Factors that mediate the relationships between household socio-economic status and childhood Attention Deficit Hyperactivity Disorder (ADHD) in children and adolescents: A systematic review

Wolfgang A. Markham*, Nicholas Spencer

Division of Health Sciences, Warwick Medical School, University of Warwick, Coventry, United Kingdom
* wolfgang.markham@warwick.ac.uk

Abstract

Background
ADHD is one of the most prevalent mental health disorders among children and adolescents. Household socio-economic status (SES) in early childhood is inversely related to ADHD later in childhood or adolescence. We conducted a systematic review to examine psychological, social and behavioural factors that mediate these relationships (PROSPERO Registration number: CRD42020182832).

Methods and findings
We searched Medline, EMBASE, PsychINFO, and Web of Science from inception until May 2020. Both authors independently reviewed abstracts and identified papers for inclusion. We sought primary observational studies (cohort, cross-sectional and case control studies) of general population-based samples of children and adolescents aged 18 and under that investigated potential mediators of the relationships between SES and ADHD. Studies based upon non-general population-based samples, twins or biochemical/physiological changes were excluded. Direct and indirect effects derived from standard validated mediation analysis were extracted for potential mediators. We assessed risk of bias using a modified NIH tool and synthesised quantitative data without meta-analysis according to the (SWiM) protocol because of heterogeneity between included studies.

Family adversity, paternal and maternal ADHD symptoms, Home Learning Environment, breastfeeding duration and a combined fine motor and language score at age 2 may lie on the SES-ADHD pathway. Evidence concerning the influence of maternal depression/anxiety and adverse parenting was inconsistent across studies. There was no evidence that mother’s health-related behaviour, family characteristics, child’s consumption of fizzy drinks or other developmental characteristics at birth/during infancy lie on the SES-ADHD pathway. Publication bias may have been introduced by our decision not to search grey literature, not to approach study authors and limit the search to the English language.
Conclusions
Evidence for mediation of the SES-ADHD pathway in childhood/adolescence is under-researched. Maternal mental health, family adversity, parenting and health-related behaviours warrant further research based on longitudinal data and employing the most advanced mediation analysis methods.

Introduction
Attention Deficit Hyperactivity Disorder (ADHD) is among the commonest mental health disorders in childhood. The prevalence of children diagnosed with ADHD increased in the USA between 2003 and 2011 [1] and more children were in receipt of prescription drugs for the condition in the United States of America (USA) and western Europe in 2012 compared with 2005/6 [2]. Whether these trends reflect a true increase in prevalence or improved methods of data collection and case ascertainment remains unclear [3,4]. A meta-regression of over 100 studies, conducted in 2017, spanning the globe identified that prevalence rates hover around 5% [5].

The aetiology of ADHD is complex resulting from a range of biological, psychological and social conditions that can act individually or synergistically [6]. The association with household socioeconomic status (SES) is well-established [7]; however, explanations for the association vary from social conditions as causal [8], through reverse causality due to loss of earnings and relationship instability [9] to confounding by genetic factors that play a part in the aetiology of ADHD which may influence SES in indirect ways [10]. In considering the potential causal role of SES, a range of socially related prenatal, perinatal and early childhood risk factors probably interacting with genetic influences have been identified [11].

Russell AE et al. [12] suggest that these socially related risk factors may be on the causal pathway from SES to ADHD and mediate the relationship. Mediators are associated with both the exposure (SES) and the outcome (ADHD) and intervene between them accounting for some or all of the effect of the exposure on the outcome (Fig 1). Confounding variables, while causally related with both exposure and outcome [13], do not lie on the causal pathway between the exposure and the outcome but falsely obscure or accentuate the exposure/outcome relationship [14]. Moreover, treating mediators as confounding variables in regression models may also falsely lead to attenuation or elimination of the effect of the exposure on the outcome. Co-variates are related to the outcome but not the exposure and adjustment aims to improve the precision of the effect estimate [15]. In studies with SES as the exposure variable, adjusting in regression analysis for psychological, social or behaviour related variables which are potential mediators is likely to reduce the direct effect of SES on ADHD [16].

Mediation analysis aims to distinguish the total effect of the exposure on the outcome, the indirect effect of the potential mediator and the direct effect of the exposure. Mediation analysis is illustrated in Fig 1 where \( a = \beta \) coefficient of the path from exposure E to mediator M, \( b = \beta \) coefficient of the path from mediator M to outcome O, and \( c' = \beta \) coefficient of the path from the exposure E to outcome O. The direct effect of exposure E on outcome O is represented by \( c' \) and the indirect effect of mediator M on the path from exposure E to outcome O is the product of the unstandardised or standardised coefficients \( a \times b \) [17].

This systematic review aims to assess published evidence from cohort, cross-sectional and case control studies regarding factors that mediate the SES-ADHD pathway in childhood and adolescence.
Methods

Protocol registration and reporting

We conducted a systematic review according to a protocol that was registered in PROSPERO, an open access registry (registration number: CRD42020182832) [18]. We followed the PRISMA checklist [19] (S2 File) and SWiM (Synthesis Without Meta-Analysis) reporting guidelines [20] when reporting our findings.

Search strategy and selection criteria

A health science librarian developed a search strategy to identify eligible investigations of mediators between household SES in early childhood and ADHD or high scores for hyperactivity/inattention in standard psychometric tests later in childhood or adolescence. We systematically searched Medline, EMBASE, PsychINFO and Web of Science from inception until 1st May 2020. A combination of indexed terms, free text words and MeSH headings were used (S3 File). We also manually searched reference lists of papers identified in the electronic database search including reviews. We did not search grey literature and study investigators were not contacted for unreported data or additional details. Included studies were restricted to the English language and were deduplicated. Both authors independently assessed article abstracts and full texts of studies that passed the initial screening phase. We included primary observational studies (cohort, cross-sectional and case control studies) of general population-based samples of children and adolescents under 19 years of age that used a recognised method for assessing mediation to investigate any psychological, social or behavioural factor that potentially mediated the relationship between household SES and childhood/adolescent ADHD or childhood/adolescent hyperactivity/inattention disorder assessed using standard psychometric tests. We excluded reviews and studies that were based upon non-general population samples, twins, adults over 18 years and studies where the investigated mediating factor was biochemical or physiological e.g. brain morphology or the investigated outcome was externalising behaviour including conduct problems or conduct disorder alone. At each stage disagreements between reviewers were resolved by consensus.
Data analysis

Both authors independently extracted the following information from each included study—country, study type, population, sample size, attrition (%), SES, child’s/adolescent’s age at SES measurement, ADHD or hyperactivity/inattention measure and prevalence, child’s/adolescent’s age at ADHD measurement, mediators studied, mediation analysis method, covariates/confounders included in analyses, direct effects of SES and indirect effects of mediators (pathway coefficients), significant mediators, and mediators that were not significant. For each study, we estimated the proportion mediated derived by dividing indirect effects of mediators by the total effect of SES (indirect + direct effect) on ADHD expressed as a percentage [21]. Investigated psychological, social or behavioural factors that were identified as not meeting criteria for potential mediation because they were not significantly associated with either the exposure or the outcome were also identified. We then independently used a modified version of the NIH assessment tool for observational, cohort and cross-sectional studies [22] to assess the risk of bias (RoB) based on the quality of each included study’s methodology (S1 Table). Additional questions were added to this NIH assessment tool namely:

- Was the study population representative of the whole target population?
- Was the mediation analysis clearly specified and defined?
- Was the choice of mediators clearly specified and justified?
- Were results of the mediation analyses clearly presented allowing direct and indirect effects to be distinguished?

Studies were allocated high, moderate or low RoB based on methodological criteria (S1 Table). We synthesised quantitative data without meta-analysis according to the (SWiM) protocol [20] because of anticipated and confirmed sources of diversity and thus, heterogeneity between studies. These anticipated sources of diversity were:

- statistical diversity i.e. diversity in the methods for identifying the direct and indirect effects of household SES on ADHD and hyperactivity/inattention.
- methodological diversity i.e. included primary studies may be longitudinal studies, cross-sectional studies or case control studies.
- clinical diversity i.e. the outcome could be based upon a medical diagnosis of ADHD or having scores equal to or over the accepted cut-off point for hyperactivity/inattention on standard psychometric tests as reported by doctor, teacher, parent or self-reported.
- diversity in the measures of household SES i.e. based upon income, education, and other accepted measures of SES.
- diversity in the mediating factors investigated
- diversity in the measurement of potential mediators
- diversity in child’s/adolescent’s age at outcome measurement.

The prioritisation of results was informal and based upon RoB assessments i.e. low RoB studies are prioritized over other studies and study design i.e. cohort studies are prioritized over other studies.

Results

We identified n = 1130 citations from bibliographic databases and n = 5 citations from reference lists. After removing duplicates we screened n = 626 titles and abstracts for eligibility.
Reasons for excluding articles at the titles and abstract stage included: not ADHD & SES; non-general population/clinical sample; not ADHD; adult sample; review/opinion. We then assessed n = 82 full text articles and excluded n = 74 articles (Fig 2). Reasons for excluding full text articles included: no mediation analysis; SES was a co-variate not an exposure; the outcome was not ADHD or hyperactivity/inattention using standard psychometric tests; outcome was externalising behaviour including conduct problems or conduct disorder alone; the investigation was based upon a non-general population based sample; review article; the

![Fig 2. Prisma flow chart [19].](https://doi.org/10.1371/journal.pone.0262988.g002)
investigated mediator was biochemical or physiological. Eight papers were identified that examined potential psychological, social or behavioural mediators of the relationship between household SES (exposure) and ADHD or hyperactivity/inattention in childhood/adolescence (outcome).

Characteristics of included studies
The characteristics of included studies are summarized in Table 1. All included studies were conducted in North America (Canada and USA (2)) or Europe (France, Germany, Norway and UK (2)). Five studies were cohort studies, two were cross-sectional studies and one was a case-control study. The populations studied were children aged 3 [23,24], aged 7 [12,25], aged 7–8 [26], aged 7–14 [27], aged 6–17 [28], and adolescents aged 17/18 [29].

Household SES was based upon maternal education when youngest child was 2 months old [24]; financial hardship when child was 0–2 (parent reported difficulty in affording heating, clothing, rent/mortgage, food and/or things for the study child) [12]; family household income at 17/18 [29]; and parental occupation when child was aged 4/5 [26]. Combined SES measures were used in four studies: parental education and household income pre-pregnancy [23]; parental education and household income when child was 7–14 years [27]; parental education, household income and child’s health care insurance status when child was 6–17 years [28]; paternal education, maternal education, fathers’ social class, mothers’ social class at 9 months [25].

The measures of ADHD or hyperactivity/inattention using reliable and valid psychometric tests varied. Miller et al. [27] relied upon teacher-reported ADHD Rating Scale, Version VI (ADHD-RS-IV) [30], the Conners ADHD Rating Scale [31], and the Strengths and Difficulties Questionnaire (SDQ) [32]. Foulon et al. [23] used mother-reported SDQ questionnaires [33,34] and Schmiedeler et al. [26] used teacher-reported SDQ [33]. Meunier et al. [24] utilised the revised Ontario Child Health Study questionnaires [35] and recorded both parents’ assessments. Russell AE et al. [12] used parent and teacher assessments based upon the Development and Well-Being Assessment (DAWBA) questionnaire [36]. Boe et al. [29] used adolescent-reported assessments based upon the WHO adult ADHD self-report ASRS scale [37]. Russell G et al. [25] and Nguyen et al. [28] relied upon parent-reporting of a health care provider diagnosis of ADHD.

The methods for investigating mediation varied between studies. Boe et al. [29], Miller et al. [27] and Nguyen et al. [28] conducted Structural Equation Modelling using Mplus version 7. Schmiedeler et al. [26] used Structural Equation Modelling in AMOS software along with Full Information Maximum Likelihood estimation for latent variable interactions. Foulon et al. [23] employed the MacArthur moderator-mediator path analysis approach [38]. Meunier et al. [24] used Baron and Kenny’s traditional sequential framework for testing direct and indirect effects along with the multilevel modelling plus framework proposed by Edwards and Lambert [39] and Preacher et al. [40]. Two studies [12, 25] drew upon a mediation analysis method that adopts a products of coefficients approach [41].

Two of the included studies [25,29] did not adjust for covariates/confounders. All the other studies adjusted for the covariate, child’s gender. Four studies [24,26–28] also adjusted for the covariate, child’s age. In addition, Meunier et al. [24] adjusted for covariate, sibling gender composition. Schmiedeler et al. [26] adjusted for teacher reported ADHD at age 4 treating it as a covariate. Nguyen et al. [28] adjusted for the covariate, child’s ethnicity, and a potential confounder, diagnosis of conduct problems.

Risk of bias (RoB) assessment
Assessments of the quality of each included study’s methodology are shown in S1 Table. None of the included studies were judged to have a low RoB. Four of the five cohort studies [12,23–
| Author | Year | Country | Study type; population; sample size; Attrition(%) | SES measure/s Age at SES measurement | ADHD measure Age at ADHD measurement Number (%) | Risk of bias | Mediators studied & method | Adjusted for covariates/ confounders | Direct & Indirect effects (95% CI) | Non-significant mediators |
|--------|------|---------|-------------------------------------------------|-------------------------------------|-----------------------------------------------|-------------|---------------------------|-------------------------------------|-------------------------------|-------------------------|
| Boe et al. 2018 [29] | Norway | Cross-sectional population-based study Working sample n = 9151 Attrition = 52.9% Mean age 17.47 years Standard deviation 0.84 years | Income to needs ratio = (Family household income adjusted for size)/(60% median threshold for family household income adjusted for size) | Study participant-reported hyperactivity-Inattention continuous measure based on adult ADHD self-report ASRS scale Proportion with hyperactivity/inattention not reported | High | Structural equation modelling Mplus Version 7.4 Mediator tested: Adolescents perceived economic status (poorer than others, equal to others, better than others) | No | Unstandardised β coefficients Direct Effect: -0.208 – (-0.20 0) = -0.008 (calculated from data) Indirect Effect: Via Perceived economic status -0.200 (-0.253, -0.150) Proportion of total effect -0.208 (95% CI -0.315, -0.095) explained by the indirect pathway (0.200/0.200+0.008) = 9% |
| Foulon et al. 2015 [23] | France | Cohort: EDEN project–women recruited in pregnancy from hospital maternity units in 2 large cities 1311 Attrition 31% | Pre-pregnancy monthly household income, paternal education, maternal education combined into a single measure | Mother-reported SDQ as continuous variable collected when child aged 3 yrs.– mean score = 3.5 Proportion with hyperactivity/inattention not reported | Moderate | MacArthur moderator-mediator path analysis approach proposed by Kraemer et al. (2001) Mediators tested at 4 periods: Before pregnancy; prenatal/birth; infancy; toddlerhood. Foetal exposures Child’s temperament Child’s neurodevelopmental status Psychosocial environment | Child gender | Standardised β coefficients Direct Effect: -0.18 p<0.05 Indirect Effects: Via Breastfeeding duration (0.25 p<0.5, -0.06 p<0.5) Total indirect effect via this pathway β = (0.25 X -0.06) = -0.015 Via Breastfeeding duration and Child neuro-developmental status (Combined score of fine motor score and language score at 2 years) (0.25 p<0.5, 0.08 p<0.5, -0.15 p<0.5) Total indirect effect via this pathway β = (0.25 X 0.08 X -0.15) = -0.003 Via Maternal depression & anxiety combined assessed at 6 month of pregnancy and mother and Infant distress and dysregulation measured at 4-8-12 months (-0.15 p<0.5, 0.32 p<0.5, 0.14 p<0.5) Total indirect effect via this pathway was β = (-0.15 X 0.32 X 0.14) = -0.007 Proportion of total effects explained by the indirect pathways (0.015+0.003+0.007)/(0.015+0.003 +0.007) + 0.18) = 12% |
| Author Year | Country | Study type; population; sample size; Attrition(%) | SES measure/s Age at SES measurement | ADHD measure Age at ADHD measurement Number (%) | Risk of bias | Mediators studied & method | Adjusted for covariates/confounders | Direct & Indirect effects (95% CI) | Non-significant mediators |
|-------------|---------|-------------------------------------------------|--------------------------------------|-------------------------------------------------|-------------|-----------------------------|---------------------------------|---------------------------------|-----------------------------|
| Meunier et al. 2013 [24] Canada | Cohort: 920 children from 397 families with 2 or > children < 4 years Attrition = 20.8% | SES measure reported when youngest child was 2 months old: Number of years maternal education completed | Mean ADHD score measured by scale with well-established reliability and validity completed by both parents at 3 yrs. | Moderate | Baron and Kenny sequential framework and multilevel modelling plus framework proposed by Edwards and Lambert (2008). Mediators tested: Mother reported differential negativity Mother reported differential positivity Observed differential negativity in home Observed differential positivity measured in home | Age Child genderSibling gender composition | Standardised β coefficients in separate single mediating risk factor models Positive and negative differential parenting included in separate models Via Observed differential negativity Direct Effect: -0.17 (p<0.001) Indirect Effect: -0.016 (p < .05). Via Mother reported Differential positivity Direct effect: -0.16 (p<0.001) Indirect Effect: -0.015 (p < .05) Proportion of total effects explained by the indirect pathways Via Observed differential negativity (-0.016/(-0.17) + (-0.016)) = 9% Via mother-reported differential positivity (-0.015/(-0.015) + (0.16) = 9% | Mother reported Differential negativity Direct Effect: -0.18 (p<0.001) Indirect Effect: 0.002 |
| Miller et al. 2016 [27] USA | Case-control study: n = 931 children 7–14 years Attrition N/A | A latent SES construct created from: Highest parental education, highest parental occupation, Family household income Mean age of child = 9.3 years | Teacher ratings of hyperactivity/inattention ADHD 521 Controls 335 Sub-threshold 75 | High | Structural Equation Modelling using Mplus 7.4 Mediators tested: Self-rated or spouse rated Paternal ADHD symptoms Self-rated or spouse rated Maternal ADHD symptoms | Age Child gender | Unstandardised β coefficients Direct Effect: -0.09 (p<0.05) Indirect Effects: Paternal ADHD symptoms -0.06 (p<0.001) Maternal ADHD symptoms -0.05 (p<0.01) Proportion of total effects explained by the indirect pathways (0.05 +0.06)/(0.05 +0.06) +0.11) = 55% | (Continued) |
| Author               | Year           | Country | Study type; population; sample size; Attrition(%) | SES measure/s Age at SES measurement | ADHD measure Age at ADHD measurement Number (%) | Risk of bias | Mediators studied & method | Adjusted for covariates/ confounders | Direct & Indirect effects (95% CI) | Non-significant mediators |
|---------------------|----------------|---------|---------------------------------------------------|--------------------------------------|-------------------------------------------------|-------------|---------------------------|-------------------------------------|----------------------------------|-----------------------------|
| Nguyen MN et al. 2019 [28] | USA           | Cross-sectional study: n = 65680 children 6–17 yrs. Mean age and standard deviation not reported Attrition N/A | SES was a latent variable made up of: Household income; Parent education; Parent employment; Child’s health care insurance status | Parent report of a health care provider diagnosis of ADHD based upon 6560 (10% weighted) | High | Structural Equation Modelling using Mplus Mediators tested: Adverse Childhood Experiences (ACE) School engagement Neighbourhood safety Neighbourhood amenities | Age Child gender Child ethnicity Diagnosis of conduct problems. | Standardised β coefficients Direct Effect: No direct effect of SES on ADHD Indirect Effects: Total indirect effect (as reported in article) (β = − 0.03; p = 0.002) mostly via ACE and school engagement but also included neighbourhood safety in model Specific significant indirect effects SES to safety to school engaged to ACE to ADHD (β = − 0.08; p < 0.001) SES to safety to ACE to school engaged to ADHD (β = − 0.01; p < 0.001) Proportion of total effects explained by the indirect pathways = 100% | Neighbourhood amenities |
| Russell AE et al. 2015 [12] | UK            | ALSPAC Birth cohort n = 8132 Attrition 45% | Main SES measure in analysis Financial Hardship when child was 0–2. | Parent/carer and teachers reported ADHD based on DAWBA 7 years 172 (2.1%) | Moderate | Multiple mediation analysis method that adopts a products of coefficients approach using the products of coefficients approach (Preacher and Hayes, 2008) Mediators tested: Maternal involvement Paternal involvement Parental psychopathology (maternal depression) Child fizzy drinks consumption at 3 years Family adversity (Rutter score) | Child gender | Unstandardised β coefficients Direct Effect: 0.113(0.03,0.19) Indirect Effects: Via Mother involved 0.003(0.000– 0.009) Via Partner involved 0.008 (0.001,0.015) Via Family adversity 0.028(0.012,0.050) Proportion of total effects explained by the indirect pathways (0.003+0.008+0.028)/(0.003+0.008 +0.028 + 0.113) = 26% | Maternal depression |
| Author Year | Country | Study type; population; sample size; Attrition(%) | SES measure/s Age at SES measurement | ADHD measure Age at ADHD measurement Number (%) | Risk of bias | Mediators studied & method | Adjusted for covariates/ confounders | Direct & Indirect effects (95% CI) | Non-significant mediators |
|-------------|---------|-------------------------------------------------|--------------------------------------|-----------------------------------------------|-------------|---------------------------|----------------------------------|-------------------------------------|-----------------------------|
| Russell G et al. 2014 [25] UK | | Cohort: UK-wide Millennium Cohort Study at 9 months 13305 Attrition 31.8% | SES Index based upon Fathers' social class, mothers' social class, paternal education, maternal education all measured at 9 months | Parent report of ADHD diagnosis by health professional at any time up to 7 years of age 187(weighted % 1.5) | Moderate | Multiple mediation analysis method that adopts a products of coefficients approach using the products of coefficients approach (Preacher and Hayes, 2008) Mediators tested: Smoking in pregnancy Family conflict/distant parenting | No | Unstandardised β coefficients Direct Effect: 0.108 (0.003,0.205) p<0.05 Indirect Effect: Via Family conflict/distant parenting 0.045 (0.032,0.056) p<0.05 Proportion of total effects explained by the indirect pathway (0.45/(0.45+1.08)= 29% | | |
| Schmiedeler et al. 2014 [26] Germany | | Cohort: children attending mean age 4 yrs. n = 468 Attrition 49.4% Imputation used & final sample 924 | Wegener prestige scale for parental occupation assessed 4–5.25 yrs. States SES was assessed during kindergarten period (T1-T3), T1 mean age was 4 and T3 which was approximately 14 months later | Teacher report of SDQ at ages 7&8 | High | Structural Equation Modelling in AMOS software along with Full Information Maximum Likelihood estimation for latent variable interactions Mediators tested: Home learning environment TV exposure | Age Child gender Teacher-reported ADHD at mean age 4. Proportion with ADHD 5.6% (teachers’ reports) | Unstandardised β coefficients Direct Effect: No direct effect of SES on hyperactivity/inattention Indirect effect: Via Home Learning Environment 0.08 p<0.05 Proportion of total effects explained by the indirect pathway = 100% | TV exposure |
had a moderate RoB. The single cohort study to be rated as high RoB [26] overcontrolled for SES. The other studies had a high RoB primarily because their design prevented assessments of the direction of causality as they were either cross-sectional studies [28,29] or a case control study [27].

Additional factors contributing to the RoB assessments included: not adjusting for covariates/confounders [25,29]; not clearly defining exposure [12]; non-representative sample [23,24,27]; high attrition rate (>20%) in all the cohort studies except for Schmiedeler et al. [26]; conflation between the exposure and the mediator [29] and overcontrolling for SES [26,28].

Potential mediators of the SES-ADHD pathway in childhood and adolescence

Table 1 shows the significant and non-significant indirect effects that were investigated and Table 2 highlights the potential mediators that were investigated but did not meet the criteria for potential mediation as they were not associated with the exposure and/or the outcome.

Table 2. Factors identified as not meeting criteria for potential mediation.

| Factors Identified as not meeting criteria for potential mediation |
|---------------------------------------------------------------|
| **Foulon et al. 2015 [23]** |
| **Mother’s health-related behaviour** |
| Mean number of alcohol glasses/week (Measured at first trimester and third trimester during pregnancy) |
| Cannabis consumption (During pregnancy) |
| Maternal psychoactive drugs intake (When baby 4–12 months old and when baby 24 months old) |
| **Mother’s psychological well being** |
| Maternal history of hospitalisation in psychiatry (pre-pregnancy) |
| Psychiatrist or psychologist consultation in the year before pregnancy (pre-pregnancy) |
| Number of psychiatrist or psychological consultations (When baby 4–12 months old and when baby 24 months old) |
| **Mother’s characteristics** |
| Maternal age at first child |
| **Baby’s characteristics at birth** |
| Birth weight, Gestational age at delivery, Apgar score at 5 minutes |
| Child required resuscitation at birth |
| **Baby/Infant/characteristics** |
| Baby unpredictable (When baby 4–12 months old) |
| Baby Inadaptable (When baby 4–12 months old) |
| Baby Dull (When baby 4–12 months old) |
| Child gross motor (When baby 24 months old) |
| **Family life** |
| Number of children with whom the child is cared (When baby 4–12 months old) |
| Number of stressful life events (When baby 4–12 months old) |
| Number of siblings (When baby 4–12 months old) |
| Parents living together (When baby 4–12 months old and when baby 24 months old) |
| Paternal involvement (When baby 4–12 months old) |
| Maternal child care (When baby 24 months old) |
| **Meunier et al. 2013 [24]** |
| **Family life** |
| Observed maternal differential positivity |
| **Russell AE et al. 2015 [12]** |
| **Mother’s health-related behaviour** |
| Substance use (use of hard drugs or alcohol consumption of more than 3 glasses a day for more than 10 days) (At age 2–4) |
| **Infant/Toddler health-related behaviour** |
| Fizzy drinks/caffeine consumption at age 3 years |
| **Family life** |
| Partner cruelty (physical or emotional) (At age 2–4) |

https://doi.org/10.1371/journal.pone.0262988.t002
Parental psychological factors

Maternal depression. Foulon et al. [23] showed that one of the significant mediating pathways between pre-pregnancy household SES and inattention-hyperactivity at age 3 had two steps. The first step was via a combined perinatal maternal depression and anxiety factor that was based on scores from the Center for Epidemiologic Studies Depression Scale (CES-D) and the State Trait Inventory Anxiety (STAI) [42,43] (standardised β coefficient for indirect effect = -0.15). The second step was via impaired mother-child relationships based on post-partum depression symptoms [44] and infant’s difficult temperament (standardised β coefficients for indirect effects = 0.32, 0.14). Thus, the total indirect effect via this pathway was β = (-0.15 X 0.32 X 0.14 = -0.007). However, other factors related to maternal psychological well-being were found not to meet the criteria for potential mediation. These factors included pre-pregnancy maternal history of psychiatric hospitalisation, psychiatrist or psychologist consultation in the year before pregnancy and number of psychiatrist or psychological consultations when the baby was 4–12 months old and 24 months old.

Russell AE et al. [12] used a different method for assessing mediation than Foulon et al. [23] and reported in their cohort study that maternal depression was not a significant mediating factor in the relationship between household SES and ADHD at age 7. In contrast to Foulon et al.’s 2-step pathway, Russell AE et al. [12] assessed maternal depression based upon a score of 13 or more on the Edinburgh Postnatal Depression Scale [45] measured when the child was 2 years and 9 months of age.

Parental ADHD symptoms

Miller et al. [27] concluded that both paternal and maternal ADHD symptoms were significant mediating factors between household SES and ADHD among 7–14 year olds (β coefficient for indirect effect: Paternal ADHD symptoms −0.06 (p<0.001); Maternal ADHD symptoms −0.05 (p<0.01)). Their measure of parental ADHD was based upon self-reported/spouse-reported current and recalled ADHD symptoms using the Conners Adult ADHD rating scale (CAARS) ADHD index [46] and the Barkley Adult ADHD rating scale (BAARS) [47]. Parental ADHD is likely to precede both household SES and the child’s ADHD. The temporal relationship of parental ADHD and its relationship with the child’s ADHD suggests it is a confounder although it is theoretically possible for parental ADHD to be both a confounder and a mediator [48]. However, Miller et al. [27] acknowledge that parental ADHD “could statistically or mechanistically explain both social disadvantage (due to downward drift) and the child’s ADHD” (p.2) which suggests treating parental ADHD as a mediator rather than a confounder is problematic.

Parenting

Issues related to adverse parenting were investigated as potential mediating factors between household SES and childhood/adolescent ADHD or high scores for hyperactivity/inattention in four studies.

Russell G et al. [25] observed that family conflict/attachment, based upon the Child-Parent Relationship Scale [49] and measured when the child was 3 years old, was a significant mediating factor between household SES and ADHD when the child was 7 years old (β coefficient for indirect effect: 0.045 (95% CI (0.032,0.056)).

Russell AE et al. [12] identified that both maternal and paternal involvement/engagement in activities with their child at 6 years of age were significant mediators of the relationship between household SES and ADHD when the child was aged 7 (β coefficient for indirect effect: Mother involved 0.003 (95% CI 0.000–0.009); Partner involved 0.008 (95% CI 0.001,0.015)).
However, these findings did not echo the findings of Foulon et al. (2014) [23] who reported that both paternal involvement when the baby was 4–12 months old and maternal child care when the child was 24 months old did not meet the criteria for mediation between household SES and childhood inattention-hyperactivity when the child was aged 3.

**Differential parenting.** Differential parenting (favourable/unfavourable parental treatment of one sibling compared to another) was investigated as a potential mediating factor between household SES and ADHD when the child was aged 3 by Meunier et al. [24]. Observed differential negativity [50] and mother-reported differential positivity [51,52] were significant mediators of this relationship ($\beta$ coefficients for indirect effect respectively: -0.016 ($p < .05$) and -0.015 ($p < .05$)) in separate single mediating risk factor models. Their study did however, report contradictory findings as mother-reported differential negativity [51,52] was not significant and observed maternal differential positivity [53] did not meet the criteria for potential mediation. These contradictory findings make the findings regarding parental differential negativity and positivity, in families with more than one child, [24] difficult to interpret.

**Family life**

**Home Learning Environment.** Schmiedeler et al. [26] found that the relationship between household SES and hyperactivity-inattention when children were aged 7 and 8 was fully mediated by Home Learning Environment ($\beta$ coefficient for indirect effect: -0.08 $p<0.05$) as the direct effect of household SES was non-significant. Home Learning Environment was measured by 11 questions including parents reading to their children, possessing books and daily newspapers, playing dice games with their children and owning a library card and visiting the library. High level of television viewing did not mediate the SES ADHD relationship. Television viewing (TV) focused on how many hours the child watched TV per day and how many hours the parent watched TV per day. However, Schmiedeler et al. [26] overcontrolled for SES by including as a covariate ADHD at a younger age.

**General family life characteristics.** Foulon et al. [23] reported that a number of other factors related to family life were not potential mediators between household SES and hyperactivity-inattention of 3 year old children as they did not meet the criteria for potential mediation. These factors included the number of children cared for when the child was 4–12 months old, number of siblings when the child was 4–12 months old and parents living together when the child was 4–12 months old and 24 months old.

**Family adversity including financial stress**

Russell AE et al. [12] reported that family adversity when the child was 2–4 years was a significant mediating factor between household SES and ADHD at age 7 ($\beta$ coefficient for indirect effect: 0.028 (95% CI (0.012,0.050)). Their family adversity index [54] was based on Rutter’s original indicators of adversity [55] and included exposure to the following factors, lack of partner affection, partner cruelty (physical or emotional), family major problems, psychopathology of mother, substance use and trouble with the police. The authors also investigated partner cruelty and substance use on their own as potential mediators but neither of these factors on their own met the criteria for potential mediation. Stressful life events are more commonly experienced by low SES households [56]. However, Foulon et al. [23] reported that the number of stressful life events when the baby was 4–12 months old did not meet the criteria for potential mediation.

Nguyen et al. [28] reported that household SES had no direct effect on ADHD when the child was aged between 6 and 17 years after accounting for indirect effects of multiple mediators (Adverse Childhood Experiences (ACEs), school engagement, neighbourhood safety and
neighbourhood amenities). Household SES was a latent variable in their Structural Equation Model made up of household income, parent education, parent employment and child’s health care insurance status. Nonetheless, household SES did have an indirect effect on ADHD when the child was aged between 6 and 17 years mostly via ACEs and school engagement (Total indirect effect standardised \( \beta \) coefficient = -0.03; \( p = 0.002 \)). ACE was represented by nine hardships including financial stress, having lived with divorced/separated parent, lived with a parent who died or served time in jail, having witnessed domestic violence, having been a victim or witnessed violence in the neighbourhood, having lived with someone with mental health problems or a substance use problems and having experienced racial discrimination. School engagement focused on caring about doing well at school and doing all the required homework. Neighbourhood safety focussed on feeling safe in their neighbourhood and feeling safe at school. Neighbourhood amenities focussed on having pavements, a park, a recreational centre or a library. However, these authors over controlled for SES in the mediation analysis because ACEs included financial stress which is a recognised measure of SES. [12] This is likely to diminish the direct effect of SES.

**Economic factors.** Boe et al. [29] concluded that adolescents’ perceived economic status at mean age 17.5 years was a significant mediating factor between household SES based upon adjusted family household income and adolescent hyperactivity/inattention (\( \beta \) coefficient for indirect effect: -0.200 (95% CI (-0.253, -0.150))). However, adolescents’ perceived economic status and household parental SES are likely to be conflated. Moreover, the temporal relationship between adolescent perceived economic status and ADHD in the cross-sectional study by Boe et al. [29] is unclear. The assumption in this study is that adolescents’ perceived economic status precedes hyperactivity/inattention measured once at 17.5 years; however, the natural history of hyperactivity/inattention which commonly starts in early childhood suggests the reverse i.e. the condition in these adolescents would precede their perception of economic status.

**Behavioural factors that mediate the relationship between household SES and childhood/adolescent ADHD or hyperactivity/inattention**

Breastfeeding duration during the period (4-8-12 months) was reported by Foulon et al. [23] to mediate the relationship between pre-pregnancy household SES and childhood inattention/hyperactivity at age three via two pathways. The first one-step pathway was via breastfeeding duration only (standardised \( \beta \) coefficients for indirect effect = 0.25 and -0.06; Total indirect effect via this pathway \( \beta = -0.015 \)). The second two-step pathway was via breastfeeding duration during the period (4-8-12 months) (Step 1) (standardised \( \beta \) coefficients for indirect effect = 0.25,) followed by child neuro-developmental status (combined score of fine motor and language score at 2 years) (Step 2) (standardised \( \beta \) coefficient for indirect effects = 0.08, -0.15). Thus, the total indirect effect via this pathway \( \beta = (0.25 \times 0.08 \times -0.15) = -0.003 \).

Two studies [23,25] reported that smoking during pregnancy was not a significant mediator of the SES-ADHD relationship. Foulon et al. [23] also reported that other health-related behaviours did not meet the criteria for potential mediation such as mean number of alcohol glasses/week (measured at first and third trimester during pregnancy), cannabis consumption during pregnancy, maternal psychoactive drugs intake when the baby was 4–12 months old and 24 months old. Moreover, Russell AE et al. (2015) [12] found that substance use (use of hard drugs or alcohol consumption of more than 3 glasses a day for more than 10 days) during the period that the baby was aged 2–4 did not meet the criteria for potential mediation. This study also found that the child’s consumption of fizzy drinks/caffeine when it was 3 years old did not meet the criteria for potential mediation.
Socio-biological characteristics of mother and baby that mediate the relationship between household SES and childhood/adolescent ADHD or hyperactivity/inattention

As highlighted above, Foulon et al [23] observed a two-step pathway between household SES and childhood inattention-hyperactivity that focussed on breastfeeding duration (Step 1) and child neuro-developmental status (combined score of fine motor and language score at 2 years). However, other developmental characteristics investigated by the authors did not meet the criteria for potential mediation including the baby being unpredictable when it was 4–12 months old, the baby being unadaptable when it was 4–12 months old, the baby being dull when it was 4–12 months old and gross motor score when the child was 24 months old [57].

These authors also reported that mother’s age at birth was not a significant mediator of the relationship between household SES and childhood inattention-hyperactivity. Additionally, mother’s age when she had her first child and baby’s birth weight, gestational age, Apgar Score at 5 minutes and requiring resuscitation at birth did not meet the criteria for potential mediation.

Direct effects of household SES on child/adolescent ADHD or hyperactivity/inattention

The proportion of the total effects of household SES on child/adolescent ADHD or hyperactivity/inattention that was mediated varied between studies. The proportions mediated were smaller in the four cohort studies we assessed as having a moderate RoB than in the studies we assessed as having a high RoB. In the moderate RoB cohort studies, mediation accounted for 9% of the total SES effect in both of the separate models of Meunier et al. [24], 12% in Foulon et al. [23], 26% in Russell AE et al. [12] and 29% in Russell G et al. [25]. In the high RoB studies, mediation accounted for 100% of the total SES effect in two studies [26,28], 96% in one [29] and 55% in another [27] (Table 1).

Meunier et. [24] reported direct effects of household SES on hyperactivity/attention among 3 year old children of -0.17 (p<0.001) and -0.16 (p<0.001).

Foulon et al. [23] reported a direct effect of household SES of -0.18 (standardised β coefficient) (p<0.05) on ADHD among 3 year olds.

Russell AE et al. [12] found a direct effect of household SES on ADHD among 7 year olds of 0.113 (95% CI (0.03,0.19)) after accounting for the effects of maternal depression, family adversity, mother involvement and father involvement as potential mediators.

Russell G et al. [25] reported a direct effect of household SES on ADHD among 7 year olds of 0.108 (95% CI (0.003,0.205).

Regarding the four studies we assessed as having a high RoB, Boe et al. [29] do not report the direct effect of objective household SES. However, calculation from study data identified a small direct effect of household SES (0.008) on self-reported hyperactivity/inattention among adolescents aged 17/18 years. The only case-control study we included in this review [27] reported a relatively small direct effect of household SES on teacher-rated hyperactivity/inattention among 7–14 year olds of -0.09 (p<0.05).

Two studies we assessed as having a high RoB [26,28] reported no direct effect of household SES on ADHD or hyperactivity/inattention but both of these studies overcontrolled for household SES. Nguyen et al. [28] focussed on parent-reported health care provider diagnosis of ADHD when the child was aged between 6 and 17 years. Schmiedeler et al. [26] focussed on teacher-reported hyperactivity/inattention among 7–8 year olds.
Discussion

Our review shows that evidence for mediation of the pathway between household SES and ADHD in childhood/adolescence is sparse and under-researched. We only identified eight studies that met the inclusion criteria which examined a range of psychological, social and behavioural risk factors as potential mediators. There were no studies of child populations outside North America and northern Europe. We synthesised the quantitative data using the SWiM guidelines for narrative synthesis without meta-analysis as diversity of mediators studied, study designs, population samples, exposure and outcome measures, and study methods precluded meta-analysis. For example, maternal depression was measured differently at different times in the foetal/infant life course and potential mediation was tested by different methods.

Main findings

When indirect effects of mediators were accounted for in the four cohort studies we assessed as having a moderate RoB, the direct effects of household SES on ADHD were robust with proportions mediated less than 30%. These results are likely to have greater validity than those from the high RoB studies which reported mediation of all or a high proportion of the household SES effect.

The review found supporting evidence for mediation of the SES-ADHD pathway by parenting behaviours, including parental conflict/attachment [25], parental engagement [12], parental differential negativity and positivity [24], and Home Learning Environment [26]. Breast feeding was the only health behaviour shown to mediate the SES-ADHD pathway. [23] One study [12] reported mediation by maternal anxiety and depression, known risk factors for ADHD in children [58]; however, a study using a different methodology did not support mediation. [23] Mediation by Adverse Childhood Experiences (ACEs), which are strongly correlated with SES [59], was reported by two studies [12,28] but a further study did not support mediation [23].

The review found no evidence of a mediating role for smoking in pregnancy, alcohol and/or cannabis consumption during pregnancy, maternal substance abuse of hard or psychoactive drugs during the child’s early years and child’s consumption of fizzy drinks. Known socio-biological risk factors for ADHD in children [11], including maternal age at the child’s birth and the birth of her first child, the baby’s birth weight, gestational age at delivery, and Apgar score at 5 minutes, all failed to meet the criteria for potential mediation [23].

Methodological limitations of the studies included in this review. The included papers had substantial methodological limitations. An essential prerequisite of mediation analysis is that the exposure precedes the outcome and potential mediators temporally lie between the exposure and the outcome. These temporal relationships are verifiable in longitudinal studies but are more difficult to verify in cross-sectional and case-control studies. Nguyen et al. [28] acknowledge this limitation but suggest, without supporting evidence, that using SEM in their study allows directionality of the variables to be examined in cross-sectional data. The temporal relationship between ADHD and both adolescent perceived economic status and parental ADHD makes interpretation of the findings of respectively Boe et al. [29] Miller et al. [27] problematic.

All the included studies had limitations related to study samples. Foulon et al. [23] recruited pregnant women from hospital maternity units in two large French cities and it is not clear if this hospital-based sample is representative of the target population. The cohort recruited by Schmiedeler et al. [26] was embedded in a national longitudinal study but the representativeness of the sub-sample is unclear. Meunier et al. [24] excluded families with a single child from
their cohort sample and Miller et al. [27] recruited both their cases and controls using community mailout lists and public advertisements which were unlikely to be representative of the target population.

Non-participation rates exceeded 50% in three studies [24,28,29] and were not reported in four studies [12,25–27] potentially introducing significant selection bias. Attrition is a universal problem in cohort studies. Sample weighting and imputation can be used to minimise the bias associated with differential loss to follow by socially related factors. [60] Attrition rates in all the included cohorts exceeded 20% and only one study [25] reported using weighting for analysis of the association of SES with ADHD but used unweighted data in the mediation analysis. The authors justify the use of unweighted data citing evidence that unweighted regression models are often robust in large datasets [61].

Two studies [25,29] did not adjust for covariates/confounders introducing a source of potential bias as confounders may falsely obscure or accentuate the exposure/outcome relationship. Two studies introduced socially related variables which are likely to have resulted in over-controlling for SES and reducing the total and direct effects of SES on ADHD [26,28].

**Validity.** Mediation by parenting behaviour reported by two moderate RoB studies [12,25] are more likely to be valid finding. The findings that maternal health-related behaviours in pregnancy or in the child’s first three years of life and peri-natal sociobiological factors either did not mediate the SES-ADHD pathway or failed to meet the criteria for mediation are likely to be valid as they are reported by moderate RoB studies [12,23]. The finding that the Home Learning Environment mediates the SES-ADHD pathway [26] is open to question as the study carries a high RoB and it is not clear if the measure of Home Learning Environment has been properly validated. Failure to include evidence of the validity of the methods used to derive adolescents’ perceived SES in Boe et al [29] and school engagement and neighborhood safety/amenities in Nguyen et al [28], also threatens the validity of these study results.

**Methods of mediation analysis.** All the included studies employed recognized, valid mediation analysis methods; however, these methods have limitations as they do not take account of models with interactions and non-linearities [62]. As a consequence, the methods used in the included papers may be subject to bias due to the mediator being affected by the exposure which, in turn, confounds the relationship between the mediator and the outcome—the exposure-induced mediator-outcome confounder effect. This effect becomes more likely the longer the period between the exposure and the outcome [62] suggesting that the cohort studies included in the review are susceptible to this limitation.

**Strengths and limitations of the systematic review**
The review protocol was registered in the PROSPERO review registry following revisions. We

- employed a robust method of risk of bias assessment modified to include potential sources of bias specific to studies of mediation
- only included studies in which the outcome was ADHD or hyperactivity/inattention as externalizing behaviour in standard psychometric tests includes conduct disorder which has a different relationship with SES
- followed the methodologically robust SWiM protocol [20] for narrative synthesis of the included studies

Grey literature databases were not included in the search strategy and additional studies may have been missed. However, this is a narrow field of academic interest and studies meeting the inclusion criteria are very likely to have been published in international peer-reviewed
journals. Publication bias may have been increased by our decision not to approach study authors and limit the search to English language publication. Exclusion of studies examining potential biochemical and physiological mediators from the review may have limited assessment of the full range of mediators reported in the literature; however, we sought to identify modifiable psychological, social and behavioural mediators and factors such as brain morphology are unlikely to be modifiable.

**Implications for future research and policy development**

Future research should be based on methodologically robust longitudinal studies with comparable measures of psychological, social and behavioural factors which meet the criteria for mediation. Maternal mental health, family adversity, parenting and health-related behaviours warrant further research. Studies will need to be sufficient in number and quality to enable meta-analysis to estimate robust pooled indirect effects of potential mediators on the SES-ADHD pathway. Studies should employ the most advanced mediation analysis methods which account for potential exposure-induced mediator outcome confounding [62]. More robust, methodologically sound research is important for future policy development aimed at minimising the link between low household SES in early childhood and later ADHD.

**Conclusions**

A range of psychological, social and behavioural risk factors have been studied as potential mediators of the SES-ADHD pathway; however, reliable conclusions on their effects on the pathway are limited by the small number of studies combined with the moderate to high risk of bias in these studies and diversity of study design, mediators studied and measurement SES and ADHD. However, no evidence of effect is not the same as evidence of no effect. Hence, we propose maternal mental health, family adversity, parenting and health-related behaviours warrant further research.

**Supporting information**

S1 Table. Risk of bias (quality) assessment.

(SDOCX)

S1 File. Prospero systematic review protocol (registration no. CRD42020182832).

(SDOCX)

S2 File. PRISMA-2009 checklist.

(SDOC)

S3 File. Search strategies for electronic databases.

(SDOCX)

**Acknowledgments**

We are most grateful to Mrs Samantha Johnson, Senior Health Science Librarian, University of Warwick for her expert advice and assistance in preparing the search strategy and running the literature searches. We are also grateful to Claire New for administrative support.

**Author Contributions**

**Conceptualization:** Wolfgang A. Markham, Nicholas Spencer.

**Data curation:** Nicholas Spencer.
Formal analysis: Wolfgang A. Markham, Nicholas Spencer.
Investigation: Wolfgang A. Markham, Nicholas Spencer.
Writing – original draft: Wolfgang A. Markham.
Writing – review & editing: Wolfgang A. Markham, Nicholas Spencer.

References

1. Xu G, Strathearn L, Liu B, Yang B, Bao W. Twenty-Year Trends in Diagnosed Attention-Deficit/Hyperactivity Disorder Among US Children and Adolescents, 1997–2016. JAMA Netw Open. 2018 Aug 3; 1 (4):e181471. https://doi.org/10.1001/jamanetworkopen.2018.1471 PMID: 30646132

2. Bachmann C, Wijlaars L, Kalverdijk L, Burcu M, Glaeske G, Schuling-Veninga CCM, et al. Trends in ADHD medication use in children and adolescents in five western countries, 2005–2012. Eur Neuropsychopharmacol. 2017 May; 27(5):484–493. https://doi.org/10.1016/j.euroneuro.2017.03.002 Epub 2017 Mar 20. PMID: 28336088

3. Davidovich M, Koren G, Fund N, Shrem M, Porath A. Challenges in defining the rates of ADHD diagnosis and treatment: trends over the last decade. BMC Pediatr. 2017 Dec 29; 17(1):218. https://doi.org/10.1186/s12887-017-0971-0 PMID: 29284437

4. Thomas R, Sanders S, Doust J, Beller E, Glasziou P. Prevalence of attention deficit/hyperactivity disorder: a systematic review and meta-analysis. Pediatrics. 2015 Apr; 135(4):e994–1001. https://doi.org/10.1542/peds.2014-3482 Epub 2015 Mar 2. PMID: 25733754

5. Polanczyk G, de Lima MS, Horta BL, Biederman J, Rohde LA. The worldwide prevalence of ADHD: a systematic review and meta-regression analysis. Am J Psychiatry. 2007 Jun; 164(6):942–8. https://doi.org/10.1176/ajp.2007.164.6.942 PMID: 17541055

6. Schachar R, Tannock R. Syndromes of hyperactivity and attention deficit, In: Rutter M, Taylor E, editors. Child and adolescent psychiatry. Oxford: Blackwell; 2002. p.399–418.

7. Russell AE, Ford T, Williams R, Russell G. The Association Between Socioeconomic Disadvantage and Attention Deficit/Hyperactivity Disorder (ADHD): A Systematic Review. Child Psychiatry Hum Dev. 2016 Jun; 47(3):440–50. https://doi.org/10.1007/s10578-015-0578-5 PMID: 26266467

8. Larsson H, Sariastian A, Längström N, D’Onofrio B, Lichtenstein P. Family income in early childhood and subsequent Attention-Deficit/Hyperactivity Disorder: A quasi-experimental study. J Child Psychol Psychiatry. 2014 May; 55(5):428–435. https://doi.org/10.1111/jcpp.12140 PMID: 24111650

9. Kvist AP, Nielsen HS, Simonsen M. The importance of children’s ADHD for parents’ relationship stability and labor supply. Soc Sci Med. 2013 Jul; 88:30–8. https://doi.org/10.1016/j.socscimed.2013.04.001 PMID: 23702207

10. Lasky-Su J, Faraone SJ, Lange C, Tsuang MT, Doyle AE, Smoller JW et al. A Study of how Socioeconomic Status Moderates the Relationship between SNPs Encompassing BDNF and ADHD Symptom Counts in ADHD Families. Behav Genet. 2007 May; 37(3):487–97. https://doi.org/10.1007/s10519-006-9136-x Epub 2013 Apr 10. PMID: 17216343

11. Thapar A, Cooper M, Jefferies R, Stergiakouli E. What causes attention deficit hyperactivity disorder? Arch Dis Child. 2012 Mar; 97(3):260–265. https://doi.org/10.1136/archdischild-2011-300482 PMID: 21903599

12. Russell AE, Ford T, Russell G. Socioeconomic Associations with ADHD: Findings from a Mediation Analysis. PLoS One. 2015 Jun 1; 10(6):e0128248. https://doi.org/10.1371/journal.pone.0128248 PMID: 26030626

13. Babylak MA. Understanding confounding and mediation. Evid Based Ment Health. Aug; 12(3):68–71. https://doi.org/10.1136/ebmh.12.3.68 PMID: 19633239

14. MacKinnon DP, Krull JL, Lockwood CM. Equivalence of the Mediation, Confounding and Suppression Effect. Prev Sci. 2000 Dec; 1(4):173–81. https://doi.org/10.1023/a:1026595011371 PMID: 11523746

15. Jiang H, Kulikarni P, Mallinckrodt CH, Shurzinsky L, Molenberghs G, Lipkovich I. Covariate Adjustment for Logistic Regression Analysis of Binary Clinical Trial Data. Stat Biopharm Res. 2017 Sep; 9(1):126–134. https://doi.org/10.1080/19466315.2016.1234973

16. Christenfeld NJ, Sloan RP, Carroll D, Greenland S. Risk Factors, Confounding, and the Illusion of Statistical Control. Psychosom Med. 2004 Nov-Dec; 66(6):688–75. https://doi.org/10.1097/01.psy.0000140008.70939.41 PMID: 15564351
Mediating factors between socio-economic status and childhood ADHD

17. Rijnhart JJM, Twisk JWR, Eekhout I, Heymans MW. Comparison of logistic-regression based methods for simple mediation analysis with a dichotomous outcome variable. BMC Med Res Methodol 2019 Jan; 19:19. https://doi.org/10.1186/s12874-018-0654-z PMID: 30665353

18. Markham W, Spencer N. Systematic review of factors that mediate the relationships between parental socio-economic status and childhood ADHD. PROSPERO CRD42020182832. 2020. Available from https://www.crd.york.ac.uk/prospero/display_record.php?ID=CRD42020182832.

19. Moher D., Shamseer L., Clarke M, Ghersi D, Liberati A, Petticrew M. Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 statement. Syst Rev 2015 Jan 2; 350:g7647. https://doi.org/10.1186/2046-4053-4-1 PMID: 25554246

20. Campbell M, McKenzie JE, Sowden A, Katikireddi SV, Brennan SE, Ellis S, et al. Synthesis without meta-analysis (SWIM) in systematic reviews: reporting guideline. BMJ. 2020; 368, 16890. https://doi.org/10.1136/bmj.l6890 PMID: 31948937

21. Meunier JC, Boyle M, O'Connor TG, Jenkins JM. Multilevel Mediation: Cumulative Contextual Risk, Maternal Differential Treatment, and Children's Behavior Within Families. Child Dev. 2013 Sep/Oct; 84(5):1594–1615. https://doi.org/10.1111/cdev.12066 PMID: 23397960

22. Russell G, Ford T, Rosenberg R, Kelly S. The association of attention deficit hyperactivity disorder with socioeconomic disadvantage: alternative explanations and evidence. J Child Psychol Psychiatry. 2014 May; 55(5):436–445. https://doi.org/10.1111/jcpp.12170 PMID: 24274762

23. Meulders S, Niklas F, Schneider W. Symptoms of attention-deficit hyperactivity disorder (ADHD) and home learning environment (HLE): a longitudinal study. Eur J Psychol Educ. 2014 Jun; 29:467–482. https://doi.org/10.1177/1359104513489979

24. Miller LL, Gustafsson HC, Tipsord J, Song M, Nousen E, Dieckmann N, et al. Is the association of ADHD with socio-economic disadvantage explained by child comorbid externalizing problems or parent ADHD? J Abnorm Child Psychol. 2018 Jul; 46(5): 951–963. https://doi.org/10.1007/s10802-017-0356-8 PMID: 29128953

25. Nguyen MN, Watanabe-Galloway S, Hill JL, Siahpush M, Tibbits MK, Wichman C. Ecological model of school engagement and attention-deficit/hyperactivity disorder in school-aged children. Eur Child Adolesc Psychiatry 2019 Jun; 28(6):795–805. https://doi.org/10.1007/s00787-018-1248-3 PMID: 30390147

26. Bee T, Dearing E, Stormark KM, Zachrisson HD. Subjective Economic Status in Adolescence: Determinants and Associations with Mental Health in the Norwegian Youth@Hordaland Study. J Fam Econ Iss. 2018(2); 39:323–336. https://doi.org/10.1016/j.jfei.2017.09.005

27. DuPaul GJ, Power TJ, Anastopoulos AD, Redd R. ADHD Rating Scale IV; Checklists, norms, and clinical interpretation. New York: Guildford Publications; 1998.

28. Conners C.K. 3rd Edition Manual. North Tonawanda (NY): Multi-Health Systems Inc.; 2008.

29. Goodman R. Psychometric properties of the Strength and Difficulties Questionnaire. J Am Acad Child Adolesc Psychiatry. 2001 Nov; 40(11):1337–45. https://doi.org/10.1097/00004583-200111000-00015 PMID: 11699809

30. Goodman R. The Strengths and difficulties Questionnaire: a research note. J Child Psychol Psychiatry. 1997 Jul; 38:581–586. https://doi.org/10.1111/j.1469-7610.1997.tb01545.x PMID: 9255702

31. Theunissen MHC, Vogels AGC, de Wolff MS, Reijneveld SA. Characteristics of the Strengths and Difficulties Questionnaire I Preschool Children. Pediatrics 2013 Feb; 131(2):e446–54. https://doi.org/10.1542/peds.2012-0089 PMID: 23296429

32. Boyle MH, Offord DR, Racine YA, Fleming JE, Szatmari P, Sanford M. Evaluation of the revised Ontario Child Health Study scales. J Child Psychol Psychiatry 1993 Feb; 34(2):189–213. https://doi.org/10.1111/1469-7610.1993.tb00979.x PMID: 8444992

33. Goodman R, Ford T, Richards H, Gatward R, Meltzer H. The Development and Well-Being Assessment: description and initial validation of an integrated assessment of child and adolescent psychopathology. J Child Psychol Psychiatry 2000 Jul; 41(5):645–55. PMID: 10946756

34. Kessler RC, Adler L, Ames M, Demler O, Faraone S, Hiripi E et al. The World Health Organization adult ADHD self-report scale (ASRS): A short screening scale for use in the general population. Psychol Med 2005 Feb; 35(2):245–56. https://doi.org/10.1017/s0033291704002892 PMID: 15841682
38. Kraemer HC, Stice E, Kazdin A, Offord D, Kupfer D. How do risk factors work together? Mediators, moderators, and independent, overlapping and proxy risk factors. Am J Psychiatry. 2001 Jun; 158(6):848–56. https://doi.org/10.1176/appi.ajp.158.6.848 PMID: 11384888

39. Edwards JR, Lambert LS. Methods for integrating moderation and mediation: A general analytic framework using moderated path analysis. Psychol Methods. 2007 Mar; 12(1):1–22. https://doi.org/10.1037/1082-989X.12.1.1 PMID: 17402809

40. Preacher KJ, Rucker DD, Hayes AF. Addressing moderated mediation hypotheses: theory, methods and prescriptions. Multivariate Behav Res 2007 Jan-Mar; 42(1):185–227. https://doi.org/10.1080/00273170701341316 PMID: 26821081

41. Preacher KJ, Rucker DD, Hayes AF. Asymptotic and resampling strategies for assessing and comparing indirect effects in multiple mediator models. Behav Res Methods 2008 Aug; 40(3):879–91. https://doi.org/10.3758/brm.40.3.879 PMID: 18697684

42. Van Dam NT, Earleywine M. Validation of the Center for Epidemiologic Studies Depression Scale–Revised (CESD-R): pragmatic depression assessment in the general population. Psychiatry Res. 2011 Mar 30; 186(1):128–32. https://doi.org/10.1016/j.psychres.2010.08.018 Epub 2010 Sep 16. PMID: 20843557

43. Meades R, Ayers S. Anxiety measures validated in perinatal populations: a systematic review. J Affect Disord. 2011 Sep; 133(1–2):1–15. https://doi.org/10.1016/j.jad.2010.10.009 Epub 2010 Nov 13. PMID: 21078523

44. Gibson J, McKenzie-Mcharg K, Shakespeare J, Price J, Gray R. A systematic review of studies validating the Edinburgh Postnatal Depression Scale in antenatal and postpartum women. Acta Psychiatr Scand. 2009 May; 119(5):350–64. https://doi.org/10.1111/j.1600-0447.2009.01363.x Epub 2009 Mar 2. PMID: 19298573

45. Eberhard-Gran M, Eskild A, Tambs K, Opjordsmoen S, Ove Samuelson S. Review of validation studies of the Edinburgh Postnatal Depression Scale. Acta Psychiatr Scand. 2001 Oct; 104(4):243–9. https://doi.org/10.1034/j.1600-0447.2001.00187.x PMID: 11722298

46. Conners CK, Erhart D, Sparrow E. Conners Adult ADHD Rating Scales (CAARS) Technical Manuel North Tonawanda (NY): Multi-Health Systems Inc.; 1999.

47. Barkley RA. Barkley Adult ADHD Rating Scale-IV (BAARS-IV). New York: Guildford Publications; 2011.

48. VanderWeele TJ. Principles of confounder selection. Eur J Epidemiology 2019 Mar; 34(3):211–219. https://doi.org/10.1007/s10654-019-00494-6 Epub 2019 Mar 6. PMID: 30840181

49. Pianta RC. Child-Parent Relationship Scale. Charlottesville (VA): University of Virginia Press; 1995.

50. Matias C, Scott S, O’Connor TG. Coding of Attachment-Related Parenting (CARP). London: Kings College Institute of Psychiatry; 2006.

51. National Longitudinal Survey of Children and Youth. Users handbook and microdata guide. Ottawa: Statistics Canada and Human Resources Development Canada; 1995.

52. Stayhorn JM, Weidman CS. A parent practices scale, and its relation to parent and child mental health. J Am Acad Child Adolesc Psychiatry. 1998 Sep; 37(5):613–8. https://doi.org/10.1097/00004583-199809000-00016 PMID: 3182627

53. Deater-Deckard K. Parenting and child behavioral adjustment in early childhood: A quantitative approach to studying family processes in the National Longitudinal Survey of Children and Youth. J Fam Psychol. 2000; 14(2):243–59. https://doi.org/10.1037/0893-3200.14.2.243 Epub 2000 Mar 1. PMID: 10834478

54. Steer C, Wolke D, the ALSPAC study team. An Index of Family Adversity [poster]. presented at the 3rd Conference of Epidemiological Longitudinal Studies in Europe. University of Bristol, UK; 2004.

55. Rutter M, Quinton D. Psychiatric disorder: ecological factors and concepts of causation. In Ecological factors in human development, North Holland Publishing Company: International Society for the Study of Behavioral Development; 1977. p. 173–187.

56. Reiss F, Meyrose A-K, Otto C, Lampert T, Klase R, Ravens-Sieberer U. Socioeconomic status, stressful life situations and mental health problems in children and adolescents: Results of the German BELLA cohort-study. PLoS One. 2019; 14(3): e0213700. doi: https://doi.org/10.1371/journal.pone.0213700 PMID: 30865713

57. Bertras S, Laroque B, Bouvier-Colbe MH, Kaminski M. [Infant temperament at 6–9 months old: validation on the French version of the Infant Characteristics Questionnaire and factors associated with measurement. Rev. Epidemiol Sante Publique. 1999 Jun; 47(3):263–77. PMID: 10422120.

58. Thomas Ray G, Croen LA, Habel LA. Mothers of Children Diagnosed with Attention-Deficit/Hyperactivity Disorder: Health Conditions and Medical Care Utilization in Periods Before and After Birth of the Child. Med Care. 2009 Jan; 47(1):105–114. https://doi.org/10.1097/MLR.0b013e31817e18c0 PMID: 19106738
59. Walsh D, McCartney G, Smith M, Armour G. Relationship between childhood socioeconomic position and adverse childhood experiences (ACEs): a systematic review. J Epidemiol Community Health. 2019 Dec; 73(12):1087–1093. https://doi.org/10.1136/jech-2019-212738 PMID: 31563897

60. How LD, Tilling K, Galobardes B, Lawlor DA. Loss to Follow-up in Cohort Studies: Bias in Estimates of Socioeconomic Inequalities. Epidemiology 2013 Jan; 24(1):1–9. https://doi.org/10.1097/EDE.0b013e31827623b1 PMID: 23211345

61. Wolke D, Wayle A, Smara M, Steer C, Goodman R, Fird T, et al. Selective drop-out in longitudinal studies and non-biased prediction of behaviour disorders. Br J Psychiatry. 2009 Sep; 195(3):249–56. https://doi.org/10.1192/bjp.bp.108.053751 PMID: 19721116

62. VanderWeele TJ, Vansteelandt S, Robins JM. Effect decomposition in the presence of an exposure-induced mediator-outcome confounder. Epidemiology. 2014 Mar; 25(2):300–6. https://doi.org/10.1097/EDE.000000000000034 PMID: 24487213