

## CLDRC MASTER PUBLICATIONS LIST

### 2016 and In Press

- Ayorech, Z., Selzam, S., Smith-Woolley, E., Knopik, V. S., Neiderhiser, J. M., DeFries, J. C., & Plomin, R. (in press). Publication Trends Over 55 Years of Behavioral Genetic Research. *Behavior Genetics*.
- Byrne, B., Olson, R.K., & Samuelsson, S. (in press). The longitudinal perspective on developmental disorders. In C. Marshal (Ed.), *Current issues in developmental disorders*. Hove, UK: Psychology Press.
- Cirino, P. A., & Willcutt, E. G. (in press). The importance of executive function in cognitive models of learning disabilities. *Journal of Learning Disabilities*.
- Gialluisi, A., Visconti, A., Willcutt, E. G., Smith, S. D., Pennington, B. F., Falchi, M., . . . Fisher, S. E. (in press). Investigating the effects of copy number variants on reading and language performance. *Journal of Neurodevelopmental Disorders*. PMCID: PMC4868026.
- Godinez, D. A., Willcutt, E. G., Pearson, K., Depue, B. E., Burgess, G. C., Andrews-Hanna, J., . . . Banich, M. T. (in press). Familial risk and ADHD specific neural activity revealed by a case-control, discordant twin pair design. NIHMS: NIHMS714497.
- Grasby, K. L., Coventry, W. L., Byrne, B., Olson, R. K., & Medland, S. E. (In press). Genetic and environmental influences on literacy and numeracy performance in Australian school children in grades 3, 5, 7, and 9. *Behavior Genetics*.
- Huibregtse, B. M., Corley, R. P., Wadsworth, S. J., Vandever, J. M., DeFries, J. C., & Stallings, M. C. (in press). A longitudinal adoption study of substance use behavior in adolescence. *Twin Research and Human Genetics*.
- Jacobson, L. A., Koriakin, T., Lipkin, P., Boada, R., Frijters, J., Lovett, M., . . . Mahone, E. M. (in press). Executive functions contribute uniquely to reading competence in minority youth. *Journal of Learning Disabilities*.
- Johnson, E. P., Pennington, B. F., Lowenstein, L. H., & Nittrouer, S. (in press). Sensitivity to structure in the speech signal by children with speech sound disorder and reading disability. *Journal of Communication Disorders*.
- Keenan, J. M. (in press). Measure for measure: Challenges in assessing reading comprehension. In J. Sabatini & E. Albro (Eds.), *Assessing Reading in the 21st Century: Aligning and Applying Advances in the Reading and Measurement Sciences*. Lanham, MD: R & L Education.
- Leopold, D. R., Christopher, M. E., Burns, G. L., Becker, S. P., Olson, R. K., & Willcutt, E. G. (in press). ADHD and sluggish cognitive tempo throughout childhood: Temporal invariance and stability from preschool through ninth grade. *Journal of Child Psychology and Psychiatry*. PMCID: PMC4334688.
- McGrath, L. M., Braaten, E. B., Doty, N. D., Willoughby, B. L., Wilson, H. K., O'Donnell, E. D., . . . Doyle, A. E. (in press). Executive functions predict dimensions of three neuropsychiatric conditions in youth. *Journal of Child Psychology and Psychiatry*.
- Peterson, R. L., Boada, R., McGrath, L., Willcutt, E. G., Olson, R. K., & Pennington, B. F. (in press). Cognitive prediction of reading, math, and attention: Shared and unique influences. *Journal of Learning Disabilities*.
- Treiman, R., Kessler, B., Pollo, T. C., Byrne, B., & Olson, R. K. (in press). Measures of kindergarten spelling and their relations to later spelling performance. *Scientific Studies of Reading*.
- Wang, Z., Soden, B., Deater-Deckard, K., Lukowski, S., Schenker, V. J., Willcutt, E., . . . Petrill, S. (in press). Development in Reading and Math in Children from Different SES Backgrounds: The Moderating Role of Child Temperament. *Developmental Science*.
- Willcutt, E. G. (in press). Genetics of ADHD. In D. E. Barch (Ed.), *Cognitive and Affective Neuroscience of Psychopathology*. Oxford, UK: Oxford University Press.
- Willcutt, E. G., Leopold, D. R., Christopher, M. E., Olson, R. K., & Petrill, S. A. (in press). ADHD and Learning Disabilities. In A. Zuddas, D. Coghill & T. Banaschewski (Eds.), *Oxford Textbook of ADHD*. Oxford, UK: Oxford Press.
- Knopik, V. S., Niederhiser, J. M., DeFries, J. C., & Plomin, R. (2017). *Behavioral genetics*. New York: Worth Publishers.
- Becker, S. P., Willcutt, E. G., Marshall, S. A., Leopold, D. R., Burns, G. L., Jarrett, M. A., . . . McBurnett, K. (2016). A critical review of the diagnostic validity of sluggish cognitive tempo in child and adolescent psychiatry. *Journal of the American Academy of Child and Adolescent Psychiatry*, 55, 163-178.
- Hartung, C. M., Lefler, E. K., Canu, W. H., Stevens, A. E., Jaconis, M., LaCount, P. A., . . . Willcutt, E. G. (2016). DSM-5 and Other Symptom Thresholds for ADHD: Which Is the Best Predictor of Impairment in College Students? *Journal of Attention Disorders*.
- Plomin, R., DeFries, J. C., Knopik, V. S., & Neiderhiser, J. M. (2016). Top 10 Replicated Findings From Behavioral Genetics. *Perspectives on Psychological Science*, 11, 3-23.
- Powers, N. R., Eicher, J. D., Miller, L. R., Kong, Y., Smith, S. D., Pennington, B. F., . . . Gruen, J. R. (2016). The regulatory element READ1 epistatically influences reading and language, with both deleterious and protective alleles. *Journal of Medical Genetics*, 53, 163-171. PMCID: PMC4789805.

### 2015

- Arnett, A. B., Pennington, B. F., Willcutt, E. G., DeFries, J. C., & Olson, R. K. (2015). Sex differences in ADHD symptom severity. *Journal of Child Psychology and Psychiatry*, 56, 632-639. PMCID: PMC4385512.
- Arnett, A. B., Pennington, B. F., Young, J. F., & Hankin, B. L. (2015). Links between within-person fluctuations in hyperactivity/attention problems and subsequent conduct problems. *Journal of Child Psychology and Psychiatry*.
- Christopher, M. E., Hulsander, J., Keenan, J., DeFries, J., Pennington, B. F., Byrne, B., . . . Wadsworth, S. (2015). The genetic and environmental etiologies of the longitudinal relations between pre-reading skills and reading at the end of first and fourth grades. *Child Development*, 86, 342-361. PMCID: PMC4375099.
- Eicher, J. D., & Gruen, J. R. (2015). Language impairment and dyslexia genes influence language skills in children with autism

- spectrum disorders. *Autism Res*, 8, 229-234. PMCID: PMC4412753.
- Eicher, J. D., Montgomery, A. M., Akshoomoff, N., Amaral, D. G., Bloss, C. S., Libiger, O., . . . Gruen, J. R. (2015). Dyslexia and language impairment associated genetic markers influence cortical thickness and white matter in typically developing children. *Brain Imaging Behav*. PMCID: PMC4639472.
- Eicher, J. D., Stein, C. M., Deng, F., Ciesla, A. A., Powers, N. R., Boada, R., . . . Gruen, J. R. (2015). The DYX2 locus and neurochemical signaling genes contribute to speech sound disorder and related neurocognitive domains. *Genes, Brain, and Behavior*, 14, 377-385. PMCID: PMC4492462.
- Elwér, Å., Gustafson, S., Byrne, B., Olson, R. K., Keenan, J. M., & Samuelsson, S. (2015). A retrospective longitudinal study of cognitive and language skills in poor reading comprehension. *Scandinavian Journal of Psychology*, 56, 157-166. PMCID: PMC4356634.
- Elwér, Å., Gustafson, S., Byrne, B., Olson, R. K., Keenan, J. M., & Samuelsson, S. (2015). Oral language deficits in poor reading comprehension with adequate word decoding *Scandinavian Journal of Psychology*, 56, 157-166. PMCID: PMC4356634.
- Grasby, K. L., Byrne, B., & Olson, R. K. (2015). Validity of large scale reading tests: A phenotypic and behavior-genetic analysis. *Australian Journal of Education*, 59, 5-21.
- Green, T., Bade Shrestha, S., Chromik, L. C., Rutledge, K., Pennington, B. F., Hong, D. S., & Reiss, A. L. (2015). Elucidating X chromosome influences on Attention Deficit Hyperactivity Disorder and executive function. *Journal of Psychiatric Research*, 68, 217-225. PMCID: PMC4528918.
- Keenan, J. M. (2015). Psychology of inferences. In J. D. Wright (Ed.), *International Encyclopedia of the Social & Behavioral Sciences*, 2nd Edition (Vol. 19, pp. 394-399). Oxford, UK: Elsevier.
- Leopold, D. R., Bryan, A. D., Willcutt, E. G., & Pennington, B. F. (2015). Evaluating the construct validity of adult ADHD and SCT among college students: A multitrait-multimethod analysis of convergent and discriminant validity. *Journal of Attention Disorders*, 19, 200-210. PMCID: PMC4334688.
- Lukowski, S. L., DiTrapani, J. B., Jeon, M., Wang, Z., Schenker, V. J., Doran, M. M., . . . Petrill, S. A. (in press). Multidimensionality in the measurement of math-specific anxiety and its relationship with mathematical skill. *Learning and Individual Differences*.
- MacDonald, B., Pennington, B. F., Willcutt, E. G., Dmitrieva, J., Samuelsson, S., Byrne, B., & Olson, R. K. (2015). *Understanding cross-country differences in rates of attention deficit hyperactivity disorder*. Manuscript submitted for publication.
- Peterson, R. L., & Pennington, B. F. (2015). Developmental dyslexia. *Annu Rev Clin Psychol*, 11, 283-307.
- Soden, B., Christopher, M. E., Hulslander, J., Olson, R. K., Cutting, L., Keenan, J. M., . . . Petrill, S. A. (2015). Longitudinal stability in reading comprehension is largely heritable from grades 1 to 6. *PLoS One*, 10, e0113807. PMCID: PMC4300224.
- Wadsworth, S. J., DeFries, J. C., Willcutt, E. G., Pennington, B. F., & Olson, R. K. (2015). The Colorado longitudinal twin study of reading difficulties and ADHD: Etiologies of comorbidity and stability. *Journal of Twin Research and Human Genetics*. PMCID: PMC4684790.
- Willcutt, E. G. (2015). Theories of ADHD. In R. Barkley (Ed.), *Attention Deficit Hyperactivity Disorder: A Clinical Handbook* (4th ed., pp. 391-404). New York: Guilford.
- Willcutt, E. G., Leopold, D. R., Petrill, S. A., Christopher, M., & Olson, R. K. (2015). *Functional and neuropsychological correlates of sluggish cognitive tempo and attention-deficit hyperactivity disorder: A ten year longitudinal study* Manuscript submitted for publication.

## 2014

- Bidwell, L. C., Henry, E. A., Willcutt, E. G., Kinnear, M. K., & Ito, T. A. (2014). Childhood and current ADHD symptom dimensions are associated with risk for more severe cannabis outcomes in college students. *Drug and Alcohol Dependence*, 135, 88-94. PMCID: PMC3904106.
- Brodsky, K., Willcutt, E. G., Davalos, D., & Ross, R. G. (2014). Neuropsychological functioning in childhood-onset psychosis and attention-deficit/hyperactivity disorder. *Journal of Child Psychology and Psychiatry*, 55, 811-818. PMCID: PMC4065620.
- Eicher, J. D., Powers, N. R., Miller, L. L., Mueller, K. L., Mascheretti, S., Marino, C., . . . Gruen, J. R. (2014). Characterization of the DYX2 locus on chromosome 6p22 with reading disability, language impairment, and IQ. *Human Genetics*, 133. PMCID: PMC4053598.
- Gialluisi, A., Newbury, D. F., Willcutt, E. G., Olson, R. K., DeFries, J. C., Bandler, W. M., . . . Fisher, S. E. (2014). Genome-wide screening for DNA variants associated with reading and language traits. *Genes, Brain, and Behavior*. PMCID: PMC4165772.
- Greven, C. U., Kovas, Y., Willcutt, E. G., Petrill, S. A., & Plomin, R. (2014). Evidence for shared genetic risk between ADHD symptoms and reduced mathematics ability: a twin study. *Journal of Child Psychology and Psychiatry*, 55, 39-48. PMCID: PMC3865138.
- Hoffmann, M. S., Polanczyk, G. V., Kieling, C., dos Santos, I. P., Willcutt, E. G., Rohde, L. A., & Salum, G. A. (2014). Attention-Deficit/Hyperactivity Disorder and Solar Irradiance: a cloudy perspective. *Biological Psychiatry*, 76, e19-e20.
- Hua, A. N., & Keenan, J. M. (2014). The role of text memory in inferencing and in comprehension deficits. *Scientific Studies of Reading*, 18, 415-431. PMCID: PMC4196386.
- Keenan, J. M. (2014). Assessment of reading comprehension. In E. R. S. C. A. Stone, B. J. Ehren, & G. P. Wallach (Ed.), *Handbook of Language and Literacy: Development and Disorders* (2nd Edition). New York: Guilford Press.
- Keenan, J. M., Hua, A. N., Meenan, C. E., Olson, R. K., Pennington, B. F., & Willcutt, E. G. (2014). Issues in identifying poor comprehenders. *Topics in Cognitive Psychology*. PMCID: PMC4414263.
- Keenan, J. M., & Meenan, C. E. (2014). Test differences in diagnosing reading comprehension deficits. *Journal of Learning Disabilities*, 47, 125-135. PMCID: PMC3383937.
- Olson, R. K., Keenan, J. M., Byrne, B., & Samuelsson, S. (2014). Why do children differ in their development of reading and related skills? *Scientific Studies of Reading*, 18, 38-54. PMCID: PMC4120985.
- Peterson, R. L., Pennington, B. F., Olson, R. K., & Wadsworth, S. (2014). Longitudinal stability of phonological and surface subtypes of developmental dyslexia. *Scientific Studies of Reading*, 18, 347-362. PMCID: PMC4241299.
- Polanczyk, G., Willcutt, E. G., Salum, G. A., Kieling, C., & Rohde, L. A. (2014). ADHD prevalence estimates across three decades: an updated systematic review and meta-regression analysis. *International Journal of Epidemiology*, 43, 434-442.

- Power, R. A., Nagoshi, C., DeFries, J. C., Plomin, R., & Wellcome Trust Case Control Consortium 2. (2014). Genome-wide estimates of inbreeding in unrelated individuals and their association with cognitive ability. *European Journal of Human Genetics*, 22, 386-390.
- Tamm, L., Epstein, J. N., Denton, C. A., Vaughn, A., Peugh, J., & Willcutt, E. (2014). Reaction time variability associated with reading skills in poor readers with ADHD. *Journal of the International Neuropsychological Society*, 20, 292-301. PMCID: PMC3963392.
- Treiman, R., Gordon, J., Boada, R., Peterson, R. L., & Pennington, B. F. (2014). Statistical learning, letter reversals, and reading. *Scientific Studies of Reading*.
- Wadsworth, S. J., Corley, R. P., & DeFries, J. C. (2014). Cognitive abilities in childhood and adolescence. In D. Finkel & C. A. Reynolds (Eds.), *Behavior Genetics of Cognition Across the Lifespan*. New York, NY: Springer.
- Willcutt, E. G. (2014). Using behavior genetic methods to understand the etiology of comorbidity. In S. Rhee & A. Ronald (Eds.), *Behavior Genetics of Psychopathology* (pp. 231-252). New York: Springer.
- Willcutt, E. G., Chhabildas, N., Kinnear, M., Defries, J. C., Olson, R. K., Leopold, D. R., . . . Pennington, B. F. (2014). The internal and external validity of sluggish cognitive tempo and its relation with DSM-IV ADHD. *Journal of Abnormal Child Psychology*, 42, 21-35. PMCID: PMC3947432.
- Wu, S. S., Mittal, V., Pennington, B. F., & Willcutt, E. G. (2014). Mathematics achievement scores and early psychosis in school-aged children. *Schizophrenia Research*, 133-134.
- Wu, S. S., Willcutt, E. G., Escobar, E., & Menon, V. (2014). Mathematics achievement and anxiety and their relation to internalizing and externalizing behaviors. *Journal of Learning Disabilities*, 47, 503-514. PMCID: PMC3883980.

## 2013

- Arnett, A. B., Pennington, B. F., Friend, A., Willcutt, E. G., Byrne, B., Samuelsson, S., & Olson, R. K. (2013). The SWAN captures variance at both the negative and positive ends of the ADHD symptom spectrum. *Journal of Attention Disorders*, 17, 152-162. PMCID: PMC333013.
- Brant, A. M., Munakata, Y., Boomsma, D. I., DeFries, J. C., Haworth, C. M., Keller, M. C., . . . Hewitt, J. K. (2013). The nature and nurture of high IQ: an extended sensitive period for intellectual development. *Psychological Science*, 24, 1487-1495. PMCID: PMC4511162.
- Byrne, B., Samuelsson, S., & Olson, R. K. (2013). Reading and reading acquisition in European languages. In P. P. a. H. Winskel (Ed.), *South and Southeast Asian Psycholinguistics* (pp. 159-170). Cambridge University Press.
- Byrne, B., Samuelsson, S., & Olson, R. K. (2013). Dyslexia. In A. J. Holliman (Ed.), *The Routledge international companion to educational psychology* (pp. 297-306). Oxford, UK: Routledge.
- Byrne, B., Samuelsson, S., & Olson, R. K. (2013). Reading and reading acquisition in European languages. In P. Padakanaya & H. Winskel (Eds.), *South and Southeast Asian Psycholinguistics* (pp. 159-170). Cambridge, UK: Cambridge University Press.
- Byrne, B., Wadsworth, S. J., Boehme, K., Talk, A. C., Coventry, W. L., Olson, R. K., . . . Corley, R. (2013). Multivariate genetic analysis of learning and early reading development. *Scientific Studies of Reading*, 17, 224-242. PMCID: PMC3633536.
- Christopher, M. E., Hulslander, J., Byrne, B., Samuelsson, S., Keenan, J. M., Pennington, B. F., . . . Olson, R. K. (2013). The genetic and environmental etiologies of individual differences in early reading growth in Australia, the United States, and Scandinavia. *Journal of Experimental Child Psychology*, 125, 453-467. PMCID: PMC3661747.
- Christopher, M. E., Hulslander, J., Byrne, B., Samuelsson, S., Keenan, J. M., Pennington, B. F., . . . Olson, R. K. (2013). Modeling the etiology of individual differences in early reading development: Evidence for strong genetic influences. *Scientific Studies of Reading*, 17, 350-358. PMCID: PMC3905458.
- Eicher, J. D., & Gruen, J. R. (2013). Imaging-genetics in dyslexia: connecting risk genetic variants to brain neuroimaging and ultimately to reading impairments. *Mol Genet Metab*, 110, 201-212. PMCID: PMC3800223.
- Eicher, J. D., Powers, N. R., Cho, K., Miller, L. L., Mueller, K. L., Ring, S. M., . . . Gruen, J. R. (2013). Associations of prenatal nicotine exposure and the dopamine related genes ANKK1 and DRD2 to verbal language. *PLoS One*, 8, e63762. PMCID: PMC3655151.
- Eicher, J. D., Powers, N. R., Miller, L. L., Akshoomoff, N., Amaral, D. G., Bloss, C. S., . . . Gruen, J. R. (2013). Genome-wide association study of shared components of reading disability and language impairment. *Genes, Brain, and Behavior*, 12, 792-801. PMCID: PMC3904347.
- Elwer, S., Keenan, J. M., Olson, R. K., Byrne, B., & Samuelsson, S. (2013). Longitudinal stability and predictors of poor oral comprehenders and poor decoders. *Journal of Experimental Child Psychology*, 115, 497-516. PMCID: PMC3661702.
- Fujisawa, K. K., Wadsworth, S. J., Kakihana, S., Olson, R. K., DeFries, J. C., Byrne, B., & Ando, J. (2013). A multivariate twin study of early literacy in Japanese Kana. *Learning and Individual Differences*, 24, 160-167. PMCID: PMC3753202.
- Miller, A. C., Keenan, J. M., Betjemann, R. S., Willcutt, E. G., Pennington, B. F., & Olson, R. K. (2013). Reading comprehension in children with ADHD: Cognitive underpinnings of the centrality deficit. *Journal of Abnormal Child Psychology*, 41, 473-483. PMCID: PMC3121914.
- O'Brien, B. A., Van Orden, G., & Pennington, B. F. (2013). Do Dyslexics Misread a ROWS for a ROSE? *Read Writ*, 26, 381-402. PMCID: PMC4004072.
- Olson, R. K., Hulslander, J., Christopher, M., Keenan, J. M., Wadsworth, S. J., Willcutt, E. G., . . . DeFries, J. C. (2013). Genetic and environmental influences on writing and their relations to language and reading. *Annals of Dyslexia*. PMCID: PMC2182316.
- Peterson, R. L., Pennington, B. F., & Olson, R. K. (2013). Subtypes of developmental dyslexia: testing the predictions of the dual-route and connectionist frameworks. *Cognition*, 126, 20-38. PMCID: PMC3491101.
- Peterson, R. L., Pennington, B. F., Samuelsson, S., Byrne, B., & Olson, R. K. (2013). Shared etiology of phonological memory and vocabulary deficits in school-age children. *Journal of Speech, Language, and Hearing Research*, 56, 1249-1259. PMCID: PMC3615120.
- Petrill, S. A., Bartlett, C. W., & Blair, C. (2013). Gene-environment interplay in child psychology and psychiatry--challenges and ways forward. *Journal of Child Psychology and Psychiatry*, 54, 1029.
- Plomin, P., DeFries, J. C., Knopik, V. S., & Neiderhiser, J. M. (2013). *Behavioral Genetics (Sixth Edition)*. New York, NY: Worth

Publishers.

- Powers, N. R., Eicher, J. D., Butter, F., Kong, Y., Miller, L. L., Ring, S. M., . . . Gruen, J. R. (2013). Alleles of a polymorphic ETV6 binding site in DCDC2 confer risk of reading and language impairment. *American Journal of Human Genetics*, 93, 19-28. PMCID: PMC3710765.
- Rhea, S. A., Bricker, J. B., Corley, R. P., DeFries, J. C., & Wadsworth, S. J. (2013). Design, utility, and history of the Colorado Adoption Project: Examples involving adjustment interactions. *Adoption Quarterly*, 16, 17-39.
- Trzaskowski, M., Davis, O. S. P., DeFries, J. C., Yang, J., Visscher, P. M., & Plomin, R. (2013). DNA evidence for strong genome-wide pleiotropy of cognitive and learning abilities. *Behavior Genetics*, 43, 267-273.
- Willcutt, E. G., Petrill, S. A., Wu, S., Boada, R., DeFries, J. C., Olson, R. K., & Pennington, B. F. (2013). Implications of comorbidity between reading and math disability: Neuropsychological and functional impairment. *Journal of Learning Disabilities*, 46, 500-516. PMCID: PMC3749272.
- Wolock, S. L., Yates, A., Petrill, S. A., Bohland, J. W., Blair, C., Li, N., . . . Bartlett, C. W. (2013). Gene  $\times$  smoking interactions on human brain gene expression: finding common mechanisms in adolescents and adults. *Journal of Child Psychology and Psychiatry*, 54, 1109-1119. PMCID: PMC3809890.

**2012**

- Anthoni, H., Sucheston, L. E., Lewis, B. A., Fan, X., Zucchelli, M., Tapia-Páez, I., . . . Kere, J. (2012). The aromatase gene CYP19A1: Several genetic and functional lines of evidence supporting a role in reading, speech, and language. *Behavior Genetics*, 42, 509-527. PMCID: PMC3375077.
- Arnett, A. B., Pennington, B. F., Willcutt, E. G., Dmitrieva, J., Byrne, B., Samuelsson, S., & Olson, R. K. (2012). A cross-lagged model of the development of ADHD inattention symptoms and rapid naming speed. *Journal of Abnormal Child Psychology*, 40, 1313-1326. PMCID: PMC3546520.
- Astrom, R. L., Wadsworth, S. J., Olson, R. K., Willcutt, E. G., & DeFries, J. C. (2012). Genetic and environmental etiologies of reading difficulties: DeFries-Fulker analysis of reading performance data from twin pairs and their non-twin siblings. *Learning and Individual Differences*. PMCID: PMC3423974.
- Boada, R., Pennington, B. F., & Willcutt, E. G. (2012). Understanding the comorbidity between dyslexia and attention deficit/hyperactivity disorder. *Topics in Language Disorders*, 32, 264-284.
- Christopher, M. E., Miyake, A., Keenan, J. M., Pennington, B. F., DeFries, J. C., Wadsworth, S. J., . . . Olson, R. K. (2012). Predicting word reading and comprehension with executive function and speed measures: A latent variable analysis. *Journal of Experimental Psychology: General*, 141, 470-488. PMCID: PMC3360115.
- Cope, N., Eicher, J. D., Meng, H., Gibson, C. J., Hager, K., Lacadie, C., . . . Gruen, J. R. (2012). Variants in the DYX2 locus are associated with altered brain activation in reading-related brain regions in subjects with reading disability. *Neuroimage*, 63, 148-156. PMCID: PMC3518451.
- Keenan, J. M. (2012). Measure for measure: Challenges in assessing reading comprehension. In E. A. J. Sabatini, & T. O'Reilly (Ed.), *Measuring Up: Advances in How We Assess Reading Ability*. Lanham, MD: Rowman & Littlefield.
- Pennington, B. F., Santerre-Lemmon, L., Rosenberg, J., MacDonald, B., Boada, R., Friend, A., . . . Olson, R. K. (2012). Individual prediction of dyslexia by single versus multiple deficit models. *Journal of Abnormal Psychology*, 121, 212-224. PMCID: PMC3270218.
- Peterson, R. L., & Pennington, B. F. (2012). Developmental dyslexia. *Lancet*, 379, 1997-2007. PMCID: PMC3465717.
- Petrill, S., Logan, J., Hart, S., Vincent, P., Thompson, L., Kovas, Y., & Plomin, R. (2012). Math fluency is etiologically distinct from untimed math performance, decoding fluency, and untimed reading performance: evidence from a twin study. *Journal of Learning Disabilities*, 45, 371-381. PMCID: PMC3413280.
- Rosenberg, J., Pennington, B. F., Willcutt, E. G., & Olson, R. K. (2012). Gene by environment interactions influencing reading disability and the inattentive symptom dimension of attention deficit/hyperactivity disorder. *Journal of Child Psychology and Psychiatry*, 53, 243-251. PMCID: PMC3235245.
- Wadsworth, S. J., Olson, R. K., Willcutt, E. G., & DeFries, J. C. (2012). Multiple regression analysis of reading performance data from twin pairs with reading difficulties and nontwin siblings: The augmented model. *Twin Research and Human Genetics*, 15, 116-119. PMCID: PMC3398740.
- Willcutt, E. G. (2012). The prevalence of DSM-IV attention-deficit/hyperactivity disorder: a meta-analytic review. *Neurotherapeutics*, 9, 490-499. PMCID: PMC3441936.
- Willcutt, E. G., Nigg, J. T., Pennington, B. F., Solanto, M. V., Rohde, L. A., Tannock, R., . . . Lahey, B. B. (2012). Validity of DSM-IV attention-deficit/hyperactivity disorder dimensions and subtypes. *Journal of Abnormal Psychology*, 121, 991-1010. PMCID: PMC3622557.

**2011**

- Astrom, R. L., Wadsworth, S. J., Olson, R. K., Willcutt, E. G., & DeFries, J. C. (2011). DeFries-Fulker analysis of longitudinal reading performance data from twin pairs ascertained for reading difficulties and from their nontwin siblings. *Behavior Genetics*, 41, 660-667. PMCID: PMC3085049.
- Betjemann, R. S., Keenan, J. M., Olson, R. K., & DeFries, J. C. (2011). Choice of reading comprehension test influences the outcomes of genetic analyses. *Scientific Studies of Reading*, 15, 363-382. PMCID: PMC3143485.
- Bidwell, L. C., Willcutt, E. G., McQueen, M. B., DeFries, J. C., Olson, R. K., Smith, S. D., & Pennington, B. F. (2011). A family based association study of DRD4, DAT1, and 5HTT and continuous traits of attention-deficit hyperactivity disorder. *Behavior Genetics*, 41, 165-174. PMCID: PMC3674022.
- Coventry, W. L., Byrne, B., Olson, R. K., Corley, R., & Samuelsson, S. (2011). Dynamic and static assessment of phonological awareness in preschool: A behavior genetic study. *Journal of Learning Disabilities*, 44, 322-329. PMCID: PMC3158276.
- Johnson, E. P., Pennington, B. F., Lowenstein, L. H., & Nittrouer, S. (2011). Sensitivity to structure in the speech signal by children with

- speech sound disorder and reading disability. *Journal of Communication Disorders*, 44, 294-314. PMCID: PMC3095671.
- Marino, C., Mascheretti, S., Riva, V., Cattaneo, F., Rigoleto, C., Rusconi, M., . . . Molteni, M. (2011). Pleiotropic effects of DCDC2 and DYX1C1 genes on language and mathematics traits in nuclear families of developmental dyslexia. *Behavior Genetics*, 41, 67-76. PMCID: PMC3939676.
- McGrath, L. M., Pennington, B. F., Shanahan, M. A., Santerre-Lemmon, L. E., Barnard, H. D., Willcutt, E. G., . . . Olson, R. K. (2011). A multiple deficit model of reading disability and attention-deficit/hyperactivity disorder: Searching for shared cognitive deficits. *Journal of Child Psychology and Psychiatry*, 52, 547-557. PMCID: PMC3079018.
- Meng, H., Powers, N. R., Tang, L., Cope, N. A., Zhang, P. X., Fulkehan, R., . . . Gruen, J. R. (2011). A dyslexia-associated variant in DCDC2 changes gene expression. *Behavior Genetics*, 41, 58-66. PMCID: PMC3053575.
- Miller, A. C., & Keenan, J. M. (2011). Understanding the centrality deficit: insight from foreign language learners. *Memory & Cognition*, 39, 873-883. PMCID: PMC3121914.
- Nilsson, R., Rhee, S. H., Corley, R. P., Rhea, S. A., Wadsworth, S. J., & DeFries, J. C. (2011). Conduct Problems in Adopted and Non-Adopted Adolescents and Adoption Satisfaction as a Protective Factor. *Adoption Quarterly*, 14, 181-198. PMCID: PMC3259118.
- Olson, R. K. (2011). Genetic and environmental influences on phonological abilities and reading achievement. In S. Brady, D. Braze & C. Fowler (Eds.), *Explaining individual differences in reading: theory and evidence*. New York: Psychology Press / Taylor Publishing.
- Olson, R. K. (2011). Evaluation of Fast ForWord Language effects on language and reading. *Perspectives on Language and Literacy*, 37, 11-15. NIHMSID: NIHMS287175.
- Olson, R. K., Keenan, J. M., Byrne, B., Samuelsson, S., Coventry, W. L., Corley, R., . . . Hulslander, J. (2011). Genetic and environmental influences on vocabulary and reading development from pre-kindergarten through grade 4. *Scientific Studies of Reading*, 15, 26-46. PMCID: PMC3019615.
- Priebe, S. J., Keenan, J. M., & Miller, A. C. (2011). How prior knowledge affects word identification and comprehension. *Reading and Writing*, 7, 581-586. PMCID: PMC3142886.
- Rivkees, S. A., Hager, K., Hosono, S., Wise, A., Li, P., Rinder, H. M., & Gruen, J. R. (2011). A highly sensitive, high-throughput assay for the detection of Turner syndrome. *Journal of Clinical Endocrinology and Metabolism*, 96, 699-705. PMCID: PMC3047225.
- Willcutt, E. G., & Bidwell, L. C. (2011). Etiology of ADHD: Implications for assessment and treatment. In B. Hoza & S. W. Evans (Eds.), *Treating Attention Deficit Hyperactivity Disorder* (pp. 6-2 - 6-18). Kingston, NJ: Civic Research Institute.
- Willcutt, E. G., Boada, R., Riddle, M. W., Chhabildas, N., DeFries, J. C., & Pennington, B. F. (2011). Colorado Learning Difficulties Questionnaire: Validation of a parent-report screening measure. *Psychological Assessment*, 23, 778-791. PMCID: PMC21574721.

## 2010

- Betjemann, R. S., Johnson, E. P., Barnard, H., Boada, R., Filley, C. M., Filipek, P. A., . . . Pennington, B. F. (2010). Genetic covariation between brain volumes and IQ, reading performance, and processing speed. *Behavior Genetics*, 40, 135-145. PMCID: PMC3608477.
- Burgess, G. C., Depue, B. E., Ruzic, L., Willcutt, E. G., Du, Y. P. P., & Banich, M. T. (2010). Attentional control activation relates to working memory in Attention-Deficit/Hyperactivity Disorder. *Biological Psychiatry*, 67, 632-640. PMCID: PMC2953472.
- Byrne, B., Coventry, W. L., Olson, R. K., Wadsworth, S. J., Samuelsson, S., Petrill, S. A., . . . Corley, R. (2010). "Teacher Effects" in early literacy development: Evidence from a study of twins. *Journal of Educational Psychology*, 102, 32-42. PMCID: PMC2830009.
- Byrne, B., Khlebtzos, D., Olson, R. K., & Samuelsson, S. (2010). Evolutionary and genetic perspectives on educational achievement. In K. Littleton, C. Wood & J. K. Staarman (Eds.), *Handbook of educational psychology* (pp. 3-34). Bingley, UK: Emerald Press.
- DeFries, J. C. (2010). Haseman and Elston sib-pair linkage analysis: a brief historical note. *Behavior Genetics*, 40, 1-2.
- Depue, B. E., Burgess, G. C., Bidwell, L. C., Willcutt, E. G., & Banich, M. T. (2010). Behavioral performance predicts grey matter reductions in the right inferior frontal gyrus in young adults with combined type ADHD. *Psychiatry Research-Neuroimaging*, 182, 231-237. PMCID: PMC2914826.
- Depue, B. E., Burgess, G. C., Willcutt, E. G., Bidwell, L. C., Ruzic, L., & Banich, M. T. (2010). Symptom-correlated brain regions in young adults with combined-type ADHD: Their organization, variability, and relation to behavioral performance. *Psychiatry Research: Neuroimaging*, 182, 96-102. PMCID: PMC2878757.
- Depue, B. E., Burgess, G. C., Willcutt, E. G., Ruzic, L., & Banich, M. T. (2010). Inhibitory control of memory retrieval and motor processing associated with the right lateral prefrontal cortex: evidence from deficits in individuals with ADHD. *Neuropsychologia*, 48, 3909-3917. PMCID: PMC2979319.
- Ebejer, J. L., Coventry, W. L., Byrne, B., Willcutt, E. G., Olson, R. K., Corley, R., & Samuelsson, S. (2010). Genetic and environmental influences on inattention, hyperactivity-impulsivity, and reading: Kindergarten to grade 2. *Scientific Studies of Reading*, 14, 293-316. PMCID: PMC2930267.
- Edgin, J. O., Pennington, B. F., & Mervis, C. B. (2010). Neuropsychological components of intellectual disability: the contributions of immediate, working, and associative memory. *Journal of Intellectual Disabilities Research*, 54, 406-417. PMCID: PMC3088787.
- Friend, A., & Olson, R. K. (2010). Phonological spelling and reading deficits in children with spelling disabilities. *Scientific Studies of Reading*, 12, 90-105. PMCID: PMC2892167.
- Gabel, L. A., Gibson, C. J., Gruen, J. R., & LoTurco, J. J. (2010). Progress towards a cellular neurobiology of reading disability. *Neurobiology of Disease*, 38, 173-180. PMCID: PMC2854314.
- Hart, S. A., Petrill, S. A., Willcutt, E. G., Thompson, L. A., Schatschneider, C., Deater-Deckard, K., & Cutting, L. E. (2010). Exploring how ADHD symptoms are related to reading and mathematics performance: General genes, general environments. *Psychological Science*, 21, 1708-1715. PMCID: PMC3708699.
- Haworth, C. M. A., Wright, M. J., Luciano, M., Martin, N. G., de Geus, E. J. C., van Beijsterveldt, C. E. M., . . . Plomin, R. (2010). The

- heritability of general cognitive ability increases linearly from childhood to young adulthood. *Molecular Psychiatry*, 15, 1112-1120. PMCID: PMC2889158.
- Hulslander, J., Olson, R. K., Willcutt, E. G., & Wadsworth, S. J. (2010). Longitudinal stability of reading-related skills and their prediction of reading development. *Scientific Studies of Reading*, 14, 111-136. PMCID: PMC2885806.
- Lahey, B. B., & Willcutt, E. G. (2010). Predictive validity of a continuous alternative to nominal subtypes of attention-deficit hyperactivity disorder in DSM-IV. *Journal of Clinical Child and Adolescent Psychology*, 39, 761-775. PMCID: PMC2979319.
- Miller, A. C., & Keenan, J. M. (in press). Understanding the centrality deficit: Insight from foreign language learners. *Memory and Cognition*. NIHMSID: NIHMS287169.
- Nittrouer, S., & Pennington, B. F. (2010). New approaches to the study of childhood language disorders. *Current Directions in Psychological Science*, 19, 308-313. PMCID: PMC3374334.
- Peterson, R. L., & Pennington, B. F. (2010). Reading disability. In K. O. Yeates, G. Taylor, D. O. Ris & B. Pennington, F. (Eds.), *Pediatric Neuropsychology: Research, theory, and practice* (pp. 324-362). New York: Guilford Press.
- Raitano Lee, N., Pennington, B. F., & Keenan, J. M. (2010). Verbal short term memory deficits in Down syndrome: phonological, semantic or both? *Journal of Neurodevelopmental Disorders*, 2, 9-25.
- Scerri, T. S., Paracchini, S., Morris, A., MacPhie, I. L., Talcott, J., Stein, J., . . . Richardson, A. J. (2010). Identification of candidate genes for dyslexia susceptibility on chromosome 18. *PLoS One*, 5, e13712. PMCID: PMC2965662.
- Sergeant, J., Willcutt, E. G., & Nigg, J. T. (2010). How clinically functional are executive function measures of ADHD? In D. Shaffer, E. Leibenluft, L. A. Rohde, P. Sirovatka & D. A. Regier (Eds.), *Externalizing disorders of childhood: Refining the research agenda for DSM-V*. Arlington, VA: American Psychiatric Association.
- Smith, S. D. (2010). Human genetic contributions to the neurobiology of dyslexia. In M. McCardle, J. Ren & O. Tzeng (Eds.), *Dyslexia across languages: Orthography and the brain-gene-behavior link*. Baltimore, MD: Brookes Publishing Company.
- Smith, S. D. (2010). Learning Disabilities. In J. Nurnberger & W. Berrettini (Eds.), *Psychiatric Genetics*. Cambridge, UK: Cambridge University Press.
- Smith, S. D., Grigorenko, E., Willcutt, E. G., Pennington, B. F., & Olson, R. K. (2010). Etiologies and molecular mechanisms of communication disorders. *Journal of Developmental and Behavioral Pediatrics*, 31, 555-563. PMCID: PMC2943674.
- Wadsworth, S. J., Olson, R. K., & DeFries, J. C. (2010). Differential genetic etiology of reading difficulties as a function of IQ: An update. *Behavior Genetics*, 40, 751-758.
- Willcutt, E. G., Betjemann, R. S., McGrath, L. M., Chhabildas, N. A., Olson, R. K., DeFries, J. C., & Pennington, B. F. (2010). Etiology and neuropsychology of comorbidity between RD and ADHD: The case for multiple-deficit models. *Cortex*, 46, 1345-1361. PMCID: PMC2993430.
- Willcutt, E. G., & McQueen, M. B. (2010). Genetic and environmental risk factors for bipolar spectrum disorders. In D. J. Miklowitz & D. V. Cicchetti (Eds.), *The Developmental Psychopathology of Bipolar Disorder* (pp. 225-258).
- Willcutt, E. G., Pennington, B. F., Duncan, L., Smith, S. D., Keenan, J. M., Wadsworth, S. J., & DeFries, J. C. (2010). Understanding the complex etiology of developmental disorders: Behavioral and molecular genetic approaches. *Journal of Developmental and Behavioral Pediatrics*, 31, 533-544. PMCID: PMC2953861.
- Yeates, K. O., Ris, M. D., Taylor, H. G., & Pennington, B. F. (2010). *Pediatric Neuropsychology: Research, theory, and practice*. New York: Guilford.
- Zhang, H., Gelernter, J., Gruen, J. R., Kranzler, H. R., Herman, A. I., & Simen, A. A. (2010). Functional impact of a single-nucleotide polymorphism in the OPRD1 promoter region. *Journal of Human Genetics*, 55, 278-284. PMCID: PMC2876206.

## 2009

- Banich, M. T., Burgess, G. C., Depue, B. E., Ruzic, L., Bidwell, L. C., Hitt-Laustsen, S., . . . Willcutt, E. G. (2009). The neural basis of sustained and transient attentional control in young adults with ADHD. *Neuropsychologia*, 47, 3095-3104. PMCID: PMC3703501.
- Boada, R., Janusz, J., Hutaff-Lee, C., & Tartaglia, N. (2009). The cognitive phenotype in Klinefelter syndrome: A review of the literature including genetic and hormonal factors. *Developmental Disabilities Research Reviews*, 15, 284-294. PMCID: PMC3056507.
- Brant, A. M., Haberstick, B. C., Corley, R. P., Wadsworth, S. J., DeFries, J. C., & Hewitt, J. K. (2009). The developmental etiology of high IQ. *Behavior Genetics*, 39, 393-405. NIHMSID: NIHMS284517.
- Byrne, B., Coventry, W. L., Olson, R. K., Samuelsson, S., Corley, R., Willcutt, E. G., . . . DeFries, J. C. (2009). Genetic and environmental influences on aspects of literacy and language in early childhood: Continuity and change from preschool to grade 2. *Journal of Neurolinguistics*, 22, 219-236. PMCID: PMC2724015.
- Cardoso-Martins, C., Pennington, B. F., Peterson, R., & Olson, R. K. (2009). Component reading skills in Down syndrome. *Reading and Writing*, 22, 277-292.
- Coventry, W. L., Byrne, B., Coleman, M., Olson, R. K., Corley, R., Willcutt, E., & Samuelsson, S. (2009). Does classroom separation affect twins' reading ability in the early years of school? *Twin Research and Human Genetics*, 12, 455-461. PMCID: PMC19803773.
- Friend, A., DeFries, J. C., Olson, R. K., Pennington, B. F., Harlaar, N., Byrne, B., . . . Keenan, J. M. (2009). Heritability of high reading ability and its interaction with parental education. *Behavior Genetics*, 39, 427-436. PMCID: PMC3387983.
- Hawke, J. L., Olson, R. K., Willcutt, E. G., Wadsworth, S. J., & DeFries, J. C. (2009). Gender ratios for reading difficulties. *Dyslexia*, 15, 239-242. PMCID: PMC2739722.
- Johnson, E. P., Pennington, B. F., Lee, N. R., & Boada, R. (2009). Directional effects between rapid auditory processing and phonological awareness in children. *Journal of Child Psychology and Psychiatry*, 50, 902-910.
- Keenan, J. M., Olson, R. K., & Betjemann, R. S. (2009). Assessment and etiology of individual differences in reading comprehension. In R. Wagner, C. Schatschneider & C. Phythian-Sence (Eds.), *Beyond decoding: The biological and behavioral foundations of reading comprehension* (pp. 227-245). New York: Guilford Press.
- Miller, A. C., & Keenan, J. M. (2009). How word decoding skill impacts text memory: The centrality deficit and how domain knowledge can compensate. *Annals of Dyslexia*, 59, 99-113. PMCID: PMC2887733.

- Olson, R. K., Byrne, B., & Samuelsson, S. (2009). Reconciling strong genetic and strong environmental influences on individual differences and deficits in reading ability. In K. R. Pugh & P. McCardle (Eds.), *How Children Learn To Read: Current Issues and New Directions in the Integration of Cognition, Neurobiology and Genetics of Reading and Dyslexia Research and Practice* (pp. 215-233). New York: Lawrence Erlbaum/Taylor-Francis.
- Pennington, B. F. (2009). How neuropsychology informs our understanding of developmental disorders. *Journal of Child Psychology and Psychiatry*, 50, 72-78.
- Pennington, B. F. (2009). *Diagnosing learning disorders* (2nd ed.). New York, NY: Guilford Press.
- Pennington, B. F., & Bishop, D. V. M. (2009). Relations Among Speech, Language, and Reading Disorders. *Annual Review of Psychology*, 60, 283-306.
- Pennington, B. F., McGrath, L. M., Rosenberg, J., Barnard, H., Smith, S. D., Willcutt, E. G., . . . Olson, R. K. (2009). Gene X environment interactions in reading disability and attention-deficit/hyperactivity disorder. *Developmental Psychology*, 45, 77-89. PMCID: PMC2743891.
- Pennington, B. F., McGrath, L. M., & Smith, S. D. (2009). Genetics of dyslexia: Cognitive analysis, candidate genes, comorbidities, and etiologic interactions. In T. E. Goldberg & D. Weinberger (Eds.), *Genetics of cognitive neuroscience* (pp. 177-193). Cambridge, MA: MIT Press.
- Peterson, R. L., Pennington, B. F., Shriberg, L. D., & Boada, R. (2009). What influences literacy outcome in children with speech sound disorder? *Journal of Speech, Language, and Hearing Research*, 52, 1175-1188. PMCID: PMC3608470.
- Phinney, E., Pennington, B. F., Raitano-Lee, N., & Boada, R. (2009). Directional effects between rapid auditory processing and phonological awareness in children. *Journal of Clinical Child and Adolescent Psychology*, 50, 902-910.
- Rice, M., Smith, S. D., & Gayán, J. (2009). Convergent genetic linkage and associations to language, speech and reading measures in families of probands with Specific Language Impairment. *Journal of Neurodevelopmental Disorders*, 1, 264-282. PMCID: PMC2788915.
- Willcutt, E. G. (2009). ADHD. In K. O. Yeats, D. O. Ris, G. Taylor & B. F. Pennington (Eds.), *Pediatric Neuropsychology: Research, Theory, and Practice* (pp. 393-417). New York: Guilford Press.
- Wise, B. W., Rogan, L., & Sessions, L. (2009). Training teachers in evidence-based intervention: The story of Linguistic Remedies. In S. Rosenfield & V. Berninger (Eds.), *Translating science-supported instruction into evidence-based practices: Understanding and applying the implementation process*. Oxford: Oxford University Press.
- Young, S. E., Friedman, N. P., Miyake, A., Willcutt, E. G., Corley, R. P., Haberstick, B. C., & Hewitt, J. K. (2009). Behavioral disinhibition: liability for externalizing spectrum disorders and its genetic and environmental relation to response inhibition across adolescence. *Journal of Abnormal Psychology*, 118, 117-130. PMCID: PMC2775710.
- Zhang, H., Kranzler, H. R., Poling, J., Gruen, J. R., & Gelernter, J. (2009). Cognitive flexibility is associated with KIBRA variant and modulated by recent tobacco use. *Neuropsychopharmacology*, 34, 2508-2516. PMCID: PMC2898508.
- Betjemann, R. S., & Keenan, J. M. (2008). Phonological and semantic priming in children with reading disability. *Child Development*, 79, 1086-1102. PMCID: Accepted for publication before April 7, 2008.
- Betjemann, R. S., Willcutt, E. G., Olson, R. K., Keenan, J. M., DeFries, J. C., & Wadsworth, S. J. (2008). Word reading and reading comprehension: stability, overlap, and independence. *Reading and Writing*, 21, 539-558. PMCID: Accepted for publication before April 7, 2008.
- Brojde, C., & Wise, B. W. (2008). An evaluation of the testing effect with third grade students. In B. C. Love, K. McRae & V. M. Sloutsky (Eds.), *Proceedings of the 30th Annual Conference of the Cognitive Science Society* (pp. 1362-1367). Washington, DC: Cognitive Science Society.
- Byrne, B., Coventry, W. L., Olson, R. K., Hulslander, J., Wadsworth, S., DeFries, J. C., . . . Samuelsson, S. (2008). A behavior-genetic analysis of orthographic learning, spelling, and decoding. *Journal of Research in Reading*, 31, 8-21. PMCID: Accepted for publication before April 7, 2008.
- Coventry, W. L., Byrne, B., Olson, R. K., Samuelsson, S., Corley, R., Wadsworth, S., & DeFries, J. C. (2008). Do the genetic effects for literacy in early childhood differ across sex or across the disabled and normal range? [Abstract]. *Behavior Genetics*, 38, 619.
- Doyle, A. E., Ferreira, M. A., Sklar, P. B., Lasky-Su, J., Petty, C., Fusillo, S. J., . . . Faraone, S. V. (2008). Multivariate genome-wide linkage scan of neurocognitive traits and ADHD symptoms: suggestive linkage to 3q13. *American Journal of Medical Genetics Part B (Neuropsychiatric Genetics)*, 147B, 1399-1411. PMCID: PMC4002289.
- Friedman, N. P., Miyake, A., Young, S. E., DeFries, J. C., Corley, R. P., & Hewitt, J. K. (2008). Individual differences in executive functions are almost entirely genetic in origin. *Journal of Experimental Psychology: General*, 137, 201-225.

## 2008

- Friend, A., DeFries, J. C., & Olson, R. K. (2008). Parental education moderates genetic influences on reading disability. *Psychological Science*, 19, 1124-1130. PMCID: PMC2605635.
- Friend, A., & Olson, R. K. (2008). Phonological spelling and reading deficits in children with spelling disabilities. *Scientific Studies of Reading*, 12, 90-105. PMCID: Accepted for publication before April 7, 2008.
- Gibson, C. J., & Gruen, J. R. (2008). The human lexinome: genes of language and reading. *Journal of Communication Disorders*, 41, 409-420. PMCID: PMC2488410.
- Hawke, J. L., Stallings, M. C., Wadsworth, S. J., & DeFries, J. C. (2008). DeFries-Fulker and Pearson-Aitken model-fitting analyses of reading performance data from selected and unselected twin pairs. *Behavior Genetics*, 38, 101-107. PMCID: Accepted for publication before April 7, 2008.
- Keenan, J. M. (2008). How the components of reading comprehension differ with age and test. In E. Albro, J. Sabatini, T. O'Reilly & M. Ventura (Eds.), *Working papers on assessing reading in the 21st century*. Princeton, NJ: Educational Testing Services.
- Keenan, J. M., Betjemann, R. S., & Olson, R. K. (2008). Reading comprehension tests vary in the skills they assess: Differential dependence on decoding and oral comprehension. *Scientific Studies of Reading*, 12, 281-300. PMCID: Accepted for publication before April 7, 2008.
- Ludwig, K. U., Schumacher, J., Schulte-Körne, G., König, I. R., Warnke, A., Plume, E., . . . Hoffmann, P. (2008). Investigation of the

- DCDC2 intron 2 deletion/compound short tandem repeat polymorphism in a large German dyslexia sample. *Psychiatric Genetics*, 18, 310-312.
- Meda, S. A., Gelernter, J., Gruen, J. R., Calhoun, V. D., Meng, H., Cope, N. A., & Pearlson, G. D. (2008). Polymorphism of DCDC2 reveals differences in cortical morphology of healthy individuals - a preliminary voxel based morphometry study. *Brain Imaging and Behavior*, 2, 21-26.
- Pennington, B. F. (2008). *Diagnosing Learning Disorders (2nd Edition)*. New York: Guilford Press.
- Plomin, R., DeFries, J. C., McClearn, G. E., & McGuffin, P. (2008). *Behavioral Genetics* (Vol. Fifth). New York: Worth Publishers.
- Rhee, S. H., Willcutt, E. G., Hartman, C. A., Pennington, B. F., & DeFries, J. C. (2008). Test of alternative hypotheses explaining the comorbidity between attention-deficit/hyperactivity disorder and conduct disorder. *Journal of Abnormal Child Psychology*, 36, 29-40. PMCID: Accepted for publication before April 7, 2008.
- Samuelsson, S., Byrne, B., Olson, R. K., Hulslander, J., Wadsworth, S., Corley, R., . . . DeFries, J. C. (2008). Response to early literacy instruction in the United States, Australia, and Scandinavia: A behavioral-genetic analysis. *Learning and Individual Differences*, 18, 289-295. PMCID: PMC2570222.
- Schmidt, G. L., Kimel, L. K., Winterrowd, E., Pennington, B. F., Hepburn, S. L., & Rojas, D. C. (2008). Impairments in phonological processing and nonverbal intellectual function in parents of children with autism. *Journal of Clinical and Experimental Neuropsychology*, 30, 557-567.
- Shanahan, M. A., Pennington, B. F., & Willcutt, E. G. (2008). Do motivational incentives reduce the inhibition deficit in ADHD? *Developmental Neuropsychology*, 33, 137-159. PMCID: Accepted for publication before April 7, 2008.
- Sonuga-Barke, E. J., Sergeant, J. A., Nigg, J., & Willcutt, E. (2008). Executive dysfunction and delay aversion in attention deficit hyperactivity disorder: nosologic and diagnostic implications. *Child and Adolescent Psychiatric Clinics of North America*, 17, 367-384, ix. PMCID: Accepted for publication before April 7, 2008.
- Treiman, R., Pennington, B. F., Shriberg, L. D., & Boada, R. (2008). Which children benefit from letter names in learning letter sounds? *Cognition*, 106, 1322-1338. PMCID: PMC2267370.
- Willcutt, E. G., Sonuga-Barke, E. J. S., Nigg, J. T., & Sergeant, J. A. (2008). Recent developments in neuropsychological models of childhood disorders. *Advances in Biological Psychiatry*, 24, 195-226.
- Wise, B. W., Cole, R., Van Vuuren, S., Schwartz, S., Snyder, L., Ngampatipatpong, N., . . . Pellom, B. (2008). Learning to read with a virtual tutor. In C. Kinzer & L. Verhoeven (Eds.), *Interactive Literacy Education: facilitating literacy environments through technology* (pp. 31-75). Mahwah, NJ: Lawrence Erlbaum.
- Wu, S. S., Meyer, M. L., Maeda, U., Salimpoor, V. N., Tomiyama, S., Geary, D. C., & Menon, V. (2008). Standardized Assessment of Strategy Use and Working Memory in Early Mental Arithmetic Performance. *Developmental Neuropsychology*, 33, 365-393.

## 2007

- Astrom, R. L., Wadsworth, S. J., & DeFries, J. C. (2007). Etiology of the stability of reading difficulties: the longitudinal twin study of reading disabilities. *Twin Research and Human Genetics*, 10, 434-439.
- Bidwell, L. C., Willcutt, E. G., DeFries, J. C., & Pennington, B. F. (2007). Testing for neuropsychological endophenotypes in siblings discordant for attention-deficit/hyperactivity disorder. *Biological Psychiatry*, 62, 991-998. PMCID: PMC2687149.
- Byrne, B., Samuelsson, S., Wadsworth, S. J., Hulslander, J., Corley, R., DeFries, J. C., . . . Olson, R. K. (2007). Longitudinal twin study of early literacy development: preschool through grade 1. *Reading and Writing*, 20, 77-102.
- Cole, R., Wise, B., & Van Vuuren, S. (2007). How Marni teaches children to read. *Educational Technology*, 24, 14-18.
- Frank, M. J., Santamaria, A., O'Reilly, R. C., & Willcutt, E. (2007). Testing computational models of dopamine and noradrenaline dysfunction in attention deficit/hyperactivity disorder. *Neuropsychopharmacology*, 32, 1583-1599.
- Friedman, N. P., Haberstick, B. C., Willcutt, E. G., Miyake, A., Young, S. E., Corley, R. P., & Hewitt, J. K. (2007). Greater attention problems during childhood predict poorer executive functioning in late adolescence. *Psychological Science*, 18, 893-900.
- Friend, A., DeFries, J. C., Wadsworth, S. J., & Olson, R. K. (2007). Genetic and environmental influences on word recognition and spelling deficits as a function of age. *Behavior Genetics*, 37, 477-486.
- Hartman, C. A., Rhee, S. H., Willcutt, E. G., & Pennington, B. F. (2007). Modeling rater disagreement for ADHD: are parents or teachers biased? *Journal of Abnormal Child Psychology*, 35, 536-542.
- Hawke, J. L., Wadsworth, S. J., Olson, R. K., & DeFries, J. C. (2007). Etiology of reading difficulties as a function of gender and severity. *Reading and Writing*, 20, 13-25.
- Keenan, J. M., & Betjemann, R. S. (2007). Comprehension of single words: The role of semantics in reading and reading disability. In E. Grigorenko & A. Naples (Eds.), *Single Word Reading* (pp. 191-210). London: Taylor & Francis.
- McGrath, L. M., Hutaff-Lee, C., Scott, A., Boada, R., Shriberg, L. D., & Pennington, B. F. (2007). Children with comorbid Speech Sound Disorder and Specific Language Impairment are at increased risk for Attention-Deficit/Hyperactivity Disorder. *Journal of Abnormal Child Psychology*, 36, 151-163.
- McGrath, L. M., Pennington, B. F., Willcutt, E. G., Boada, R., Shriberg, L. D., & Smith, S. D. (2007). Gene x environment interactions in speech sound disorder predict language and preliteracy outcomes. *Development and Psychopathology*, 19, 1047-1072.
- Olson, R. K. (2007). Genetic and environmental influences on word reading skills. In E. Grigorenko & A. Naples (Eds.), *Single word reading: Behavioral and biological approaches* (pp. 233-253). Mahwah, N.J.: Laurence Erlbaum Associates.
- Olson, R. K. (2007). Introduction to the special issue on genes, environment, and reading. *Reading and Writing*, 20, 1-11.
- Olson, R. K., Keenan, J. M., Byrne, B., Samuelsson, S., Coventry, W. L., Corley, R., . . . Hulslander, J. (2007). Genetic and environmental influences on vocabulary and reading development. *Scientific Studies of Reading*, 20, 51-75. PMCID: PMC3019615.
- Pennington, B. F. (2007). Integrating cognition and motivation in our understanding of ADHD [Commentary]. In D. E. Barch (Ed.), *Cognitive and affective neuroscience of psychopathology*. New York: Oxford University Press.
- Pennington, B. F., Snyder, K. A., & Roberts, R. J. (2007). Developmental cognitive neuroscience: Origins, issues, and prospects. *Developmental Review*, 27, 428-441.
- Peterson, R. L., McGrath, L. M., Smith, S. D., & Pennington, B. F. (2007). Neuropsychology and genetics of speech, language, and

- literacy disorders. *Pediatric Clinics of North America*, 54, 543-561, vii.
- Phinney, E., Pennington, B. F., Olson, R., Filley, C. M., & Filipek, P. A. (2007). Brain structure correlates of component reading processes: implications for reading disability. *Cortex*, 43, 777-791.
- Samuelsson, S., Olson, R., Wadsworth, S., Corley, R., DeFries, J. C., Willcutt, E., . . . Byrne, B. (2007). Genetic and environmental influences on prereading skills and early reading and spelling development in the United States, Australia, and Scandinavia. *Reading and Writing*, 20, 51-75.
- Shaywitz, S. E., Gruen, J. R., & Shaywitz, B. A. (2007). Management of dyslexia, its rationale, and underlying neurobiology. *Pediatric Clinics of North America*, 54, 609-623, viii.
- Smith, S. D. (2007). Genes, language development, and language disorders. *Mental Retardation and Developmental Disabilities Research Reviews*, 13, 96-105.
- Smith, S. D., & Gilger, J. W. (2007). Dyslexia and Other Specific Learning Disorders. In D. L. Rimoin, J. M. Connor, R. E. Pyeritz, B. R. Korf & A. E. H. Emery (Eds.), *Emery and Rimoin's principles and practice of medical genetics*, 4th Edition (pp. 2548-2568). New York: Elsevier.
- Wadsworth, S. J., DeFries, J. C., Olson, R. K., & Willcutt, E. G. (2007). Colorado longitudinal twin study of reading disability. *Annals of Dyslexia*, 57, 139-160.
- Willcutt, E. G., Betjemann, R. S., Pennington, B. F., Olson, R. K., DeFries, J. C., & Wadsworth, S. J. (2007). Longitudinal study of reading disability and attention-deficit/hyperactivity disorder: Implications for education. *Mind, Brain, and Education*, 4, 181-192. PMCID: Accepted for publication before April 7, 2008
- Willcutt, E. G., Betjemann, R. S., Wadsworth, S. J., Samuelsson, S., Corley, R., DeFries, J. C., . . . Olson, R. K. (2007). Preschool twin study of the relation between attention-deficit/hyperactivity disorder and prereading skills. *Reading and Writing*, 20, 103-125. PMCID: Accepted for publication before April 7, 2008
- Willcutt, E. G., Pennington, B. F., Olson, R. K., & DeFries, J. C. (2007). Understanding comorbidity: a twin study of reading disability and attention-deficit/hyperactivity disorder. *American Journal of Medical Genetics Part B (Neuropsychiatric Genetics)*, 144B, 709-714.
- Wise, B. (2007). Turning reading research into policy. *Reading Research Quarterly*, 42, 407-411.
- Wise, B. W., & Raskind, M. (2007). Technology and Reading Difficulties: Introduction to the special issue. *Perspectives of the International Dyslexia Association*, 7-8.
- Wise, B. W., & Van Vuuren, S. (2007). Choosing Software Gems to Improve Your Child's Reading. *Perspectives of the International Dyslexia Association*, 33, 34-38.
- Yerys, B. E., Hepburn, S. L., Pennington, B. F., & Rogers, S. J. (2007). Executive function in preschoolers with autism: Evidence consistent with a secondary deficit. *Journal of Autism and Developmental Disorders*, 37, 1068-1079.
- Zhang, H., Liu, L., Wang, X., & Gruen, J. R. (2007). Guideline for data analysis of genomewide association studies. *Cancer Genomics Proteomics*, 4, 27-34.

## 2006

- Boada, R., & Pennington, B. (2006). Deficient implicit phonological representations in children with dyslexia. *Journal of Experimental Child Psychology*, 95, 153-193.
- Bricker, J. B., Stallings, M. C., Corley, R. P., Wadsworth, S. J., Bryan, A., Timberlake, D. S., . . . DeFries, J. C. (2006). Genetic and environmental influences on age at sexual initiation in the Colorado Adoption Project. *Behavior Genetics*, 36, 820-832.
- Byrne, B., Olson, R. K., Samuelsson, S., Wadsworth, S., Corley, R., DeFries, J. C., & Willcutt, E. (2006). Genetic and environmental influences on early literacy. *Journal of Research in Reading*, 29, 33-49.
- Caspi, A., Hofer, S. M., Rhea, S. A., & DeFries, J. C. (2006). Genetic and environmental influences on age at sexual initiation in the Colorado Adoption Project. *Behavior Genetics*, 36, 820-832.
- Deupree, J. D., Smith, S. D., Kratochvil, C. J., Bohac, D., Ellis, C. R., Polaha, J., & Bylund, D. B. (2006). Possible involvement of alpha-2A adrenergic receptors in attention deficit hyperactivity disorder: radioligand binding and polymorphism studies. *American Journal of Medical Genetics Part B (Neuropsychiatric Genetics)*, 141B, 877-884.
- Friedman, N. P., Miyake, A., Corley, R. P., Young, S. E., DeFries, J. C., & Hewitt, J. K. (2006). Not all executive functions are related to intelligence. *Psychological Science*, 17, 172-179.
- Hawke, J. L., Wadsworth, S. J., & DeFries, J. C. (2006). Genetic influences on reading difficulties in boys and girls: the Colorado twin study. *Dyslexia*, 12, 21-29.
- Keenan, J. M., & Betjemann, R. S. (2006). Comprehending the Gray Oral Reading Test without reading it: Why comprehension tests should not include passage-independent items. *Scientific Studies of Reading*, 10, 363-380.
- Keenan, J. M., Betjemann, R. S., Wadsworth, S. J., DeFries, J. C., & Olson, R. K. (2006). Genetic and environmental influences on reading and listening comprehension. *Journal of Research in Reading*, 29, 79-91.
- McGrath, L. M., Smith, S. D., & Pennington, B. F. (2006). Breakthroughs in the search for dyslexia candidate genes. *Trends in Molecular Medicine*, 12, 333-341.
- Olson, R. K. (2006). Genes, environment, and dyslexia - The 2005 Norman Geschwind memorial lecture. *Annals of Dyslexia*, 56, 205-238.
- Olson, R. K., & Wise, B. (2006). Computer-based remediation for reading and related phonological disabilities. In M. McKenna, L. Labbo, R. Kieffer & D. Reinking (Eds.), *Handbook of literacy and technology*, vol. 2 (pp. 57-74). Mahwah, N.J.: Lawrence Erlbaum.
- Ozonoff, S., Pennington, B. F., & Solomon, M. (2006). Neuropsychological perspectives on developmental psychopathology. In D. V. Cicchetti & D. Cohen (Eds.), *Developmental Psychopathology* (2nd Edition) (pp. 332-380). New York: Wiley and Sons.
- Pennington, B. F. (2006). From single to multiple deficit models of developmental disorders. *Cognition*, 101, 385-413.
- Pennington, B. F., & Chhabildas, N. (2006). Attention deficit hyperactivity disorder. In M. Farah & T. E. Todd (Eds.), *Patient-based approaches to cognitive neuroscience* (2nd Edition) (pp. 407-418). Cambridge, MA: MIT Press.
- Pennington, B. F., Williams, J., & Rogers, S. J. (2006). Conclusions. In S. J. Rogers & J. Williams (Eds.), *Imitation and the development*

- of the social mind: Lessons from typical development and autism* (pp. 431-453). New York: Guilford Press.
- Rutherford, M. D., Pennington, B. F., & Rogers, S. J. (2006). The perception of animacy in young children with autism. *Journal of Autism and Developmental Disorders*, 36, 983-992.
- Shanahan, M. A., Pennington, B. F., Yerys, B. E., Scott, A., Boada, R., Willcutt, E. G., . . . DeFries, J. C. (2006). Processing speed deficits in attention deficit/hyperactivity disorder and reading disability. *Journal of Abnormal Child Psychology*, 34, 585-602.
- Wadsworth, S. J., Corley, R., Plomin, R., Hewitt, J. K., & DeFries, J. C. (2006). Genetic and environment influences on the continuity and change in reading achievement in the Colorado Adoption Project. In A. C. Huston & M. N. Ripke (Eds.), *Developmental contexts in middle childhood* (pp. 87-106). New York, NY: Cambridge University Press.

## 2005

- Banaschewski, T., Hollis, C., Oosterlaan, J., Roeyers, H., Rubia, K., Willcutt, E., & Taylor, E. (2005). Towards an understanding of unique and shared pathways in the psychopathophysiology of ADHD. *Developmental Science*, 8, 132-140.
- Biederman, J., Pennington, B. F., Nigg, J. T., Willcutt, E. G., & Doyle, A. E., . . . & Sklar, P. (2005). Attention-deficit/hyperactivity disorder: A selective overview. *Biological Psychiatry*, 57, 1215-1220.
- Byrne, B., Wadsworth, S., Corley, R., Samuelsson, S., Quain, P., DeFries, J. C., . . . Olson, R. K. (2005). Longitudinal twin study of early literacy development: Preschool and kindergarten phases. *Scientific Studies of Reading*, 9, 219-235.
- Doyle, A. E., Faraone, S. V., Seidman, L. J., Willcutt, E. G., Nigg, J. T., Waldman, I. D., . . . Biederman, J. (2005). Are endophenotypes based on measures of executive functions useful for molecular genetic studies of ADHD? *Journal of Child Psychology and Psychiatry*, 46, 774-803.
- Doyle, A. E., Willcutt, E. G., Seidman, L. J., Biederman, J., Chouinard, V. A., Silva, J., & Faraone, S. V. (2005). Attention-deficit/hyperactivity disorder endophenotypes. *Biological Psychiatry*, 57, 1324-1335.
- Edgin, J. O., & Pennington, B. F. (2005). Spatial cognition in autism spectrum disorders: superior, impaired, or just intact? *Journal of Autism and Developmental Disorders*, 35, 729-745.
- Friedman, N. P., & Miyake, A. (2005). Comparison of four scoring methods for the reading span test. *Behavioral Research Methods*, 37, 581-590.
- Gayan, J., Willcutt, E. G., Fisher, S. E., Francks, C., Cardon, L. R., Olson, R. K., . . . DeFries, J. C. (2005). Bivariate linkage scan for reading disability and attention-deficit/hyperactivity disorder localizes pleiotropic loci. *Journal of Child Psychology and Psychiatry*, 46, 1045-1056.
- Lahey, B. B., Pelham, W. E., Loney, J., Lee, S. S., & Willcutt, E. (2005). Instability of the DSM-IV Subtypes of ADHD from preschool through elementary school. *Archives of General Psychiatry*, 62, 896-902.
- Meng, H., Hager, K., Held, M., Page, G. P., Olson, R. K., Pennington, B. F., . . . Gruen, J. R. (2005). TDT-association analysis of EKN1 and dyslexia in a Colorado twin cohort. *Human Genetics*, 118, 87-90.
- Meng, H., Smith, S. D., Hager, K., Held, M., Liu, J., Olson, R. K., . . . Gruen, J. R. (2005). DCDC2 is associated with reading disability and modulates neuronal development in the brain. *Proceedings of the National Academy of Sciences of the United States of America*, 102, 17053-17058. PMCID: PMC1278934.
- Nigg, J. T., Willcutt, E. G., Doyle, A. E., & Sonuga-Barke, E. J. (2005). Causal heterogeneity in attention-deficit/hyperactivity disorder: do we need neuropsychologically impaired subtypes? *Biological Psychiatry*, 57, 1224-1230.
- Olson, R. K., & Byrne, B. (2005). Genetic and environmental influences on reading and language ability and disability. In H. W. Catts & A. Kamhi (Eds.), *The connections between language and reading disabilities* (pp. 173-200). Mahwah, NJ: Laurence Erlbaum Associates.
- Pennington, B. F. (2005). Toward a new neuropsychological model of attention-deficit/hyperactivity disorder: subtypes and multiple deficits. *Biological Psychiatry*, 57, 1221-1223.
- Pennington, B. F., & Olson, R. K. (2005). Genetics of dyslexia. In M. J. Snowling & C. Hulme (Eds.), *The science of reading: A handbook* (pp. 453-472). Oxford: Blackwell Publishing.
- Pennington, B. F., Willcutt, E., & Rhee, S. H. (2005). Analyzing comorbidity. *Advances in Child Development and Behavior*, 33, 263-304.
- Raha-Chowdhury, R., Andrews, S. R., & Gruen, J. R. (2005). CAT 53: a protein phosphatase 1 nuclear targeting subunit encoded in the MHC Class I region strongly expressed in regions of the brain involved in memory, learning, and Alzheimer's disease. *Brain Research: Molecular Brain Research*, 138, 70-83.
- Rhee, S. H., Hewitt, J. K., Corley, R. P., Willcutt, E. G., & Pennington, B. F. (2005). Testing hypotheses regarding the causes of comorbidity: Examining the underlying deficits of comorbid disorders. *Journal of Abnormal Psychology*, 114, 346-362.
- Samuelsson, S., Byrne, B., Quain, P., Wadsworth, S., Corley, R., DeFries, J. C., . . . Olson, R. (2005). Environmental and genetic influences on prereading skills in Australia, Scandinavia, and the United States. *Journal of Educational Psychology*, 97, 705-722.
- Smith, S. D., & Morris, C. A. (2005). Planning studies of etiology. *Applied Psycholinguistics*, 26, 97-110.
- Smith, S. D., Pennington, B. F., Boada, R., & Shriberg, L. D. (2005). Linkage of speech sound disorder to reading disability loci. *Journal of Child Psychology and Psychiatry*, 46, 1057-1066.
- Snyder, L., Caccamise, D., & Wise, B. (2005). The assessment of reading comprehension: Considerations and cautions. *Topics in Language Disorders*, 25, 33-50.
- Stevenson, J., Asherson, P., Hay, D., Levy, F., Swanson, J., Thapar, A., & Willcutt, E. (2005). Characterizing the ADHD phenotype for genetic studies. *Developmental Science*, 8, 115-121.
- Todd, R. D., Huang, H., Smalley, S. L., Nelson, S. F., Willcutt, E. G., Pennington, B. F., . . . Neuman, R. J. (2005). Collaborative analysis of DRD4 and DAT genotypes in population-defined ADHD subtypes. *Journal of Child Psychology and Psychiatry*, 46, 1067-1073.
- Wadsworth, S. J., & DeFries, J. C. (2005). Genetic etiology of reading difficulties in boys and girls. *Twin Research and Human Genetics*, 8, 594-601.
- Weaver, C. A., Keenan, J. M., Perrig, W., & Patel, V. (2005). Walter Kintsch: A psychology beyond words. In A. Healy (Ed.),

- Experimental cognitive psychology and its applications* (pp. 17-29). Washington, DC: American Psychological Association.
- Willcutt, E. G., Brodsky, K., Chhabildas, N. A., Shanahan, M., Yerys, B. E., Scott, A., & Pennington, B. F. (2005). The neuropsychology of ADHD: Validity of the executive function hypothesis. In D. Gozal & D. L. Molfese (Eds.), *Attention deficit hyperactivity disorder: From genes to patients* (pp. 185-213). Totowa, NJ: Humana Press.
- Willcutt, E. G., & Carlson, C. L. (2005). Diagnostic validity of attention-deficit/hyperactivity disorder. *Clinical Neuroscience Review*, 5, 219-232. PMCID: Accepted for publication before April 7, 2008
- Willcutt, E. G., Doyle, A. E., Nigg, J. T., Faraone, S. V., & Pennington, B. F. (2005). Validity of the executive function theory of attention-deficit/hyperactivity disorder: A meta-analytic review. *Biological Psychiatry*, 57, 1336-1346.
- Willcutt, E. G., Pennington, B. F., Olson, R. K., Chhabildas, N., & Hulslander, J. (2005). Neuropsychological analyses of comorbidity between reading disability and attention deficit hyperactivity disorder: In search of the common deficit. *Developmental Neuropsychology*, 27, 35-78.

## 2004

- Cardoso-Martins, C., & Pennington, B. F. (2004). The relationship between phoneme awareness and rapid serial naming skills and literacy acquisition: The role of developmental period and reading ability. *Scientific Studies of Reading*, 8, 27-52.
- de Jong, P. F., & Olson, R. K. (2004). Early predictors of letter knowledge. *Journal of Experimental Child Psychology*, 88, 254-273.
- Deffenbacher, K. E., Kenyon, J. B., Hoover, D. M., Olson, R. K., Pennington, B. F., DeFries, J. C., & Smith, S. D. (2004). Refinement of the 6p21.3 quantitative trait locus influencing dyslexia: linkage and association analyses. *Human Genetics*, 115, 128-138.
- Flint, J., DeFries, J. C., & Henderson, N. D. (2004). Little epistasis for anxiety-related measures in the DeFries strains of laboratory mice. *Mammalian Genome: Genes and Phenotypes*, 15, 77-82.
- Francks, C., Paracchini, S., Smith, S. D., Richardson, A. J., Scerri, T. S., Cardon, L. R., . . . Monaco, A. P. (2004). A 77-kilobase region of chromosome 6p22.2 is associated with dyslexia in families from the United Kingdom and from the United States. *American Journal of Human Genetics*, 75, 1046-1058. PMCID: PMC1182140.
- Friedman, N. P., & Miyake, A. (2004). The relations among inhibition and interference control functions: a latent-variable analysis. *Journal of Experimental Psychology: General*, 133, 101-135.
- Hartman, C. A., Willcutt, E. G., Rhee, S. H., & Pennington, B. F. (2004). The relation between sluggish cognitive tempo and DSM-IV ADHD. *Journal of Abnormal Child Psychology*, 32, 491-503.
- Henderson, N. D., Turri, M. G., DeFries, J. C., & Flint, J. (2004). QTL analysis of multiple behavioral measures of anxiety in mice. *Behavior Genetics*, 34, 267-293.
- Hulslander, J., Talcott, J., Witton, C., DeFries, J., Pennington, B., Wadsworth, S., . . . Olson, R. (2004). Sensory processing, reading, IQ, and attention. *Journal of Experimental Child Psychology*, 88, 274-295.
- Keenan, J. M., & Simon, J. A. (2004). Inference deficits in women with Fragile X Syndrome: A problem in working memory. *Cognitive Neuropsychology*, 21, 579-596.
- Lahey, B. B., Pelham, W. E., Loney, J., Kipp, H., Ehrhardt, A., Lee, S. S., . . . Massetti, G. (2004). Three-year predictive validity of DSM-IV attention deficit hyperactivity disorder in children diagnosed at 4-6 years of age. *American Journal of Psychiatry*, 161, 2014-2020.
- Lowe, N., Kirley, A., Hawi, Z., Sham, P., Wickham, H., Kratochvil, C. J., . . . Gill, M. (2004). Joint analysis of the DRD5 marker concludes association with attention-deficit/hyperactivity disorder confined to the predominantly inattentive and combined subtypes. *American Journal of Human Genetics*, 74, 348-356.
- Ma, J., Cole, R., Pellom, B., Ward, W., & Wise, B. (2004). Accurate automatic visible speech synthesis of arbitrary 3D models based on concatenation of diviseme motion capture data. *Computer animation and virtual worlds*, 15, 485-500.
- Olson, R. K. (2004). SSSR, environment, and genes. *Scientific Studies of Reading*, 8, 111-124.
- Ozonoff, S., Cook, I., Coon, H., Dawson, G., Joseph, R. M., Klin, A., . . . Wrathall, D. (2004). Performance on Cambridge Neuropsychological Test Automated Battery subtests sensitive to frontal lobe function in people with autistic disorder: evidence from the Collaborative Programs of Excellence in Autism network. *Journal of Autism and Developmental Disorders*, 34, 139-150.
- Petrill, S. A., Hewitt, J. K., Cherny, S. S., Lipton, P. A., Plomin, R., Corley, R., & DeFries, J. C. (2004). Genetic and environmental contributions to general cognitive ability through the first 16 years of life. *Developmental Psychology*, 40, 805-812.
- Petrill, S. A., Lipton, P. A., Hewitt, J. K., Plomin, R., Cherny, S. S., Corley, R., & DeFries, J. C. (2004). Genetic and environmental contributions to general cognitive ability through the first 16 years of life. *Developmental Psychology*, 40, 805-812.
- Raitano, N. A., Pennington, B. F., Tunick, R. A., Boada, R., & Shriberg, L. D. (2004). Pre-literacy skills of subgroups of children with speech sound disorders. *Journal of Child Psychology and Psychiatry*, 45, 821-835. PMCID: PMC3164030.
- Smith, S. D. (2004). Localization and identification of genes influencing language and learning disorders. In M. L. Rice & S. F. Warren (Eds.), *Developmental Language Disorder: From phenotype to etiologies* (pp. 329-354). Mahwah, NJ: Lawrence Erlbaum Associates.
- Smith, S. D., & Taggart, R. T. (2004). Nonsyndromic Hearing Impairment. In H. Toriello, W. Reardon & R. J. Gorlin (Eds.), *Heredity hearing loss and its syndromes* (pp. 37-82). New York: Oxford University Press.
- Tiu, R. D., Jr., Wadsworth, S. J., Olson, R. K., & DeFries, J. C. (2004). Causal models of reading disability: a twin study. *Twin Research and Human Genetics*, 7, 275-283.
- Turri, M. G., DeFries, J. C., Henderson, N. D., & Flint, J. (2004). Multivariate analysis of quantitative trait loci influencing variation in anxiety-related behavior in laboratory mice. *Mammalian Genome*, 15, 69-76.
- Willcutt, E. G. (2004). Behavior Genetics. In W. E. Craighead & C. Nemerooff (Eds.), *Corsini Concise Encyclopedia of Psychology and Behavioral Sciences* (pp. 105-107). New York, NY: Wiley & Sons.
- Willcutt, E. G., & Gaffney-Brown, R. (2004). Etiology of dyslexia, ADHD, and related difficulties: Using genetic methods to understand comorbidity. *Perspectives of the International Dyslexia Association*, 30, 12-15.
- Williams, R. W., Bennett, B., Lu, L., Gu, J., DeFries, J. C., Carosone-Link, P. J., . . . Johnson, T. E. (2004). Genetic structure of the LXS panel of recombinant inbred mouse strains: a powerful resource for complex trait analysis. *Mammalian Genome*, 15, 637-647.

Wise, B. W. (2004). Facts, Fictions, and Facts in the Reading Wars. In M. Joshi (Ed.), *Dyslexia: Myths, misconceptions, and some practical applications*. Mahwah, NJ: Lawrence Erlbaum.

## 2003

- Alarcón, M., Plomin, R., Corley, R. P., & DeFries, J. C. (2003). Multivariate parent-offspring analyses of specific cognitive abilities. In S. A. Petrill, R. Plomin, J. C. DeFries & J. K. Hewitt (Eds.), *Nature, nurture, and the transition to early adolescence*. New York: Oxford University Press.
- Bishop, E. G., Cherny, S. S., Corley, R., Plomin, R., DeFries, J. C., & Hewitt, J. K. (2003). Development genetic analysis of general cognitive ability from 1 to 12 years in a sample of adoptees, biological siblings, and twins. *Intelligence*, 31, 31-49.
- Bodensteiner, J. B., Smith, S. D., & Schaefer, G. B. (2003). Hypotonia, congenital hearing loss, and hypoactive labyrinths. *Journal of Child Neurology*, 18, 171-173.
- Friedman, M. C., Chhabildas, N., Budhiraja, N., Willcutt, E. G., & Pennington, B. F. (2003). Etiology of the comorbidity between RD and ADHD: Exploration of the non-random mating hypothesis. *American Journal of Medical Genetics Part B (Neuropsychiatric Genetics)*, 120B, 109-115.
- Gayán, J. (2003). Genetic and environmental influences on individual differences in printed word recognition. *Journal of Experimental Child Psychology*, 84, 97-123.
- Gayán, J. (2003). Genetic and environmental influences on individual differences in printed word recognition. *Journal of Experimental Child Psychology*, 84, 97-123.
- Gayán, J., & Olson, R. K. (2003). Genetic and environmental influences on individual differences in printed word recognition. *Journal of Experimental Child Psychology*, 84, 97-123.
- Kaakinen, J. K., Hyona, J., & Keenan, J. M. (2003). How prior knowledge, working memory capacity, and relevance of information affect eye fixations in expository text. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, 29, 447-457.
- Londin, E. R., Meng, H., & Gruen, J. R. (2003). A transcription map of the 6p22.3 reading disability locus identifying candidate genes. *BMC Genomics*, 4, 25. PMCID: PMC166143.
- O'Connor, T. G., Caspi, A., DeFries, J. C., & Plomin, R. (2003). Genotype-environment interaction in children's adjustment to parental separation. *Journal of Child Psychology and Psychiatry*, 44, 849-856.
- Pennington, B. F. (2003). Acceptance of the Samuel Torrey Orton Award - November 15, 2002, Atlanta, Georgia - Understanding the comorbidity of dyslexia. *Annals of Dyslexia*, 53, 15-22.
- Pennington, B. F., & Chhabildas, N. (2003). Attention deficit hyperactivity disorder. In T. E. Feinberg & M. Farah (Eds.), *Behavioral neurology and neuropsychology* (pp. 831-842). New York: McGraw Hill.
- Pennington, B. F., Moon, J., Edgin, J., Stedron, J., & Nadel, L. (2003). The neuropsychology of Down syndrome: Evidence for hippocampal dysfunction. *Child Development*, 74, 75-93.
- Petrill, S. A., Plomin, R., DeFries, J. C., & Hewitt, J. K. (2003). Nature, nurture, and adolescent development. In S. A. Petrill, R. Plomin, J. C. DeFries & J. K. Hewitt (Eds.), *Nature, nurture, and the transition to early adolescence* (pp. 3-12). New York: Oxford University Press.
- Petrill, S. A., Plomin, R., DeFries, J. C., & Hewitt, J. K. (2003). Conclusions. In S. A. Petrill, R. Plomin, J. C. DeFries & J. K. Hewitt (Eds.), *Nature, nurture, and the transition to early adolescence* (pp. 310-316). New York, NY: Oxford University Press.
- Petrill, S. A., Plomin, R., DeFries, J. C., & Hewitt, J. K. (2003). *Nature, Nurture and the Transition to Early Adolescence*. Oxford, NY: Oxford University Press.
- Plomin, R., DeFries, J. C., Craig, I. W., & McGuffin, P. (2003). Behavioral genetics. In R. Plomin, J. C. DeFries, I. W. Craig & P. McGuffin (Eds.), *Behavioral genetics in the postgenomic era* (pp. 3-15). Washington, DC: American Psychological Association.
- Plomin, R., DeFries, J. C., Craig, I. W., & McGuffin, P. (2003). *Behavioral genetics in the postgenomic era*. Washington, DC: American Psychological Association.
- Smith, S. D., Deffenbacher, K. E., & Brower, A. M. (2003). Reading and Dyslexias. In N. P. Group (Ed.), *Encyclopedia of the Human Genome*: Nature Publishing Group.
- Wadsworth, S. J., & DeFries, J. C. (2003). Etiology of the stability of reading performance from 7 to 12 years of age and its possible mediation by IQ. In S. A. Petrill, R. Plomin, J. C. DeFries & J. K. Hewitt (Eds.), *Nature, nurture, and the transition to early adolescence* (pp. 49-61). New York: Oxford University Press.
- Willcutt, E. G., DeFries, J. C., Pennington, B. F., Olson, R. K., Smith, S. D., & Cardon, L. R. (2003). Genetic etiology of comorbid reading difficulties and ADHD. In R. Plomin, J. C. DeFries, P. McGuffin & I. Craig (Eds.), *Behavioral Genetics in a Postgenomic Era* (pp. 227-246). Washington DC: American Psychological Association.

## 2002

- Ahn, J., Won, T. W., Kaplan, D. E., Londin, E. R., Kuzmic, P., Gelernter, J., & Gruen, J. R. (2002). A detailed physical map of the 6p reading disability locus, including new markers and confirmation of recombination suppression. *Human Genetics*, 111, 339-349.
- Barkley, R. A., Cook, E. H., Diamond, A., Zametkin, A., Thapar, A., Teeter, A., . . . Pelham, W. E. (2002). International consensus statement on ADHD. January 2002. *Clinical Child and Family Psychology Review*, 5, 89-111.
- Boada, R., Willcutt, E. G., Tunick, R., Chhabildas, N. A., Oglane, J., Olson, R. K., . . . Pennington, B. F. (2002). A twin study of the etiology of high reading ability. *Reading and Writing*, 15, 683-707.
- Byrne, B., Delaland, C., Fielding-Barnsley, R., Quain, P., Samuelsson, S., Hoien, T., . . . Olson, R. K. (2002). Longitudinal twin study of early reading development in three countries: Preliminary results. *Annals of Dyslexia*, 52, 49-73.
- Carpenter, M., Pennington, B. F., & Rogers, S. J. (2002). Interrelations among social-cognitive skills in young children with autism. *Journal of Autism and Developmental Disorders*, 32, 91-106.
- Compton, D. L., Olson, R. K., DeFries, J. C., & Pennington, B. F. (2002). Comparing the relationships among two different versions of alphanumeric rapid automatized naming and word level reading skills. *Scientific Studies of Reading*, 6, 343-368.

- De Leenheer, E. M., Ensink, R. J., Kunst, H. P., Marres, H. A., Talebizadeh, Z., Declau, F., . . . Cremers, C. W. (2002). DFNA2/KCNQ4 and its manifestations. *Advances in Otorhinolaryngology*, 61, 41-46.
- Fisher, S. E., & DeFries, J. C. (2002). Developmental dyslexia: genetic dissection of a complex cognitive trait. *Nature Reviews: Neuroscience*, 3, 767-780.
- Fisher, S. E., Francks, C., Marlow, A. J., MacPhie, I. L., Newbury, D. F., Cardon, L. R., . . . Monaco, A. P. (2002). Independent genome-wide scans identify a chromosome 18 quantitative-trait locus influencing dyslexia. *Nature Genetics*, 30, 86-91.
- Francks, C., Fisher, S. E., Olson, R. K., Pennington, B. F., Smith, S. D., DeFries, J. C., & Monaco, A. P. (2002). Fine mapping of the chromosome 2p12-16 dyslexia susceptibility locus: quantitative association analysis and positional candidate genes SEMA4F and OTX1. *Psychiatric Genetics*, 12, 35-41.
- Hartung, C. M., Willcutt, E. G., Lahey, B. B., Pelham, W. E., Loney, J., Stein, M. A., & Keenan, K. (2002). Sex differences in young children who meet criteria for attention deficit hyperactivity disorder. *Journal of Clinical Child and Adolescent Psychology*, 31, 453-464.
- Kaakinen, J., Hyona, J., & Keenan, J. M. (2002). Individual differences in perspective effects on on-line text processing. *Discourse Processes*, 33, 159-173.
- Kaplan, D. E., Gayan, J., Ahn, J., Won, T. W., Pauls, D., Olson, R. K., . . . Gruen, J. R. (2002). Evidence for linkage and association with reading disability on 6p21.3-22. *American Journal of Human Genetics*, 70, 1287-1298. PMCID: PMC447603.
- Keenan, J. M. (2002). Inferences. In N. Smelser & P. Balter (Eds.), *International Encyclopedia of the Social & Behavioral Sciences* (Vol. 11, pp. 7432 – 7435). Amsterdam: Pergamon.
- Knopik, V. S., Smith, S. D., Cardon, L., Pennington, B., Gayán, J., Olson, R. K., & DeFries, J. C. (2002). Differential genetic etiology of reading component processes as a function of IQ. *Behavior Genetics*, 32, 181-198.
- Koczat, D. L., Rogers, S. J., Pennington, B. F., & Ross, R. G. (2002). Eye movement abnormality suggestive of a spatial working memory deficit is present in parents of autistic probands. *Journal of Autism and Developmental Disorders*, 32, 513-518.
- Lahey, B. B., & Willcutt, E. G. (2002). Validity of the diagnosis and dimensions of attention deficit hyperactivity disorder. In P. J. Jensen & J. R. Cooper (Eds.), *Attention Deficit Hyperactivity Disorder: State of the Science* (pp. 1-23). New York: Civic Research Institute.
- Nigg, J. T., John, O. P., Blaskey, L. G., Huang-Pollock, C. L., Willcutt, E. G., Hinshaw, S. P., & Pennington, B. (2002). Big five dimensions and ADHD symptoms: links between personality traits and clinical symptoms. *Journal of Personality and Social Psychology*, 83, 451-469.
- Olson, R. K. (2002). Dyslexia: nature and nurture. *Dyslexia*, 8, 143-159.
- Pennington, B. F. (2002). *The Development of Psychopathology*. New York, NY: Guilford Press.
- Pritchard, M., & Keenan, J. M. (2002). Does jury deliberation really improve jurors' memories? *Journal of Applied Cognitive Psychology*, 16, 589-601.
- Rawson, K. A., & Miyake, A. (2002). Does relocating information in text depend on verbal or visuospatial abilities? An individual-differences analysis. *Psychonomics Bulletin and Review*, 9, 801-806.
- Rende, B., Ramsberger, G., & Miyake, A. (2002). Commonalities and differences in the working memory components underlying letter and category fluency tasks: a dual-task investigation. *Neuropsychology*, 16, 309-321.
- Smith, S. D., Gilger, J. W., & Pennington, B. F. (2002). Dyslexia and other language/learning disorders. In D. L. Rimoin, J. M. Conner & R. Pyeritz (Eds.), *Emery and Rimoin's principles and practice of medical genetics, 4th Edition* (pp. 2827-2865). New York: Churchill Livingstone.
- Tunick, R. A., & Pennington, B. F. (2002). The etiological relationship between reading disability and phonological disorder. *Annals of Dyslexia*, 52, 75-97.
- Wadsworth, S. J., Corley, R. P., Hewitt, J. K., Plomin, R., & DeFries, J. C. (2002). Parent-offspring resemblance for reading performance at 7, 12 and 16 years of age in the Colorado Adoption Project. *Journal of Child Psychology and Psychiatry*, 43, 769-774.
- Wadsworth, S. J., Davis, C. J., Knopik, V. S., Willcutt, E. G., & DeFries, J. C. (2002). Genetics of reading disabilities *Proceedings of the 12th Postgraduate Course in Pediatric Neurology: Learning Disabilities in Reading and Writing* (pp. 23-36). Milan: FrancoAngeli.
- Wadsworth, S. J., J., D. C., Knopik, V. S., Willcutt, E. G., & DeFries, J. C. (2002). Genetics of reading disabilities. In I. Arcolini & G. Zardini (Eds.), *I disturbi di apprendimento della lettura e della scrittura*. Milan, Italy: FrancoAngeli.
- Willcutt, E. G., Pennington, B. F., Smith, S. D., Cardon, L. R., Gayán, J., Knopik, V. S., . . . DeFries, J. C. (2002). Quantitative trait locus for reading disability on chromosome 6p is pleiotropic for attention-deficit/hyperactivity disorder. *American Journal of Medical Genetics*, 114, 260-268.
- Wise, B. W., & Snyder, L. (2002). Clinical judgments in language-based learning disabilities. In R. Bradley, L. Danielson & D. Hallahan (Eds.), *Identification of Learning Disabilities: Research to practice*. Mahwah, NJ: Lawrence Erlbaum.
- Young, S. E., Smolen, A., Corley, R. P., Krauter, K. S., DeFries, J. C., Crowley, T. J., & Hewitt, J. K. (2002). Dopamine transporter polymorphism associated with externalizing behavior problems in children. *American Journal of Medical Genetics*, 114, 144-149.
- Ahn, J., Won, T. W., Zia, A., Reutter, H., Kaplan, D. E., Sparks, R., & Gruen, J. R. (2001). Peaks of linkage are localized by a BAC/PAC contig of the 6p reading disability locus. *Genomics*, 78, 19-29.

## 2001

- Bennetto, L., Pennington, B. F., Porter, D., Taylor, A. K., & Hagerman, R. J. (2001). Profile of cognitive functioning in women with the fragile X mutation. *Neuropsychology*, 15, 290-299.
- Carpenter, M., Pennington, B. F., & Rogers, S. J. (2001). Understanding of others' intentions in children with autism. *Journal of Autism and Developmental Disorders*, 31, 589-599.
- Cherny, S. S., Saudino, K. J., Fulker, D. W., Plomin, R., Corley, R. P., & DeFries, J. C. (2001). The development of observed shyness from 14 to 20 months. In R. N. Emde & J. K. Hewitt (Eds.), *Infancy to early childhood: Genetic and environmental influences*

- on developmental change (pp. 269-282). Oxford, NY: Oxford University Press.
- Chhabildas, N., Pennington, B. F., & Willcutt, E. G. (2001). A comparison of the neuropsychological profiles of the DSM-IV subtypes of ADHD. *Journal of Abnormal Child Psychology*, 29, 529-540.
- Compton, D. L., Davis, C. J., DeFries, J. C., Gayán, J., & Olson, R. K. (2001). Genetic and environmental influences on reading and RAN: An overview of results from the Colorado Twin Study. In M. Wolf (Ed.), *Dyslexia, fluency, and the brain* (pp. 277-303). Timonium, NJ: York Press.
- Compton, D. L., DeFries, J. C., & Olson, R. K. (2001). Are RAN- and phonological awareness-deficits additive in children with reading disabilities? *Dyslexia*, 7, 125-149.
- Davis, C. J., Gayán, J., Knopik, V. S., Smith, S. D., Cardon, L. R., Pennington, B. F., . . . DeFries, J. C. (2001). Etiology of reading difficulties and rapid naming: the Colorado Twin Study of Reading Disability. *Behavior Genetics*, 31, 625-635.
- Davis, C. J., Knopik, V. S., Olson, R. K., Wadsworth, S. J., & DeFries, J. C. (2001). Genetic and environmental influences on rapid naming and reading ability: A twin study. *Annals of Dyslexia*, 51, 231-247.
- Fisher, S. E., & Smith, S. D. (2001). Progress towards the identification of genes influencing developmental dyslexia. In A. J. Fawcett (Ed.), *Dyslexia: Theory and Good Practice* (pp. 39-64). London: Whurr Publishing.
- Gayán, J., & Olson, R. K. (2001). Genetic and environmental influences on orthographic and phonological skills in children with reading disabilities. *Developmental Neuropsychology*, 20, 483-507.
- Kaakinen, J., Hyona, J., & Keenan, J. M. (2001). Individual differences in perspective effects on text memory. *Current Psychology Letters: Behavior, Brain, & Cognition*, 5, 21-32.
- Miyake, A. (2001). Individual differences in working memory: introduction to the special section. *Journal of Experimental Psychology-General*, 130, 163-168.
- Miyake, A., Witzki, A. H., & Emerson, M. J. (2001). Field dependence-independence from a working memory perspective: A dual-task investigation of the Hidden Figures Test. *Memory*, 9, 445-457.
- Pennington, B. F., Cardoso-Martins, C., Green, P. A., & Lefly, D. L. (2001). Comparing the phonological and double deficit hypotheses for developmental dyslexia. *Reading and Writing*, 14, 707-755.
- Pennington, B. F., & Lefly, D. L. (2001). Early reading development in children at family risk for dyslexia. *Child Development*, 72, 816-833.
- Pennington, B. F., Mohr, J., Ogle, J., Stedron, J., & Nadel, L. (2001). Neuropsychology of Down syndrome. *Cytogenetics and Cell Genetics*, 92, 16-16.
- Plomin, R., DeFries, J. C., McClearn, G. E., & McGuffin, P. (2001). *Behavioral genetics (4th edition)*. New York: Worth Publishing.
- Plomin, R., Emde, R. N., Hewitt, J. K., Kagan, J., & DeFries, J. C. (2001). An experiment in collaborative science. In R. N. Emde & J. K. Hewitt (Eds.), *Infancy to early childhood: Genetic and environmental influences on developmental change* (pp. 355-381). New York: Oxford University Press.
- Polk, T. A., Reed, C. L., Keenan, J. M., Hogarth, P., & Anderson, C. A. (2001). A dissociation between symbolic number knowledge and analogue magnitude information. *Brain and Cognition*, 47, 545-563.
- S., C. S., Fulker, D. W., Emde, R. N., Plomin, R., Corley, R. P., & DeFries, J. C. (2001). Continuity and change in general cognitive ability from 14 to 36 months. In R. N. Emde & J. K. Hewitt (Eds.), *Infancy to early childhood: Genetic and environmental influences on developmental change* (pp. 206-220). Oxford, NY: Oxford University Press.
- Simon, J. A., Keenan, J. M., Pennington, B. F., Taylor, A. K., & Hagerman, R. J. (2001). Discourse processing in women with fragile X syndrome: evidence for a deficit establishing coherence. *Cognitive Neuropsychology*, 18, 1-18.
- Smith, S. D. (2001). Relationships between neurologic disorders and hereditary hearing loss. *Seminars in Pediatric Neurology*, 8, 147-159.
- Smith, S. D., Kelley, P. M., Askew, J. W., Hoover, D. M., Deffenbacher, K. E., Gayán, J., . . . Olson, R. K. (2001). Reading disability and chromosome 6p21.3: evaluation of MOG as a candidate gene. *Journal of Learning Disabilities*, 34, 512-519.
- Turri, M. G., Datta, S. R., DeFries, J., Henderson, N. D., & Flint, J. (2001). QTL analysis identifies multiple behavioral dimensions in ethological tests of anxiety in laboratory mice. *Current Biology*, 11, 725-734.
- Turri, M. G., Henderson, N. D., DeFries, J. C., & Flint, J. (2001). Quantitative trait locus mapping in laboratory mice derived from a replicated selection experiment for open-field activity. *Genetics*, 158, 1217-1226.
- Van Orden, G. C., Pennington, B. F., & Stone, G. O. (2001). What do double dissociations prove? *Cognitive Science*, 25, 111-172.
- Wadsworth, S. J., Corley, R. P., Hewitt, J. K., & DeFries, J. C. (2001). Stability of genetic and environmental influences on reading performance at 7, 12, and 16 years of age in the Colorado Adoption Project. *Behavior Genetics*, 31, 353-359.
- Willcutt, E. G., Chhabildas, N. A., & Pennington, B. F. (2001). Validity of the DSM-IV subtypes of ADHD. *The ADHD Report*, 9, 2-5.
- Willcutt, E. G., Pennington, B. F., Boada, R., Ogle, J. S., Tunick, R. A., Chhabildas, N. A., & Olson, R. K. (2001). A comparison of the cognitive deficits in reading disability and attention-deficit/hyperactivity disorder. *Journal of Abnormal Psychology*, 110, 157-172.
- Wise, B. W. (2001). The indomitable dinosaur builder (and how she overcame her phonological deficit and learned to read instructions, and other things). *Journal of Special Education*, 35, 134-144.
- Wright, M., De Geus, E., Ando, J., Luciano, M., Posthuma, D., Ono, Y., . . . Boomsma, D. (2001). Genetics of cognition: outline of a collaborative twin study. *Twin Research*, 4, 48-56.

## 2000

- Alarcon, M., Knopik, V. S., & DeFries, J. C. (2000). Covariation of mathematics achievement and general cognitive ability in twins. *Journal of School Psychology*, 38, 63-77.
- Alarcon, M., Pennington, B. F., Filipek, P. A., & DeFries, J. C. (2000). Etiology of neuroanatomical correlates of reading disability. *Developmental Neuropsychology*, 17, 339-360.
- Crnic, L. S., & Pennington, B. F. (2000). Down syndrome: Neuropsychology and animal models. In C. Rovee-Collier, L. P. Lipsitt & H. Hayne (Eds.), *Progress in infancy research* (pp. 69-111). Mahwah, NJ: Lawrence Erlbaum Associates.
- Davis, C. J., Knopik, V. S., Wadsworth, S. J., & DeFries, J. C. (2000). Self-reported reading problems in parents of twins with reading

- difficulties. *Twin Research and Human Genetics*, 3, 88-91.
- DeFries, J. C. (2000). Quantitative behavioral genetic aspects in twin research. In B. Smedby, I. Lundberg & T. Sorenson (Eds.), *Scientific evaluation of the Swedish Twin Registry* (pp. 80-89). Stockholm, Sweden: Swedish Council for Planning and Coordination of Research.
- Friedman, N. P., & Miyake, A. (2000). Differential roles for visuospatial and verbal working memory in situation model construction. *Journal of Experimental Psychology: General*, 129, 61-83.
- Lefly, D. L., & Pennington, B. F. (2000). Reliability and validity of the adult reading history questionnaire. *Journal of Learning Disabilities*, 33, 286-296.
- Miyake, A., Emerson, M. J., & Friedman, N. P. (2000). Assessment of executive functions in clinical settings: problems and recommendations. *Seminars in Speech and Language*, 21, 169-183.
- O'Connor, T. G., Caspi, A., DeFries, J. C., & Plomin, R. (2000). Are associations between parental divorce and children's adjustment genetically mediated? An adoption study. *Developmental Psychology*, 36, 429-437.
- Oliver, A., Johnson, M. H., Karmiloff-Smith, A., & Pennington, B. F. (2000). Deviations in the emergence of representations: A neuroconstructivist framework for analyzing developmental disorders. *Developmental Science*, 3, 1-40.
- Pennington, B. F., Filipek, P. A., Lefly, D., Chhabildas, N., Kennedy, D. N., Simon, J. H., . . . DeFries, J. C. (2000). A twin MRI study of size variations in human brain. *Journal of Cognitive Neuroscience*, 12, 223-232.
- Smith, S. D., Kelley, P. M., Kenyon, J. B., & Hoover, D. (2000). Tietz syndrome (hypopigmentation/deafness) caused by mutation of MITF. *Journal of Medical Genetics*, 37, 446-448.
- Wadsworth, S. J., Knopik, V. S., & DeFries, J. C. (2000). Reading disability in boys and girls: No evidence for a differential genetic etiology. *Reading and Writing*, 13, 133-145.
- Wadsworth, S. J., Olson, R. K., Pennington, B. F., & DeFries, J. C. (2000). Differential genetic etiology of reading disability as a function of IQ. *Journal of Learning Disabilities*, 33, 192-199.
- Welsh, M., & Pennington, B. F. (2000). Phenylketonuria. In K. O. Yeates, M. D. Ris & H. G. Taylor (Eds.), *Pediatric neuropsychology: Research, theory, and practice* (pp. 275-299). New York, NY: Guilford Press.
- Willcutt, E. G., & Pennington, B. F. (2000). Comorbidity of reading disability and attention-deficit/hyperactivity disorder: differences by gender and subtype. *Journal of Learning Disabilities*, 33, 179-191.
- Willcutt, E. G., & Pennington, B. F. (2000). Psychiatric comorbidity in children and adolescents with reading disability. *Journal of Child Psychology and Psychiatry*, 41, 1039-1048.
- Willcutt, E. G., Pennington, B. F., & DeFries, J. C. (2000). Twin study of the etiology of comorbidity between reading disability and attention-deficit/hyperactivity disorder. *American Journal of Medical Genetics Part B (Neuropsychiatric Genetics)*, 96, 293-301.
- Willcutt, E. G., Pennington, B. F., & DeFries, J. C. (2000). Etiology of inattention and hyperactivity/impulsivity in a community sample of twins with learning difficulties. *Journal of Abnormal Child Psychology*, 28, 149-159.
- Wise, B. W., Ring, J., & Olson, R. K. (2000). Individual differences in gains from computer-assisted remedial reading. *Journal of Experimental Child Psychology*, 77, 197-235.
- Young, S. E., Stallings, M. C., Corley, R. P., Krauter, K. S., & Hewitt, J. K. (2000). Genetic and environmental influences on behavioral disinhibition. *American Journal of Medical Genetics*, 96, 684-695.

## 1999

- Alarcón, M., Plomin, R., Fulker, D. W., Corley, R., & DeFries, J. C. (1999). Molarity not modularity: Multivariate genetic analysis of specific cognitive abilities in parents and their 16-year-old children in the Colorado Adoption Project. *Cognitive Development*, 14, 175-193.
- Castles, A., Datta, H., Gayán, J., & Olson, R. K. (1999). Varieties of developmental reading disorder: genetic and environmental influences. *Journal of Experimental Child Psychology*, 72, 73-94.
- Coucke, P. J., Van Hauwe, P., Kelley, P. M., Kunst, H., Schatteman, I., Van Velzen, D., . . . Van Camp, G. (1999). Mutations in the KCNQ4 gene are responsible for autosomal dominant deafness in four DFNA2 families. *Human Molecular Genetics*, 8, 1321-1328.
- DeFries, J. C., Knopik, V. S., & Wadsworth, S. J. (1999). Colorado Twin Study of Reading Disability. In D. D. Duane (Ed.), *Reading and attention disorders: Neurobiological correlates* (pp. 17-41). Baltimore, MD: York Press.
- Eslinger, P. J., Biddle, K., Pennington, B. F., & Page, R. B. (1999). Cognitive and behavioral development up to 4 years after early right frontal lobe lesion. *Developmental Neuropsychology*, 15, 157-171.
- Filipek, P. A., Pennington, B. F., Simon, J. H., Filley, C. M., & DeFries, J. C. (1999). Structural and functional neuroanatomy in reading disorder. In D. D. Duane (Ed.), *Reading and attention disorders: Neurobiological correlates*. Baltimore, MD: York Press.
- Gayán, J., & Olson, R. K. (1999). Reading disability: evidence for a genetic etiology. *European Child and Adolescent Psychiatry*, 8 Suppl 3, 52-55.
- Gayán, J., & Olson, R. K. (1999). Reading disability: evidence for a genetic etiology. *European Child and Adolescent Psychiatry*, 8 Suppl 3, 52-55.
- Gayán, J., Smith, S. D., Cherny, S. S., Cardon, L. R., Fulker, D. W., Brower, A. M., . . . DeFries, J. C. (1999). Quantitative-trait locus for specific language and reading deficits on chromosome 6p. *American Journal of Human Genetics*, 64, 157-164. PMCID: PMC1377713.
- Griffith, E. M., Pennington, B. F., Wehner, E. A., & Rogers, S. J. (1999). Executive functions in young children with autism. *Child Development*, 70, 817-832.
- Kelley, P. M., Abe, S., Askew, J. W., Smith, S. D., Usami, S. I., & Kimberling, W. J. (1999). Human connexin 30 (GJB6), a candidate gene for nonsyndromic hearing loss: molecular cloning, tissue-specific expression, and assignment to chromosome 13q12. *Genomics*, 62, 172-176.
- Kintsch, W., Healy, A. F., Hegarty, M., Pennington, B. F., & Salthouse, T. A. (1999). Models of working memory: Eight questions and some general issues. In A. Miyake & P. Shah (Eds.), *Models of working memory* (pp. 412-441). Cambridge, MA: Cambridge University Press.

- Knopik, V. S., & DeFries, J. C. (1999). Etiology of covariation between reading and mathematics performance: a twin study. *Twin Research and Human Genetics*, 2, 226-234.
- McBurnett, K., Pfiffner, L. J., Willcutt, E., Tamm, L., Lerner, M., Ottolini, Y. L., & Furman, M. B. (1999). Experimental cross-validation of DSM-IV types of attention-deficit/hyperactivity disorder. *Journal of the American Academy of Child and Adolescent Psychiatry*, 38, 17-24.
- Olson, R. K., Datta, H., Gayán, J., & DeFries, J. C. (1999). A behavioral-genetic analysis of reading disabilities and component processes. In R. M. Klein & P. A. McMullen (Eds.), *Converging methods for understanding reading and dyslexia* (pp. 133-151). Cambridge, MA: the MIT Press.
- Pennington, B. F. (1999). Toward an integrated understanding of dyslexia: Genetic, neurological, and cognitive mechanisms. *Development and Psychopathology*, 11, 629-654.
- Pennington, B. F., Filipek, P. A., Lefly, D., Churchwell, J., Kennedy, D. N., Simon, J. H., . . . DeFries, J. C. (1999). Brain morphometry in reading-disabled twins. *Neurology*, 53, 723-729.
- Pennington, B. F., Filipek, P. A., Lefly, D., Churchwell, J., Kennedy, D. N., Simon, J. H., . . . DeFries, J. C. (1999). Brain morphometry in reading-disabled twins. *Neurology*, 53, 723-729.
- Schmitz, S., Fulker, D. W., Plomin, R., Zahn-Waxler, C., Emde, R. N., & DeFries, J. C. (1999). Temperament and problem behaviour during early childhood. *International Journal of Behavioral Development*, 23, 333-355.
- Talebizadeh, Z., Kelley, P. M., Askew, J. W., Beisel, K. W., & Smith, S. D. (1999). A novel mutation in the KCNQ4 gene in a large kindred with dominant progressive hearing loss. *Human Mutation*, 14, 493-501.
- Van Hauwe, P., Coucke, P. J., Declau, F., Kunst, H., Ensink, R. J., Marres, H. A., . . . Van Camp, G. (1999). Deafness linked to DFNA2: one locus but how many genes? *Nature Genetics*, 21, 263.
- Wadsworth, S. J., Fulker, D. W., & DeFries, J. C. (1999). Stability of genetic and environmental influences on reading performance at 7 and 12 years of age in the Colorado Adoption Project. *International Journal of Behavioral Development*, 23, 319-332.
- Willcutt, E. G., Hartung, C. M., Lahey, B. B., Loney, J., & Pelham, W. E. (1999). Utility of behavior ratings by examiners during assessments of preschool children with attention-deficit/hyperactivity disorder. *Journal of Abnormal Child Psychology*, 27, 463-472.
- Willcutt, E. G., Pennington, B. F., Chhabildas, N. A., Friedman, M. C., & Alexander, J. (1999). Psychiatric comorbidity associated with DSM-IV ADHD in a nonreferred sample of twins. *Journal of the American Academy of Child and Adolescent Psychiatry*, 38, 1355-1362.
- Wise, B. W., Ring, J., & Olson, R. K. (1999). Training phonological awareness with and without explicit attention to articulation. *Journal of Experimental Child Psychology*, 72, 271-304.

## 1998

- Alarcón, M., Plomin, R., Fulker, D. W., Corley, R., & DeFries, J. C. (1998). Multivariate path analysis of specific cognitive abilities data at 12 years of age in the Colorado Adoption Project. *Behavior Genetics*, 28, 255-264.
- Aman, C. J., Roberts, R. J., Jr., & Pennington, B. F. (1998). A neuropsychological examination of the underlying deficit in attention deficit hyperactivity disorder: frontal lobe versus right parietal lobe theories. *Developmental Psychology*, 34, 956-969.
- Brookhouser, P. E., & Smith, S. D. (1998). Genetic hearing loss. In B. J. Bailey (Ed.), *Head and Neck Surgery-Otolaryngology, 2nd Edition* (pp. 1311-1327). Philadelphia: Lippincott-Raven.
- Gilger, J. W., Pennington, B. F., Harbeck, R. J., DeFries, J. C., Kotzin, B., Green, P., & Smith, S. (1998). A twin and family study of the association between immune system dysfunction and dyslexia using blood serum immunoassay and survey data. *Brain and Cognition*, 36, 310-333.
- Kelley, P. M., Harris, D. J., Comer, B. C., Askew, J. W., Fowler, T., Smith, S. D., & Kimberling, W. J. (1998). Novel mutations in the connexin 26 gene (GJB2) that cause autosomal recessive (DFNB1) hearing loss. *American Journal of Human Genetics*, 62, 792-799.
- Knopik, V. S., Alarcón, M., & DeFries, J. C. (1998). Common and specific gender influences on individual differences in reading performance: a twin study. *Personality and Individual Differences*, 25, 269-277.
- Knopik, V. S., & DeFries, J. C. (1998). A twin study of gender-influenced individual differences in general cognitive ability. *Intelligence*, 26, 81-89.
- Lahey, B. B., Pelham, W. E., Stein, M. A., Loney, J., Trapani, C., Nugent, K., . . . Baumann, B. (1998). Validity of DSM-IV attention-deficit/hyperactivity disorder for younger children. *Journal of the American Academy of Child and Adolescent Psychiatry*, 37, 695-702.
- Light, J. G., DeFries, J. C., & Olson, R. K. (1998). Multivariate behavioral genetic analysis of achievement and cognitive measures in reading-disabled and control twin pairs. *Human Biology*, 70, 215-237.
- O'Connor, T. G., Deater-Deckard, K., Fulker, D., Rutter, M., & Plomin, R. (1998). Genotype-environment correlations in late childhood and early adolescence: antisocial behavioral problems and coercive parenting. *Developmental Psychology*, 34, 970-981.
- Pennington, B. F., & Bennetto, L. (1998). Toward a neuropsychology of mental retardation. In J. A. Burack, R. M. Hodapp & E. Zigler (Eds.), *Handbook of mental retardation and development* (pp. 80-114).
- Plomin, R., Corley, R., Caspi, A., Fulker, D. W., & DeFries, J. (1998). Adoption results for self-reported personality: evidence for nonadditive genetic effects? *Journal of Personality and Social Psychology*, 75, 211-218.
- Plomin, R., & DeFries, J. C. (1998). The genetics of cognitive abilities and disabilities. *Scientific American*, 278, 62-69.
- Riddle, J. E., Cheema, A., Sobesky, W. E., Gardner, S. C., Taylor, A. K., Pennington, B. F., & Hagerman, R. J. (1998). Phenotypic involvement in females with the FMR1 gene mutation. *American Journal of Mental Retardation*, 102, 590-601.
- Smith, S. D., Brower, A. M., Cardon, L. R., & DeFries, J. C. (1998). Genetics of reading disability: Further evidence for a gene on Chromosome 6. In B. K. Shapiro, P. J. Accardo & A. J. Capute (Eds.), *Specific reading disability: A view of the spectrum* (pp. 63-74). Timonium, MD: York Press, Inc.
- Smith, S. D., & Harker, L. A. (1998). Single gene influences on radiologically-detectable malformations of the inner ear. *Journal of Communication Disorders*, 31, 391-408.

- Smith, S. D., Kelley, P. M., & Brower, A. M. (1998). Molecular approaches to the genetic analysis of specific reading disability. *Human Biology*, 70, 239-256.
- Smith, S. D., Kimberling, W. J., Schaefer, G. B., Horton, M. B., & Tinley, S. T. (1998). Medical genetic evaluation for the etiology of hearing loss in children. *Journal of Communication Disorders*, 31, 371-389.
- Smith, S. D., Schaefer, G. B., Horton, M. B., Tinley, S. T., & Kimberling, W. J. (1998). Medical Genetic Evaluation for the Etiology of Hearing Loss in Children. *Journal of Communication Disorders*, 31, 371-388.

## 1997

- Alarcon, M., & DeFries, J. C. (1997). Reading performance and general cognitive ability in twins with reading difficulties and control pairs. *Personality and Individual Differences*, 22, 793-803.
- Alarcon, M., DeFries, J. C., Light, J. G., & Pennington, B. F. (1997). A twin study of mathematics disability. *Journal of Learning Disabilities*, 30, 617-623.
- Brown, M. R., Tomek, M. S., Van Laer, L., Smith, S., Kenyon, J. B., Van Camp, G., & Smith, R. J. H. (1997). A novel locus for autosomal dominant nonsyndromic hearing loss, DFNA13, maps to chromosome 6p. *American Journal of Human Genetics*, 61, 924-927.
- DeFries, J. C., Alarcón, M., & Olson, R. K. (1997). Genetic aetiologies of reading and spelling deficits: Developmental differences. In C. H. Hulme & M. J. Snowling (Eds.), *Dyslexia: Biology, cognition, and intervention* (pp. 20-37). London: Whurr Publishers Ltd.
- DeFries, J. C., Filipek, P. A., Fulker, D. W., Olson, R. K., Pennington, B. F., Smith, S. D., & Wise, B. W. (1997). Colorado Learning Disabilities Research Center. *Learning Disabilities: A Multidisciplinary Journal*, 8, 7-19.
- Knopik, V. S., Alarcon, M., & DeFries, J. C. (1997). Comorbidity of mathematics and reading deficits: evidence for a genetic etiology. *Behavior Genetics*, 27, 447-453.
- Knopik, V. S., Alarcón, M., & DeFries, J. C. (1997). Comorbidity of mathematics and reading deficits: evidence for a genetic etiology. *Behavior Genetics*, 27, 447-453.
- Pennington, B. F. (1997). Using genetics to dissect cognition. *American Journal of Human Genetics*, 60, 13-16. PMCID: PMC1712539.
- Pennington, B. F. (1997). Dimensions of executive functions in normal and abnormal development. In N. A. Krasnegor & G. R. Lyon (Eds.), *Development of the prefrontal cortex* (pp. 265-281). New York, NY: Paul H. Brookes Publishing Company.
- Pennington, B. F., Rogers, S. J., Bennetto, L., Griffith, E. M., Reed, D. T., & Shyu, V. (1997). Validity tests of the executive dysfunction hypothesis of autism. In J. Russell (Ed.), *Autism as an executive disorder* (pp. 147-178). Oxford: Oxford University Press.
- Pennington, B. F., & Smith, S. D. (1997). Genetic analysis of dyslexia and other complex behavioral phenotypes. *Current Opinion in Pediatrics*, 9, 636-641.
- Plomin, R., DeFries, J. C., McClearn, G. E., & Rutter, M. (1997). *Behavioral genetics (3rd edition)*. New York: Freeman and Company.
- Plomin, R., Fulker, D. W., Corley, R., & DeFries, J. C. (1997). Nature, nurture, and cognitive development from 1 to 16 years: A parent-offspring adoption study. *Psychological Science*, 8, 442-447.
- Sherman, S. L., DeFries, J. C., Gottesman, I. I., Loehlin, J. C., Meyer, J. M., Pelias, M. Z., . . . Waldman, I. (1997). ASHG statement - Recent developments in human behavioral genetics: Past accomplishments and future directions. *American Journal of Human Genetics*, 60, 1265-1275.
- Van Camp, G. V., Coucke, P. J., Kinst, H., Schatteman, I., Van Velzen, D., Marres, H., . . . Willems, P. J. (1997). Linkage analysis of progressive hearing loss in five extended families maps the DFNA2 gene to a 1.25 Mb region on chromosome 1p. *Genomics*, 41.
- Wadsworth, S. J., Corley, R. P., DeFries, J. C., Fulker, D. W., Carey, G., & Plomin, R. (1997). Substance experimentation in the Colorado Adoption Project. *Personality and Individual Differences*, 23, 463-471.
- Wise, B. W., Ring, J., Sessions, L., & Olson, R. K. (1997). Phonological awareness with and without articulation: A preliminary study. *Learning Disability Quarterly*, 20, 211-225.

## 1996

- Bennetto, L., Pennington, B. F., & Rogers, S. J. (1996). Intact and impaired memory functions in autism. *Child Development*, 67, 1816-1835.
- Boetsch, E. A., Green, P. A., & Pennington, B. F. (1996). Psychosocial correlates of dyslexia across the life span. *Development and Psychopathology*, 8, 539-562.
- Casto, S. D., Pennington, B. F., Light, J. G., & DeFries, J. C. (1996). Differential genetic etiology of reading disability as a function of mathematics performance. *Reading and Writing*, 8, 295-306.
- DeFries, J. C., & Alarcón, M. (1996). Genetics of specific reading disability. *Mental Retardation and Developmental Disabilities Research Reviews*, 2, 39-47.
- DeFries, J. C., & Light, J. G. (1996). Twin studies of reading disability. In J. H. Beitchman, N. J. Cohen, M. M. Konstantareas & R. Tannoux (Eds.), *Language, learning, and behavior disorders* (pp. 272-292). New York, NY: Cambridge University Press.
- Gilger, J. W., Borecki, I. B., Smith, S. D., DeFries, J. C., & Pennington, B. F. (1996). The etiology of extreme scores for complex phenotypes: An illustration using reading performance. In C. H. Chase, D. G. Rosen & G. F. Sherman (Eds.), *Developmental dyslexia* (pp. 63-85). Baltimore, MD: York Press.
- Gilger, J. W., Hanebuth, E., Smith, S. D., & Pennington, B. F. (1996). Differential risk for developmental reading disorders in the offspring of compensated versus noncompensated parents. *Reading and Writing*, 8, 407-417.
- Knopik, V. S., DeFries, J. C., & Alarcón, M. (1996). Gender differences in cognitive abilities of opposite-sex and same-sex twin pairs with reading disability. *Annals of Dyslexia*, 46, 241-257.
- Lefly, D. L., & Pennington, B. F. (1996). Longitudinal study of children at high family risk for dyslexia: The first two years. In M. L. Rice (Ed.), *Toward a genetics of language* (pp. 49-75). Mahwah, NJ: Lawrence Erlbaum.
- Markel, P. D., Fulker, D. W., Bennett, B., Corley, R. P., DeFries, J. C., Erwin, V. G., & Johnson, T. E. (1996). Quantitative trait loci for ethanol sensitivity in the LS x SS recombinant inbred strains interval mapping. *Behavior Genetics*, 26, 447-458.

- Pennington, B. F., & Gilger, J. W. (1996). How is dyslexia transmitted? In C. H. Chase, G. D. Rosen & G. F. Sherman (Eds.), *Developmental dyslexia: Neural, cognitive, and genetic mechanisms* (pp. 41-61). Parkton, MD: York Press.
- Pennington, B. F., & Ozonoff, S. (1996). Executive functions and developmental psychopathology. *Journal of Child Psychology and Psychiatry*, 37, 51-87.
- Roberts, R. J., & Pennington, B. F. (1996). An interactive framework for examining prefrontal cognitive processes. *Developmental Neuropsychology*, 12, 105-126.
- Rogers, S. J., Bennetto, L., McEvoy, R., & Pennington, B. F. (1996). Imitation and pantomime in high-functioning adolescents with autism spectrum disorders. *Child Development*, 67, 2060-2073.
- Saudino, K. J., Plomin, R., & DeFries, J. C. (1996). Tester-rated temperament at 14, 20 and 24 months: Environmental change and genetic continuity. *British Journal of Developmental Psychology*, 14, 129-144.
- Schmitz, S., Saudino, K. J., Plomin, R., Fulker, D. W., & DeFries, J. C. (1996). Genetic and environmental influences on temperament in middle childhood: Analyses of teacher and tester ratings. *Child Development*, 67, 409-422.
- Smith, S. D., Pennington, B. F., & DeFries, J. C. (1996). Linkage analysis with complex behavioral traits. In M. L. Rice (Ed.), *Towards a genetics of language*. Mahwah, NJ: Lawrence Erlbaum.
- Sobesky, W. E., Taylor, A. K., Pennington, B. F., Bennetto, L., Porter, D., Riddle, J., & Hagerman, R. J. (1996). Molecular/clinical correlations in females with fragile X. *American Journal of Medical Genetics*, 64, 340-345.

## 1995

- Alarcon, M., DeFries, J. C., & Fulker, D. W. (1995). Etiology of individual differences in reading performance: a test of sex limitation. *Behavior Genetics*, 25, 17-23.
- Braungartreker, J., Rende, R. D., Plomin, R., DeFries, J. C., & Fulker, D. W. (1995). Genetic mediation of longitudinal associations between family environment and childhood behavior problems. *Development and Psychopathology*, 7, 233-245.
- Cardon, L. R., Smith, S. D., Fulker, D. W., Kimberling, W. J., Pennington, B. F., & DeFries, J. C. (1995). Quantitative trait locus for reading disability: correction. *Science*, 268, 1553.
- Cardon, L. R., Smith, S. D., Fulker, D. W., Kimberling, W. J., Pennington, B. F., & DeFries, J. C. (1995). Response. *Science*, 268, 787-788.
- Cardon, L. R., Smith, S. D., Fulker, D. W., Kimberling, W. J., Pennington, B. F., & DeFries, J. C. (1995). Quantitative trait locus for reading disability. *Science*, 268, 1553-1553.
- Casto, S. D., DeFries, J. C., & Fulker, D. W. (1995). Multivariate genetic analysis of Wechsler Intelligence Scale for Children--Revised (WISC-R) factors. *Behavior Genetics*, 25, 25-32.
- Flint, J., Corley, R., DeFries, J. C., Fulker, D. W., Gray, J. A., Miller, S., & Collins, A. C. (1995). A simple genetic basis for a complex psychological trait in laboratory mice. *Science*, 269, 1432-1435.
- Fukushima, K., Ramesh, A., Srikumari Srisailapathy, C. R., Ni, L., Chen, A., O'Neill, M., . . . Smith, R. J. H. (1995). Consanguineous nuclear families used to identify a new locus for recessive non-syndromic hearing loss on 14q. *Human Molecular Genetics*, 4, 1643-1648.
- Fukushima, K., Ramesh, A., Srisailapathy, C. R., Ni, L., Wayne, S., O'Neill, M. E., . . . Smith, R. J. H. (1995). An autosomal recessive non-syndromic form of sensorineural hearing loss maps to 3p-DFNB6. *Genome Research*, 5, 305-308.
- Gilger, J. W., & Pennington, B. F. (1995). Why associations among traits do not necessarily indicate their common etiology: a comment on the Geschwind-Behan-Galaburda model. *Brain and Cognition*, 27, 89-93.
- Hyona, J., & Olson, R. K. (1995). Eye fixation patterns among dyslexic and normal readers: Effects of word length and word frequency. *Journal of Experimental Psychology-Learning Memory and Cognition*, 21, 1430-1440.
- Light, J. G., & DeFries, J. C. (1995). Comorbidity of reading and mathematics disabilities: genetic and environmental etiologies. *Journal of Learning Disabilities*, 28, 96-106.
- Light, J. G., Pennington, B. F., Gilger, J. W., & DeFries, J. C. (1995). Reading Disability and Hyperactivity Disorder - evidence for a common genetic etiology. *Developmental Neuropsychology*, 11, 323-335.
- Markel, P. D., DeFries, J. C., & Johnson, T. E. (1995). Ethanol-induced anesthesia in inbred strains of long-sleep and short-sleep mice: a genetic analysis of repeated measures using censored data. *Behavior Genetics*, 25, 67-73.
- Markel, P. D., DeFries, J. C., & Johnson, T. E. (1995). Use of repeated measures in an analysis of ethanol-induced loss of righting reflex in inbred long-sleep and short-sleep mice. *Alcoholism: Clinical and Experimental Research*, 19, 299-304.
- Pennington, B. F. (1995). Genetics of learning disabilities. *Journal of Child Neurology*, 10 Suppl 1, S69-77.
- Pennington, B. F., Bennetto, L., McAleer, O. K., & Roberts, R. J., Jr. (1995). Executive functions and working memory: Theoretical measurement issues. In G. R. Lyon & N. A. Krasnegor (Eds.), *Attention, memory, and executive function* (pp. 327-348). New York, NY: Paul H. Brookes Publishing Company.
- Pennington, B. F., & Welsh, M. C. (1995). Neuropsychology and developmental psychopathology. In D. V. Cicchetti & D. J. Cohen (Eds.), *Manual of developmental psychopathology* (pp. 254-290). New York, NY: Wiley and Sons.
- Schaefer, G. B., Novak, K., Steele, D., Buehler, B., Smith, S. D., Zaleski, D., . . . Sanger, W. (1995). Familial inverted duplication 7p. *American Journal of Medical Genetics*, 56, 184-187.
- Smith, S. D. (1995). Overview of genetic auditory syndromes. *Journal of the American Academy of Audiology*, 6, 1-14.
- Sobesky, W. E., Porter, D., Pennington, B. F., & Hagerman, R. J. (1995). Dimensions of shyness in fragile X females. *Developmental Brain Dysfunction*, 8, 280-292.
- Van Camp, G., Coucke, P., Balemans, W., Van Velzen, D., Van de Bilt, C., Van Laer, L., . . . Willemse, P. J. (1995). Localization of a gene for non-syndromic hearing loss (DFNA5) to chromosome 7p15. *Human Molecular Genetics*, 4, 2159-2163.
- Wadsworth, S. J., DeFries, J. C., Fulker, D. W., Olson, R. K., & Pennington, B. F. (1995). Reading performance and verbal short-term memory - A twin study of reciprocal causation. *Intelligence*, 20, 145-167.
- Wadsworth, S. J., DeFries, J. C., Fulker, D. W., & Plomin, R. (1995). Cognitive ability and academic achievement in the Colorado Adoption Project: a multivariate genetic analysis of parent-offspring and sibling data. *Behavior Genetics*, 25, 1-15.
- Wadsworth, S. J., DeFries, J. C., Fulker, D. W., & Plomin, R. (1995). Covariation among measures of cognitive ability and academic

achievement in the Colorado Adoption Project - Sibling Analysis. *Personality and Individual Differences*, 18, 63-73.  
Wise, B. W., & Olson, R. K. (1995). Computer-based phonological awareness and reading instruction. *Annals of Dyslexia*, 45, 99-122.

### 1994

- Alarcon, M., DeFries, J. C., & Gillis, J. J. (1994). Familial resemblance for measures of reading performance in families of reading disabled and control twins. *Reading and Writing*, 6, 93-101.
- Cardon, L. R., Smith, S. D., Fulker, D. W., Kimberling, W. J., Pennington, B. F., & DeFries, J. C. (1994). Quantitative trait locus for reading disability on chromosome 6. *Science*, 266, 276-279.
- Cherny, S. S., Fulker, D. W., Corley, R. P., Plomin, R., & DeFries, J. C. (1994). Continuity and change in infant shyness from 14 to 20 months. *Behavior Genetics*, 24, 365-379.
- Cherny, S. S., Fulker, D. W., Emde, R. N., Robinson, J., Corley, R. P., Reznick, J. S., . . . DeFries, J. C. (1994). A developmental-genetic analysis of continuity and change in the Bayley Mental Development Index from 14 to 24 Months: the MacArthur Longitudinal Twin Study. *Psychological Science*, 5, 354-360.
- Coucke, P. J., Van Camp, G., Djayodiharjo, B., Smith, S. D., Frants, R. R., Padberg, G. W., . . . Willems, P. J. (1994). Linkage of autosomal dominant hearing loss to the short arm of chromosome 1 in two families. *New England Journal of Medicine*, 331, 425-431.
- DeFries, J. C., Plomin, R., & Fulker, D. W. (1994). *Nature and nurture during middle childhood*. Oxford: Blackwell Publishers.
- Gilger, J. W., Borecki, I. B., DeFries, J. C., & Pennington, B. F. (1994). Commingling and segregation analysis of reading performance in families of normal reading probands. *Behavior Genetics*, 24, 345-355.
- Mazzocco, M. M. M., Nord, A. M., Vandoornicck, W., Greene, C. L., Kovar, C. G., & Pennington, B. F. (1994). Cognitive development among children with early-treated phenylketonuria. *Developmental Neuropsychology*, 10, 133-151.
- Mazzocco, M. M. M., Pennington, B. F., & Hagerman, R. J. (1994). Social cognition skills among females with Fragile X. *Journal of Autism and Developmental Disorders*, 24, 473-485.
- Olson, R. K. (1994). Language deficits in "specific" reading disability. In M. Gernsbacher (Ed.), *Handbook of Psycholinguistics* (pp. 895-916). New York, NY: Academic Press.
- Olson, R. K., Forsberg, H., & Wise, B. (1994). Genes, environment, and the development of orthographic skills. In V. W. Berninger (Ed.), *The varieties of orthographic knowledge I: Theoretical and developmental issues* (pp. 27-72). Dordrecht, Netherlands: Kluwer Academic Publishers.
- Olson, R. K., Forsberg, H., Wise, B., & Rack, J. (1994). Measurement of word recognition, orthographic, and phonological skills. In G. R. Lyon (Ed.), *Frames of reference for the assessment of learning disabilities: New views on measurement issues* (pp. 243-277). Baltimore, MD: Paul H. Brookes Publishing Company.
- Smolen, A., Marks, M. J., DeFries, J. C., & Henderson, N. D. (1994). Individual differences in sensitivity to nicotine in mice: response to six generations of selective breeding. *Pharmacology Biochemistry and Behavior*, 49, 531-540.
- Sobesky, W. E., Pennington, B. F., Porter, D., Hull, C. E., & Hagerman, R. J. (1994). Emotional and neurocognitive deficits in fragile X. *American Journal of Medical Genetics*, 51, 378-385.
- Wise, B. W. (1994). Beginning to spell: a study of 1st-Grade children. *Contemporary Psychology*, 39, 844-845.

### 1993

- Coon, H., Carey, G., Fulker, D. W., & DeFries, J. C. (1993). Influences of school environment on the academic achievement scores of adopted and nonadopted children. *Intelligence*, 17, 79-104.
- DeFries, J. C., & Gillis, J. J. (1993). Genetics of reading disability. In R. Plomin & G. E. McClearn (Eds.), *Nature, nurture and psychology* (pp. 121-145). Washington, DC: American Psychological Association.
- DeFries, J. C., Gillis, J. J., & Wadsworth, S. J. (1993). Genes and genders: A twin study of reading disability. In A. M. Galaburda (Ed.), *Dyslexia and development: Neurobiological aspects of extra-ordinary brains* (pp. 187-204). Cambridge, MA: Harvard University Press.
- Mazzocco, M. M. M., Pennington, B. F., & Hagerman, R. J. (1993). The neurocognitive phenotype of female carriers of Fragile X: Additional evidence for specificity. *Journal of Developmental and Behavioral Pediatrics*, 14, 328-335.
- McEvoy, R. E., Rogers, S. J., & Pennington, B. F. (1993). Executive function and social communication deficits in young autistic children. *Journal of Child Psychology and Psychiatry*, 34, 563-578.
- Mitton, J. B., Schuster, W. S. F., Cothran, E. G., & Defries, J. C. (1993). Correlation between the Individual Heterozygosity of Parents and Their Offspring. *Heredity*, 71, 59-63.
- Ozonoff, S., Rogers, S. J., Farnham, J. M., & Pennington, B. F. (1993). Can standard measures identify subclinical markers of autism? *Journal of Autism and Developmental Disorders*, 23, 429-441.
- Pennington, B. F., & Bennetto, L. (1993). Main Effects Or Transactions in the Neuropsychology of Conduct Disorder - the Neuropsychology of Conduct Disorder - Commentary. *Development and Psychopathology*, 5, 153-164.
- Pennington, B. F., Groisser, D., & Welsh, M. C. (1993). Contrasting cognitive deficits in attention-deficit hyperactivity disorder versus reading disability. *Developmental Psychology*, 29, 511-523.
- Plomin, R., Emde, R. N., Braungart, J. M., Campos, J., Corley, R., Fulker, D. W., . . . DeFries, J. C. (1993). Genetic change and continuity from 14 to 20 months - the MacArthur Longitudinal Twin Study. *Child Development*, 64, 1354-1376.
- Rack, J. P., & Olson, R. K. (1993). Phonological deficits, IQ, and individual differences in reading disability: Genetic and environmental influences. *Developmental Review*, 13, 269-278.
- Stevenson, J., Pennington, B. F., Gilger, J. W., DeFries, J. C., & Gillis, J. J. (1993). Hyperactivity and spelling disability: testing for shared genetic aetiology. *Journal of Child Psychology and Psychiatry*, 34, 1137-1152.
- Swanson, J. M., McBurnett, K., Wigal, T., Pfiffner, L., Lerner, M., Williams, L., . . . Crinella, F. (1993). Effect of stimulant medication on children with Attention Deficit Disorder: A "review of reviews". *Exceptional Children*, 60, 154-162.
- Wadsworth, S. J., DeFries, J. C., & Fulker, D. W. (1993). Cognitive abilities of children at 7 and 12 years of age in the Colorado

adoption project. *Journal of Learning Disabilities*, 26, 611-615.

## 1992

- Braungart, J. M., Plomin, R., DeFries, J. C., & Fulker, D. W. (1992). Genetic influence on tester-rated infant temperament as assessed by Bayley Infant Behavior Record: Nonadoptive and adoptive siblings and twins. *Developmental Psychology*, 28, 40-47.
- Cardon, L. R., Corley, R. P., DeFries, J. C., Plomin, R., & Fulker, D. W. (1992). Factorial validation of a telephone test battery of specific cognitive abilities. *Personality and Individual Differences*, 13, 1047-1050.
- Cardon, L. R., Fulker, D. W., DeFries, J. C., & Plomin, R. (1992). Multivariate genetic analysis of specific cognitive abilities in the Colorado Adoption Project at Age 7. *Intelligence*, 16, 383-400.
- Cardon, L. R., Fulker, D. W., DeFries, J. C., & Plomin, R. (1992). Continuity and change in general cognitive ability from 1 to 7 Years of Age. *Developmental Psychology*, 28, 64-73.
- Cherny, S. S., Cardon, L. R., Fulker, D. W., & DeFries, J. C. (1992). Differential heritability across levels of cognitive ability. *Behavior Genetics*, 22, 153-162.
- Cherny, S. S., DeFries, J. C., & Fulker, D. W. (1992). Multiple regression analysis of twin data: a model-fitting approach. *Behavior Genetics*, 22, 489-497.
- Emde, R. N., Plomin, R., Robinson, J. A., Corley, R., DeFries, J., Fulker, D. W., . . . Zahn-Waxler, C. (1992). Temperament, emotion, and cognition at fourteen months: the MacArthur Longitudinal Twin Study. *Child Development*, 63, 1437-1455.
- Gilger, J. W., Pennington, B. F., & DeFries, J. C. (1992). A twin study of the etiology of comorbidity: attention-deficit hyperactivity disorder and dyslexia. *Journal of the American Academy of Child and Adolescent Psychiatry*, 31, 343-348.
- Gilger, J. W., Pennington, B. F., Green, P., Smith, S. M., & Smith, S. D. (1992). Reading disability, immune disorders and non-right-handedness: twin and family studies of their relations. *Neuropsychologia*, 30, 209-227.
- Gillis, J. J., DeFries, J. C., & Fulker, D. W. (1992). Confirmatory factor analysis of reading and mathematics performance: a twin study. *Acta Geneticae Medicae et Gemellologiae*, 41, 287-300.
- Gillis, J. J., Gilger, J. W., Pennington, B. F., & DeFries, J. C. (1992). Attention deficit disorder in reading-disabled twins: evidence for a genetic etiology. *Journal of Abnormal Child Psychology*, 20, 303-315.
- Johnson, T. E., DeFries, J. C., & Markel, P. D. (1992). Mapping quantitative trait loci for behavioral traits in the mouse. *Behavior Genetics*, 22, 635-653.
- Mazzocco, M. M. M., Hagerman, R. J., Cronister-Silverman, A., & Pennington, B. F. (1992). Specific frontal-lobe deficits among women with the Fragile-X gene. *Journal of the American Academy of Child and Adolescent Psychiatry*, 31, 1141-1148.
- Mazzocco, M. M. M., Hagerman, R. J., & Pennington, B. F. (1992). Problem-solving limitations among cytogenetically expressing Fragile-X women. *American Journal of Medical Genetics*, 43, 78-86.
- Mazzocco, M. M. M., Yannicelli, S., Nord, A. M., Vandoornicx, W., Davidsonmundt, A. J., Pennington, B. F., & Greene, C. L. (1992). Cognition and Tyrosine Supplementation Among School-Aged Children with Phenylketonuria. *American Journal of Diseases of Children*, 146, 1261-1264.
- Olson, R. K., & Wise, B. W. (1992). Reading on the computer with orthographic and speech feedback: an overview of the Colorado Remediation Project. *Reading and Writing*, 4, 107-144.
- Pennington, B. F., & Gilger, J. W. (1992). Dyslexia. *New England Journal of Medicine*, 327, 280; author reply 280-281.
- Pennington, B. F., Gilger, J. W., Olson, R. K., & DeFries, J. C. (1992). The external validity of age- versus IQ-discrepancy definitions of reading disability: lessons from a twin study. *Journal of Learning Disabilities*, 25, 562-573.
- Plomin, R., Corley, R., DeFries, J. C., & Fulker, D. W. (1992). Childrens Television Viewing - Response. *Psychological Science*, 3, 75-76.
- Rack, J. P., Snowling, M. J., & Olson, R. K. (1992). The nonword reading deficit in developmental dyslexia - a review. *Reading Research Quarterly*, 27, 28-53.
- Smith, S. D. (1992). Identification of genetic influences. *Journal of Communication Disorders*, 2, 73-85.
- Smith, S. D. (1992). Genetic Counseling. In J. G. Clark & F. N. Martin (Eds.), *Effective Counseling In Audiology: Perspectives and Practice* (pp. 70-91). New York: Prentice Hall.
- Smith, S. D., Kimberling, W. J., & Pennington, B. F. (1992). Screening for multiple genes influencing dyslexia. In B. F. Pennington (Ed.), *Reading Disabilities: Genetic and Neurological Influences* (pp. 97-110). Dordrecht, the Netherlands: Kluwer Academic Publishers.
- Swanson, J. M., Willcutt, E. G., McBurnett, R. K., Cantwell, D., Lerner, M., Pfiffner, L. J., . . . Tamm, L. (1992). Treatment of ADHD: Beyond Medication. *Beyond Behavior*, 4, 12-22.
- Wadsworth, S. J., DeFries, J. C., Stevenson, J., Gilger, J. W., & Pennington, B. F. (1992). Gender ratios among reading-disabled children and their siblings as a function of parental impairment. *Journal of Child Psychology and Psychiatry*, 33, 1229-1239.
- Waldman, I. D., DeFries, J. C., & Fulker, D. W. (1992). Quantitative genetic analysis of IQ development in young children: Multivariate multiple-regression with orthogonal polynomials. *Behavior Genetics*, 22, 229-238.
- Wise, B. W. (1992). Whole words and decoding for short-term learning: Comparisons on a talking-computer system. *Journal of Experimental Child Psychology*, 54, 147-167.
- Wise, B. W., & Olson, R. K. (1992). How poor readers and spellers use interactive speech in a computerized spelling program. *Reading and Writing*, 4, 145-163.

## 1991

- DeFries, J. C., & Gillis, J. J. (1991). Etiology of reading deficits in learning disabilities: Quantitative genetic analysis. In J. E. Obrzut & G. W. Hynd (Eds.), *Neuropsychological foundations of learning disabilities: A handbook of issues, methods and practice* (pp. 29-47). Orlando, FL: Academic Press.
- DeFries, J. C., Olson, R. K., Pennington, B. F., & Smith, S. D. (1991). Colorado Reading Project: Past, present, and future. *Learning Disabilities: A Multidisciplinary Journal*, 2, 37-46.

- DeFries, J. C., Olson, R. K., Pennington, B. F., & Smith, S. D. (1991). Colorado Reading Project: An update. In D. D. Duane & D. B. Gray (Eds.), *The reading brain: The biological basis of dyslexia* (pp. 53-87). Parkton, MD: York Press.
- DeFries, J. C., Stevenson, J., Gillis, J. J., & Wadsworth, S. J. (1991). Genetic etiology of spelling deficits in the Colorado and London Twin Studies of Reading Disability. *Reading and Writing*, 3, 271-283.
- Fulker, D. W., Cardon, L. R., DeFries, J. C., Kimberling, W. J., Pennington, B. F., & Smith, S. D. (1991). Multiple-regression analysis of sib-pair data on reading to detect quantitative trait loci. *Reading and Writing*, 3, 299-313.
- Fulker, D. W., Cardon, L. R., DeFries, J. C., Kimberling, W. J., Pennington, B. F., & Smith, S. D. (1991). Multiple regression analysis of sib-pair data on reading to detect quantitative trait loci. *Reading and Writing: An Interdisciplinary Journal*, 3, 299-313.
- Gilger, J. W., Pennington, B. F., & DeFries, J. C. (1991). Risk for reading disability as a function of parental history in 3 family studies. *Reading and Writing*, 3, 205-217.
- Olson, R. K., Gillis, J. J., Rack, J. P., DeFries, J. C., & Fulker, D. W. (1991). Confirmatory factor analysis of word recognition and process measures in the Colorado Reading Project. *Reading and Writing*, 3, 235-248.
- Olson, R. K., Rack, J. P., Conners, F. A., DeFries, J. C., & Fulker, D. W. (1991). Genetic etiology of individual differences in reading disability. In L. V. Feagans, E. J. Short & L. J. Meltzer (Eds.), *Subtypes of learning disabilities: Theoretical perspectives and research*. Hillsdale, NJ: Lawrence Erlbaum Associates.
- Ozonoff, S., Pennington, B. F., & Rogers, S. J. (1991). Executive function deficits in high-functioning autistic individuals: relationship to theory of mind. *Journal of Child Psychology and Psychiatry*, 32, 1081-1105.
- Ozonoff, S., Rogers, S. J., & Pennington, B. F. (1991). Asperger's syndrome: evidence of an empirical distinction from high-functioning autism. *Journal of Child Psychology and Psychiatry*, 32, 1107-1122.
- Pennington, B. F. (1991). Genetic and neurological influences on reading disability: An overview. *Reading and Writing*, 3, 191-201.
- Pennington, B. F. (1991). Genetics of learning disabilities. *Seminars in Neurology*, 11, 28-34.
- Pennington, B. F., Gilger, J. W., Pauls, D., Smith, S. A., Smith, S. D., & DeFries, J. C. (1991). Evidence for major gene transmission of developmental dyslexia. *Journal of the American Medical Association*, 266, 1527-1534.
- Plomin, R., Coon, H., Carey, G., DeFries, J. C., & Fulker, D. W. (1991). Parent-offspring and sibling adoption analyses of parental ratings of temperament in infancy and childhood. *Journal of Personality*, 59, 705-732.
- Rogers, S. J., & Pennington, B. F. (1991). A theoretical approach to the deficits in infantile autism. *Development and Psychopathology*, 3, 137-162.
- Smith, S. D. (1991). Recurrence Risks. *Annals of the New York Academy of Sciences*, 603, 203-211.
- Smith, S. D., Kimberling, W. J., & Pennington, B. F. (1991). Screening for multiple genes influencing dyslexia. *Reading and Writing*, 3, 285-298.
- Stevenson, J., & DeFries, J. C. (1991). Twin studies of spelling. In L. Richards & P. Jeffrey (Eds.), *Research information for teachers* (pp. 1-4). Wellington, New Zealand: Australian Council for Educational Research and the New Zealand Council for Educational Research.
- Welsh, M. C., Pennington, B. F., & Groisser, D. B. (1991). A normative developmental study of executive function: A window on prefrontal function in children. *Developmental Neuropsychology*, 7, 131-149.