Qualitative investigation of the factors that generate ambivalent feelings in women who give birth after receiving negative results from non-invasive prenatal testing

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Abstract
Background: Women who receive negative results from non-invasive prenatal genetic testing (NIPT) may find that they later have mixed or ambivalent feelings, for example, feelings of accepting NIPT and regretting undergoing the test. This study aimed to investigate the factors generating ambivalent feelings among women who gave birth after having received negative results from NIPT.
Methods: A questionnaire was sent to women who received a negative NIPT result, and a contents analysis was conducted focusing on ambivalent expressions for those 1562 women who responded the questionnaire. The qualitative data gathered from the questionnaire were analyzed using the N-Vivo software package.
Results: Environmental factors, genetic counseling-related factors, and increased anticipatory anxiety, affected the feeling of ambivalence among pregnant women. Furthermore, pregnant women desired more information regarding the detailed prognosis for individuals with Down syndrome and living with them and/or termination, assuming the possibility that they were positive.
Conclusions: Three major interrelated factors affected the feeling of ambivalence in women. Highlighting and discussing such factors during genetic counseling may resolve some of these ambivalences, thereby enhancing the quality of decisions made by pregnant women.

Background
A prenatal test for fetal abnormalities may cause maternal anxiety [1, 2]. Pregnant women who made an uninformed choice for non-invasive prenatal genetic testing (NIPT) felt deeper decisional regret associated with prolonged anxiety [3, 4].
Women typically have two conflicting thoughts at the early stage of pregnancy: the desire to know about the neonate via ultrasound examination and the reluctance to receive negative news. Many women reportedly take the NIPT to relieve their ambivalence regarding negative results [5].
Furthermore, certain women feel like their pregnancy is provisional or temporary until they learn about their NIPT results, i.e., although they may feel physical changes, they may not acknowledge their pregnancy until they see their results [6]. Ambivalence has been identified among the attitudes of the general public regarding prenatal testing; however, the origins of the ambivalence remain
unclear [7, 8]. Lewis et al. reported on the ambivalence of the women taking the NIPT had ambivalent feelings, e.g., desire for the information about the baby by the test and additional anxiety whilst waiting, concerns around “over-testing” and “pressure by the health professionals”. [9]. Ambivalence can be defined as the “simultaneous existence of positive and negative evaluations of an attitude” [10].

In our previous study, we conducted an “awareness survey of NIPT in Japan”, and reported that women with negative results had a higher NIPT rating than women with positive results, but with respect to the ethical aspects of NIPT, women with negative results responded unclearly [11]. In the presence of negative NIPT results, diseases other than the three chromosomal aneuploidies may be identified during later stages of life [12] then they are expected to have mixed feelings about the NIPT.

Based on these findings, we thought that even women with negative test results could have ambivalent feelings later. This study aimed to investigate factors affecting ambivalent feelings in pregnant women after undergoing NIPT.

Methods

Design

The present study is based on a qualitative research methodology, involving content analysis using the free-form description of the first year after questionnaire answered by women who received negative results for NIPT.

Research procedure

All pregnant women who took the NIPT were asked to participate in a questionnaire-based survey conducted by the Japan NIPT Consortium. The survey was in two parts: (i) pre-and immediately after the NIPT test, from April 2013 to March 2014 and (ii) a year- after the NIPT test, from April 2014 to March 2015. We have already reported on part (i) [11], and the present study is for part (ii). A year- after research was conducted by a mail-in survey to determine whether the assessment of the NIPT and genetic counseling was changed after the test and one year after. The contents of the
questionnaire consisted of a one-year-after evaluation and free description for NIPT and genetic counseling, and the question of the free description part was an open-ended question, such as, “Please show your opinion or feedback of the NIPT.” In the present study, we conducted content analysis for 1562 respondents (20.6%) who provided the free-form description to the open-ended question (Fig. 1).

Participants
Pregnant women who were identified as being at an increased risk of chromosomal disorders due to advanced maternal age, the results of ultrasound marker or maternal serum marker tests, or a family history of chromosomal abnormalities or those who had a high risk of being a translocation carrier, were eligible for NIPT. In most cases, the indication for NIPT was advanced maternal age (96.5%). In addition, 14.4% of the women underwent NIPT based on a recommendation from their family. Because the responses in the first year after the questionnaire were anonymous, and it was impossible to obtain the background data, the personal background of the respondents was unknown; however, the available necessary background information of the respondents who underwent NIPT was shown in the previous study [11] of 7740 women, whereas the present study was limited to those who received negative NIPT results among the 7740 women, and therefore, the background data were not same.

Data Analysis
The questionnaires were subsequently mailed to a third-party NIPT data center for data entry. Qualitative methods are required in situations warranting detailed analysis and an understanding of the process to determine the nature of the issue being investigated. We undertook a combined content analysis (quantitative and qualitative) via text mining approaches, using N-Vivo Qualitative software for the free-form description part of the questionnaire focusing on ambivalent expressions. The advantage of computer-assisted content analysis of textual data is the coding reliability that helps to generate comparable results [13]. The text in the free-form description part was entered into
the computer and used as input for the N-vivo software. N-vivo was used for line-by-line coding of the texts, to identify emerging themes. Furthermore, all the text was read, and the coding process was discussed with two researchers. The sub-categories extracted by N-vivo were assessed by two reviewers. The code of those lower categories determined after the evaluation was extracted inductively into the above category by two reviewers. The coding process is shown below.

Step 1 Line-by-line coding with N-vivo, using an inductive approach.

Step 2 Thematic grouping. The transversal analysis showed new themes by two researchers.

Step 3 Creating higher-level categories and evaluating relationships

Step 4 Reviewing these analyses. Thematic stabilization. All authors.

Ethics

To conduct this study, the participating centers obtained approval from their respective ethics committees.

Results

We identified three primary categories generating feelings of ambivalence among women: (1) factors related to genetic counseling; (2) environmental factors, and (3) increased anticipatory anxiety.

Furthermore, the former two factors caused the anticipatory anxiety (Table.1, Fig.2).

Factors related to genetic counseling

This category Factors related to genetic counseling comprised two subcategories: “Lack of information” and “Lack of psychological care adapted to individual needs.” “Lack of information” further comprised two mid-level categories: “Inadequate support by medical staff up to taking NIPT” and “Lack of information in case of positive result.” “Lack of psychological care adapted to individual needs comprised “Lack of psychological care at genetic counseling,” and “Lack of genetic counseling at the appropriate moment.”

Regarding “Lack of information,” “support by medical staff up to taking NIPT” referred to the lack of support from the family physician during NIPT. Many family physicians were not only not helpful in referring their patient for genetic counseling, but they also themselves were unaware of it. Owing to
the limited number of NIPT facilities for pregnant women in Japan, some respondents also reported receiving a reproachful response from their family physician upon requesting information regarding NIPT or referrals.

Furthermore, certain women were criticized for taking the NIPT during genetic counseling, and many women wished that genetic counseling would be performed assuming that NIPT could give positive results regarding “Lack of information in case of positive result.” They wished to receive detailed information regarding the methods of terminating their pregnancy as one of the alternatives in case of chromosomal abnormalities in the fetus.

_I don't think it hurts to take some time to explain about the options for getting an artificial termination of pregnancy in the case of a positive result. There is this idea that an abortion bad, but if the chance that you can love your child unconditionally is 0%, then I believe that it is an important option, even though it's not easy to discuss..._

_(if the pregnancy was terminated)_

However, some women who considered continuing their pregnancy despite receiving positive results upon NIPT were seeking information regarding the exact quality of life for individuals living with such illnesses and the social resources available for them.

_What I really want to see added is what to do if the results do come out positive. I want to hear during the counseling session the exact types of social support that I could get if my child is born with Down syndrome or some other illness to help me live with and raise the child with the illness, for example at school, daycare, at home, names of organizations or municipal centers where I can get support._

_(if the pregnancy was continued)_

These data indicated that respondents were seeking information during genetic counseling that would support the decision to continue or terminate their pregnancy. These women desired information regarding the daily lives of children with disabilities and other related materials to guide their decision of whether they could raise a child with disabilities; furthermore, they desired follow-up support for women reporting with positive results in the NIPT.

_Environmental factors_
**Environmental factors** included the comfort levels of women for raising a child with disabilities; this aspect comprised three subcategories: “Lack of awareness and education about diversity,” “Insecurities on raising the child,” and “Lack of social support systems for people with disabilities.” “Insecurities on raising the child” comprised “Insecurities for the future,” “Vague insecurity,” “Insecurities for the child-rearing environment,” and “Familiarity with challenges of living with a disabled person.”

The reason why many people have the image of prenatal testing as an "immoral thing" in Japan may come from views about ethics on this subject in Japan, or prejudice and closed-mindedness against people with Down syndrome and other chromosomal abnormalities. I just sometimes wish that those who are debating and giving their input on ethics would also take part in discussing how to change Japan into a more livable society for children with Down syndrome and their families.

*(Lack of awareness and education about diversity)*

I want to give birth even if child with Down syndrome, if possible ... The word “life sorting” is not good, but we feel that social systems are forced to do so. I think we must create a welfare, education and social system that everyone can give birth and raise without worrying, even if a child has a disability.

*(Lack of social support systems for people with disabilities)*

Regarding “Insecurities on raising the child”, older parents had a sense of responsibility to raise the child; they were concerned regarding the life of the child after their death owing to their age at pregnancy, and they were concerned regarding burdening the siblings of the child and, thus, had “Insecurities for the future.”

Furthermore, certain respondents also exhibited “Vague insecurity” because raising a disabled child is unfathomable and difficult to imagine, owing to the lack of experience. Others, however, already had “Familiarity with challenges of living with a disabled person,” because they knew somebody or had family members with disabilities, thereby fostering the understanding of the merits and demerits and having discerned that it would be challenging. They also faced “Insecurities for the child-rearing environment” owing to their concern regarding an inadequate financial support or the social
environment for raising a child with disabilities.

As an actual mother of a child with chromosomal abnormalities, I'm glad that I had my baby, but raising my child is full of challenges and would be absolutely impossible without the cooperation of people around me, so it's not something that can be glossed over. I got tested for my second pregnancy, but I don't think I would've had the child if the results were positive. (Familiarity with challenges of living with a person with disabilities).

**Increased anticipatory anxiety**

**Increased anticipatory anxiety** included the two subcategories “Time-related anxiety” and “Anxiety about test accuracy.” Among the pregnant women who opted for NIPT, there were some who were so anxious during the two weeks until they got the results that they could not sleep until finding out that their results were not positive. Also, knowing that test accuracy of NIPT was not 100%, some people were anxious that their result might be a false-negative, and they could not eliminate this anxiety until the birth of the child.

**Components of Ambivalence**

**Ambivalent feeling** comprised five subcategories: “Options in the case of a positive result,” “Guilt towards the child,” “Criticisms on NIPT from others,” “Denial of disabled people”, and “How to tell the child.”

Some respondents “Stated their decision” with regard to the course of action in case of a positive NIPT result, whereas others expressed “Difficulty stating their decision” regarding continuing or terminating their pregnancy. In both cases, respondents seemed agonized over their decision in the case of positive results (Fig. 3).

I had decided that if the results came out positive, I would not give birth to the baby. However, even though I now know that the results were negative, I haven't told my own parents that I took the test. At the same time that I feel negative about the idea that we were picking and choosing who to keep if the results were positive, I also blame myself for my lack of confidence in raising the child, even though it is because I have health problems myself. (Stated their decision)
Some women seemed to have a sense of “Guilt towards the child” after given birth and parenting the child, knowing that they were trying to decide the life of the child.

I don't think I would have taken the test if I were younger. Some people criticize that people take the test too casually because it places very little stress on the mother or child. But I believe that anybody who chose to take the test did so after putting lots of thought into it. Looking into my child's face, sometimes I remember about when I took the test, and I can't help feeling sorry towards my child. My baby's life is very valuable, and I want to give all the love it deserves.

(Guilt towards the child)

Furthermore, some women could not tell anybody that they took the NIPT owing to social pressure, “Criticisms from others on NIPT”, and others who felt guilty that having “Denial of disabled people” upon having children with disorders, which were screened during NIPT, despite obtaining negative NIPT results.

When I watch documentaries on TV, etc., and hear that the majority of opinions seem to support that “Whether or not a life should live should not be decided by parents, but should be treated as an independent life” “Even if my child has Down syndrome, I am so happy that they came to this world”. People who, for whatever reason decided not to have the baby, will be burdened with guilt for the rest of their lives. I did test negative, but whenever I look into my child's face and see a person with Down syndrome, I criticize myself because “I decided on her life.”

(Denial of disabled people)

Discussion

The results of this study also showed that factors related to genetic counseling and environmental factors greatly influence women’s decision-making regarding taking the NIPT. Because of the impact of factors related to adequate information, i.e., Factors related to genetic counseling before undergoing NIPT and Environmental factors, which describe an inadequate social environment wherein the ability of independent decision making among women is perturbed, Anticipatory anxiety is increased. The decision-making among pregnant women who receive a prenatal diagnosis is affected not only by individuals close to them, e.g., family members or friends, but also by the
opinions of medical staff, the social environment, and uncertainty regarding fetal health conditions [13-16].

Furthermore, the guilt that women experience when deciding to give birth or abort the fetus, or the guilt they experience regarding decision-making related to disabled individuals and contrary sentiments associated with ideas that promote prenatal testing and that having to face this moral opposition induces a state of Ambivalence among some women even one year after taking the NIPT, whenever the unresolved feelings associated with the thought “What if the result had been positive” resurfaced in their minds.

There are problems related to genetic counseling about assuming positive NIPT results. Genetic counselors themselves are concerned that talking to parents about Down syndrome during prenatal testing is likely to elicit fear among them. Overly optimistic or negative information from medical staff, who largely influence the parents’ ultimate decision-making upon receiving positive NIPT results can also introduce a bias [17, 18], similar to the genetic counselors themselves who lack the necessary skills and knowledge [3]. Perhaps, this is focused on avoiding any negative effects of providing information while assuming a positive NIPT result on the mental status of pregnant women, who seek emotional relief from undergoing the test. However, many women also seek information regarding the precise living conditions of children with the screened genetic disorders and regarding the types of social resources available to them.

Statistics have revealed that in England, a few more women who received positive NIPT results chose to have an abortion compared to those who tested positive upon amniocentesis [3]. Concurrently, in Japan, the possibility of continuing the pregnancy existed among many women, despite obtaining positive NIPT results. The present results also show that it is important to offer genetic counseling under the assumption of a potentially positive NIPT result.

Among numerous pregnant women, the motivation to undergo prenatal testing stems from their need to "feel relieved [5]." However, among the women who inquire about prenatal diagnosis, many request information regarding not only about the actual test, but also regarding the many anxieties they may have about their children during pregnancy. Pregnant women attended NIPT genetic
counseling sessions at a time when they could openly speak and be listened to; hence, it is very important to provide psychological support to pregnant women in addition to information regarding testing methods or disorders that can be screened for during the test [19].

Fears regarding the potential discrimination faced by the child, prejudice, the lack of understanding among family members or other people, and the lack of societal support for raising the child are some of the reasons that motivate women to have a "healthy" child. There is a cultural feature in Japan that is sensitive to the surroundings and is easily affected. Furthermore, the present results suggest that the society is not very open and livable for individuals with disabilities. Simultaneously, numerous individuals strongly felt the need to adapt to the social support systems for people giving birth to children with disabilities. These results suggest a positive and supportive perspective towards disabled individuals, fostering the hope that in the future, diversity will be widely accepted and that society will be welcoming to all children, irrespective of being born with disabilities. Many of the present study subjects were mothers with late-life pregnancies. We believe the potential impacts on family and siblings, societal prejudices on disabilities, and attachment towards the unborn child were weighed after careful consideration and understanding of societal prejudices and the increased chance of having a child with an abnormal chromosomal number owing to their age before choosing to undergo NIPT.

Similar to other types of prenatal testing, we believe that NIPT may also increase the pregnant women's worries and anxieties about the unborn child. The rate of infertility treatment was high among the women who underwent NIPT (42.2%). Furthermore, women undergoing infertility treatment have strong fears regarding chromosomal abnormalities among their children [20McMahon, 2013 #895, 21]; however, informing women that results are usually negative for 98% of older mothers may as well reduce the anxiety among women undergoing infertility treatment [22, 23].

Offering detailed information to pregnant women during genetic counseling sessions to enable its juxtaposition with the mothers’ own experiences and values to make an informed decision whether or not to undergo NIPT lowers the amount of uncertainty, stress, and anxiety they experience during the decision-making [24] [25].
The present study indicates that numerous individuals seek detailed information regarding the survival of children with positive results upon NIPT, including social support services, which indicates the lack of information and its accuracy regarding these disorders. As such, administration of NIPT is accompanied by high levels of stress among mothers, thereby potentially explaining their increased levels of anxiety until receiving the test results or their anxieties regarding the accuracy of the test [26].

Pregnant women taking the NIPT are compelled to gain adequate knowledge and understanding of the NIPT and the disorders screened therewith and to decide between continuing or terminating their pregnancy upon receiving a positive result within a very limited time. It is normal for all pregnant women to wish for a healthy child, which motivates them to take the NIPT to be “relieved [27].” However, even when they are relieved upon receiving negative results, many women were still ambivalent about their decision, feeling guilty about having tried to decide their child’s life, or having felt denial towards individuals with disabilities, but also having to endorse the thoughts about the test. This is probably what made them recognize their ambivalence and internal conflict at having these unresolved complex emotions.

**Conclusion**

The present study shows that even negative results can induce ambivalent feelings among pregnant women, and it is important to recognize that these feelings may be due to interrelated factors concerning genetic counseling, environmental factors, and increased anticipatory anxiety. Moreover, pregnant women have requested additional information regarding the detailed prognosis of individuals with Down syndrome or other congenital disorders and/or artificial abortion, assuming positive NIPT results. Therefore, genetic counseling and appropriate follow-up sessions are essential for the well-being of pregnant women taking the NIPT.

**Limitation and future direction**

There may be some possible limitations to this study. Firstly, a possible bias due to only women filling in the free description column of the questionnaire after one year; the background of these women is
also not clear. Further prospective studies are needed to investigate the psychosocial situations of women undergoing NIPT before and after the examination and after the passage of time. Even in the case of a negative NIPT result, women may have various conflicts and ambivalent feelings, and recognizing the factors that elicit these feelings may help future genetic counseling.

**Practice Implications**

To reduce internal conflict in pregnant women due to ambivalent feelings, it is critical to actively provide them detailed information about living with a child affected by one of the disorders screened for during the genetic counseling sessions and to discuss this information, assuming a potentially positive result, including specifics regarding medical pregnancy termination. Importantly, we believe that pregnant women should receive personalized psychological care so that they can make independent decisions based on factual knowledge. Thus, even if pregnant women experience feelings of ambivalence, they should be reassured and accept that undergoing NIPT was their decision after careful consideration, thereby reducing their potential internal conflicts.

**Abbreviations**

NIPT: Non-invasive prenatal testing

**Declarations**

**Ethics approval and consent to participate**

Subjects have given their written informed consent.

The study protocol has been approved by the ethics committee of Center for Maternal-Fetal, Neonatal and Reproductive Medicine, National Center for Child Health and Development (No 602), and each research institute’s committee on human research, as follows.

Ethics committee of Showa University, Ethics committee of Nagoya City University, The Ethical committee of the Jikei University School of Medicine, Nagasaki University Hospital Clinical Research Ethics Committee, The ethical committee of Hyogo College of Medicine, Ethics committee of Miyagi Children’s Hospital, Yokohama City University Certified Institutional Review Board, Ethics committee of Saitama Medical School, Ethics committee of Sapporo Medical University, Ethics committee of Iwate
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Consent for publication
Not applicable.

Availability of data and materials
The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Competing interests
The authors have no conflicts of interest to declare.

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Haruhiko Sago: Study planning, data collection, data review, revising the manuscript.

All authors have read and have approved the final manuscript.

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Tables
Please see supplementary files.

Figures
Figure 1

Flowchart of the respondent selection
Figure 2

The relationship between women's ambivalent emotions and their factors

Factors related to genetic counseling

Environmental factors

Increased anticipatory anxiety

Women’s ambivalent feelings
Components of the ambivalence Footnote: n = Number of occurrences of each category.
Categories may partially overlap.

Supplementary Files
This is a list of supplementary files associated with this preprint. Click to download.
renamed_a972e.pptx
revise Sep_2019__table__.pdf
The Questionnaire.docx