

Estimation of the frequency of involuntary infertility on a nation-wide basis

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1 ABSTRACT

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3 BACKGROUND: Assessing the couple fecundity on a nation-wide basis without excluding 4 couples who eventually remain infertile is challenging. Our aim was to describe the couple 5 fecundity (in terms of frequency of involuntary infertility) among the general population living in France. METHODS: We used a current-duration design. A random sample of 64,262 6 7 households was selected in 2007-2008, allowing us to identify 15,810 women aged 18-44 8 years. Eligible women (n=1089) were those having regular sexual intercourse with a male 9 partner, not using any method of contraception and not having delivered in the previous 3 10 months. These women reported information on the current duration of unprotected 11 intercourse (CDUI, the time elapsed between the start of the period of unprotected 12 intercourse and the time of inclusion in the study). The CDUI distribution was used to 13 estimate the frequency of involuntary infertility, using a newly developed statistical technique 14 that does not require couples to be followed-up until the end of the period of unprotected 15 intercourse. RESULTS: CDUI was defined for 867 women. An estimated 46% of couples had 16 no detected pregnancy conceived during the first 6 months of unprotected intercourse (95% 17 confidence interval, 36-56%). The proportions of couples with no detected pregnancy within 18 12 and 24 months were 24% (19-30%) and 11% (8-14%), respectively. CONCLUSIONS: 19 These results constitute one of the few descriptions of the fecundity of a nation-wide 20 representative sample of couples from the general population, not limited to couples who 21 eventually conceived or to those resorting to medical help.

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23 Key words: Fecundability; Fecundity; Infertility; Involuntary infertility; Pregnancy;

INTRODUCTION 1

2 Few descriptions of the couples' ability to conceive a pregnancy (fecundity) exist in non-3 selected groups of the general population. Fertility, the actual number of children per woman, 4 is monitored in most countries and is statistically associated with having a time to pregnancy 5 longer than 12 months (Joffe, et al., 2009). However, outside the specific context of populations not using contraceptive methods (Larsen and Vaupel, 1993), it constitutes a very 6 7 indirect and probably not very sensitive indicator of the occurrence of impaired fecundity. 8 Data on the use of assisted reproduction technologies (ART) are widely available (de 9 Mouzon, et al., 2010, de Mouzon, et al., 2009), but information on the number of couples 10 resorting to ART is difficult to interpret without knowledge of the population at risk (those 11 trying to become pregnant), and because involuntarily infertile couples seek for medical help 12 after variable durations of unprotected intercourse (Moreau, et al., 2010) and do not always 13 resort to ART.

14 Fecundity studies relying on a pregnancy-based design have been conducted in several 15 countries (Jensen, et al., 2001, Juul, et al., 1999, Muller, et al., 2006) (reviewed by Leridon, 16 2007), but this design excludes couples who do not conceive finally, thus overestimating the 17 fecundity level. Few surveys including couples who remain infertile have been conducted. 18 The European Studies of Infertility and Subfecundity (Karmaus, et al., 1999), a set of 19 retrospective studies conducted in Denmark, Germany, Poland, Italy and Spain in 1991-93, 20 considered unsuccessful attempts at pregnancy. Some cohort studies of pregnancy planners 21 have been conducted, but generally for etiologic or biologic purposes rather than to describe 22 fecundity at the population level, and in selected populations (Bonde, et al., 1998, Buck, et 23 al., 2004, reviewed by Guzick and Swan, 2006, Tietze, 1968, Weinberg, et al., 1994, Wilcox, 24 et al., 1988, Zinaman, et al., 2000). Demographers have also studied perceived fecundity or 25 impaired fecundity in the general population, for example in France (Leridon, 1992, Leridon, 26 2007) or in the US National Survey of Family Growth (Stephen and Chandra, 2006); these 27 studies, usually relying on large and well-defined population based samples are difficult to 28 compare to the above-mentioned "time to pregnancy" studies, either because they did not use an objective cut-off (e.g., 12 or 24 months) to define involuntary infertility (Leridon, 1992)
or, in the case of the National Survey of Family Growth, because the number of cases of
(current) 12-month infertility is not divided by the number of couples (currently) at risk of
pregnancy, but by the size of the (larger) group of married women, regardless of whether
they were "at risk" of pregnancy, leading to a much lower rate of involuntary infertility than in
other studies (Guzick and Swan, 2006).

7 Our aim was to describe the frequency of involuntary infertility among the general population 8 of France without excluding couples involuntarily infertile. In terms of approach and study 9 design, the need to efficiently include infertile couples (Olsen and Rachootin, 2003, Sallmen, et al., 2000, Slama, et al., 2004) implied the exclusion of retrospective designs. Among 10 11 prospective designs, the main options are the incident cohort design (Bonde, et al., 1998), 12 the prevalent cohort design (Keiding, 1992, Wise, et al., 2010), and the current duration 13 approach (Keiding, et al., 2002, Slama, et al., 2006). The very low eligibility rate of the incident cohort design -probably around 1% of women of reproductive age (Bonde, et al., 14 15 1998, Slama, et al., 2006) - made it in our eyes little suited for descriptive studies. The first 16 stage of the prevalent cohort design requires one to recruit a cross-sectional sample of 17 couples not using contraception and to collect the time elapsed since the start of the period 18 at risk of pregnancy to account for left-censoring in the analysis; this first stage also 19 corresponds to a current duration approach, which is therefore embedded into the prevalent 20 cohort design. We have recently demonstrated the feasibility of this design (Slama, et al., 21 2006). For these reasons, we chose to use a current duration approach (Keiding, et al., 22 2002, Slama, et al., 2006, Weinberg and Gladen, 1986).

1 METHODS

The study, termed the Observatory of Fecundity in France (Obseff), aims is to describe the fecundity of couples from the general population at the nation-wide level, to describe characteristics of menstrual cycle function using urinary biomarkers and to characterize the influence of environmental factors (in particular atmospheric pollutants) on fecundity; only the first aim is presented here. Our study follows the methodology of our feasibility study on the current duration approach (Slama, et al., 2006), with slight modifications.

8

9 Population sample

Our approach was a two-stage stratified sampling (Slama, et al., 2006). At the first stage, a 10 11 random probability sample of households was selected using a stratified random sample of 12 all landline phones in mainland France (86% of households had a landline phone at the time 13 of the survey). Subjects who only had a mobile phone or no phone were not included in the 14 sample. Households in areas corresponding to urban areas of more than 100,000 inhabitants 15 were over-sampled and corresponded to about 61% of the random list of phone numbers 16 (compared to 46% in the whole French population), a feature taken into account in the 17 analysis by a reweighing approach (see below).

18 Eligibility criteria

19 Eligible households were those which were the main residence of a woman aged 18 to 44 20 years; if several women in this age range lived in the selected household, we randomly drew one, without selecting another one if she turned out not to be eligible. Women aged 18 to 44 21 years (i.e. between their 18th and 45th birthdays) answered an eligibility questionnaire lasting 22 23 about 4 minutes. Eligible women were those living with a male partner or engaged in a 24 regular relationship with a male partner, who were not regularly using any method to avoid 25 pregnancy (nor was their partner) at the time of the study. The interviewer asked questions 26 about all types of methods to avoid pregnancy, enumerating all of them. Couples sporadically 27 using contraception were not considered to be eligible, unless they simultaneously declared 28 that they were trying to become pregnant. Women who did not have sexual intercourse in the

1 month prior to interview, and women who had delivered in the 3 months prior to interview 2 were excluded. Women were asked if they were currently trying to become pregnant, but this was not an eligibility criterion. Indeed, following the logic of a previous study (Karmaus, et al., 3 4 1999), our aim was to capture the whole population theoretically at risk of pregnancy, 5 whatever their fecundity level, without excluding those who are subfertile or may consider 6 themselves sterile after some duration of involuntary infertility. For the same reason, we did 7 not exclude from the main analysis women with irregular menstrual periods or couples in 8 which a male fertility disorder (e.g., low sperm count) had been diagnosed. Sensitivity 9 analyses excluding "non-planners" (defined with respect to the start of the period of 10 unprotected intercourse) and women without menstrual bleeding in the last 12 months are 11 provided (see below).

12 Correction for population sampling and selection bias

13 We used a weighting approach to correct for possible differences between the women who 14 accepted to reply to the eligibility questionnaire and women from the general population, and 15 to correct for the over-sampling of subjects living in urban areas of more than 100,000 16 inhabitants. First, a weight correcting for the over-sampling of large urban areas was defined; 17 a weight correcting for the lower probability of inclusion of women who were not the only 18 woman aged 18 to 44 years in the home was then generated. Finally a weight allowing the 19 recruited population of women aged 18 to 44 years who answered the eligibility criteria to be 20 more similar to the general population of women in this age range (based on the distributions 21 of age, marital life, number of children, age at the end of studies observed in the national 22 census) was created. The estimation of this last weight was based on a generalized raking 23 procedure (Deville, et al., 1993) and was implemented using CALMAR macro implemented 24 on SAS statistical software (INSEE). Finally, these weights were multiplied, and the 25 corresponding final weight was used in all analyses.

26

27 Current duration of unprotected intercourse

The main outcome was the rates of 6-month (respectively 12- and 24-month) involuntary 1 2 infertility, defined as the proportion of couples without a recognized pregnancy within 6 months (respectively, 12 or 24 months) of unprotected intercourse (although the term 3 4 involuntary tends to assume that couples wished to become pregnant -at least at the start of 5 the period of unprotected intercourse- we also used it in analyses including all periods of unprotected intercourse, even those started in the absence of a pregnancy wish). These 6 7 outcomes were derived from the current duration of unprotected intercourse (CDUI), using an 8 approach outlined below. CDUI corresponds to the time elapsed between the start of the 9 period of unprotected intercourse and the interview. Women were also asked to provide the 10 duration elapsed since they last used a method to avoid pregnancy (in weeks, months and 11 years), if any, which was used if the date was missing. The starting date of the period of 12 unprotected intercourse was determined in three different ways. For the majority of women 13 (63%), it corresponded to the date of discontinuation of use of the last contraceptive method, as declared by the woman. Women were asked if, upon discontinuing use of contraception, 14 15 they had waited for 1, 2 or 3 cycles before actively trying to become pregnant; if this was the 16 case then 28, 56 or 84 days, respectively, were subtracted from CDUI. We also checked that 17 no pregnancy declared by the woman had occurred since this date, and corrected the 18 starting date accordingly if this was not the case. If the couple had not used any method to 19 avoid pregnancy since the last pregnancy (32% of women), then the starting date was 20 defined as the end of the last pregnancy, plus three months in case of a live or stillbirth; if the 21 resulting assumed starting date was after the date of interview (e.g., for women who 22 delivered a live newborn in the previous 3 months), then the couple was not considered to be 23 eligible. For couples who had never used any method to avoid pregnancy and in which the 24 woman had never been pregnant (5% of women), the starting date was that of the start of the 25 relationship.

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27 Estimation of the probability of pregnancy

1 The principle of the current duration approach is to infer the underlying distribution (or 2 survival function) corresponding to an unobserved time interval until a given event, from the distribution of the so far elapsed part of this time interval (Keiding, et al., 2002, van Es, et al., 3 4 2000, Weinberg and Gladen, 1986, Yamaguchi, 2003). Here, we are interested in total time 5 interval of unprotected intercourse before a pregnancy (if any), but we only observe couples not using contraception at a certain time period (the time of interview), without following them 6 7 up like in a cohort study. More precisely, the end of the period of unprotected intercourse, in 8 addition to successful conception, may also happen because the couple gives up (resumes 9 contraception because the couple considers it is not anymore a good time to become 10 pregnant, e.g. as a result of a change in their financial situation, or because the couple splits, 11 becomes too old, die). The possible onset of fertility treatment can also be considered as the 12 end of the period of unprotected intercourse, but another interpretation is possible. Indeed, it 13 is possible to argue *either* that fertility treatments change to some extent the probability of conception away from the primary interest, so that the relevant time is unprotected 14 15 intercourse before onset of medical infertility treatment; in other words, we then consider 16 start of fertility treatment as another way of giving up trying. Alternatively, one may consider 17 that the couple is still trying to become pregnant during fertility treatment, if our focus is the 18 actual fecundity in today's society as it is (in which case onset of such treatments can be 19 ignored in analyses). We present both results here since each one has its logic. Note that 20 this way of handling a competing risk is necessary in the current duration approach in order 21 to avoid complex hypotheses on the transition rate to infertility treatments, and that it differs 22 from classical prospective (e.g., Cox) survival modelling, in which fertility treatments can be 23 handled by censoring or as competing risks. Infertility treatments were assessed by asking 24 specifically if each type of medical infertility treatment (drugs, artificial insemination, in vitro 25 fertilization, intra-cytoplasmic sperm injection) had been used by the woman or her partner, 26 and when.

The survival function corresponding to this underlying distribution has been estimated using a parametric approach assuming a generalized gamma distribution (Keiding, et al., 2002,

1 Yamaguchi, 2003). Confidence intervals were built using a bootstrap approach. The 2 estimations were repeated restricting the population to women declaring that they were currently trying to become pregnant or that they had stopped using a contraceptive method 3 4 because they wished to become pregnant (thereafter called "pregnancy planners"). A 5 modification of the approach initially proposed (Keiding, et al., 2002, Slama, et al., 2006) 6 regards very long current durations of unprotected intercourse. Many of these long durations 7 most probably concern couples who are aware of strong subfertility or even sterility, and 8 since our interest is in the shape of the distribution of time to pregnancy for the very first 9 years, we based the estimation on all durations shorter than 36 months and replaced 10 (without excluding them) those reported as being longer than 36 months by the information 11 that they were longer than 36 months.

The resulting estimated survival function can be interpreted as an estimate of the proportion of couples who, after a given number of months, still have unprotected intercourse (that is, they did not give up the period of unprotected intercourse) and did not obtain a clinically detected pregnancy. In the case where couples with infertility treatments are excluded, it is an estimate of an alternative fecundity measure, namely the proportion of couples who, after a given number of months, still have unprotected intercourse and have neither obtained a clinically detected pregnancy nor started fertility treatment.

19

20 Sensitivity analyses

21 We repeated the estimation of the proportion of couples who still have unprotected 22 intercourse in various subgroups or imposing alternative sets of assumptions: 1) assuming 23 that couples who had not resumed contraception since their last pregnancy had a duration of 24 post-partum infertility of 6 months (instead of 3 in the main analysis); 2) excluding women 25 who declared not to have had menstrual cycles in the last 12 months; 3) excluding couples 26 who declared that they had used contraception sporadically (included in the main analysis if 27 they declared to be currently trying to become pregnant) and 4) including couples with 28 sporadic contraception mentioned in 3) above, but assigning them a reduced CDUI, to

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- 1 account for the fact that they were not exposed to pregnancy risk during all months since the
- 2 start of the pregnancy attempt; in practice, we arbitrarily halved their CDUI.

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1 RESULTS

2 Population sample

We drew a random sample of 64,262 households, 19,121 of which were the main residence of an 18-44 year old woman; 15,810 of these women accepted to reply to the eligibility questionnaire (Figure I). Among women aged 18-44 years, 4.8% were pregnant at the time of the eligibility questionnaire; 5.5% reported that they were currently trying to become pregnant and 4.4% declared that they planned to try to become pregnant within the next 12 months (Table I). Half of the women (48%) reported that they had no intention to try to become pregnant in the future.

10

11 Estimation of the frequency of infertility

12 Among the women who answered the eligibility questionnaire, 1089 (6.9%) were eligible; that 13 is, they were not using any birth control method, had a male partner and had been sexually 14 active in the previous month. Out of these, 943 accepted to participate. The estimated overall 15 participation rate was 63% (Figure I). The current duration of unprotected intercourse (CDUI) 16 could be defined for 867 participating women. The characteristics of women with a defined 17 CDUI are shown in Table II. The median duration of CDUI, which has no direct interpretation, was 13.2 months, with 25-75th percentiles equal to 3.7-41.8 months. From CDUI, we 18 19 estimated the underlying "survival" function, corresponding to the time to conception or end 20 of the period of unprotected intercourse for the source population (Figure IIA). The estimated 21 proportion of couples who would still not have conceived and would still be having 22 unprotected intercourse 6 months after the start of the interval of unprotected intercourse 23 was 46% (95% confidence interval, CI, 36-56%); it was 24% (95% CI, 19-30%) and 11% 24 (95% CI, 8-14%) within 12 and 24 months after the start of the period of unprotected 25 intercourse, respectively. Values were very slightly lower after restriction to the 708 couples 26 who declared that they had stopped using birth control methods in order to obtain a 27 pregnancy: rates were 45% (95% CI, 34-55%), 23% (18-28%) and 10% (8-12%) at 6, 12 and 28 24 months, respectively. After restriction to nulliparous couples, the proportions of couples

who had not conceived and were still having unprotected intercourse at 6, 12 and 24 months 1 2 after the start of the period of unprotected intercourse were 47% (95% CI, 26-68%), 26% (15-3 36%) and 11% (7-16%), respectively (Figure IIB). In the analysis based on the alternative fecundity measure (excluding from the initial population couples who had resorted to infertility 4 5 treatments, in order to estimate the proportion of couples who had not conceived nor started infertility treatment and were still having unprotected intercourse), the rates at 6, 12 and 24 6 7 months after the start of the period of unprotected intercourse were 43% (95% CI, 34-53%), 8 20% (95% CI, 16-25%) and 8% (95% CI, 6-10%), respectively (Figure IIA).

9

10 Sensitivity analyses

Assuming a duration of 6 months of post-partum infertility instead of 3 led to the exclusion of 11 12 9 couples but had no impact on our estimates. Twelve women declared not to have had 13 menstrual cycles over the last 12 months before inclusion, and, again, excluding them had 14 no impact either on our results. Excluding couples who declared that they were trying to 15 become pregnant and that they were using contraception sporadically (n=113) yielded very 16 slightly increased rates of involuntary infertility (47%, 26% and 12% at 6, 12 and 24 months, 17 respectively). Including these couples with a halved current duration decreased the rates of 18 involuntary infertility to 37% (95% CI, 28-47%), 20% (15-25%) and 9% (7-11%) at 6, 12 and 19 24 months, respectively.

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1 DISCUSSION

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3 Comparison with former studies

To our knowledge, our study is one of the first to estimate the frequency of involuntary
infertility in a nation wide representative population sample and not relying on a retrospective
study design.

7 Several studies using other designs have provided descriptions of the couple fecundity (e.g. 8 Bonde, et al., 1998, Jensen, et al., 2001, Joffe, 2000, Karmaus, et al., 1999, Muller, et al., 9 2006, Scheike, et al., 2008, Slama, et al., 2008). Most of these studies relied on retrospectively collected information in women or men asked to describe earlier pregnancy 10 11 attempts, using either pregnancy-based (Jensen, et al., 2001, Joffe, et al., 2005, Scheike, et 12 al., 2008) or historically prospective (Karmaus, et al., 1999, Slama, et al., 2008) designs (the 13 former design is restricted to pregnancy attempts ending in a pregnancy, while the latter also 14 includes those ending with no pregnancy). In a historically prospective study, 932 couples 15 from Denmark, Germany, Poland, Italy and Spain described their periods of unprotected 16 intercourse resulting or not in a pregnancy, provided they had started less than 5 years prior 17 to the interview (Karmaus, et al., 1999). When the analysis was restricted to the first period of 18 unprotected intercourse, 23% of women had not conceived within the first 12 months of 19 attempt (varying from 33% in Poland down to 15% in South Italy); when the most recent time 20 of unprotected intercourse was considered (including current attempts started not more than 21 5 years before interview), the proportion of women who had not conceived within 12 months 22 of unprotected intercourse was 29% (Karmaus, et al., 1999). In a historically prospective 23 study in two French rural areas conducted in 2000 that considered pregnancy attempts 24 started between 1985 and 2000, Slama et al. (2008) reported a rate of 12-month involuntary 25 infertility of 16%. In a prospective study among Danish first pregnancy planners followed-up 26 for six months, 256 couples out of 430 conceived (Bonde, et al., 1998), which, after taking 27 into account censoring, corresponds to a 6-month involuntary infertility rate of 36%. In 28 another cohort of 221 women from America who volunteered in 1983-1985 as they were

1 planning to stop using birth-control methods in order to become pregnant, the 6-month 2 cumulative pregnancy rate was 78% (Wilcox, et al., 1988). Retrospective studies relying only on periods of unprotected intercourse (or pregnancy attempts) leading to a live birth (the so-3 called "pregnancy-based" design) yield lower rates of 12-month involuntary infertility: in 4 5 England, Joffe (2000) reported that 10% of couples whose first pregnancy attempt leading to a live birth started in 1991-93 had needed more than 12 months to conceive. In another 6 7 pregnancy-based study in 4 European cities among couples who delivered in 1996-98, the 8 corresponding rate for couples recruited in Paris was 9.9% (Jensen, et al., 2001); it ranged 9 from 5 to 11% among fertile couples recruited using a similar design in four French cities in 10 2002-2003 (Muller, et al., 2006). In the Enguête Nationale Périnatale, a national sample 11 constituted of all women who delivered in all French maternity units in a given week in 2003 12 (Blondel, et al., 2006), time to pregnancy could be defined for 10,262 out of 14,482 live 13 births, and the rates of 6, 12 and 24-month infertility were 32% (95%, 31-33%), 18% (95% 14 CI, 17-19%) and 6% (95% CI, 5-6%), respectively (Slama, et al., (in press)). Although our 15 confidence intervals were relatively broad (19% to 30% for the estimated rate of 12-month 16 infertility), our estimate of the rate of 12-month involuntary infertility is somewhat higher than 17 the pregnancy-based values reported in the *Enquête Nationale Périnatale*. This is coherent 18 with what can expected from the facts that the latter study did not take couples remaining 19 infertile or giving up the pregnancy attempt into account and that couples fecund enough to 20 have many children are over-represented in such a pregnancy-based study.

21

22 Study population

Our study is based on a random sample from the general population. The estimated participation rate of eligible couples was 63%, which offers potential for bias. Our population was weighted to limit such bias. This weighing approach, consisting in making our population more similar to the general population in terms of age and age at the end of studies, is expected to correct efficiently for selection bias to the extent that factors associated with participation and fecundity level are statistically linked with these sociodemographic

1 variables. Unlike some previous studies on time to pregnancy using a pregnancy-based 2 design, inclusion was not conditioned on couples having eventually obtained a pregnancy. 3 We tried to identify all couples possibly at risk of pregnancy, without excluding sterile or 4 subfertile couples. We chose not to condition inclusion on couples being currently trying to 5 become pregnant, because subfertile couples who had started a pregnancy attempt in order to become pregnant may consider themselves subfertile or sterile after several months on 6 7 involuntary infertility, and may and consequently declare that they are not trying anymore to 8 become pregnant although they still have unprotected intercourse. We reported additional 9 results limited to couples who declared to have started the period of unprotected intercourse 10 in order to become pregnant so as to describe the fecundity of pregnancy planners, which 11 turned out to be very similar to that estimated without that exclusion.

12 Including parous women may induce bias or limit representativeness of the population 13 sample because previous reproductive history may impact persistency in trying to become 14 pregnant; this may also limit comparability with future studies because of possible temporal 15 changes in desired family size; for this reason, we also reported results restricted to 16 nulliparous women. These women had a very slightly higher rate of 12-month involuntary 17 infertility, compared to the overall population also including parous women.

18

19 Assumptions made by our design

20 Fecundity studies have several potential limitations. These relate in particular to the fact that 21 couples with unplanned or mistimed pregnancies usually have no defined duration of 22 unprotected intercourse; to variability in the delay before pregnancy detection, in access to 23 and use of contraception or in desired family size. These limitations however apply to most 24 other types of fecundity studies, and have been discussed at length elsewhere (see e.g., 25 Joffe, et al., 2005, Key, et al., 2009, Slama, et al., 2006, Weinberg, et al., 1994); we will here 26 focus on sources of bias and assumptions specific to our study design. Our cross-sectional 27 design implies that couples who have had unprotected intercourse for a long time have a 28 higher probability of inclusion than couples with a shorter waiting time; however this length-

1 biased sampling is taken into account in the statistical analysis, which provides an unbiased 2 estimate under certain assumptions (Keiding, et al., 2002, Weinberg and Gladen, 1986). A limitation of the design is that it does not allow one to distinguish couples who conceive a 3 4 pregnancy from those who abandon the pregnancy attempt, either because they split or 5 resume contraception before they conceive; that is, a period of involuntary infertility as defined in our study corresponds to the time until pregnancy occurrence or end of 6 7 unprotected intercourse with no pregnancy. Data on the frequency of couples stopping a 8 period of unprotected intercourse before pregnancy occurrence are limited; persistency in 9 trying to become pregnant may vary according to mother's age at starting date, country 10 (Basso, et al., 2000), and probably other factors such as parity. The impact of such attempts 11 on the estimates from the current duration approach is expected to correspond to an 12 overestimation of the fecundity level, whose amplitude will depend on the frequency of these 13 attempts terminated because the couple split or resumed contraception. As forcefully pointed 14 out by Basso et al. (Basso, et al., 2000), these pregnancy attempts not ending in a 15 pregnancy also constitute a source of bias in the pregnancy-based design, which does not 16 allow including them either and in which the collected time to pregnancy is conditioned on 17 couples not giving up the period of unprotected intercourse; in that design also, their 18 exclusion leads to an overestimation of the fecundity level. In a prevalent cohort based on 19 couples recruited after less than 12 months of pregnancy attempt, the proportion of couples 20 who reported to discontinue trying to become pregnant during the following year was 5% 21 (Wise, et al., 2010). Compared to what had originally been suggested (Keiding, et al., 2002, 22 Slama, et al., 2006), we report here results including couples who had initiated an infertility 23 treatment since the start of the period of unprotected intercourse (as well as analyses 24 excluding these couples, corresponding to the approach used in our previous publication). If 25 these couples with infertility treatment are excluded, then the studied event is either 26 pregnancy, or end of the period of unprotected intercourse or start of a fertility treatment; if 27 these couples with infertility treatment are included, then the event is simply either pregnancy 28 or end of the period of unprotected intercourse; both estimates are worth reporting.

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Coherently, including these couples yielded a somewhat higher frequency of 12-month
 involuntary infertility (24%, compared to 20% if couples with a treatment are excluded),
 corresponding to a longer time to event.

4

5 Relevance of the current duration approach to describe fecundity

6 Describing and possibly monitoring couple fecundity takes on importance in the context of 7 increasing use of ART and of the possible deterioration of male reproductive health. Indeed, 8 several studies have described temporal decreases in sperm concentration and motility in 9 specific areas of industrialised countries over the last decades (Auger, et al., 1995, Carlsen, et al., 1992, Nelson and Bunge, 1974, Swan, et al., 2000). These decreases cannot be 10 11 considered as certain, in particular because studies reporting temporal trends in semen 12 quality are often based on self-selected subjects, not allowing quantification of participation 13 rates and description of possible selection bias (Cohn, et al., 2002, Eustache, et al., 2004, 14 Hauser, et al., 2005, Muller, et al., 2004). If true, such a decline might have had an impact on 15 fecundability, the cycle-specific probability of pregnancy among non-contracepting couples, 16 as indicated by a simulation study (Slama, et al., 2004). This decline in fecundability may not 17 imply strong changes in the average number of children per couple, but could entail 18 increases in the proportion of couples subject to 1- to 5-year involuntary infertility (Leridon 19 and Slama, 2008).

20 Very few studies directly tried to assess temporal trends in fecundability or involuntary 21 infertility of populations (Jensen, et al., 2005, Joffe, 2000, 2008). These studies generally do 22 not cover areas where temporal declines in semen quality have been reported. Results do 23 not show decreases in couple fecundity. Their pregnancy-based design excludes couples 24 who remain involuntarily infertile, therefore limiting the statistical power of analyses trying to 25 highlight changes in fecundity, and possibly biasing the comparison towards absence of 26 change in fecundity (Slama, et al., 2004). Additionally, it has been argued that temporal 27 trends in the proportion of unwanted pregnancies and abortion rates may make it difficult to 28 describe time trends in fecundity (Sallmen, et al., 2005). Several approaches can be used to

correct for these potential biases (Joffe, et al., 2006, Joffe, et al., 2005, Key, et al., 2009) but
the efficiency of these approaches may in practice be limited in the context of a retrospective
survey design (Sallmen, et al., 2006). For these reasons, alternatives to a retrospective
design are worth investigating. These include the (incident) cohort design (Bonde, et al.,
1998), the prevalent cohort design (Keiding, 1992, Wise, et al., 2010) and the current
duration approach. Other designs have been suggested (Olsen and Andersen, 1999) but not
applied to our knowledge.

8 The eligibility rate in the current duration approach is relatively low; we had to survey 15,810 9 couples in order to recruit 867 couples (5.5%) with a defined current duration of unprotected 10 intercourse. This eligibility rate is higher than that of the main other prospective design, the 11 (classical) cohort design, in which couples are recruited before the start of the period of 12 unprotected intercourse. Indeed, less than 2% of women aged 18-44 years contacted 13 indicated that they planned to start a pregnancy attempt within the next 6 months, out of 14 which probably not all will do so within this duration. Conversely, some couples may start a 15 pregnancy attempt in the same time period without having planned it long in advance, and 16 these would be hard to identify and include in an incident cohort. These study designs are 17 actually not incompatible one with the other; indeed, couples who plan to start a pregnancy 18 attempt soon (eligible in an incident or prevalent cohort on fecundity) can also be identified 19 from the eligibility questionnaire of a current-duration study such as ours, and these can be 20 followed-up, together with couples eligible for the current duration design at the time of 21 interview. If repeated at regular time intervals, such a design could be used to prospectively 22 monitor time trends in fecundity. This could be seen as a parallel to the studies monitoring 23 semen quality in young men in Scandinavian countries (Jorgensen, et al., 2006); as 24 suggested by Olsen and Rachootin (2003), a system monitoring fecundity should monitor 25 semen quality in parallel to a measure of the couple fecundity such as involuntary infertility.

26

In conclusion, our study provides an estimate of the frequency of involuntary infertility in the general population showing that about one in four to five couples have not conceived a

18

- 1 detected pregnancy and are still having unprotected intercourse 12 month after having
- 2 stopped using a contraceptive method, and that about one couple out of ten may still be
- 3 unsuccessful in conceiving after 2 years of unprotected intercourse.

1 AUTHORS' ROLES

RS and JB initiated the study. RS, BD, NK, AB and JB designed the study and
questionnaires; AB, BD, JB and RS supervised data collection. AB, BD and RS cleaned and
prepared data. The statistical analyses were planned and supervised by RS and NK and
carried out by OH, DS and LGA. RS drafted the first version of the manuscript. BD, LGA,
MJCE, LR, JCT, NK and JB critically reviewed the manuscript.

7

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12

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3

TABLES 1

2 Table I: Characteristics of the women who replied to the eligibility questionnaire.

| | Whole pop | ulation (n=15,810) | | Current duration | on group | | |
|---|-----------|-------------------------|-------------|-------------------------|---------------|----------------|--|
| Characteristic | n | Weighted % ^a | Yes (n=867) | | No (n=14,943) | | |
| | | | n | Weighted % ^a | n | % ^a | |
| Age at interview (years) | | | - | | | | |
| 18-24 | 2,756 | 22.8 | 53 | 9.4 | 2,703 | 23.5 | |
| 25-29 | 2,601 | 17.1 | 193 | 25.0 | 2,408 | 16.7 | |
| 30-34 | 3,048 | 19.2 | 254 | 30.4 | 2,794 | 18.6 | |
| 35-39 | 3,836 | 20.5 | 222 | 21.0 | 3,614 | 20.4 | |
| 40-44 | 3,569 | 20.5 | 145 | 14.2 | 3,424 | 20.8 | |
| Currently has a male partner | , | | | | , | | |
| Yes | 12,178 | 66.9 | 867 | 100 | 11,311 | 65.1 | |
| No | 3,632 | 33.1 | 0 | 0 | 3,632 | 34.9 | |
| Number of children | , | | | | , | | |
| 0 | 5,420 | 38.7 | 360 | 42.3 | 5,060 | 38.5 | |
| 1 | 3,016 | 20.4 | 281 | 32.9 | 2,735 | 19.7 | |
| 2 | 4,693 | 24.8 | 159 | 15.2 | 4,534 | 25.3 | |
| 3 and more | 2,681 | 16.2 | 67 | 9.6 | 2,614 | 16.5 | |
| Current contraception | , | | | | , - | | |
| Yes, systematically used | 13,105 | 82.0 | 0 | 0 | 13,105 | 86.5 | |
| Yes, sporadically used | 136 | 0.9 | 113 | 13.1 | 23 | 0.2 | |
| No method to avoid pregnancy ^b | 2,150 | 14.3 | 754 | 86.9 | 1,396 | 10.4 | |
| Surgical sterilisation c | 406 | 2.8 | 0 | 0 | 406 | 2.9 | |
| Do not know | 13 | 0.1 | 0 | 0 | 13 | 0.1 | |
| Defined CDUI | | | | | | | |
| Yes | 867 | 5.2 | 867 | 100 | 0 | 0 | |
| No | 14,943 | 94.9 | 0 | 0 | 14,943 | 100 | |
| Planning to try to become pregnant ^a | | | | | | | |
| Yes, currently trying | 859 | 5.5 | 655 | 75.6 | 204 | 1.5 | |
| Yes, will start within 1 or 2 months | 27 | 0.1 | 3 | 0.3 | 24 | 0.1 | |
| Yes, in about 3 months | 60 | 0.4 | 2 | 0.1 | 58 | 0.4 | |
| Yes, within 4-6 months | 181 | 0.9 | 1 | 0.03 | 180 | 1.0 | |
| Yes, within 7-12 months | 511 | 3.0 | 12 | 1.7 | 499 | 3.1 | |
| Yes, in more than a year | 1,789 | 12.0 | 17 | 2.2 | 1,772 | 12.5 | |
| Yes, but not planned when | 3,460 | 24.9 | 32 | 4.0 | 3,428 | 26.1 | |
| No | 7,700 | 48.1 | 139 | 15.6 | 7,561 | 49.9 | |
| Does not know | 57 | 0.4 | 6 | 0.5 | 51 | 0.4 | |
| Currently pregnant | 751 | 4.8 | Õ | 0 | 751 | 5.0 | |

3 4 5 6 7 8 9

CDUI: Current duration of unprotected intercourse. ^a Percentages were corrected for possible selection bias and over-representation of urban compared to rural

areas. ^b Including pregnant women.

^c Surgical sterilisation corresponded to tubal ligation, vasectomy or hysterectomy. ^d Women with surgical sterilisation have not been asked about pregnancy planning.

10

11

- Table II: Characteristics of the 867 women with defined current duration of unprotected 1
- 2 intercourse.

| Characteristic | n | Weighted % ^a |
|--|-----|-------------------------|
| Age at interview | | |
| 18-24 years | 53 | 9.4 |
| 25-29 years | 193 | 24.9 |
| 30-34 years | 254 | 30.4 |
| 35-39 years | 222 | 21.0 |
| 40-44 years | 145 | 14.2 |
| | 145 | 14.2 |
| Age at start of period of unprotected intercourse | | 10.0 |
| 18-24 years | 101 | 16.2 |
| 25-29 years | 263 | 32.7 |
| 30-34 years | 293 | 30.1 |
| 35-39 years | 175 | 17.6 |
| 40-44 years | 35 | 3.3 |
| Number of children | | |
| 0 | 360 | 42.3 |
| 1 | 281 | 32.9 |
| 2 | 159 | 15.2 |
| 3 and more | 67 | 9.6 |
| Frequency of sexual intercourse | - | |
| 1-3 per month | 165 | 18.0 |
| 1-2 per week | 418 | 47.5 |
| >2 per week | 269 | 34.6 |
| Duration of menstrual cycle | 200 | 04.0 |
| <27 days | 161 | 24.2 |
| 27-29 days | 356 | 42.4 |
| | | |
| >29 days | 273 | 33.5 |
| Medical treatment for infertility since start of the PUI | 700 | 00.4 |
| No | 708 | 83.4 |
| Yes | 159 | 16.6 |
| Started the PUI to obtain a pregnancy | | |
| Yes | 708 | 80.8 |
| No | 159 | 19.3 |
| Currently trying to become pregnant | | |
| Yes | 655 | 75.6 |
| No | 212 | 24.4 |
| Smoking at the start of the PUI | | |
| No | 580 | 62.0 |
| Yes | 279 | 38.0 |
| Body mass index (kg/m ²) | | |
| <18.5 | 78 | 10.9 |
| 18.5-19.9 | 132 | 12.9 |
| 20-22.4 | 248 | 25.4 |
| 22.5-24.9 | 163 | 20.0 |
| 25-29.9 | 151 | 18.8 |
| | | |
| \geq 30 | 82 | 12.0 |
| Current duration of unprotected intercourse ^b | 100 | 00.0 |
| <3 months | 182 | 20.2 |
| 3-5.9 months | 113 | 12.9 |
| 6-11.9 months | 118 | 13.2 |
| 12-23.9 months | 130 | 16.2 |
| 24-35.9 months | 75 | 8.9 |
| ≥ 36 months | 249 | 28.6 |

3 PUI: Period of Unprotected Intercourse

^a Percentages were corrected for possible selection bias and over-representation of urban compared to rural 4

5 areas by a reweighting approach. This explains why the variables relative to medical treatment and to whether 6 couples started the PUI to obtain a pregnancy have different weighted percentages although the number of

7

subjects in each "Yes" and "No" categories, and hence the unweighted percentages, are the same. ^b The distribution of the current duration of unprotected intercourse declared by couples has no direct 8

9 interpretation due to the length-biased sampling and should therefore not be interpreted as an estimate of the

10 frequency of involuntary infertility (given in Figure II).

1 FIGURES

2

3 Figure I: Flow chart of study population.

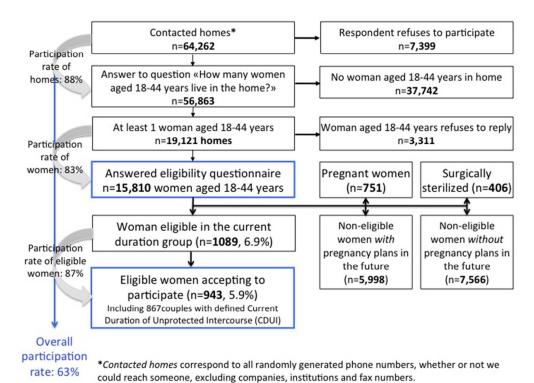
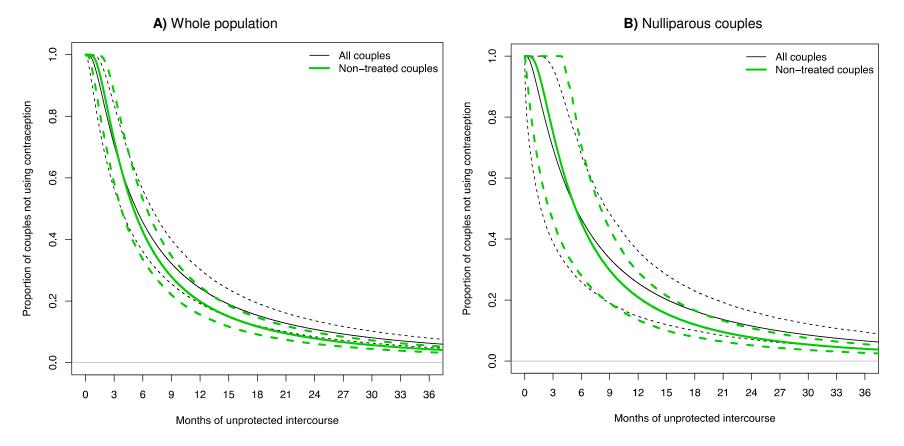


Figure II: Estimation of the proportion of sexually active couples still without detected pregnancy and still not using contraception, as a function of the number of months elapsed since the start of the period of unprotected intercourse. The usual clinical definitions of infertility would correspond to the values at 12 and 24 months. **A)** Whole eligible population (n=867, and n=708 after restriction to non treated couples); **B)** nulliparous couples only (n=360 and n=277 after restriction to non treated couples).



The survival function corresponding to time until pregnancy or end of the period of unprotected intercourse without pregnancy is indicated by the black solid line (estimated from the whole population at risk of pregnancy at inclusion). The green curve is the same estimate, but recomputed after exclusion of couples who had begun treatment for infertility. Dotted curves indicate 95% confidence intervals.