Incidence and risk factors for insufficient endometrial tissue from endometrial sampling

Objective: To assess the incidence of and factors that predict insufficient tissue after endometrial sampling.

Methods: This study reviewed the records of women undergoing endometrial sampling at Khon Kaen University’s Srinagarind Hospital between June 2014 and June 2015. It excluded cases in which the device could not be inserted into the uterine cavity due to pain intolerance or equipment failure. The criterion for diagnosing insufficient endometrial tissue was a lack of any intact tissue fragments containing both glands and stroma.

Results: Medical records of 233 women were reviewed. Insufficient tissue following endometrial sampling was noted in 67 cases (28.8%; 95% confidence interval [CI]=23.0–35.0). Histologic results in the remaining 166 women included normal pathological endometrium (121, 51.9%), endometrial polyps (7, 3.0%) endometrial hyperplasia (27, 11.6%), and endometrial cancer (11, 4.7%). According to multivariable analysis, menopausal status (odds ratio [OR] =3.60, 95% CI=1.84–7.05) and endometrial thickness of less than 8 mm (OR =3.91, 95% CI=1.49–10.21) were significant independent predictors for insufficient endometrial tissue after endometrial sampling.

Conclusion: The incidence of insufficient tissue following endometrial sampling was 28.8%. Significant independent factors associated with an increased risk of insufficient tissue were menopausal status and endometrial thickness of less than 8 mm.

Keywords: insufficient endometrial tissue, endometrial sampling, Endocell®, sample adequacy, inadequate sampling

Introduction

Endometrial sampling is a method used to obtain endometrial tissue in order to histologically examine women who have abnormal uterine bleeding.¹ This procedure is commonly performed to look for significant endometrial pathologies, including endometrial hyperplasia and endometrial carcinoma, in women who have an abnormally thick endometrium on imaging, atypical glandular cells on cervical smear, or postmenopausal bleeding.²,³ Previous studies have indicated that the performance of endometrial sampling is as good as those of dilatation and curettage (D&C) or manual vacuum aspiration.⁴,⁵ In cases of postmenopausal bleeding, endometrial sampling has been shown to have a sensitivity of 90% and specificity of 99%–100% for diagnosing endometrial hyperplasia or endometrial cancer with D&C or hysteroscopy with biopsy as a reference standard.⁶ A previous systematic review found moderate agreement with regard to tumor grade between endometrial sampling and final pathology from hysterectomy.⁷

The disadvantage of endometrial sampling relates to the small volume of endometrial tissue obtained. Endometrial sampling yields insufficient tissue for definite pathological diagnosis in 6%–33% of cases.⁸–¹³ Approximately 6% of postmenopausal women with...
nondiagnostic specimens after endometrial sampling for abnormal uterine bleeding are subsequently found to have significant endometrial lesions on further evaluation. Therefore, it is necessary to identify patients in whom endometrial sampling may yield insufficient tissue and for whom other endometrial evaluation procedures would be more suitable. The aim of this study is to assess the incidence of and factors that predict insufficient tissue after endometrial sampling.

Methods
After receiving approval from the Khon Kaen University Ethics Committee for Human Research based on the Declaration of Helsinki and the ICH Good Clinical Practice Guidelines (HE591350), we reviewed the records of women who underwent endometrial sampling as a primary treatment at Khon Kaen University (KKU)’s Srinagarind Hospital, between June 2014 and June 2015. Because this was a retrospective study and data were analyzed anonymously, the need for informed consent was waived by the ethics committee. We excluded cases in which the device could not be inserted into the uterine cavity due to intolerance to pain or equipment failure.

Methods to obtain endometrial tissues for pathology examination in our institution depended on the menopausal status. Endometrial sampling was performed in all premenopausal women with abnormal uterine bleeding. For postmenopausal women, there were two options, ie, endometrial aspiration or transvaginal ultrasound for endometrial measurement.

All endometrial sampling procedures were performed via Endocell® (disposable endometrial cell sampler tube; CooperSurgical, Inc., Trumbull, CT, USA) requiring only manual suction to extract a sample from the endometrium. This device provides excellent suction using an elastomeric seal piston plunger with an outside diameter of 3.1 mm. All sampling procedures were conducted by the residents in training or members of the gynecology staff. Abstract data included in this study were patient characteristics, indications for endometrial sampling, endometrial thickness by ultrasound (if any), types of operators (gynecology residents vs attending staffs), and detailed pathological reports. Pathological tissue reports were reviewed by a single gynecologic pathologist (PK). The criterion for diagnosis of insufficient endometrial tissue was a lack of any intact tissue fragments containing both glands and stroma.

Descriptive statistics were used for reporting the data. The chi-square and Fisher’s exact tests were used to univariately identify factors potentially related to insufficient endometrial tissue samples. Factors with a P-value less than 0.20 according to univariate analysis were subjected to multivariate analysis using a logistic model to reveal which were the independent factors. An odds ratio (OR) with a 95% confidence interval (CI) that did not include unity was considered statistically significant. Statistical analyses were carried out using SPSS version 17.0 (SPSS Inc., Chicago, IL, USA).

Results
The medical records of 233 women who underwent endometrial sampling were reviewed. Table 1 displays the baseline characteristics of the patients. The mean age of the patients was 48.3 years (range=27–86 years). Transvaginal ultrasound for evaluating endometrial thickness was performed in 108 (46.4%) women. Endometrial thickness ranged from 1.9 to 53.0 mm.

Insufficient tissue following endometrial sampling was noted in 67 women (a rate of 28.8%; 23.0–35.0). The histologic results of the remaining 166 women included normal pathological endometrium (121, 51.9%), endometrial polyps (7, 3.0%), endometrial hyperplasia (27, 11.6%), and endometrial cancer (11, 4.7%).

Details regarding endometrial thickness as measured by transvaginal ultrasound are reported in Table 1. Subsequent

| Characteristics                        | Number of women (%) |
|----------------------------------------|---------------------|
| Age                                    |                     |
| Younger than 60 years                  | 213 (91.4)          |
| 60 years or older                      | 20 (8.6)            |
| Parity                                 |                     |
| Nulliparous                            | 28 (12.0)           |
| Multiparous                            | 205 (88.0)          |
| Menopausal status                      |                     |
| Premenopause                           | 162 (69.5)          |
| Postmenopause                          | 71 (30.5)           |
| Previous vaginal birth                 |                     |
| Yes                                    | 177 (75.9)          |
| No                                     | 56 (24.1)           |
| Current hormone use                    |                     |
| Yes                                    | 55 (23.6)           |
| No                                     | 178 (76.4)          |
| Types of operator                      |                     |
| In-training resident                   | 192 (82.4)          |
| Gynecology staff                       | 41 (17.6)           |
| Indications for endometrial sampling   |                     |
| Abnormal uterine bleeding              | 190 (81.5)          |
| Abnormal glandular cells on cervical smears | 37 (15.9)          |
| Postmenopausal pyometra                | 6 (2.6)             |
| Endometrial thickness (cutoff 8 mm)    |                     |
| Less than 8 mm                         | 59 (54.6)           |
| 8 mm or more                           | 49 (45.4)           |
management of women with insufficient samples included fractional curettage (18; 26.9%), re-endometrial biopsy (5; 7.5%), hysterectomy (9; 13.4%), and no further management (35; 52.2%). In the 67 women with insufficient endometrial tissue samples, there is no evidence of endometrial carcinoma development over a median follow-up period of 36 months. Eleven (4.7%) patients were diagnosed with endometrial carcinoma (one unstaged, four stage IA, two stage IB, two stage II, one stage IIIIC1, and one stage IVB).

Eight clinical variables were analyzed, including advanced age, parity, menopausal status, operator, previous vaginal birth, current hormonal use, indications for endometrial sampling, and endometrial thickness, to significantly predict insufficient endometrial tissue among the specimens. Menopausal status, endometrial thickness, and advanced age had P-values of less than 0.20 according to univariate analysis. Multivariate analysis was performed using a logistic regression model that included three significant covariates. Only menopausal status (OR=3.60, 95% CI=1.84–7.05) and endometrial thickness less than 8 mm (OR=3.91, 95% CI=1.49–10.21) remained statistically significant predictors for insufficient endometrial tissue on disposable endometrial cell sampler tube after this analysis (Table 2).

Table 2 Factors predicting insufficient endometrial tissue

| Variables                                | Number of insufficient (%) | Univariate P-value | Multivariate OR (95% CI) |
|------------------------------------------|----------------------------|--------------------|--------------------------|
| Menopausal status                        |                            |                    |                          |
| Postmenopause (n=71)                     | 34 (47.9)                  | <0.05              | 3.60 (1.84–7.05)         |
| Premenopause (n=162)                    | 33 (20.4)                  |                    | Reference               |
| Endometrial thickness                    |                            |                    |                          |
| Less than 8 mm (n=59)                    | 25 (42.4)                  | 0.01               | 3.91 (1.49–10.21)        |
| 8 mm or more (n=49)                      | 8 (16.3)                   |                    | Reference               |
| Age                                      |                            |                    |                          |
| 60 years or older (n=20)                 | 9 (45.0)                   | 0.09               | 0.95 (0.32–2.77)         |
| Younger than 60 years (n=213)            | 58 (27.2)                  |                    | Reference               |
| Indications for endometrial sampling     |                            |                    |                          |
| AUB (n=190)                              | 58 (30.5)                  | 0.20               | Variable removed        |
| Non-AUB (n=43)                           | 9 (20.9)                   |                    |                         |
| Current hormone use                      |                            |                    |                          |
| Yes (n=55)                               | 13 (23.6)                  | 0.34               | Variable removed        |
| No (n=178)                               | 54 (30.3)                  |                    |                         |
| Parity                                   |                            |                    |                          |
| Nulliparity (n=28)                       | 10 (35.7)                  | 0.38               | Variable removed        |
| Multiparity (n=205)                      | 57 (27.8)                  |                    |                         |
| Previous vaginal birth                   |                            |                    |                          |
| Yes (n=177)                              | 53 (29.9)                  | 0.47               | Variable removed        |
| No (n=56)                                | 14 (25.0)                  |                    |                         |
| Operator                                 |                            |                    |                          |
| In-training resident (n=192)             | 55 (28.6)                  | 0.93               | Variable removed        |
| Staff member (n=41)                      | 12 (29.2)                  |                    |                         |

Abbreviations: AUB, abnormal uterine bleeding; OR, odds ratio; CI, confidence interval.

Discussion

Insufficient endometrial tissue following endometrial sampling is widely observed in gynecological practices and can often lead to critical problems in clinical management. In this study, we evaluated the magnitude of and factors related to the collection of insufficient endometrial samples. The incidence of insufficient endometrial tissue was 28.8% (95% CI=23.0–35.0). A multivariate analysis revealed that postmenopausal status and endometrial thickness less than 8 mm were significant predictors for insufficient endometrial tissue.

The rate of insufficient tissue following endometrial sampling varies widely in the literature, from 6% to 33%. This may be due to secondary differences across the studies in terms of sample characteristics, in particular the proportions of menopausal patients, diagnostic criteria for insufficient tissue, and types of provider.8,9,11,12 Bakour et al8 reported that menopausal status was associated with insufficient specimen (OR=4.49, 95% CI=2.49–8.09). Williams et al10 found that
the rate of adequate tissue after endometrial sampling was
52% in postmenopausal women, but 92% in those who were
premenopausal. Recently, Xie et al11 confirmed that meno-
pausal status increased the risk of sample inadequacy follow-
ing endometrial sampling (OR=8.324, 95% CI=3.79–18.24).
The higher rate of insufficient tissue after endometrial sam-
plying among postmenopausal women may be due to atrophic
changes to the endometrium, which could, in turn, result in
less tissue being obtained.16

Previous studies have consistently reported an association
between endometrial thickness as evaluated by transvaginal
ultrasonography and risk of insufficient tissue after sub-
sequent endometrial sampling.8,11,17,18 However, the cutoff
threshold for endometrial thickness has varied across these
studies. Bakour et al6 found that endometrial thickness above
5 mm decreased the risk of insufficient sample on endometri-
al biopsy (OR=0.19, 95% CI=0.07–0.53). Elsandabesee and
Greenwood19 stated that there was only a 27% probability
of getting an adequate endometrial sample in women with
an endometrial thickness of less than 5 mm, compared to
60% among those with an endometrial thickness ≥5 mm.
Visser et al17 showed that endometrial thickness >12 mm
independently decreased the chance of insufficient sampling
(OR=0.26, 95% CI=0.09–0.76). Recently, Xie et al11 found
that an endometrial thickness of less than 7 mm was sig-
ificantly associated with sample inadequacy (OR=6.74,
95% CI=3.04–14.90). In our study, an endometrial thick-
ness of less than 8 mm according to transvaginal ultrasound
increased the risk of inadequate endometrial sampling
(OR=3.91, 95% CI=1.49–10.21).

There has been controversy about the association between
advanced age and an adequacy of endometrial samples.8,17,19,20
Previous studies noted an approximate 4% increase in the
rate of insufficient tissue after endometrial sampling with
each year of increasing age.8,17 Kandil et al19 demonstrated
that there was a significantly higher rate of insufficient
samples among women 60 years and older compared to
younger women (14.6% vs 5.8%). However, Adambekov
et al18 showed that age over 55 years was not associated with
inadequate tissue following endometrial sampling (OR=1.95,
95% CI=0.72–5.30). In this study, women 60 years of age
and older had a higher rate of insufficient endometrial tissue
compared to younger women (45.0% vs 27.2%). However,
this difference was not statistically significant according to
multivariate analysis (OR=0.95, 95% CI=0.32–2.77).

This study found no association between insufficient
samples and other reproductive characteristics, such as
nulliparity, previous vaginal birth, or current hormonal use,
which contrast with the results of previous studies.11,17,19
Prolonged use of oral combined contraceptive pills, depo-
medoxyprogesterone acetate, or menopausal hormonal
therapy can cause endometrial atrophy, thus resulting in
higher risk of insufficient samples following endometrial
sampling.21 Additionally, Adambekov et al18 reported an
increased risk of insufficient tissue after endometrial sam-
plying among women who presented with postmeno-
pausal bleeding when compared to those who underwent
endometrial sampling with other indications (OR=7.41,
95% CI=2.27–24.14).19 However, there was no significant
difference between women who had abnormal uterine bleed-
ing and those who presented with other symptoms in terms
of the risk of insufficient tissue after endometrial sampling
in the present study.

It is debatable whether or not the training level of
operator affects the rate of insufficient tissue after endome-
trial sampling. A recent study showed that the procedure
being conducted by a nonphysician provider was associ-
ated with an increased risk of sampling failure (OR=9.15,
95% CI=2.49–33.69).19 However, this association was not
reaffirmed by other studies.11,17 In this study, there was no
significant association between the type of operator (in-
training residents vs staff members) and risk of insufficient
endometrial tissue.

In this study, 35 (52.2%) of the 67 women with insuf-
cient endometrial tissue samples did not undergo further
endometrial evaluation. Previous studies indicated that
approximately 6%–7% of women with insufficient endome-
trial tissue samples were subsequently found to have atypical
endometrial hyperplasia or endometrial cancer.12 These
findings highlight the importance of further evaluation to
ensure exclusion of significant endometrial pathology among
women with insufficient endometrial samples.

The main limitation of this study is that, due to its ret-
rospective nature, some information was unavailable. This
included sonographic results regarding endometrial thick-
ness and uterine volume, as well as previous failed attempts
at endometrial sampling, all of which have been shown to
be important factors associated with the risk of insuf-
cient endometrial tissue samples. In addition, the reasons
for not performing further diagnostic procedures among
women with insufficient samples are unknown. Despite this
limitation, the primary strength of the study was that it was
conducted in a single institution, where the pathological
specimens are reviewed by experienced gynecologic patholo-
gists, and criteria for evaluating the adequacy of tissue are
standardized.
Conclusion
The incidence of insufficient tissue following endometrial sampling was 28.8% (23.0–35.0). Significant independent factors associated with an increased risk of insufficient tissue were menopausal status and endometrial thickness less than 8 mm.

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Disclosure
The authors report no conflicts of interest in this work.

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