

University of Warwick institutional repository: http://go.warwick.ac.uk/wrap

This paper is made available online in accordance with publisher policies. Please scroll down to view the document itself. Please refer to the repository record for this item and our policy information available from the repository home page for further information.

To see the final version of this paper please visit the publisher's website. Access to the published version may require a subscription.

 $\label{eq:Author} \mbox{Author(s): A Waylen , T Ford , R Goodman , M Samara , D Wolke Article Title: Can early intake of dietary omega-3 predict childhood}$

externalizing behaviour? Year of publication: 2009

Link to published version: http://dx.doi.org/_10.1111/j.1651-

2227.2009.01434.x

Publisher statement: None



Can early intake of dietary omega-3 predict childhood externalizing behavior?

| Journal: | Acta Paediatrica |
|-------------------------------|---|
| Manuscript ID: | SPAE-2008-0875.R2 |
| Manuscript Type: | Regular Article |
| Date Submitted by the Author: | |
| Complete List of Authors: | Waylen, Andrea; Bristol Dental School, Oral and Dental Science Ford, Tamsin; Peninsula Medical School Goodman, Robert; Institute of Psychiatry, Child and Adolescent Psychiatry Samara, Muthanna; University of Warwick, Department of Psychology Wolke, Dieter; University of Warwick, Department of Psychology; Warwick Medical School, Health Sciences Research Institute |
| Keywords: | ALSPAC, breastfeeding, externalising behaviour, longitudinal study, omega-3 |
| | |



Can early intake of dietary omega-3 predict childhood externalizing behaviour?

A. Waylen¹, T. Ford², R. Goodman³, M. Samara⁴ and D. Wolke^{4,5}

Short title: Dietary omega-3 and childhood externalising behaviour

Corresponding author: Dr Andrea Waylen, Department of Oral and Dental Science, Bristol Dental School, Lower Maudlin Street, Bristol BS1 2LY

Email: andrea.waylen@bristol.ac.uk

Tel: +44 (0)117 342 4281 Fax: +44 (0)117 929 9898

¹ Bristol Dental School, Department of Oral and Dental Science

² Peninsula Medical School

³ Kings College, London, Department of Child and Adolescent Psychiatry

⁴University of Warwick, Department of Psychology

⁵ University of Warwick, Warwick Medical School

ABSTRACT

Aim: To determine whether maternal and child intake of dietary omega-3 fatty acids, together with the presence or absence of breastfeeding, predicted psychiatric diagnosis of externalising disorders in childhood.

Methods: Data concerning childhood externalising disorders was collected from 8242 children aged 7.9 years in a large British cohort. Intake of omega-3 fatty acids was measured for the study mother during pregnancy and for the child at three years. Duration of breastfeeding was examined to account for moderating effects. Adjustment was made for a variety of potential confounders.

Results: Maternal intake of omega-3 and breastfeeding predicted oppositional / conduct disorder and comorbid externalising disorder before adjustment for confounding factors. However, there was no association between intake of omega-3 by mother or child and any type of externalising disorder once socio-demographic factors were taken into account.

Conclusions: Any association between intake of omega-3 and childhood externalising disorders appears to be strongly confounded with sociodemographic factors. This is important to note given the current popularity of omega-3 as a possible treatment for behaviour problems related to inattention and impulsivity. Care must be taken that studies investigating this relationship account fully for factors associated with both behaviour and diet.

Keywords: ALSPAC, breastfeeding, externalising behaviour, longitudinal study, omega-3

Nutrition in infancy or early childhood is associated with childhood behaviours such as inattentiveness, hyperactivity, impulsivity and anti-social behaviour; these, in turn are associated with childhood externalising disorders including ADHD [1]. One group of nutrients associated with externalising disorders in children is that of essential fatty acids (EFAs). Results from two recent randomised control trials conclude that nutritional interventions using a combined fatty acid supplement reduce problems such as inattention, hyperactivity and impulsivity in middle childhood and that these effects can be maintained over time [2, 3] but other studies report no effects [4].

EFAs are essential for both optimal development [5] and the functioning of the central nervous system (CNS) [6]. During the last trimester of pregnancy and up to two years of age, they accumulate rapidly in the CNS of the infant, transferring from mother to child via the placenta and in breast-milk. EFAs must be obtained from the maternal diet from foods like oily fish as they cannot be synthesised by humans [7]. However early neurodevelopment may be compromised if maternal levels of EFAs are not supplemented by diet during pregnancy and lactation [5] or if infant levels are not maintained post-natally via breast or supplemented feeding and complementary foods [8].

The current paper focuses upon one type of EFA, the omega-3 series of long-chain polyunsaturated fatty acids (LC-PUFAs). LC-PUFAs are found primarily in cell membrane phospholipids, are important in the regulation of cell function and affect gene transcription [5]. They are necessary during periods of rapid growth (pre-natal development, early childhood and adolescence) and nutrient loss (pregnancy and lactation) [6]. Neonates with neurological dysfunction have lower indices of both LC-PUFAs and EFAs [8] and there is increased risk of suboptimal cognitive, social and motor skills in children at eight years of age as maternal intake of FAs during pregnancy declines [9]. On the other hand, increased CNS maturity is observed in infants who are breastfed [10] possibly because of the transfer of LC-PUFAs and other fatty acids in breast milk [11]. The aim of the present study was to determine whether maternal intake of omega-3 LC-PUFAs at 32 weeks gestation and by the child aged three years predicted diagnosis of externalizing disorders overall and specific sub-types of disorders in middle childhood. We also examined the moderating effects of breastfeeding in the relationship between diagnosis of externalising disorders and intake of omega-3 in pregnancy and childhood.

MATERIALS AND METHODS

Patients: The data analysed here were collected for the Avon Longitudinal Study of Parents and Children (ALSPAC: see www.bristol.ac.uk/alspac/), reported in detail elsewhere [12]. Within the former county of Avon, UK, 14,541 pregnant women expected to deliver between April 1991 and December 1992 enrolled into the

study. By the time children entered primary school (aged 5), around 11,500 families were still participating.

Mothers consented to join the study at recruitment and were free to withdraw at any time; all aspects of the study conform to the ethical regulations of both the ALSPAC Law and Ethics Committee and local research ethics committees.

As with many longitudinal studies there was attrition from ALSPAC during the data collection period. When the study children were aged 7.9 years (SD 1.4 months) behavioural data was returned by parents for 8242 children (59%) and by teachers for 5115 study children (39%). Mothers who dropped out from the study (non-responders) were younger than those who remained (p<.001), were more likely to leave school earlier (p<.001), were more likely to live in rented accommodation (p<.001) and were more likely to still be smoking in pregnancy (p<.001). These families were also exposed to higher levels of family adversity during the pregnancy (p<.001). Babies born to non-responders were more likely to have been born prematurely (p<.001) and to weigh less than 2500g (p<.001). These babies were less likely to have been breastfed (p<.001). Non-responder mothers had lower levels of omega-3 in their diet than those who remained in the study (p<.001) but there was no difference in the omega-3 intake of the children at 3 years of age. (Actual data for this comparison are provided in a supplementary file). Diet Assessment: Information on mother's diet at 32 weeks gestation and child's diet at 3 years was collected using food-frequency questionnaires; details are provided elsewhere [13]. The transfer of EFAs during the final trimester of pregnancy is important for the infant's neurological development and although the CNS is no longer so demanding of omega-3 for development at 3 years of age, intake at this stage might be considered as a maintenance dose. Three questions on seafood consumption were used to estimate omega-3 intake per week for mothers and children and intake was categorised as either high or low [9, 14]. Mothers and children with high intake were used as the reference group in analyses. Infant-feeding data were taken from the 15-month questionnaire with the reference group being children who were breastfed for any period of time. Confounding factors associated with diet were: maternal age and education, family adversity during pregnancy (an aggregate measure of 18 variables measuring adverse family circumstances including maternal psychopathology, financial difficulties and standard of housing) [15] and whether the mother smoked cigarettes during pregnancy (never smoked, stopped smoking in pregnancy, still smoking in pregnancy).

Behavioural data: Information about child behaviour at 7.9 years was collected from parents and teachers using a questionnaire version of the Development and Well-Being Assessment (DAWBA) [16]. This is a validated measure consisting of structured and semi-structured questions. An experienced clinician (TF) combined all information about symptoms and their impact using a computerised heuristic to make DSM-IV diagnoses of

childhood psychiatric disorders [17]. Data from teacher questionnaires were available for 66% of children with parent data. Confounding factors associated with behaviour were gender, gestational age (<37 weeks or >=37 weeks) and birth weight (<2500g or >=2500g).

Statistical methods: All analyses were undertaken using STATA 9.0[18]. Diagnoses of externalizing disorders were categorised into groups: oppositional defiant disorder or conduct disorder without any type of comorbid disorder ("pure ODD/CD": N = 189), attention-deficit hyperactive disorder without any type of comorbid disorder ("pure ADHD": N = 103) or ADHD plus ODD/CD (comorbid disorder, N = 72). We focussed on "pure" disorders and ADHD plus ODD / CD as a separate group so that children with comorbid disorders were not double-counted. Descriptive statistics were calculated by diagnostic category (Table 1). (Table 1 about here)

Unadjusted independent logistic regressions were carried out to predict diagnosis using mother and child omega-3 FA consumption and whether or not the child was breast-fed, together with an interaction term for breast-feeding by maternal intake of omega-3 FAs. The interaction term allowed the identification of a) mothers who breastfed but did not eat oily fish during pregnancy and b) those who ate oily fish during pregnancy but did not breastfeed. Multivariate step-wise models were built adjusting for all confounding factors. To account for multiple analyses, a conservative value of p<.001 was used.

RESULTS

Dietary factors: Children with "pure ODD / CD" were less likely to have been breastfed (p<.001). However, the evidence for differences in omega-3 intake by either mother or child between children diagnosed with a disorder and those without was weak (p>=.001) (Table 1).

Externalising disorders: For every type of externalizing disorder, more boys were diagnosed than girls (p<.001). Mothers of children with externalizing disorders were younger than those with no disorder (p<.05) but there was no difference in their education (p=.68). The families of children with externalising disorders experienced more family adversity during pregnancy than those with no disorder (p<.001). With respect to specific disorders, children diagnosed with "pure ADHD" were more likely to have been born before 37 weeks than those without (p<.001). Mothers of children with "pure ODD / CD" and "ADHD plus ODD / CD were more likely to continue to smoke throughout pregnancy (p<.001).

Dietary factors and externalising disorders: Neither maternal nor child intake of omega-3 nor breastfeeding predicted "pure ADHD" in the unadjusted analyses (p>=.38; odds ratios and confidence intervals in Table 2). Formula-feeding (never breastfed) predicted "pure ODD/CD" in the unadjusted analyses (p<.001) but after adjustment for confounding factors, there was no evidence for formula feeding as a predictor (p>.40).

There was weak evidence that maternal omega-3 (p = .01), formula feeding (p = .02) and the interaction between the two (p = 0.18) predicted the combination of ADHD plus ODD/CD in the unadjusted model but after adjustment for confounding factors these relationships disappeared (p > = .34).

DISCUSSION

This study aimed to investigate the ability of maternal intake of omega-3 during pregnancy and child intake of omega-3 at 3 years to predict externalising disorders in middle childhood. Children with externalising disorders were more likely to be boys and to be born into families where there was strong evidence of relatively high levels of family adversity during pregnancy. Children diagnosed with ADHD were more likely to have been born before 37 weeks.

Conclusions from previous research regarding the effects of omega-3 on externalising disorders have been inconsistent: some studies report positive effects of omega-3 [2, 3] while other work has been less conclusive [5]. Our results showed no overall benefit of omega-3 intake by either mother or child on the diagnosis of externalising disorders after adjustment. There were no associations between omega-3 intake and diagnosis of "pure ADHD" in this British cohort, consistent with some [4] but not other research [19]. Children who were breastfed had a lower rate of "pure ODD/CD" than children who were formula-fed before adjustment [20]. Similarly, when maternal intake of omega-3 was low during pregnancy and the baby was subsequently formula-fed, there was an increased risk of ADHD plus ODD/CD in unadjusted analyses. However, the predictive ability of these variables disappeared after adjustment, perhaps because the unadjusted relationships were a function of residual confounding eg breast-feeding may be easier to manage for mothers who, for example, are not depressed or have more time to spend supporting and supervising their children. Alternatively, we may have over-adjusted in some circumstances and the unadjusted relationships may be accurate eg omega-3 intake may be reduced in families exposed to adversity and reduced intake during pregnancy may be associated with an increased risk of oppositional and / or conduct disorders in childhood. In order to examine this hypothesis, the effects of omega-3 on such disorders should be further investigated. We are unable to rule out the possibility that other processes (not measured here) may influence the risk of externalising disorders in childhood [21] eg inefficient metabolism as opposed to a dietary deficiency of omega-3 [4] or the ratio of different fatty acids to one another [11] or variation in the nutritional content of breast-milk [22]. Further research is needed to improve our understanding of the influence of absolute levels of nutrients in diet on externalizing behaviours, particularly ADHD. While the strengths of the study are large sample size and observer-rated psychiatric diagnoses, there are limitations. Only 59% of the original cohort provided DAWBA data and fewer than 5% of this group met the

criteria for diagnosis of an externalising disorder. Given the lack of teacher data on 34% of the sample, we will have misclassified some children as having no disorder [17], reducing our chances of detecting statistical associations. Other work on this cohort [23] has suggested that a lower proportion of families remaining in the study have children with externalising disorders compared to those who dropped out but bias will only occur if the association between omega-3 and externalising symptoms varies between responders and non-responders and we have no reason to believe that this is the case. In statistical simulations this type of selection bias has been shown to attenuate rather than alter the relationship between predictors [24].

We were restricted to using dietary intake of omega-3 FAs and breast-feeding practice as proxies for physiological measures of omega-3. This is problematic regarding the transfer of omega-3 via breast-milk given that the omega-3 content of breast-milk can vary ten-fold [22]. Similarly, other factors in human milk may affect neurodevelopment so studies such as this, comparing breast- and formula-fed infants, cannot completely resolve the role of EFAs in development. Also, the dichotomisation of each of these variables may have resulted in a loss of statistical power to detect differences between groups but given that the outcome of interest was diagnosis rather than symptom counts, we believe that categorisation allows the possibility of capturing non-linear effects in the relationship with externalising disorders.

In conclusion, neither maternal nor child intake of omega-3 nor breastfeeding predicted any type of externalising disorder after adjusting for social factors associated with diet and behaviour Our results are important to consider in light of the current tendency to promote omega-3 as a possible treatment for ADHD-related problems [2]. If studies do not adjust sufficiently for confounding factors such as family adversity and pre-term delivery then they may misrepresent the relationship between externalising disorders and omega-3 [25]. Stronger evidence is required from randomised control trials to clarify the role of omega-3 alone and in conjunction with other LC-PUFAs in childhood externalising disorders.

Dietary omega-3 and childhood externalising behaviour

ACKNOWLEDGEMENTS

We are extremely grateful to all the families who took part in this study, the midwives for their help in recruiting them, and the whole ALSPAC team, which includes interviewers, computer and laboratory technicians, clerical workers, research scientists, volunteers, managers, receptionists and nurses. The UK Medical Research Council, the Wellcome Trust and the University of Bristol provide core support for ALSPAC. This publication is the work of the authors and Dr Waylen will serve as guarantor for the contents of this paper. This research was specifically funded by The Health Foundation (Grant 265/1981). We thank Colin Steer (ALSPAC, University of Bristol) for his guidance with statistical analysis and Imogen Rogers for help with the interpretation of the omega-3 data. We would also like to thank the reviewers for their helpful comments during the construction of this manuscript.

ABBREVIATIONS

EFA, essential fatty acid; CNS, central nervous system; LC-PUFA, long-chain polyunsaturated fatty acid; FA, fatty acid; IQ, intelligence quotient; ALSPAC, Avon Longitudinal Study of Parents and Children; UK, United Kingdom; DAWBA, Development and Well-Being Assessment; ODD/CD, oppositional development disorder / conduct disorder; ADHD, attention-deficit hyperactive disorder; OR, odds ratio; CI, confidence interval; SES, socio-economic status; FAI, family adversity index; n-3, omega-3

REFERENCES

- [1] Schnoll R, Burshteyn D, Cea-Aravena J. Nutrition in the treatment of attention-deficit hyperactivity disorder: a neglected but important aspect. Applied Psychophysiology and Biofeedback. 2003;28(1):63-75.
- [2] Sinn N, Bryan J. Effect of supplementation with polyunsaturated fatty acids and micronutrients on learning and behavior problems associated with child ADHD. Journal of Developmental and Behavioral Pediatrics. 2007;28(2):82-91.
- [3] Richardson AJ, Montgomery P. The Oxford-Durham study: A randomized, controlled trial of dietary supplementation with fatty acids in children with developmental coordination disorder. Pediatrics. 2005;115(5):1360-6.
- [4] Richardson AJ, Puri BK. The potential role of fatty acids in attention- deficit/hyperactivity disorder. Prostaglandins Leukotrienes and Essential Fatty Acids. 2000 Jul-Aug;63(1-2):79-87.
- [5] Heird W, Lapillone A. The role of essential fatty acids in development. Annual Review of Nutrition. 2005;25:549-71.
- [6] Muskiet FAJ, van Goor SA, Kuipers RS, Velzing-Aarts FV, Smit EN, Bouwstra H, et al. Long-chain polyunsaturated fatty acids in maternal and infant nutrition. Prostaglandins Leukotrienes and Essential Fatty Acids. 2006;75(3):135-44.
- [7] Koletzko B, Agostini C, Carlson SE, Clandinin T, Hornstra G, Neuringer M, et al. Long chain polyunsaturated fatty acids (LC-PUFA) and perinatal development. Acta Paediatrica. 2001;90:460-4.
- [8] Loosemore ED, Judge MP, Lammi-Keefe CJ. Dietary intake of essential and long-chain polyunsaturated fatty acids in pregnancy. Lipids. 2004;39(5):421-4.
- [9] Hibbeln JR, Davis JM, Steer C, Emmett P, Rogers I, Williams C, et al. Maternal seafood consumption in pregnancy and neurodevelometral outcomes in childhood (ALSPAC study): an observational cohort study. The Lancet. 2007;369:578-85.
- [10] Willatts P, Forsyth JS. The role of long-chain polyunsaturated fatty acids in infant cognitive development. Prostaglandins Leukotrienes and Essential Fatty Acids. 2000 Jul-Aug;63(1-2):95-100.
- [11] Gustafsson P, Duchen K, Birberg U, Karlsson T. Breastfeeding, very long polyunsaturated fatty acids(PUFA) and IQ at 61/2 years of age. Acta Paediatrica. 2004;93:1280-7.
- [12] Golding J, Pembrey M, Jones R. ALSPAC--the Avon Longitudinal Study of Parents and Children. I. Study methodology. Paediatr Perinat Epidemiol. 2001;15(1):74-87.

- [13] North K, Emmett P, Team TAS. Multivariate analysis of diet among three-year-old children and associations with socio-demographic characterisitics. European Journal of Clinical Nutrition. 2000;54:73-80.
- [14] Australian Government. Nutrient Reference values for Australia and New Zealand: Ministry of Health; 2006.
- [15] Bowen E, Heron J, Waylen A, Wolke D. Domestic violence risk during and after pregnancy: findings from a British longitudinal study. BJOG: An International Journal of Obstetrics and Gynaecology. 2005;112(8):1083-9.
- [16] 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. Journal of Child Psychology & Psychiatry. 2000;41:645-55.
- [17] Ford T, Goodman R, Meltzer H. The British Child and Adolescent Mental Health Survey 1999: the prevalence of DSM-IV disorders. Journal of the American Academy of Child and Adolescent Psychiatry. 2003 Oct;42(10):1203-11.
- [18] STATA 9.0. College Station, TX: Statacorp 1985.
- [19] Raine A, Mellingen K, Liu JH, Venables P, Mednick SA. Effects of environmental enrichment at ages 3-5 years on schizotypal personality and antisocial behavior at ages 17 and 23 years. American Journal of Psychiatry. 2003 Sep;160(9):1627-35.
- [20] Jacobson S, Jacobson J. Breastfeeding and intelligence in children. BMJ. 2006;333:929-30.
- [21] Feldman R, Eidelman AI. Direct and indirect effects of breast milk on the neurobehavioural and cognitive development of premature infants. Developmental Psychobiology. 2003;43:109-19.
- [22] Innis S, Uauy R. Mechanisms of action of LCPUFA effects on infant growth and neurodevelopment: perinatal biochemistry and physiology of LCPUFA discussion. The Journal of Pediatrics. 2003 2003/10;143(4, Supplement 1):96-109.
- [23] Wolke D, Waylen A, Samara M, Steer C, Goodman R, Ford T, et al. Does selective dropout in longitudinal studies lead to biased prediction of behaviour disorders? British Journal of Psychiatry. Submitted.
- [24] Berk R. An Introduction to Sample Selection Bias in Sociological Data. American Sociological Review. 1983;48(3):386-98.
- [25] Dangour A, Allen E, Elbourne D, Fletcher A, Richards M, Uauy R. Fish consumption and cognitive function among older people in the UK: baseline data from the OPAL study. Journal of Nutrition, Health and Aging 2009;13(3):198-202.

Table 1: Descriptive data for responders according to DSM-IV diagnosis (SD)^a

| | N (%) | Maternal | Maternal | FAI [§] in | % Still | Child | Maternal n3 ^x | % < 37 | % <2500g | Whether | Child n3 ⁿ |
|---------------|--------|-------------|----------------------|---------------------|------------|---------|--------------------------|-----------|--------------|-----------|-----------------------|
| | | age | education | pregnancy | smoking in | gender | (32 weeks gestation) | weeks | birth-weight | breastfed | (3 years) |
| | | | % O levels | | pregnancy | % male | % Low intake | gestation | | % Never | % Low intake |
| | | | or less [±] | | | | <340g / week | | | | <280mg / week |
| No disorder | 7654 | 29.0 (4.6) | 58.3 | 1.1 (1.4) | 15.2 | 50.1 | 78.3 | 4.8 | 4.5 | 22.2 | 49.6 |
| | (92.9) | | | | | | | | | | |
| Any | 364 | 28.2 (5.1)* | 61.5 | 1.8 (1.9)*** | 24.2 *** | 75.5 | 82.2 * | 7.2 | 5.7 | 32.2*** | 50.5 |
| externalizing | (4.4) | | | | | *** | | | | | |
| disorder | | | | | | | | | | | |
| Pure ODD/CD | 189 | 28.3 (5.0)* | 61.4 | 1.9 (1.9)*** | 29.4*** | 68.6*** | 82.3 ** | 4.3 | 4.8 | 33.9*** | 57.1** |
| | (2.3) | | | | | | | | | | |
| Pure ADHD | 103 | 29.0 (5.1) | 56.1 | 1.5 (1.8)*** | 14.4 | 81.6*** | 76.1 | 13.6*** | 8.7 * | 26.1 | 48.9 |
| | (1.3) | | | | | | | | | | |
| ADHD plus | 72 | 27.4 (5.2)* | 72.3 | 2.1 (2.0)*** | 29.4** | 88.6*** | 88.9 * | 5.7 | 5.7 | 34.8* | 47.5 |
| ODD/CD | (0.9) | | | | | | | | | | |

^a the total number of children in this table with externalising disorders only is 8018; 224 children from the complete sample of 8242 were diagnosed with an internalising disorder but these were excluded from all analyses.

[±]O levels: educational qualification taken at 16 years of age § Family Adversity Index ⁿ n3 = omega-3 fatty acids

^{*} p<.05 ** p<.01 *** p<.001 (strength of evidence for difference from those with no disorder)

Table 2: Odds ratios and 95% CI for omega-3 and breastfeeding factors in the prediction of DSM-IV externalizing behaviors

| DSM-IV diagnosis | | Reference group | Unadjust | ed | | Adjuste | ed § | |
|------------------|------------------------------|-----------------|-------------------|------------------------------|------------------|------------|-----------------------|----------|
| ODD/CD | Maternal n3 | None | OR 1.73 | 95% CI 1.03 – 2.90 | р 0.04 | OR 0.98 | 95% CI 0.64 – 1.49 | р .92 |
| N = 189 | Child n3 | None | 1.62 | 0.92 - 2.85 | 0.09 | 1.39 | 0.99 - 1.96 | .06 |
| | Never breastfed | | 1.83 | 1.30 – 2.56 | <.001 | 1.16 | 0.57 - 2.36 | .68 |
| | Maternal omega-3 * breastfed | None*never | 1.20 | 0.40 - 3.61 | 0.74 | 1.42 | 0.63 - 3.23 | .40 |
| ADHD | Maternal n3 | None | 1.17 | 0.60 - 2.28 | 0.65 | 0.89 | 0.53 - 1.49 | .66 |
| N = 103 | Child n3 | None | 1.31 | 0.62 - 2.79 | 0.48 | 0.77 | 0.49 - 1.21 | .25 |
| | Never breastfed | | 1.24 | 0.77 - 1.98 | 0.38 | 0.83 | 0.32 - 2.17 | .70 |
| | Maternal omega-3 * breastfed | None*never | 0.60 | 0.11 - 3.32 | 0.56 | 1.40 | 0.45 - 4.40 | .56 |
| ADHD plus ODD/CD | Maternal n3 | None | 3.56 | 1.41 – 8.95 | 0.01 | 1.41 | 0.67 - 2.98 | .37 |
| N = 72 | Child n3 | None | 1.02 | 0.36 - 2.89 | 0.98 | 0.92 | 0.52 - 1.63 | .78 |
| | Never breastfed | | 1.87 | 1.13 – 3.12 | 0.02 | 1.76 | 0.55 - 5.59 | .34 |
| | Maternal omega-3 * breastfed | None*never | 2.06 | 1.13 – 3.73 | .018 | 0.99 | 0.26 - 3.79 | .99 |

[§] Adjustment factors: Maternal age and education level, family adversity (including maternal psychopathology, financial difficulties and standard of housing during pregnancy), gestational age, birthweight, breastfeeding and smoking

Table 1: Descriptive data for responders vs. non-responders

| | Responders | Non-responders | p |
|---|-------------|----------------|-------|
| | (N = 8242) | (N = 3098) | |
| Mean maternal age (SD) | 29.0 (4.6) | 26.7 (5.1) | <.001 |
| Maternal education: O levels or less ^a | 58% | 76% | <.001 |
| Home owner | 82% | 61% | <.001 |
| Mean family adversity score (SD) | 1.1 (1.4) | 1.6 (1.8) | <.001 |
| Still smoking in pregnancy | 16% | 29% | <.001 |
| Gestational age <37 weeks | 5% | 15% | <.001 |
| Birth weight <2500g | 5% | 8.4% | <.001 |
| Never breastfed | 23% | 35% | <.001 |
| Mean maternal intake (g) of omega-3 (SD) | 1.4 (0.5) | 1.3 (0.5) | <.001 |
| Mean child intake (g) of omega-3 (SD) | 0.51 (0.50) | 0.51 (0.50) | .082 |
| Gender of study child | 51% | 52% | .17 |

^a In England and Wales, O levels are a national qualification taken at 16 years of age