



HAL
open science

Elderly persons with elderly fathers - do they face additional risks?

Karen Ritchie

► **To cite this version:**

Karen Ritchie. Elderly persons with elderly fathers - do they face additional risks?. *International Psychogeriatrics*, 2008, 20 (3), pp.427-9. 10.1017/s1041610207006242 . inserm-00170307

HAL Id: inserm-00170307

<https://inserm.hal.science/inserm-00170307>

Submitted on 6 Jun 2008

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Elderly persons with elderly fathers – do they face additional risks ?

Karen Ritchie

French National Institute of Medical Research (INSERM)

Research Unit U888 “Nervous System Pathologies”

La Colombière Hospital

Montpellier

France

Psychogeriatric research has explored many factors likely to influence our mental health in later life, but one which has received surprisingly little attention given the current interest in genetic determinants, has been paternal age. We now know that both delayed motherhood and delayed fatherhood may have a significant detrimental effect on the mental health of the offspring, but by different mechanisms. While delayed motherhood has been associated with higher rates of obstetric and perinatal problems, delayed fatherhood has been associated with higher risk of new inheritable-mutation disorders. Tarin et al. (1998) have postulated this is because ageing leads to a reduction in the activities of antioxidant enzymes within the seminal plasma and spermatozoa making them more vulnerable to mutational changes. In most syndromes the mutation rate increases with paternal age at an exponential rate; a rate much higher than in elderly women perhaps due to the greater number of cell divisions in the male germ line (Crow 1997).

High paternal age has been found to have a robust association with numerous neuropsychiatric disorders including psychoses (El-Saadi et al; 2004), in particular schizophrenia (Tsuchiya et al. 2005; Malaspina et al. 2001), and to a lesser extent with epilepsy (Vestergaard et al. 2005) and autism (Reichenberg et al. 2006). Together these findings suggest the role of an increase in paternal germ cell mutation with age which may at least partly explain the persistence of these disorders within the human population given that persons with neuropsychiatric disorders have lowered fertility and reduced chances of marriage.

The impact of paternal age on cognitive functioning is of particular interest in relation to ageing populations given that genetic determinants of IQ are now thought to exercise a greater influence than environmental factors with ageing. Given the substantial and complex impact of genetic factors on intelligence, it might be expected that *de novo* genetic events in male

germ cells may be likely to have a detrimental effect on information processing abilities in offspring, and the effects of this more visible with ageing. Evidence from a birth cohort suggests that there is a U-shaped relationship between paternal age and IQ at adolescence controlling for maternal age, education, social class, sex, birth order, birth weight and obstetric complications (Malaspina et al. (2005). The adverse effect of paternal age was found in this study to be greater in relation to performance than verbal IQ.

Most studies of paternal age and intellectual functioning have been carried out on children and young adults, so that very little is known about how this factor may impact on mental health and cognitive functioning at higher ages. The relationship between paternal age and Alzheimer's disease is an old book which has been opening and shutting for over a quarter of a century with conflicting results ranging from increased risk (Urakami et al. 1989), no relationship (Corkin et al. 1983; Hofman et al. 1990; Fratiglioni et al. 1993), or even increased risk with younger paternal age (Farrer et al. 1991). The principal limitations of most of these studies, however, has been their small sample sizes (mostly case-control studies), naïve statistical analyses (often univariate), failure to adequately take into account confounding factors, and mistakenly confounding sex adjustment with sex stratification. More recent studies based on larger population cohorts, and able to control for multiple interacting and confounding factors suggest a more complex link. Re-analyzing previous case-control studies by meta-analysis, van Dijn et al. (1994) found an increased risk with higher paternal age. Whalley et al. (2001) conducted an integrative analysis of Scottish data and found increased paternal age to be a significant risk factor for both AD and vascular dementia in men only. These researchers also found increased paternal age to be related to pre-senile dementia in men only (Whalley et al. 1995). A further refinement of the question is proposed by Bertram et al. (1998) who assessed probability of carrying a major gene for AD in a series of 206 AD cases and found fathers of patients with a low major gene probability were significantly older than high probability groups and controls. These observations are consistent with the *de novo* mutation hypothesis. On the other hand there has been little interest in looking at paternal age and late-life cognitive functioning in the absence of dementia.

And what of fragility for other disorders ? There are in fact very few studies of mental health and paternal age other than AD. Ptok et al. (2000) reported no relationship with either dementia or depression in a short article in *International Psychogeriatrics* in 2000, but this study was based on a simple ANOVA comparison of clinical groups adjusting only for sex.

Moreover the groups did not contain many subjects whose fathers were over 40 (the high risk group identified in studies of other neurological disorders in younger subjects). There are a number of difficulties inherent in evaluating the question of mental health and parental age, not the least of which is the fact that more recent population genetic studies have incidentally found that second and third children in a family are commonly by another father without the child's knowledge (personal communication).

Birth cohort data is of course the best source for tracking the relationship between paternal age and late life mental health, limiting errors due to mortality bias, inaccurate recall of father's age and correct identification of the father. Rodgers (1990) has already reported that older father's age in a UK birth cohort was associated with higher offspring PSE scores at 36 years in men. Evaluations of more recent examinations of the cohort will reveal whether this vulnerability continues into old age, and how it may interact with other ageing-related factors. From a longitudinal population study of psychiatric disorder in elderly conducted in Montpellier in the south of France we have found a relationship between older fathers and probability of depressive disorders in late life (unpublished observations). Given current interest in genetic determinants of mental health in late life, it is perhaps timely to look again at the role of paternal age, especially given the availability of large population data sets with DNA banks and increasingly better phenotype data. While women have long been concerned about the effect of increasing maternal age on the well-being of their offspring, the question has received little publicity in regard to men. The question needs to be asked as to whether older fathers are not only putting their children at greater risk of childhood neurological disorders but also of a sadder old age.

References

Bertram L, Busch R, Spiegl M, Lautenschlager NT, Müller U, Kurz A. Paternal age is a risk factor for Alzheimer's disease in the absence of a major gene. **Neurogenetics** 1998 1: 277-280

Corkin S, Growdon JH, Rasmussen SL. Parental age as a risk factor for Alzheimer's disease. **Ann Neurol** 1983; 13: 674-676

Crow, JF. The high spontaneous mutation rate: is it a health risk ? **Proc Nat Acad Sci USA** 1997; 94: 8380-8386

El-Saadi O, Pedersen CB, McNeil TF, Saha S, Welham J, O'Callaghan E, Cantor-Graae E, Chant D, Mortensen PB, McGrath J. Paternal and maternal age as risk factors for psychosis : findings from Denmark, Sweden and Australia. *Schizophr Res* 2004; 67: 227-236

Farrer LA, Cupples LA, Connor L, Wolf PA, Growdon JH. Association of decreased paternal age and late-onset Alzheimer's disease. An example of genetic imprinting ? **Arch Neurol** 1991; 48: 599-604

Fratiglioni L, Ahlbom A, Viitanen M, Winblad B. Risk factors for late-onset Alzheimer's disease: a population-based, case control study. **Ann Neurol** 1993; 33: 258-266

Hofman A, van Dijn CM, Schulte W, Tanja TA, Haaxma R, Lameris AJ, Saan RJ. Is parental age related to the risk of Alzheimer's disease. **Br J Psychiatry** 1990; 157: 273-275

Malaspina D. Paternal factors and schizophrenia risk: *de novo* mutations and imprinting. *Schizophrenia Bull* 2001; 3: 379-393

Malaspina D, Reichenberg A, Weiser M, Fennig S, Davidson M, Harlap S, Wolitzky R, Rabinowitz J, Susser E, Knobler HY. Paternal age and intelligence: implications for age-related genomic changes in male germ cells. *Psychiatr Genet* 2005; 15: 117-125

Ptok U, Papassotiropoulos A, Maier W, Heun R. Advanced parental age: a risk factor for Alzheimer's disease or depression in the elderly ? *Int Psychogeriatr* 2000; 12: 445-451

Reichenberg A, Gross R, Weiser M, Bresnahan M, Silverman J, Harlap S, Rabinowitz J, Shulman C, Malaspina D, Lubin G, Knobler HY, Davidson M, Susser E. Advancing paternal age and autism. *Arch Gen Psychiatry* 2006; 63: 1026-1032

Rodgers B. Adult affective disorder and early environment. **Br J Psychiat** 1990;157:539-550

Tarin JJ, Brines J, Cano A. Long-term effects of delayed parenthood. **Human Reproduction** 1998; 13: 2371-2376

Tsuchiya KJ, Takagai S, Kawai M, Matsumoto H, Nakamura K, Minabe Y, Mori N, Takei N. Advanced paternal age associated with an elevated risk for schizophrenia in offspring in a Japanese population. **Schizophr Res** 2005 76: 337-342

Urakami K, Adachi Y, Takahashi K. A community-based study of parental age at the birth of patients with dementia of the Alzheimer type. **Arch Neurol** 1989; 46: 38-39

Van Dijn CM, Clayton DG, Chandra V, Fratiglioni L, Graves AB. Interaction between genetic and environmental risk factors for Alzheimer's disease : a re-analysis of case-control studies. EURODEM Risk Factors Research Group. **Genetic Epidemiology** 1994; 11: 539-551

Vestergaard M, Mork A, Madsen KM, Olsen J. Paternal age and epilepsy in the offspring. *Eur J Epidemiol* 2005; 20: 1003-1005

Whalley LJ. Early-onset Alzheimer's disease in Scotland: environmental and familial factors. *Br J Psychiatry Suppl* 2001; 40: s53-59

Whalley LJ, Thomas BM, Starr JM. Epidemiology of presenile Alzheimer's disease in Scotland (1974-1988)II. Exposures to possible risk factors. **Br J Psychiatry** 1995; 167: 732-738