Application of a Novel Vibrating Device for Fine-Needle Aspiration Cytology

Mami Morita¹, Ryusuke Hori¹*, Shintaro Fujimura², Yusuke Okanoue², Tsuyoshi Kojima², Koichi Omori³ and Kazuhiko Shoji³

Abstract

Objective: Fine-needle aspiration cytology (FNAC) is a valuable diagnostic technique. However, the procedure involves back-and-forth motions of a needle within a mass, which can lead to unexpected complications. We have developed a novel device and sampling techniques that use vibration and rotation instead of back-and-forth motions.

Methods: The new device consists of a vibrating motor fixed to the stopper of a 5 ml syringe with its piston. A 22-gauge needle attached to the 5 ml syringe is used for FNAC. Samples were obtained from resected specimens using the following four procedures: suction only; suction and vibration for 5 seconds; suction and 180° rotation of the syringe; and suction, vibration for 5 seconds, and 180° rotation of the syringe. Samples were also obtained using the conventional technique. The numbers of well-visualized follicular groups on glass slides were counted to compare the amounts of cellular material obtained using the five different procedures. Next, 415 patients with thyroid nodules underwent ultrasound-guided FNAC to evaluate the rate of inadequacy.

Results: Sufficient amounts of material were obtained from resected specimens using suction, vibration, and rotation, and using the conventional technique. Inadequate thyroid FNAC material was obtained in 12.3% of cases.

Conclusions: The new device and sampling techniques for thyroid FNAC collected sufficient amounts of adequate material and allowed safe and precise control of the device. Our device and sampling techniques are expected to be widely used not only for thyroid FNAC sampling but also for sampling from other anatomical sites.

Keywords
Fine-needle aspiration cytology; Vibration; Rotation; Back-and-forth motion; 5 ml syringe; Inadequacy rate; Bethesda system

Introduction

Due to its simplicity and accuracy combined with minimal discomfort and complications, fine-needle aspiration (FNAC) is a valuable diagnostic technique to evaluate suspicious masses in the head and neck region [1]. Recently, ultrasound has become widely used in conjunction with FNAC, as ultrasound guidance improves the ability to obtain a cytologic diagnosis from small lesions as well as those that are inaccessible to FNAC by palpation [2]. FNAC is usually performed using a 21- to 25-gauge needle attached to a 10- or 20 ml syringe, with or without a pistol-type holder (Figure 1). Samples are obtained by back-and-forth cutting motions of the needle within a mass, with vigorous suctioning. Although this technique is relatively straightforward to perform, there are some serious safety concerns to consider. First, these back-and-forth motions within lesions are sometimes associated with unexpected complications [3]. Several important structures and organs are present in the head and neck, including blood vessels, nerves, the trachea, and the thyroid gland. If the tip of the needle accidentally contacts these structures, complications may result. Second, a 10- or 20 ml syringe equipped with a holder is so large that it is difficult to control precisely under strong suction in the range of 10–20 ml negative pressure, particularly when acquiring samples from a small mass, so at least two people are needed to obtain samples safely during FNAC. Moreover, some patients are afraid of large syringes and needles, leading to excessive anxiety when the needle penetrates the skin overlying a lesion. Therefore, we developed a novel device that is smaller than the usual 10- or 20 ml syringe with its holder, and a new sampling technique that uses vibration and rotation to obtain samples instead of back-and-forth motions. Using this new device and novel techniques, we were able to collect sufficient samples and perform the procedure safely with precise control. In this paper, we introduce the design of our new device and sampling techniques, and describe their utility. To evaluate their efficacy, the amounts of cellular material obtained from surgically removed thyroid specimens were compared among five procedures: four different procedures with our new device and sampling techniques, and the conventional technique of back-and-forth motions using a 20 ml syringe equipped with a pistol-type holder. We also evaluated whether the thyroid FNAC materials obtained using our new device and sampling techniques in a consecutive series of patients were diagnostic or adequate based on the Bethesda system for reporting thyroid cytopathology [4,5].

Materials and Methods

Design of the new device

The new device consists of a vibrating motor attached to the stopper of a 5 ml syringe and its piston. The vibrating motor is disk shaped, 12 mm in diameter, and 3.4 mm thick, and vibrates at a frequency of 220 Hz. The design of the stopper made from aluminum plate is shown in Figure 2. It is approximately 30 × 30 × 1.5 mm, with a protrusion next to the 5 ml syringe and a slit next to the piston. When the stopper is attached to a 5 ml syringe and its piston, negative pressure with 1 ml suction is maintained (Figure 3). As the vibrating motor is glued to the stopper, this device is small and weighs only 13 g. The vibrating motor and aluminum plate are widely available, so anyone can make this device. Optimization of the frequency of vibration was not assessed in this study.

Experimental materials and patients

From 2009 to 2011, a consecutive series of patients underwent thyroid ultrasound and ultrasound-guided FNAC at Tenri Hospital.
Six of these patients (four females and two males) who were diagnosed with papillary thyroid carcinoma underwent hemi-thyroidectomy surgery. Within 1 h after resection, the six resected specimens (which were not preserved in formalin) were used to examine amounts of cellular material obtained by the various methods. To evaluate whether thyroid FNAC samples obtained using our new device and sampling techniques were diagnostic, 415 patients (316 females and 99 males; mean age, 61.5 ± 14.1 years) who had undergone ultrasound-guided FNAC for thyroid nodules ≥ 3 mm in diameter, without thyroid cysts, were enrolled in this study. Specifically, the thyroid nodules were 3-62 mm in size. All patients provided informed consent prior to ultrasound-guided FNAC and hemi-thyroidectomy. We compared the results obtained using our device and sampling techniques with those of 588 patients with thyroid nodules who underwent ultrasound-guided FNAC using the conventional technique in 2007–2009. This study was approved by the institutional review board of Tenri Hospital.

**Sampling techniques**

A 22-gauge needle attached to a 5 ml syringe was used for FNAC. To perform sampling on the resected specimens, the needle was passed through the capsule of specimens and then inserted into the cancerous sites by palpation. The device, consisting of a stopper glued to a vibrating motor, was attached to the syringe and its piston. Negative pressure was obtained by applying 1 ml suction. Subsequently, samples were obtained using four different procedures: suction only; suction and vibration for 5 seconds; suction and 180° rotation of the syringe; and suction, vibration for 5 seconds, and 180° rotation of the syringe. All suction, vibration, and rotation were halted before the needle was removed. For the conventional technique, samples were obtained by back-and-forth cutting motions in the cancer site with vigorous suction using a 20 ml disposable syringe equipped with a pistol-type holder. These five procedures were performed three times per specimen.
In our study, ultrasound-guided FNAC procedures were performed with patients in a supine position and their neck slightly extended by a pillow placed under their shoulders. The nodule was localized by ultrasound scanning (General Electric LOGIQ 500). Local anesthesia was not applied routinely, but was given (as 0.5% lidocaine) to patients with needle fear and/or excessive anxiety. During FNAC, a needle and attached 5 ml syringe were connected to the protrusion of the new device. The needle then penetrated the overlying skin and was inserted into the target nodule (Figure 4A). One hand was used to position the piston against the slit in the device and then 1 mL negative pressure was applied (Figure 4B). The vibrating motor was then switched on with the other hand (Figure 4C), and samples were obtained by the various techniques using suction, vibration for 5 seconds, and 180° rotation of the syringe (Figure 4D). All suction, vibration, and rotation were halted before the needle was removed. Importantly, no back-and-forth motions were used.

Cytological assessment and criteria for adequacy

After aspiration from the six surgically removed thyroid specimens, samples were suspended in a small amount of physiological saline. The cell suspension was then centrifuged at 1,000 rpm for 3 min (AutoSmear). Next, the cells were fixed on glass slides using AutoSmear with 95% ethanol fixation and stained with Papanicolaou. The numbers of well-preserved and well-stained glass slides using AutoSmear with 95% ethanol fixation and stained with Papanicolaou. The numbers of well-preserved and well-stained follicular cells, with at least 10 cells per group, preferably on a single slide, were counted with Papanicolaou. The numbers of well-preserved and well-stained glass slides using AutoSmear with 95% ethanol fixation and stained with Papanicolaou. The numbers of well-preserved and well-stained follicular cells, with at least 10 cells per group, preferably on a single slide, were counted with Papanicolaou. The numbers of well-preserved and well-stained glass slides using AutoSmear with 95% ethanol fixation and stained with Papanicolaou. The numbers of well-preserved and well-stained follicular cells, with at least 10 cells per group, preferably on a single slide, were counted with Papanicolaou.

Significant differences were also identified between inadequate samples of 415 patients obtained using our new device and sampling techniques and those obtained using the conventional technique using the Chi-square test.

Results

The mean numbers of follicular groups (containing at least 10 cells per group) obtained with suction only, suction and vibration for 5 seconds, and suction and 180° rotation of the syringe were 10.7, 16.0, and 19.9, respectively. The mean numbers of follicular groups obtained using our techniques and the conventional technique were 86.8 and 113.1, respectively; the difference between the two was not statistically significant (p=0.919). Variance was not equal among the five groups as determined by Bartlett’s test. The overall differences among the five groups were significant (p=0.003). The numbers of follicular groups obtained using suction, vibration for 5 seconds, and 180° rotation of the syringe were significantly larger than with suction only (p=0.036; Figure 5). Significantly larger numbers were also obtained using the conventional technique than with suction only, suction and vibration for 5 seconds, and suction and 180° rotation of the syringe (p=0.014, 0.014, and 0.018, respectively; Figure 5).

The thyroid FNAC materials obtained using our new device and sampling techniques were considered inadequate in 12.3% of 415 patients and in 16.9% of 130 patients with nodules ≤ 10 mm. There were no samples with extensive obscuring blood that hindered the evaluation of the follicular cells. No serious complications occurred in any of the 415 patients. On the other hand, using the conventional technique, the FNAC materials obtained from 18.2% of 588 patients and 35.2% of 88 patients with nodules ≤ 10 mm were considered inadequate. Significant differences were identified between inadequate samples of 415 patients obtained using our new device and sampling techniques and those of 588 patients obtained using the conventional technique (p<0.001). Significant differences were also identified between inadequate samples of 130 patients with nodules 10 mm obtained using our new device and sampling techniques and those of 88 patients with nodules ≤ 10 mm obtained using the conventional technique (p<0.001).

Figure 4: View during FNAC. (A) The needle attached to a 5 ml syringe, against which the protrusion on the device is placed, is passed through the overlying skin and inserted into a target nodule. (B) One hand is used to position the piston against the slit in the device (arrow) and suction with 1 ml negative pressure is applied. (C) The vibrating motor is turned on with the other hand. (D) Samples are obtained immediately using suction, vibration for 5 seconds, and 180° rotation of the syringe. Asterisks indicate switching of the vibrating motor fixed to the stopper.
attached to a 5 ml syringe and its piston, the instrument is smaller
samples (Figure 6). Since the stopper fixed to the vibrating motor is
syringe under 1 mL of suction is 0.85 atm, which is sufficient to obtain
is required in the needle. The calculated negative pressure in a 5 ml
sufficient vacuum because only a very small volume of dead space
stopper of the syringe and its piston are designed to maintain negative
perform FNAC did not lead to any serious complications. The
samples are not penetrated. In this study, using our new device
used instead of back-and-forth motions, ensuring that dorsal sites
is considered a safe instrument because vibration and rotation are
precisely under strong suction with 10-20 ml negative pressure.
Moreover, the 10- or 20 ml syringes that are usually used for FNAC
and-forth motion of the sampling needle within tumor masses.

One cause of these unexpected complications is likely the back-
and-forth motion of the sampling needle within tumor masses.
Moreover, the 10- or 20 ml syringes that are usually used for FNAC
are very large, and it is difficult to control the syringe and its piston
precisely under strong suction with 10-20 ml negative pressure.
Consequently, this technique can result in penetrating the dorsal
sites of nodules and damage to surrounding structures, particularly
when samples are acquired from small nodules. Our novel device
is considered a safe instrument because vibration and rotation are
used instead of back-and-forth motions, ensuring that dorsal sites
of nodules are not penetrated. In this study, using our new device
to perform FNAC did not lead to any serious complications. The
stopper of the syringe and its piston are designed to maintain negative
pressure with 1 mL of suction. Based on Boyle’s Law, this creates
sufficient vacuum because only a very small volume of dead space
is required in the needle. The calculated negative pressure in a 5 ml
syringe under 1 mL of suction is 0.85 atm, which is sufficient to obtain
samples (Figure 6). Since the stopper fixed to the vibrating motor is
attached to a 5 ml syringe and its piston, the instrument is smaller
and lighter than a 20 ml syringe equipped with a holder. This allows
our novel device to be manipulated more easily even by small hands

and provides more control during tissue sampling. In particular, our
new device is useful when sampling is performed in confined spaces
within the compact anatomy of the head and neck.

An FNAC sample of the thyroid is considered adequate for evaluation
if it contains a minimum of six groups of follicular cells, and inadequate samples are reported as non-diagnostic or
unsatisfactory [4,5]. To demonstrate the efficacy of our new device
and sampling techniques that use vibration and rotation instead
of back-and-forth motions, we compared the amounts of cellular
material obtained in thyroid specimens using five procedures, and
evaluated whether the thyroid FNAC samples were adequate. The
samples obtained using all five procedures contained more than six
groups of follicular cells. Although the numbers of follicular groups
obtained using suction only, suction and vibration, or suction and
rotation were smaller, sufficient amounts of material were obtained
with suction, vibration for 5 seconds, and 180° rotation of the syringe,
as well as using the conventional technique. There were also no
significant differences between the numbers of groups obtained using
our techniques versus the conventional technique. The obliqueness
of the needle tip may affect the acquisition of follicular groups during
these three procedures. Thyroid cells may be detached by the oblique
cutting edge of the needle tip and be aspirated into its lumen when
the combined sampling techniques of suction, vibration, and rotation
are applied; therefore, material may be obtained as sufficiently as with
the conventional technique. Furthermore, 12.3% of samples were
inadequate using our new device and sampling techniques compared
with 18.2% of samples obtained using the conventional technique, and
the difference between these two sampling groups was significant.
By contrast, previous studies have reported FNAC inadequacy rates for
thyroid tumor samples ranging from 13.4% to 33% [9-11]. The results
demonstrate the efficacy of our new device and sampling techniques.
In this study, we did not optimize suction or vibration time, but 5 secs
are sufficiently short that blood is not aspirated and this time did not
decrease the percentage of adequate samples.

Our novel device is considered a safe instrument that is easy to
control precisely. Sufficient amounts of material were obtained using
suction, vibration and rotation, as well as using the conventional
technique. Therefore, this device and sampling technique are
expected to be widely used. Furthermore, use of this technique at
other anatomical sites is also anticipated. Future studies should
include assessments of the efficacy of this device and these sampling
techniques when sampling parotid glands, submandibular glands,
lymphatic nodes, and other head and neck lesion sites.

Discussion

FNAC is a simple, accurate, and cost-effective technique that is
the best choice for diagnostic evaluation of head and neck masses.
Furthermore, it is usually a safe diagnostic procedure with rare cases
of serious complications. The most common complications, such as
local pain or discomfort and minor hematomas, occur post-FNAC.
However, serious complications following FNAC have been reported,
including massive hematomas, carotid hematomas, infections,
recurrent laryngeal nerve injury, vasovagal reactions due to severe
pain or anxiety, transient dysphagia, esophageal puncture, tracheal
puncture, and tumor dissemination. Furthermore, complications
may be underestimated due to the reluctance of those performing the
aspiration to report them [6-8].

The mean numbers of follicular groups obtained using suction
only, suction + vibration, suction + rotation, suction + vibration + rotation,
and the conventional technique. The mean numbers of follicular groups
obtained using suction only; suction and vibration, and suction and
rotation were 10.7, 16.0, and 19.9, respectively. The mean numbers
obtained using our techniques versus the conventional technique were
86.8 and 113.1, respectively. Significantly larger numbers of follicular
groups were obtained using suction, vibration, and rotation than by using
suction only (p = 0.036). The numbers obtained using the conventional
technique were also significantly larger than with suction only, suction
and vibration, and suction and rotation (p = 0.014, 0.014, and 0.018,
respectively).

Figure 5: The mean numbers of follicular groups obtained using suction
only, suction + vibration, suction + rotation, suction + vibration + rotation,
and the conventional technique. The mean numbers of follicular groups
obtained using suction only; suction and vibration, and suction and
rotation were 10.7, 16.0, and 19.9, respectively. The mean numbers
obtained using our techniques versus the conventional technique were
86.8 and 113.1, respectively. Significantly larger numbers of follicular
groups were obtained using suction, vibration, and rotation than by using
suction only (p = 0.036). The numbers obtained using the conventional
technique were also significantly larger than with suction only, suction
and vibration, and suction and rotation (p = 0.014, 0.014, and 0.018,
respectively).

One cause of these unexpected complications is likely the back-
and-forth motion of the sampling needle within tumor masses.
Moreover, the 10- or 20 ml syringes that are usually used for FNAC
are very large, and it is difficult to control the syringe and its piston
precisely under strong suction with 10-20 ml negative pressure.
Consequently, this technique can result in penetrating the dorsal
sites of nodules and damage to surrounding structures, particularly
when samples are acquired from small nodules. Our novel device
is considered a safe instrument because vibration and rotation are
used instead of back-and-forth motions, ensuring that dorsal sites
of nodules are not penetrated. In this study, using our new device
to perform FNAC did not lead to any serious complications. The
stopper of the syringe and its piston are designed to maintain negative
pressure with 1 mL of suction. Based on Boyle’s Law, this creates
sufficient vacuum because only a very small volume of dead space
is required in the needle. The calculated negative pressure in a 5 ml
syringe under 1 mL of suction is 0.85 atm, which is sufficient to obtain
samples (Figure 6). Since the stopper fixed to the vibrating motor is
attached to a 5 ml syringe and its piston, the instrument is smaller
and lighter than a 20 ml syringe equipped with a holder. This allows
our novel device to be manipulated more easily even by small hands

and provides more control during tissue sampling. In particular, our
new device is useful when sampling is performed in confined spaces
within the compact anatomy of the head and neck.

An FNAC sample of the thyroid is considered adequate for evaluation
if it contains a minimum of six groups of follicular cells, and inadequate samples are reported as non-diagnostic or
unsatisfactory [4,5]. To demonstrate the efficacy of our new device
and sampling techniques that use vibration and rotation instead
of back-and-forth motions, we compared the amounts of cellular
material obtained in thyroid specimens using five procedures, and
evaluated whether the thyroid FNAC samples were adequate. The
samples obtained using all five procedures contained more than six
groups of follicular cells. Although the numbers of follicular groups
obtained using suction only, suction and vibration, or suction and
rotation were smaller, sufficient amounts of material were obtained
with suction, vibration for 5 seconds, and 180° rotation of the syringe,
as well as using the conventional technique. There were also no
significant differences between the numbers of groups obtained using
our techniques versus the conventional technique. The obliqueness
of the needle tip may affect the acquisition of follicular groups during
these three procedures. Thyroid cells may be detached by the oblique
cutting edge of the needle tip and be aspirated into its lumen when
the combined sampling techniques of suction, vibration, and rotation
are applied; therefore, material may be obtained as sufficiently as with
the conventional technique. Furthermore, 12.3% of samples were
inadequate using our new device and sampling techniques compared
with 18.2% of samples obtained using the conventional technique, and
the difference between these two sampling groups was significant.
By contrast, previous studies have reported FNAC inadequacy rates for
thyroid tumor samples ranging from 13.4% to 33% [9-11]. The results
demonstrate the efficacy of our new device and sampling techniques.
In this study, we did not optimize suction or vibration time, but 5 secs
are sufficiently short that blood is not aspirated and this time did not
decrease the percentage of adequate samples.

Our novel device is considered a safe instrument that is easy to
control precisely. Sufficient amounts of material were obtained using
suction, vibration and rotation, as well as using the conventional
technique. Therefore, this device and sampling technique are
expected to be widely used. Furthermore, use of this technique at
other anatomical sites is also anticipated. Future studies should
include assessments of the efficacy of this device and these sampling
techniques when sampling parotid glands, submandibular glands,
lymphatic nodes, and other head and neck lesion sites.
Conclusion

In this study, we introduced a novel device and new sampling techniques for thyroid FNAC. The new procedure uses vibration and rotation instead of back-and-forth motions of the needle to obtain samples. This allows collection of sufficient amounts of diagnostic material and enables safe and precise control of the device. Our new device and sampling techniques are expected to be widely used not only for thyroid FNAC samples but also at other anatomical sites.

References


Submit your next manuscript and get advantages of SciTechnol submissions

- 80 Journals
- 21 Day rapid review process
- 3000 Editorial team
- 5 Million readers
- More than 5000 Quality and quick review processing through Editorial Manager System

Submit your next manuscript at www.scitechnol.com/submission