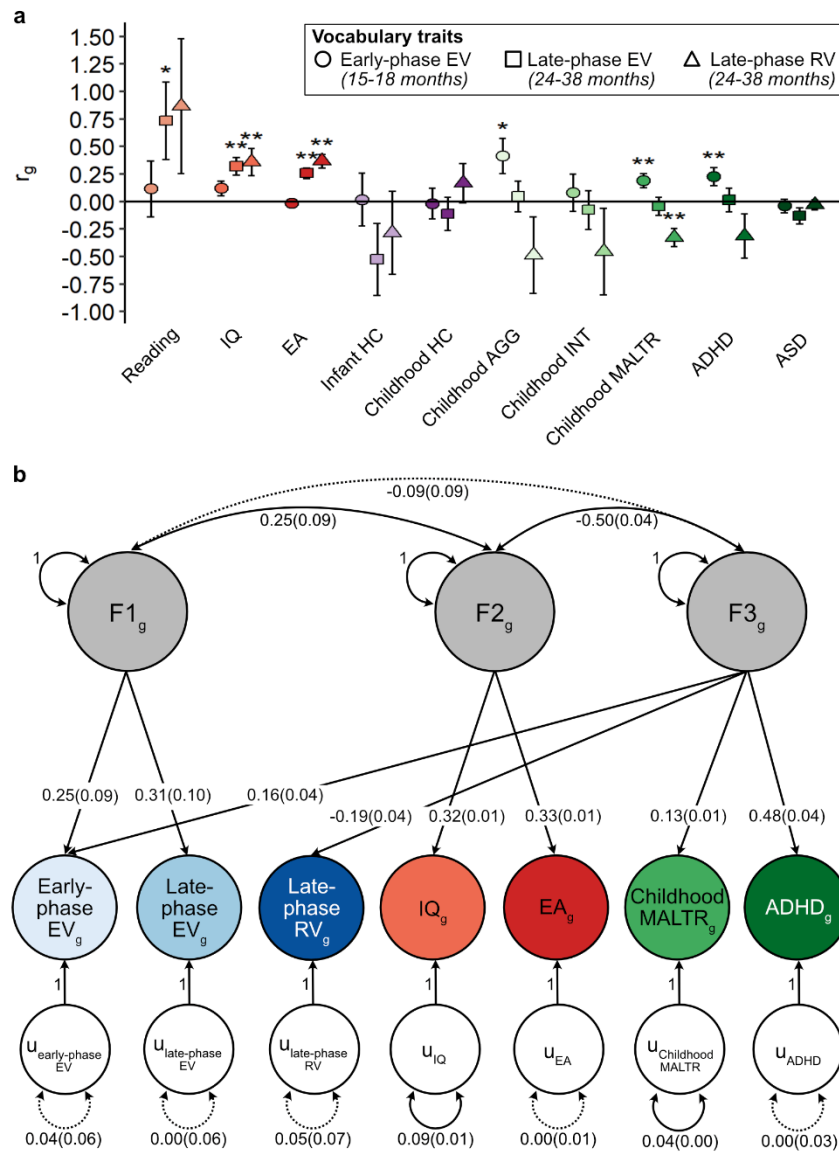


Figure 3: Genetic relationships of vocabulary with several later-life cognitive, health and behavioural outcomes

(a) Genetic correlations (r_g) were estimated using summary statistics and High-Definition Likelihood (HDL)⁴⁵. Bars represent standard errors. ** multiple-testing adjusted $P < 5.32 \times 10^{-3}$; * $P < 0.05$. **(b)** Three-factor model fitted to genetic covariance patterns of early-life vocabulary measures and genetically correlated later-life outcomes (identified with HDL, $P(r_g) < 5.32 \times 10^{-3}$) using Genomic SEM⁵⁷. Solid and dashed arrow lines represent factor loadings with $P < 0.05$ and $P \geq 0.05$, respectively. Unstandardised factor loadings are shown, with standard error in parenthesis. Model fit characteristics are provided in eTable 11.

Abbreviations: ADHD, Attention-Deficit/Hyperactivity Disorder; AGG, aggression; ASD, Autism Spectrum Disorder; EA, educational attainment; EV, expressive vocabulary; HC, head circumference; INT, internalising symptoms; IQ, general intelligence; M, months; MALTR, maltreatment; RV, receptive vocabulary; SDQ, Strengths and Difficulties Questionnaire; Y, years