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ARTICLES

Research Articles

Knowledge and attitude of men about vasectomy as a method of family planning among married men working in Babcock University, Ogun state, Nigeria  30
Owopetu Christiana, Chukwuma Sonachi and Nwozichi Chinomso

Health of women with inflammatory bowel disease: Coping with challenges from pregnancy to child rearing  36
Chisato Kimura and Toshihide Ohmori
Full Length Research Paper

Knowledge and attitude of men about vasectomy as a method of family planning among married men working in Babcock University, Ogun state, Nigeria

Owopetu Christiana, Chukwuma Sonachi and Nwozichi Chinomso

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This study investigated the level of knowledge and attitude of married male workers about vasectomy and also the factors influencing the attitude of married men working in Babcock University towards vasectomy. Stratified random sampling technique was used to select 200 participants from the academic and administrative work divisions of the University and a self-constructed questionnaire was administered and 150 were retrieved and analysed using both descriptive and inferential statistics. Findings revealed that majority (42.7%) of the participants were between the ages of 31 to 40 years, Christians (97.3%), of the Yoruba tribe (55.3%), had a bachelor’s degree (46%) and were non-academic staff (53.3%). Majority (38%) of participants had adequate knowledge and 62.7% had positive attitude towards vasectomy. There was no association between participants’ level of education and their level of knowledge of vasectomy, however, a significant association was observed between participants’ level of knowledge and their attitude towards vasectomy ($\chi^2_{\text{cal}} = 53.89$, $P \leq 0.05$). The risk of spouse’s health (54%) was the major factor influencing positive attitude and the need of more children (41.3%) was the main factor influencing negative attitude towards vasectomy. There is therefore need to develop awareness programs in order to equip the population with valid information and thus increasing their knowledge about vasectomy. Specific strategies should be developed in order to tackle the identified barriers hindering the acceptance of vasectomy among married men.

Key words: Knowledge, attitude, vasectomy, family planning, married men.

INTRODUCTION

Family planning has beneficial effects in terms of sustainable socio-economic development and protection of the environment (Alemayehu, 2012). Unfortunately, most family planning programs in Nigeria have primarily targeted women (Onasoga et al., 2013) and men often do not participate in reproductive health matters. According to Odu et al. (2006), Nigerian men still have poor knowledge and attitude towards family planning despite global move to increase the involvement of men in reproductive health matters. A concerted effort is therefore needed to enhance men’s knowledge and increase their involvement in reproductive health (Ezegwui and Enwereji, 2009).

Sterilization, which includes vasectomy, is one of the...
important options available to married men who have decided to end childbearing, however, a number of identified factors affect the acceptance of vasectomy among men. In a qualitative study conducted by Bunce et al. (2007) using a focus group discussion and in-depth interviews to assess factors affecting vasectomy acceptability in Tanzania. Their findings showed that economical factors, spousal influence, religion, provider reputation and availability, uncertainty about the future and poor vasectomy knowledge were major factors that influenced vasectomy decision making among men.

Knowledge and attitude towards vasectomy have been reported to have significant influence on its uptake among men. For example, a study conducted by Ezegwui and Enwereji (2009) reported that only 6.8% of men may accept vasectomy with the knowledge they currently have while 89% will not. This finding was also linked to the fact that majority of the men believed that sterilization should be left for women only. In the same vein, a study conducted in Pakistan reported that men felt that vasectomy could cause impotence and is exclusively meant for prisoners (Nishtar et al., 2012). These are examples of myths held by men which have affected their attitude and acceptance of Vasectomy. Worldwide, only about 3 to 6% of couples are using vasectomy as a method of contraception (Pile and Barone, 2009).

The low acceptance of vasectomy has also been reported in the developed countries. The result of a National survey in the United States of America showed that only 13.3% of married men reported having had a vasectomy done (Anderson et al., 2010). Also, another American survey reported that only 6% of married men have undergone vasectomy (Eisenberg and Lipshultz, 2010). This corroborates the opinion of Shih et al. (2014) that despite the lower risk, high cost-efficiency, and high efficacy of vasectomy compared with female sterilization, more couples from the United States of America rely on female sterilization than vasectomy. Reasons for low use of vasectomy included lack of knowledge and misconceptions about the procedure, lack of access, provider bias and patents preferences.

Healthcare professionals and public health educators have important roles to play in order to convey correct information about vasectomy to the population thereby increasing its acceptability. Unfortunately, Okunola et al. (2009) reported that majority of health care workers were unwilling to accept vasectomy and none of them had ever had vasectomy done. Similarly, Ebeigbe et al. (2011) reported that despite good knowledge about vasectomy among Nigerian resident doctors, only 41.3% would opt for it while 89.4% of doctors counselled often for bilateral tubal ligation, only 5.8% did vasectomy.

In order to increase the acceptance of vasectomy among men, awareness programmes need to be developed and conducted to enhance men's knowledge and promote positive attitude towards vasectomy. It is in light of this that the researchers sought to assess the knowledge and attitude of married men about vasectomy as a method of family planning.

**METHODOLOGY**

The study adopted descriptive survey research design in generating priority data in measuring and analysing the variables under consideration. The study area was Babcock University, Ilishan-Remo, Ogun state. Stratified random sampling technique was used to select 150 participants from the academic and administrative work divisions of the University. A self-administered 37-item close-ended questionnaire designed based on the study objectives, was administered. The questionnaire included four sections: socio-demographic data, knowledge of vasectomy, attitude towards vasectomy and factors influencing attitude towards the choice of vasectomy. Ethical clearance was obtained from the Babcock University Health Research Ethics Committee. Participants were fully informed about the nature of the study and that their participation was voluntary. Furthermore, the information provided by the participants was treated with confidentiality. The questionnaires were directly administered personally and through three trained research assistants to the respondents at their various offices and collected immediately or later as time permitted. Instructions to guide the participants in filling the questionnaire were included. 200 questionnaires were distributed, 180 were retrieved and 150 were considered appropriate for analysis which involved 180 participants.

**RESULTS**

Figure 1 shows that the risk of spouses’ health (54%) was the major factor that influenced positive attitude towards vasectomy among participants. Figure 2 shows the need for more children (41.2%), dread or surgery (40.7%), fear of risk factors (38%), unavailability of safe practices (38%) and lack of support from spouse (38%) and ability to marry more wives (37.3%) were the major factors that influenced negative attitude towards vasectomy among participants. Table 1 shows that majority of the participants were between the ages of 31 to 40 years (42.7%), Christians (97.3%), Yorubas (55.3%), of average socio-economic status (70.7%), had a bachelor's degree (46%) and were non-academic staff (53.3%).

Table 2 shows that majority of participants (38%) had adequate knowledge while 30% had inadequate knowledge about vasectomy. Table 3 shows that majority (62.7%) of participants had positive attitude while 37.3% had negative attitude towards vasectomy.

Table 4 indicates that there was no statistically significant association between participants educational level and their level of knowledge about vasectomy ($X^2_{cal} = 25.09$, $P>0.05$). Table 5 shows that there was a statistically significant association between participants knowledge level and attitude towards vasectomy ($X^2_{cal} = 53.89$, $P<0.05$).

**DISCUSSION**

Findings of this study have shown that majority of participants were between the ages of 31 to 40 years and
had at least a bachelors degree which is not synonymous with the characteristics of participants in a previous study conducted in Nigeria by Onasoga et al. (2013). The difference in educational levels could be due to the fact that the current study was conducted in an educational institution where most people are expected to be educated. However, the age group in this present study is similar to the age group in another study conducted in Ibadan-Nigeria (Okunola et al., 2009).

The result of this study reveals that majority of participants had adequate knowledge about vasectomy. This finding contradicts the findings of Akpamu et al. (2013) who reported that majority of literate men in Epoma-Nigeria had inadequate knowledge about vasectomy as a method of contraception. Although, the studies were conducted among educated men, the difference in knowledge level could be due to methodological differences and also due to the fact that the present study was an institutional study. To further describe participants’ knowledge about vasectomy, majority (60.7%) knew that a man cannot bear children once he has had a vasectomy done. This implies that majority of the participants knew that vasectomy is a permanent method of family planning. This corroborates the report of World health organization (1994) that there is a very small chance that a man’s partner will become pregnant after he has had a vasectomy. In addition, findings from this study revealed that only 10.7% of participants felt that vasectomy requires long period of hospitalization after it is done. Parallel to this, Bounce et al. (2007) opined that vasectomy...
Table 1. Socio-demographic characteristics of participants n=150.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Frequency (n= 150)</th>
<th>Percentage (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21-30</td>
<td>27</td>
<td>18.0</td>
</tr>
<tr>
<td>31-40</td>
<td>64</td>
<td>42.7</td>
</tr>
<tr>
<td>41-50</td>
<td>38</td>
<td>25.3</td>
</tr>
<tr>
<td>51-60</td>
<td>16</td>
<td>10.7</td>
</tr>
<tr>
<td>61-70</td>
<td>4</td>
<td>2.7</td>
</tr>
<tr>
<td>71-80</td>
<td>1</td>
<td>0.7</td>
</tr>
<tr>
<td>Total</td>
<td>150</td>
<td>100.0</td>
</tr>
<tr>
<td>Religion</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Christianity</td>
<td>146</td>
<td>97.3</td>
</tr>
<tr>
<td>Islam</td>
<td>4</td>
<td>2.7</td>
</tr>
<tr>
<td>Total</td>
<td>150</td>
<td>100.0</td>
</tr>
<tr>
<td>Ethnic group</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hausa</td>
<td>3</td>
<td>2.0</td>
</tr>
<tr>
<td>Igbo</td>
<td>46</td>
<td>30.7</td>
</tr>
<tr>
<td>Yoruba</td>
<td>83</td>
<td>55.3</td>
</tr>
<tr>
<td>Others</td>
<td>18</td>
<td>12.0</td>
</tr>
<tr>
<td>Total</td>
<td>150</td>
<td>100.0</td>
</tr>
<tr>
<td>Socioeconomic status</td>
<td></td>
<td></td>
</tr>
<tr>
<td>High</td>
<td>34</td>
<td>22.7</td>
</tr>
<tr>
<td>Average</td>
<td>106</td>
<td>70.7</td>
</tr>
<tr>
<td>Low</td>
<td>10</td>
<td>6.7</td>
</tr>
<tr>
<td>Total</td>
<td>150</td>
<td>100.0</td>
</tr>
<tr>
<td>Highest educational level</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NCE</td>
<td>18</td>
<td>12.0</td>
</tr>
<tr>
<td>BSc/BA/B.Ed.</td>
<td>69</td>
<td>46.0</td>
</tr>
<tr>
<td>MSc/MA/M.Ed.</td>
<td>43</td>
<td>28.7</td>
</tr>
<tr>
<td>Ph.D.</td>
<td>20</td>
<td>13.3</td>
</tr>
<tr>
<td>Total</td>
<td>150</td>
<td>100.0</td>
</tr>
<tr>
<td>Work division</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Academic</td>
<td>70</td>
<td>46.7</td>
</tr>
<tr>
<td>Non-academic</td>
<td>80</td>
<td>53.3</td>
</tr>
<tr>
<td>Total</td>
<td>150</td>
<td>100.0</td>
</tr>
</tbody>
</table>

is a safe, simple and effective but underused method of contraception. Although the safety of vasectomy has been emphasized, interestingly however, majority of the participants in this study knew that vasectomy does not protect against sexually transmitted infections and does not interfere with men’s sexual functions. This is contrary to the report of Bounce et al (2007) who stated that Tanzanian men and women believed that vasectomy would result in decreased sexual performance also in the report of Nishtar et al. (2012) reported that participants believed vasectomy could cause impotence. A positive attitude is required in decision making about vasectomy uptake. Findings from this study have shown that positive attitude towards vasectomy was observed among majority of participants. Specifically, only very few of the participants (8.7%) believed that family should be left for women alone and only 15.3% believed that vasectomy was a form of castration. This implies that negative attitude towards vasectomy could be linked to myths and misconceptions regarding the procedure. Contrary to the attitude observed among the participants in the present study, the findings of Ezegwui and Enwereji
Table 2. Overall level of knowledge about vasectomy among participants.

<table>
<thead>
<tr>
<th>Level of knowledge</th>
<th>Number (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adequate (80% and above)</td>
<td>57 (38)</td>
</tr>
<tr>
<td>Moderate (50-79%)</td>
<td>48 (32)</td>
</tr>
<tr>
<td>Inadequate (&lt;50%)</td>
<td>45 (30)</td>
</tr>
</tbody>
</table>

Table 3. Overall attitude towards vasectomy among participants.

<table>
<thead>
<tr>
<th>Attitude</th>
<th>Frequency (n= 150)</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Positive (≥ mean)</td>
<td>94</td>
<td>62.7</td>
</tr>
<tr>
<td>Negative (&lt;mean)</td>
<td>56</td>
<td>37.3</td>
</tr>
</tbody>
</table>

Table 4. Association between level of education and knowledge about vasectomy.

<table>
<thead>
<tr>
<th>Level of education</th>
<th>Level of knowledge</th>
<th>χ²</th>
<th>P -value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Adequate</td>
<td>Moderate</td>
<td>Inadequate</td>
</tr>
<tr>
<td>NCE</td>
<td>4</td>
<td>1</td>
<td>13</td>
</tr>
<tr>
<td>BSc</td>
<td>25</td>
<td>21</td>
<td>23</td>
</tr>
<tr>
<td>MSc</td>
<td>17</td>
<td>19</td>
<td>7</td>
</tr>
<tr>
<td>Ph.D.</td>
<td>11</td>
<td>7</td>
<td>2</td>
</tr>
<tr>
<td>Total</td>
<td>57</td>
<td>48</td>
<td>45</td>
</tr>
</tbody>
</table>

Table 5. Association between level of knowledge and attitude towards Vasectomy.

<table>
<thead>
<tr>
<th>Level of knowledge</th>
<th>Attitude</th>
<th>χ²</th>
<th>P -value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Positive</td>
<td>Negative</td>
<td></td>
</tr>
<tr>
<td>High</td>
<td>51</td>
<td>6</td>
<td>53.89</td>
</tr>
<tr>
<td>Moderate</td>
<td>34</td>
<td>14</td>
<td>-</td>
</tr>
<tr>
<td>Low</td>
<td>9</td>
<td>36</td>
<td>-</td>
</tr>
<tr>
<td>Total</td>
<td>94</td>
<td>56</td>
<td>-</td>
</tr>
</tbody>
</table>

(2009) and Onasoga et al. (2013) showed that 40.7 and 48.5% respectively viewed vasectomy as castration. As component of participants’ attitudinal disposition towards vasectomy, majority disagreed that vasectomy would affect men’s dignity thus opposing the opinions of participants in Nishtar et al. (2012) where vasectomy was viewed to be meant exclusively for prisoners. While assessing further on the reasons for approval and disapproval of vasectomy as a method of family planning, result of this study showed that positive attitude towards vasectomy was majorly influenced by risk of spouses’ health (54%) and stress of tubal ligation (43.3%) while negative attitude towards vasectomy was influenced by need for more children and dread for surgery. This implies that the married men in Babcock University will most likely not consider vasectomy because they are in need of more children. Similar findings were reported in other studies as major factors that influenced men’s decision regarding vasectomy (Dutta et al., 2004; Murthy and Rao, 2003). However, this result is in contrast with a similar study conducted by Onasoga et al. (2013) in which cultural acceptance 82.4% (112), religious acceptance 72.1% (98), ignorance 70.6% (96) and accessibility of family planning clinic 66.2% (90) were the major factors negatively influencing attitude of men towards acceptance of vasectomy. Another important finding of this study was that participants’ educational qualification did not have a significant association with their level of knowledge about vasectomy. This implies that knowledge of vasectomy may not necessarily be more among the highly educated. A contrary finding was reported by Soaji et al. (2013) who
found that level of knowledge about vasectomy was more among people with higher education and also Keramat et al. (2011) reported positive associations between educational levels and choosing to have a vasectomy. With regards to the relationship between knowledge and attitude towards vasectomy, findings showed a statistically significant association between participants' knowledge level and their attitude towards vasectomy. Positive attitude was observed more among participants with adequate knowledge about vasectomy which opposes the result of Onasoga et al. (2013). Also contrary to this result, a previous study among Nigerian resident gynaecologists reported that despite good knowledge about vasectomy among doctors, majority were poorly disposed towards use of vasectomy (Ebeigbe et al., 2011). This indicates a need for effective national training programmes targeted at health workers to enhance their knowledge of vasectomy as well as break barriers to personal use of and counselling for vasectomy.

Associated risks, unavailability of safe expertise, lack of support from spouse and ability to marry more wives were additional factors that had negative influence on participant's attitude towards vasectomy. There is therefore need to develop awareness programs in order to equip the population with valid information and thus increasing their knowledge. This will invariably help individuals to overcome certain barriers. This study identified that spousal support and influence have effect on the acceptability of vasectomy, which is synonymous with the findings of Bounce et al. (2007). Therefore, the women group should be targeted in order to change their attitude toward vasectomy and thus give their support.

Conclusion

Contrary to previous studies which reported inadequate knowledge about vasectomy among men in Nigeria, this study has found that men in the study population had relatively adequate knowledge and positive attitude towards vasectomy. Since knowledge level was observed to be significantly associated with individual's attitude towards vasectomy, it is however, imperative that further education and counselling be provided to improve men's knowledge thereby promoting positive attitude towards vasectomy as a method of family planning. Specific strategies should be developed in order to tackle the identified barriers hindering the acceptance of vasectomy among married men.

ACKNOWLEDGEMENT

The authors give thanks to God Almighty for giving them strength and wisdom throughout the course of this study. They also acknowledge the efforts of the participants who gave their time to be included in this study. We will not fail to appreciate Professor E.O Ajao for his encouragement and support.
Full Length Research Paper

Health of women with inflammatory bowel disease: Coping with challenges from pregnancy to child rearing

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For women with irritable bowel disease (IBD), leading a healthy life during the perinatal period is important because it may affect their quality of life and motherhood. However, little attention has been paid to their challenges and the methods of coping with them. This study therefore aims to explore the challenges women with IBD face and the methods of coping with them during the perinatal period from their viewpoint. Exploratory, qualitative interview study design was used in this study. In-depth, one-on-one, semi-structured interviews were conducted in 2010 to 2011 among eight women having IBD, who experienced pregnancy, delivery and child rearing. The results of the interviews were analyzed by inductive content analysis. The participants recognized the following as challenges: high-risk management by obstetricians in collaboration with gastroenterologists, high risk to physical, mental and social well-being due to IBD relapse, limitation of information necessary during the perinatal period, issues related to the medical profession, and breastfeeding in their own way. They were coping with these issues by self-care for IBD, and making the most of various social support. Since IBD is often diagnosed during women's reproductive years, gastroenterology and obstetric specialists should cooperate with each other to routinely provide reproductive counseling to women with IBD. Furthermore, we should establish, as early as possible, health care communities for women with IBD, where they can maintain their quality of life and increase their preparedness for parenthood before pregnancy and during the perinatal period, based on multidisciplinary collaboration.

Key words: Inflammatory bowel disease (IBD), women's health, nursing, midwifery, multidisciplinary collaboration, reproductive care, pregnancy, child rearing, chronic illness, gastroenterology, qualitative approach.

INTRODUCTION

Recently, the incidence of inflammatory bowel disease (IBD), comprising ulcerative colitis and Crohn’s disease, a chronic, refractory and debilitating disease, has been increasing around the world, and has been acknowledged as a global disease (Molodecky et al., 2012). In Japan, the age-adjusted morbidity (/100,000 people) of IBD was estimated to be 84.5 for ulcerative colitis and 26.3 for Crohn’s disease in 2009 (Kuwahara...
As the incidence of IBD has so increased in Japan as in other countries, IBD has become a major social issue. IBD is considered to involve a complex interaction of four different mechanisms – a genetic predisposition that results in immune regulatory cell derangement, defects in the mucosal barrier, susceptibility to environmental triggers including specific antigens, and commensal bacteria in the patient’s intestinal lumen. IBD is characterized by repeated periods of remission and exacerbation, accompanied by fevers, rash, frequency bowel movements, diarrhea, rectal bleeding, loss of appetite, abdominal pain and fatigue. Further, IBD affects joints, eyes, skin, bones and organs other than the intestine during the course of the disease. Relapsing IBD attacks sufferers unexpectedly, and this calls for frequent medical checks at clinics or hospitals. Consequently, IBD sufferers experience mental fatigue (Beck et al., 2013), as well as social disabilities including unemployment, sick leave, unfilled potential in the workforce (Mallett et al., 1978; Bernklev et al., 2006) and deterioration of quality of life associated with health (Guassora et al., 2000; Graff et al., 2006; Lix et al., 2008; Jäghult et al., 2011) in addition to physical distress. Most IBD sufferers are diagnosed in the second or third decade of their life (Takebayashi and Nishiwaki, 2007; Bernick and Kane, 2010), and they experience significant life events, such as entering school, employment, marriage, pregnancy and delivery, after the development of IBD. Especially, since approximately 25% of women with IBD were reported to conceive for the first time after diagnosis (Baiocco and Korelitz, 1984), special care is needed in the perinatal management of women with IBD.

Recently the public has paid more attention to the health of women with IBD, especially medical aspects such as intensified symptoms, known as ‘mini-flares’, difficulties before and/or during menstruation (Kane, 2002), fertility (Willoughby and Truelove, 1980; Steinlauf and Present, 2004), the risk of deterioration of IBD during pregnancy (Alstead, 2002; Reddy et al., 2008; Habal and Huang, 2012), effects of drugs for IBD treatment during pregnancy (Coelho et al., 2011) and breastfeeding (Kane, 2002; Habal and Kapila, 2009). On the other hand, only a few studies have reported on the transitional process to parenthood for women with IBD, their experiences in the perinatal period (Mukherjee et al., 2002), and women with aggravated IBD during pregnancy (Cooper et al., 2011).

Research to elucidate how IBD affects mothers and their families including children in the transition period to motherhood would be useful for IBD women to prepare for their parenting role and to create new families. Findings from such studies would serve as effective information for health professionals, such as physicians, nurses and midwives, when supporting IBD women and their families in the reproductive period. Thus, the aim of this study was to examine the challenges women with IBD face and how they handle them, and to focus on the role of health professionals in supporting women with IBD from pregnancy to child rearing.

**METHODOLOGY**

**Design**

An exploratory design was employed, in order to elucidate the ascribed significance and perspectives of IBD women based on their experiences from pregnancy to child rearing.

**Participants**

The inclusion criteria were women with IBD within five years after the latest childbirth without serious complications and/or the history of mental disorder, and who were in a good condition in the IBD remission stage. They were given the formal request for this study initially by email attended by the primary investigator by a gastroenterologist, the second investigator, at a gastroenterology clinic in Saitama, Japan, or through internet sites where women could link to this study via the representative of the Society of Patients with Intractable Diseases. Although, ten women with IBD expressed their will to participate in the study, two abstained from interviews due to serious health conditions, and eight participated in the study. After receiving emails from women with IBD, the primary investigator returned a compliment and scheduled interviews.

**Data collection**

In 2010 to 2011, CK (the first author) conducted in-depth, one-on-one, semi-structured interviews (for 1.5 to 2 h) with eight women with IBD, at their homes, considering their physical condition, especially bowel control, and busy life due to child rearing, in order to interview them in a quiet and calm atmosphere. The interview guide had been prepared in advance by reviewing previous research (Mukherjee et al., 2002) that handled similar topics to ours. On the day of interviews, written informed consent was obtained before the interview. The participants first completed a brief questionnaire consisting of demographics, IBD history such as diagnosis, medication and surgery, and obstetric history, followed by a series of open-ended questions (Table 1). The interviews were tape-recorded with the participants’ permission, and transcribed verbatim by an experienced transcriber.

**Ethical considerations**

Ethical permission was obtained from the Tokyo Metropolitan University Ethics Committee. We assured the participants of their voluntary participation, protection of confidentiality/privacy, and right to withdraw participation whenever they want. Moreover, we explained that they could choose the date of the interview, and could suspend the interview at any time according to their physical condition. Lastly we explained the possibility of publishing our research findings in the future.

**Data analysis**

The contents of the recorded interviews were analyzed by inductive content analysis (Graneheim and Lundman, 2004; Elo and Kyngäs, 2008). The transcription was first read through several times to obtain a sense of the whole (Burnard, 1991). Then, the text about the participants’ experiences during the perinatal period was extracted as a unit of analysis, condensed as an analytical unit, and labeled with a code (Graneheim and Lundman, 2004). Next, the
various codes were compared based on differences and similarities, and grouped into sub-categories, which were then abstracted and named using content-characteristic words. Sub-categories with similar events were grouped together as categories, and the categories were further grouped as themes. The tentative categories were examined by two researchers (the authors) and amended. Finally, the codes, sub-categories, categories and themes were tabulated, and it was reconfirmed that the classification presented the validity and totality of the original narratives of the participants. In this connection, we quoted, in our manuscript, the participants’ original narratives that seemed to most appropriately represent the phenomena or meaning based on the empirical data.

**Rigour**

The researchers examined the rigour of our qualitative research based on the following five standards: credibility, transferability, dependability, conformability and authenticity (Guba and Lincoln, 1994). Concerning credibility, the researchers conducted a peer debriefing to compare the emerging codes and themes with the contents of the transcripts and interview summaries, and verified them in order to enhance the credibility of the findings. To ensure dependability and conformability, the researchers tape-recorded all interviews, and transcribed them verbatim, while preserving the documents/memos including all the trails of procedures or decision making regarding the research methodology. For this purpose also, the researchers utilized peer debriefing. As for transferability, the researchers reviewed the research findings of other studies concerning perinatal experiences or perinatal care of women suffering from other chronic diseases or disabilities, and provided, in the discussion section, sufficient descriptive data to enable the readers to evaluate the applicability of the data to other settings by comparing our research findings with others. Finally, concerning authenticity, the researchers repeatedly read the transcripts in consideration of the preliminary themes, and provided descriptive data, using the narratives of the participants categorized from the meaningful units in the synthesized sub-categories.

**RESULTS**

Table 2 shows the characteristics of the participants. All participants were aged between 27 and 41 years (mean: 33.8 years), with children aged from 7 months to 7 years. Three participants had two children, and one had three children. All participants were diagnosed with IBD when they were 18 to 35 years old. The diagnoses they reported are shown in Table 2. Although, the severity of symptoms varied, six participants experienced a relapse of IBD during pregnancy, and four had deterioration of IBD after delivery. Three participants had cesarean sections, and three had a preterm delivery, including one with caesarean section because of IBD, which necessitated the newborn’s admission to the Neonatal Intensive Care Unit (NICU). Two women with Crohn’s disease remained in remission by using biological preparation (#7) or mesalazine (#8) from pregnancy to child-rearing. One woman, per doctor’s orders, stopped taking biological preparation during the first trimester, but resumed from the second trimester through postpartum due to relapse (#2). One started to take biological preparation after enterectomy due to her small-intestinal perforation three months postpartum (she took mesalazine from pregnancy to two month postpartum) (#1). One woman began a course of biological preparation after her caesarean section due to relapse despite steroid use during pregnancy (#3). All three women with ulcerative colitis maintained their health by using mesalazine or predonine enema in spite of slight flares during pregnancy. These three women took the above-mentioned medications during postpartum, and two had no relapse (#4, 5). One experienced relapse due to stress related to her child’s disease, West syndrome (#6). We here present the challenges women with IBD faced and the methods of coping with them based on the main themes derived from the qualitative data. The names of participants are concealed and replaced with numbers to assure anonymity.

**Experiences of women with IBD during perinatal period**

We identified two themes derived from the experiences of women with IBD during the perinatal period: challenges women with IBD faced and methods of coping with the challenges, as summarized in Table 3.

**Challenges women with IBD faced**

We identified five categories under the heading “Challenges women with IBD faced”: High-risk management by obstetricians in collaboration with gastroenterologists; high risk to physical, mental and social well-being due to IBD relapse; limitation of information necessary
Table 2. Background characteristics of participants during perinatal period.

<table>
<thead>
<tr>
<th>Participant</th>
<th>Diagnosis (reported by participants)</th>
<th>Age at interview</th>
<th>Age at diagnosis</th>
<th>Number of children</th>
<th>Main events experienced by participants</th>
<th>During pregnancy</th>
<th>During delivery</th>
<th>During postpartum period</th>
</tr>
</thead>
<tbody>
<tr>
<td>#1</td>
<td>CD</td>
<td>35</td>
<td>29</td>
<td>2</td>
<td>IBD relapse, Long-term hospitalization</td>
<td>1st child: Caesarean section, newborn management in NICU; 2nd child: Repeat Caesarean section, normal neonate</td>
<td>IBD relapse, Hospitalization, Small-intestinal perforation, Enterectomy</td>
<td></td>
</tr>
<tr>
<td>#2</td>
<td>CD</td>
<td>31</td>
<td>27</td>
<td>2</td>
<td>IBD relapse, Attended hospital for check-ups</td>
<td>1st and 2nd child: Normal vaginal delivery</td>
<td>IBD relapse, Hospitalization for check-ups, Surgery for colonic ileus and Seton drainage for anal fistulas</td>
<td></td>
</tr>
<tr>
<td>#3</td>
<td>CD</td>
<td>27</td>
<td>20</td>
<td>1</td>
<td>IBD relapse, Long-term hospitalization</td>
<td>1st child: Preterm delivery, Caesarean section, Newborn management in NICU</td>
<td>IBD relapse, Hospitalization</td>
<td></td>
</tr>
<tr>
<td>#4</td>
<td>UC</td>
<td>27</td>
<td>22</td>
<td>2</td>
<td>IBD relapse, Attended hospital for check-ups</td>
<td>1st child: Normal vaginal delivery; 2nd child: Preterm delivery, Caesarean section, Breech presentation</td>
<td>No problem</td>
<td></td>
</tr>
<tr>
<td>#5</td>
<td>UC</td>
<td>41</td>
<td>35</td>
<td>1</td>
<td>IBD relapse, Attended hospital for check-ups</td>
<td>1st child: Normal vaginal delivery</td>
<td>No problem</td>
<td></td>
</tr>
<tr>
<td>#6</td>
<td>UC</td>
<td>39</td>
<td>25</td>
<td>1</td>
<td>IBD relapse, Attended hospital for check-ups</td>
<td>1st child: Normal vaginal delivery</td>
<td>IBD relapse, Attended hospital for check-ups Child: West syndrome</td>
<td></td>
</tr>
<tr>
<td>#7</td>
<td>CD</td>
<td>38</td>
<td>22</td>
<td>3</td>
<td>No problem</td>
<td>1st child: Normal vaginal delivery; 2nd child: Intrauterine infection, Preterm vaginal delivery, Newborn/infant management in NICU/GCU; 3rd child: Normal vaginal delivery</td>
<td>No problem</td>
<td></td>
</tr>
<tr>
<td>#8</td>
<td>CD</td>
<td>32</td>
<td>18</td>
<td>1</td>
<td>No problem</td>
<td>1st child: Normal vaginal delivery</td>
<td>No problem</td>
<td></td>
</tr>
</tbody>
</table>

CD, Crohn’s disease; UC, Ulcerative colitis

during perinatal period; issues related to medical profession; breastfeeding in one’s own way.

**High-risk management by obstetricians in collaboration with gastroenterologists:** Participants in this study described three concepts: mental preparedness for high-risk management, maintenance of a normal gastroenterological condition and maintenance of normal obstetric course. They felt that these concepts influenced each other.

1. Mental preparedness for high-risk management: Though the participants felt it was a bother to receive management by both an obstetrician and a gastroenterologist, they accepted the high-risk management as a safe system.

   “My pregnancy was risky because I needed to be managed by both an obstetrician and a gastroenterologist. It was bothersome for me to have check-ups in two different departments. But I felt at ease to know that collaborative management was possible if needed.”

2. Maintenance of a normal gastroenterological condition: Most participants continued to take medication and/or elemental dietary supplement (Elental) to control IBD. However, frequent bowel movements with associated faecal urgency and anemia did not resolve. Therefore, the participants tried to lead a stress-free life with a wait-and-see approach without receiving check-ups, refraining from going out, etc.

   “I took 5-ASA (mesalazine) during my pregnancy because my IBD symptoms were severe. But my symptoms did not get better. And I had little trouble in my daily living though I felt sluggish due to anemia. The worst thing about this disease is that I have little time from when my stomach starts
Table 3. Themes, categories and sub-categories derived from experiences of participants during perinatal period.

<table>
<thead>
<tr>
<th>Theme</th>
<th>Category</th>
<th>Sub-category</th>
</tr>
</thead>
<tbody>
<tr>
<td>Challenges participants faced</td>
<td>High-risk management by obstetricians in collaboration with gastroenterologists</td>
<td>Mental preparedness for high-risk management</td>
</tr>
<tr>
<td></td>
<td>High risk to physical, mental and social well-being due to IBD relapse</td>
<td>Maintenance of a normal gastroenterological condition</td>
</tr>
<tr>
<td></td>
<td>Limitation of information necessary during perinatal period</td>
<td>Maintenance of normal obstetric course</td>
</tr>
<tr>
<td></td>
<td>Issues related to medical profession</td>
<td>Attending hospital and/or hospitalization due to IBD relapse</td>
</tr>
<tr>
<td></td>
<td>Breastfeeding in one’s own way</td>
<td>Lack of obstetric and perinatal education</td>
</tr>
<tr>
<td>Methods of coping with challenges</td>
<td>Self-care for IBD</td>
<td>Separation of mother and child, necessitating substitute child care by other family members</td>
</tr>
<tr>
<td></td>
<td>Making the most of various social support</td>
<td>Lack of information</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Inconsistency of information</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Resistance to patient becoming a mother</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ignorance about IBD</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Various styles of breastfeeding</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Early weaning due to separation of mother and baby</td>
</tr>
</tbody>
</table>

hurting to get to a restroom. So, I tried to only go to places where I could find a toilet easily.”

3. Maintenance of normal obstetric course: The participants tried to continue taking Elental during pregnancy to maintain a normal fetal body weight. When the courses of IBD and pregnancy were normal, they led lives like a healthy woman, almost forgetting about IBD.

“I tried to take Elental for growth of my baby, but I couldn’t while I was pregnant. My obstetrician, gastroenterologist and I monitored the growth of my baby with ultrasound scans. Other than that, my condition was good while I was pregnant, and I had a really happy, stress-free time. It was as if my IBD disappeared!”

High risk to physical, mental and social well-being due to IBD relapse: When IBD relapsed, participants in this study experienced three major concerns: attending hospital and/or hospitalization because of IBD relapse, lack of obstetric perinatal education and separation of mother and baby necessitating substitute childcare by other family members. All these factors negatively affected the physical, mental and social well-being of the mothers and their families.

1. Attending hospital and/or hospitalization because of IBD relapse: Six pregnant and four postpartum participants had an IBD relapse, and attended hospital and/or were hospitalized in the gastroenterology department. One participant who had to be hospitalized for a long time from pregnancy to the postpartum period and underwent abdominal surgery after caesarean section experienced intolerable abdominal stinging and mental distress including anxiety about insufficient preparedness for motherhood.

“I had an emergency admission to hospital because of inflammation of the small intestine when I was four months pregnant. I couldn’t eat anything, and received only intravenous hyperalimentation for a long time, and restarted steroid use. As my baby grew, it pressed on my small intestine, and I was having intolerable abdominal pain. I was depressed. I was worried about whether I could take care of my baby.”

2. Lack of obstetric and perinatal education: All participants who were hospitalized because of IBD relapse commonly experienced adaptation delay in postnatal child rearing because of insufficient preparedness for motherhood.

“I had no chance to prepare for delivery or childcare like mothers who were attending mothers’ classes, and so, I didn’t have any childcare skills because I was in hospital so long. After the caesarean section, I underwent abdominal surgery again, so my husband and mother-in-law had to take care of my baby for 2 months.”

3. Separation of mother and baby necessitating substitute childcare by other family members: Participants who had
to be hospitalized for IBD relapse commonly experienced separation from their baby, and were forced to discontinue breastfeeding. Their own mothers and mothers-in-law had to take care of their babies, and the grandmothers experienced refusal of bottle-feeding by the babies, resulting in weaning babies from milk and starting supplementary foods.

“I had a normal postpartum after a C-section, but one day I suddenly developed strong abdominal pain. I took my two daughters to my parents-in-law by car and left them there. I was then taken to hospital by ambulance, and had an emergency enterectomy. It was a close call, and I am glad I survived. I was hospitalized and separated from my baby, and so my mother had to wean my baby from milk and started to give her solid food, little by little. My daughter was exclusively breastfed until that time, so my mother had a lot of trouble.”

Limitation of information necessary during perinatal period: The participants searched on the internet for the effects of IBD on pregnancy and delivery while consulting their doctors at all reproductive stages. They were confused by the lack and/or discrepancy of information.

Lack of information: Most participants stated that they had little or no information from other women with IBD. They searched for information about IBD and pregnancy/delivery on the internet as the first step. They asked their gastroenterologists to provide them with information, but they could only receive information limited to medical aspects of IBD.

“I searched the web for information on how other women with IBD were dealing with difficulties during pregnancy and delivery because I couldn’t get such information from other women with IBD. My doctor just gave me some papers written by other doctors.”

Inconsistency of information: Most participants searched for the effects of medication, especially of Infliximab and/or painkillers, on the internet. However, they could not find accurate information. Moreover, they recognized the inconsistency of information given by doctors from different specialty areas, and were confused by discrepancies.

“I searched for the effects of Infliximab and painkillers on the internet. But I couldn’t find accurate information on the fetus and breastfeeding, so I was afraid of the effects. For example, I felt that the internal medicine doctor’s opinion was different from that of surgeons with regard to the use of painkillers for my anal pain.”

Issues related to medical profession

This category included two sub-categories: resistance to patient becoming a mother, and ignorance about IBD among nurses and midwives.

Resistance to patient becoming a mother

One participant encountered direct resistance to her desire to have a baby by the gastroenterologist during the pregnancy. She was greatly hurt, and continued to have feelings of inferiority about having a child even in the later child-rearing stage, five years after delivery. However, she balanced her negative feelings with the confidence that she could deliver a child and love her.

“I don’t want to mention, but the doctor said to me “Don’t you care about your husband and children and the problems they’ll have because of your IBD?”. After that, I always had a feeling of inferiority. But my children are very lovely. I am happy that I could deliver babies and can take care of them.”

Ignorance about IBD

The participants expressed their gratitude to nurses and midwives for their compassionate care, while they considered that the nursing profession had only limited knowledge about IBD. Therefore, some participants did not dare to consult nurses and midwives about their disease or concerns.

“I think that the midwives didn’t care about my disease during pregnancy and the postpartum. I had a normal pregnancy and delivery, and so, I didn’t consult them. I felt that they did not know much about IBD, anyway.”

Breastfeeding in one’s own way

The participants experienced various styles of breastfeeding including exclusive/partial breastfeeding, early weaning due to separation of the mother and baby.

Various styles of breastfeeding

Seven participants practiced and enjoyed exclusive or partial breastfeeding, and finally accepted their own style of breastfeeding.

“I wanted to exclusively breastfeed my baby. I can contact my baby warmly while breastfeeding, and it makes me feel like a mother. All my friends exclusively breastfed their babies. So I wanted to do so, too.”

Early weaning due to separation of mother and baby

Five participants, who had to introduce artificial milk or
supplementary food to their babies when they were hospitalized for IBD relapse, did not consider the change to be negative, and understood that it was a necessary decision.

“I had to stay in hospital during the early period of childcare. One week after my hospitalization, my daughter was weaned from breast milk and began to take formula milk. But I think it was a good choice because I just had to do so.”

Methods of coping with challenges

This theme was summarized into two categories: self-care for IBD and making the most of various social support.

Self-care for IBD

The participants categorized self-care for IBD into two: maintaining physical wellness and behaving as normal or usual.

Maintaining physical wellness: The participants were making efforts to maintain their physical wellness from the perspective of IBD sufferers, though they continued to avoid food unsuitable for their constitution, prevent fatigue, and monitor for bloody stools and the pattern of bowel movements.

“Oily or cold foods affect my abdominal condition, especially ice cream. So I don’t eat it. I have dairy products with low fat. I pay careful attention to the amount of sleep I have, and try not to get too tired. I always have diarrhea and frequent bowel movements when I am tired.”

Behaving as normal or usual: In social relationships, the participants adopted two ways of letting other people know about their disease; by getting people to understand IBD without concealing it, or by avoiding informing people except their family members and friends about IBD. In these ways, they felt comfortable and able to prevent stress or problems.

“I used to talk about my disease to my old friends because I didn’t want to conceal it. It was easier for me. But, now, I don’t tell my friends who are also mothers about my disease. I feel it is not necessary to tell them about my IBD.”

Making the most of various social support

Three sub-categories of social support utilized by the participants were identified: family support, peer support and professional support.

Family support

Regardless of IBD relapse, the participants commonly utilized the support of their parents (or parents-in-law) as the first choice, which covered complete or partial childcare. Partners’ support consisted of emotional or financial support.

“During my regular checkups and hospitalization for IBD relapse, my parents take care of my child. My husband always comes back home late at night, so I can’t rely on him on weekdays. My husband and children spent time with me every weekend in the hospital.”

Peer support

Most participants had already established networks with female friends with IBD before their pregnancy or in the perinatal period. They had received effective information to visualize concrete images of daily life during the perinatal period, from the viewpoint of IBD sufferers.

“My friend with IBD became pregnant, and delivered a baby a few months earlier than me. She has Crohn’s disease, and talked to me about her experience. We have been friends for a long time. I can ask her at any time about pregnancy/delivery and child-rearing, as well as about IBD.”

Professional support

All participants had established a strong relationship with their IBD specialists. The IBD specialists monitored the symptoms of the participants, guided them about ways to have a healthy life based on their clinical assessment, and provided them with appropriate information and necessary treatment. IBD specialists were the persons who offered a sense of security to IBD women and their families.

“I always depend on my doctor. If I have any problem with the disease, I go to the clinic at any time. My doctor suggested a good time to get pregnant, with consideration of my condition.”

DISCUSSION

Strategies for IBD women to overcome difficulties in perinatal period

The participants in this study had regularly monitored their IBD symptoms, such as diarrhea, by themselves,
and paid close attention to diet and keeping themselves rested to maintain physical well-being in the same way as they did before pregnancy. This is in agreement with the research finding that many IBD sufferers acquire the ability to take care of themselves in order to maintain ‘normality as an IBD sufferer’ in their own way (Fletcher and Schneider, 2006; Fletcher et al., 2008; McCormick et al., 2012). Since the perinatal period is a transitional stage, which is associated with changes in the family system and the need to acquire new parental skills including breastfeeding, all couples experience some psychological stress. As women with IBD have only limited information about pregnancy and the disease, it is important for them to pay attention, before pregnancy, to their daily diet, rest, coping with stress, symptom control by medication, and postpartum childcare, and discuss these concerns with their families.

Regarding self-care for psychological health, some participants thought it was better to pretend to be normal and experience ordinary daily life as if they were healthy women, in spite of their difficult experiences of IBD relapse in the perinatal period. Maintaining ‘health-related normality’ and ‘pretending to be normal’ are reported to be essential for IBD suffers (Hall et al., 2005). The same was true for the respondents in this study. If IBD sufferers had a relapse in the perinatal period, they would be categorized into the group of ‘pregnant women with disability’. When women with difficulties (example, disability or illness) receive reproductive care, they are forced to confront tasks different from those women with a normal course, including collection of various information about reproductive health care, despite little information being available, and to spend the perinatal period communicating effectively with multidisciplinary care providers (Wong, 2000).

However, women with disabilities are adaptive to and knowledgeable about their own conditions and difficulties, and want to be respected by healthcare professionals as individuals (Barber, 2008). Therefore, women with disabilities, in particular those with ‘a high risk’, should have the chance to access appropriate maternity-care information and consultation as easily as their ‘low risk’ or ‘normal’ counterparts enjoy, and be looked after by a health care system that supports their maternal identity and values (Walsh-Gallagher et al., 2012). Most participants in our study could overcome the challenges during the perinatal period by making use of various social support. Especially, family support during IBD relapse was essential for IBD women to become a parent while overcoming difficulties. As the perinatal period is a transitional stage not only for IBD women but also for their family members, IBD women should develop supportive communication with their families through having anticipatory guidance on the risk of relapse and coordinated childcare with their partners and/or family members.

Further, most participants had utilized gastroenterologist as one of their social supports. In particular, they had received suggestions about the appropriate time to get pregnant and information about the interaction between pregnancy and IBD when they planned to get pregnant. At the same time, they had searched for this information on the web. These findings coincide with the results of previous studies showing that the particular information sources about pregnancy for IBD women were gastroenterologists and the internet (Marri et al., 2007; Mountfield et al., 2009; Gawron et al., 2014). Although, the internet should not be the main source of information for women with IBD, as people are accustomed to seeking information on the internet, it would probably be necessary to have a reliable web-based information package for pregnant IBD patients and those planning to get pregnant including specific guidance for daily life from before pregnancy to postpartum such as self-care and/or social supports including family and/or peers.

Considering that the peak age of IBD onset is during the reproductive period, IBD women should receive appropriate reproductive health guidance before pregnancy, on an individual or group basis, as follows: appropriate time to get pregnant, various courses and risks in the perinatal period, effects of IBD medication on pregnancy/breastfeeding, necessity for high risk management, ways to get appropriate information and methods of coping with IBD relapses. At the, IBD women should have the opportunity to come in contact with peers with IBD, and establish a supportive environment, such as a ‘buddy system’ with peers, by themselves, because they usually hesitate to talk about IBD-related topics that are rarely covered in regular parents’ classes (Blackford et al., 2000).

Roles of health professionals involved in care of women with IBD

Gastroenterologists are the key sources for women with IBD of reproductive age who decide to have a baby. Therefore, they should always support IBD women in their decision-making by providing them with updated and precise information. Gastroenterology nurses are in the best position to concretely show IBD women the challenges they face and need to manage during the perinatal period. These gastroenterology specialists should integrate, as a routine care system, reproductive counseling, such as discussion about contraceptive methods and birth control, into education for IBD women of reproductive age and their families (Gawron et al., 2014). For example, gastroenterology specialists can provide IBD women and their partners with not only appropriate information about pregnancy, but also opportunities to participate in perinatal health education classes to come in contact with IBD peers, based on collaboration with other professionals including midwives and public health nurses. Such inter-professional care
would play a pivotal role in caring for IBD women, and could help to create a favorable health care community for IBD women before their babies’ conception (Barber, 2008). Such an approach coincides with the research findings that pointed out the need for specific care for women with disabilities due to other chronic medical conditions (Corbin, 1987; Blackford et al., 2000; Aukamp and Sredl, 2004; Thompson et al., 2008).

The results of this study suggest that, in the case of long-term hospitalization during pregnancy because of IBD relapse, IBD women and their families have little or no chance to learn about childcare during the maternity stage, resulting in possible insufficient preparedness for parenthood including immediate postpartum childcare and mental stress. It is speculated that gastroenterology nurses may lack the understanding that women with IBD hospitalized for relapse should simultaneously prepare for motherhood, because gastroenterology nurses tend to manage IBD women as a medical model, while midwives tend to undervalue the disease course and management of IBD. A similar situation has often been criticized when pregnant women with complications or disabilities received multidisciplinary follow-up (Thierry, 2006; Piotrowski and Snell, 2007). Recently, although medical collaboration between gastroenterology and obstetrics has become standard policy for treatment of women with IBD, collaborative care by gastroenterology nurses/midwives and community health professionals, including public health nurses, has lagged behind. Therefore, it is necessary to establish, as early as possible, a perinatal collaborative model for IBD women based on effective communication among gastroenterology nurses, midwives and community health professionals with a multidisciplinary team approach. Such an approach may include health education for IBD women such as methods of birth control, knowledge about the course and risks of IBD, breastfeeding care during hospitalization for IBD relapse, and care coordination in communities, and will improve the quality of life of women with IBD during the perinatal period.

On the other hand, previous studies suggested that IBD sufferers have a tendency to refrain from talking about their typical symptoms, such as fatigue, to health professionals and even their own family members and friends (Beck et al., 2013; Hall et al., 2005). Gastroenterology nurses and midwives should focus on these tendencies, consider the health-related and learning needs of IBD women, and offer them opportunities to express their feelings or concerns during the perinatal period.

LIMITATIONS

This study has a few limitations. First, the fact that all interviews were conducted by the first author alone might have weakened the credibility of the findings. Second, the sample size was too small to generalize the findings to other populations. In particular, participants in this study may not present an average IBD mother as six of the eight relapsed during pregnancy, and four the perinatal period affects the results. Finally, age of their children at the interviews ranged from 7 month to 7 years old. Due to limitations within the interview surveys, there is some uncertainty whether interviewees have mixed memories with similar events when the answers involve past events or feelings. Consequently, we should plan the prospective research design to include interviews at some points from pregnancy to child rearing stage in the future.

Conclusion

Since IBD is often diagnosed during a woman’s reproductive years, gastroenterologists and obstetricians should cooperate with each other to routinely provide IBD women with reproductive counseling through individual or group education. Furthermore, we should establish, as early as possible, health care communities for women with IBD and their families, where good IBD control and health of IBD women can be maintained before pregnancy and during the perinatal period, based on multidisciplinary collaboration.

Conflict of interest

The authors declare that they have no conflicts of interest.

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