Appropriate Concentration of Acetic Acid for the Preservation of Fresh Cadaveric Brains to be Used in Surgical Training: A Preliminary Study

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SUMMARY: A small amount of acetic acid (AA), a common preservative, has been shown to increase contamination in cadaveric tissue, while larger concentrations can lead to the tissue becoming hard, especially in fresh brains. This study attempted to optimize the concentration of AA to be used in the cranial cavity in order to produce the most realistic consistency and color. Six adult cadaveric heads were preserved with descending glacial AA at concentrations of 98.5\%, 80\%, 60\%, 40\%, 20\%, and 10\%. The samples were kept at 5°C for 14 days. The brain cortex was then dissected with a suction tube and forceps to reveal the underlying brain tissue for inspection. Color change, cortical firmness, pia mater stickiness, and participant satisfaction were evaluated. The color of the brains in all concentrations was slightly yellow. However, the temporal area of the brain preserved using 20\% AA was significantly more pink. The pia mater of the brain cortex of all samples was firm and difficult to pry apart, with the firmest consistency being in the brain tissue preserved using 98.5\% AA. The brain tissue in all samples had a liquid-like consistency. The brains preserved in AA at a concentration greater than 60\% yielded higher satisfaction scores. We conclude that acetic acid has a role in brain preservation for skull base surgery training and recommend AA concentrations higher than 60\% for maximal participants satisfaction.

KEY WORDS: Acetic acid; Concentration; Preservation; Fresh cadaver; Brain.

INTRODUCTION

Acetic acid (AA) is a preservative commonly used to prevent bacterial and fungal growth in wound dressings and in the food industry (Febriana \textit{et al.}, 2012; Shukla \textit{et al.}, 2017). It has been used in post-mortem cadavers to preserve the skin color, softness, and joint mobility for fresh cadaveric surgery demonstrations (Brenner, 2014; Varlet \textit{et al.}, 2018). However, using a small amount of AA has been shown to increase contaminations in cadaveric tissue (Trcek \textit{et al.}, 2015), while larger concentrations can lead to the tissue becoming hard, especially in fresh brains.

A tremendous amount of skill and attention to detail is required to adequately address the risks that accompany surgery performed on the nervous system, which means that rigorous training is necessary before a resident can become a competent neurosurgeon. Neurosurgery residency training in Thailand takes at least of 5 years to complete. The Royal College of Surgeons of Thailand (2018) recently declared that certain milestones must be reached before this training is complete, with the ability to perform a number of specific procedures being one of them. One of the many strengths of Khon Kaen University (KCU) neurosurgery residency program is the availability of cadavers, which are donated by local residents through the Department of Anatomy. The Neurosurgery Division of the Faculty of Surgery has been organizing triennial conferences with live instruction and hands-on experience using the heads of these cadavers for the past 5 years for interested residents and neurosurgeons from all over the country. Some of the benefits of practicing on cadavers include the opportunity to familiarize oneself.

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with surgical tools, the ability to become used to the configurations of the cranial vasculature, as well as the chance to feel more comfortable and confident navigating through the complicated skull base. However, there are still problems encountered with using the cadaveric brains that should be addressed. A fixative solution consisting of a mix of glycerin and formalin is routinely used for cadaver preservation at KKU. However, the glycerin makes the brain overly pulpy, rendering the samples impossible to dissect. When preserved with formalin, the brains become hard and inelastic, characteristics unsuitable for dissection. In addition to its distinct, pungent smell and color-altering properties, formalin has been classified as an occupational hazard for health personnel. It has been reported to cause contact dermatitis, upper airway inflammation, asthma, and is implicated in nasopharyngeal cancer, sinonasal cancer, and lymphohematopoietic cancer (Nelson et al., 1986; Raja & Sultana, 2012; Friis et al., 2014; Pontén & Bruze, 2015; Davies, 2016; Kim et al., 2016). Over the past 100 years, various substances have been reported as being used preservation of the cadaveric tissue. The method of preservation mostly involves bathing the whole cadaver in a certain solution, with an unquantifiable portion of said solution entering the brain. Such solutions include fompeizole, phenoxyethanol, formaldehyde, formalin, and diethylene glycol, each of which have several advantages and disadvantages (Brenner).

Acetic acid’s use in the preservation of the post-mortem cadaver has been documented as early as in the 19th century by French chemist Jean-Nicholas Gannal (Brenner). In particular, glacial acetic acid has been shown to preserve skin color, skin softness, and joint mobility in the cadaveric arm (Tschernezky, 1984). The aim of this study was to investigate the optimal concentration of acetic acid to be poured into the cadaver’s cranial cavity in order to give fresh cadaveric brain tissue the most realistic consistency and color.

MATERIAL AND METHOD

Six fresh cadaveric heads were obtained from the KKU body donation program, in accordance with the standard procedures of the Khon Kaen University Faculty of Medicine’s Department of Anatomy. All of the cadaveric heads were of adults aged more than 18 years and deceased not longer than 3 days, and were stored in a cool preserving chamber. There were no exclusion criteria. Since this was a pilot study, there were yet no standards as to the optimum number of samples. A Burr hole was made at the vertex position in each cadaveric head. The heads were then placed upside down in a 5 °C refrigerator in order to drain all residual fluids. After 48 h, industrial grade 98.5 % glacial acetic acid was diluted with tap water to 80 %, 60 %, 40 %, 20 %, and 10 % concentrations, and each solution (including one that was undiluted) was poured into a separate head until full. The heads were then placed upright and stored in a 5 °C refrigerator. After another 48 hours, the level of the acetic acid at the Burr hole was examined, as it tends to decrease, presumably because it flows into and fills up smaller spaces in the skull cavity. Acetic acid at the same concentration was then added fill the head up to the Burr hole entrance. The cadaveric heads were then left in the 5 °C refrigerator for another 14 days. A traumatic flap was used to access the brain tissue, which was then examined for color and pia mater firmness by ten participants including neurosurgical residents and neurosurgeons. The brain cortex was then dissected with a suction tube and forceps to reveal the underlying brain tissue for inspection. Color, cortical firmness (on a scale of 1-5: 5 = very hard, 4 = fairly hard, 3 = firm, 2 = fairly soft, 1 = very soft), pial stickiness, and participant satisfaction (on a scale of 1-5: 1 = Not at All Satisfied, 2 = Partly Satisfied, 3= Satisfied, 4 = More than Satisfied, 5 =Very Satisfied) were evaluated.

RESULTS

The summary of our findings is shown in Table I. The color of all of the brains was more or less retained, appearing only slightly yellower. The brain that received 20 % acetic acid had a subarachnoid hemorrhage, with the temporal area appearing significantly more pink. All participants agreed that the pial mater of the brain cortex of all samples was firmed and difficult to pry apart. The cadaveric head preserved with 98.5 % acetic had the firmest consistency, with firmness decreasing with the acid concentration (Table I). In addition, it was observed that firmness was not distributed equally in the brains preserved with concentrations lower than 60 % (Table I). Apart from the brain preserved with 40 % solution, all specimens felt firmer near the temporal area. After the pia mater was separated with a scalpel, the brain tissue in all samples was observed to have a liquid-like consistency.

DISCUSSION

Acetic acid has been used for preservation of the post-mortem cadavers early as in the 19th century (Brenner). However, this study was the first to preliminarily
Table I. Summary of acetic acid concentrations with satisfaction parameters.

| Cadaveric heads (H) | H1  | H2  | H3  | H4  | H5  | H6  |
|---------------------|-----|-----|-----|-----|-----|-----|
| Acetic acid concentration (%) | 98.5 % | 80 % | 60 % | 40 % | 20 % | 10 % |
| Initial acetic acid poured (ml) | 130 | 90 | 100 | 110 | 50 | 160 |
| Additional acetic acid at 48 hr (ml) | 130 | 120 | 125 | 130 | 80 | 200 |
| Color at 14 days | Slightly yellow | Slightly yellow | Slightly yellow | Slightly yellow, pink at temporal lobe | Slightly yellow | Slightly yellow |
| Firmness at 14 days (Scales: 1-5) | 3 | 3 | 3 | 3 | 2 | 1 |
| Pia mater stickiness | Sticky | Sticky | Sticky | Sticky | Sticky | Sticky |
| Sylvian subarachnoid hemorrhage | NA | NA | NA | NA | NA | NA |

Note: Cortical firmness (5 = Very hard, 4 = fairly hard, 3 = Firm, 2 = fairly soft, 1 = Very soft). Participant satisfactions (1 = Not at All Satisfied, 2 = Partly Satisfied, 3= Satisfied, 4 = More than Satisfied, 5 =Very Satisfied).

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RESUMEN: Se ha demostrado que una pequeña cantidad de ácido acético (AA), un preservante común, aumenta la contaminación en el tejido del cadáver, mientras que mayores concentraciones pueden endurecer el tejido, particularmente en cerebros frescos. Este estudio intentó optimizar la concentración de AA en la cavidad craneal para producir una consistencia y coloración cerebral más cercanos a la realidad. Seis cabezas cadáveres adultas se conservaron con AA glacial descendente en concentraciones de 98.5%, 80%, 60%, 40%, 20% y 10%. Las muestras se mantuvieron a 5 °C durante 14 días. Luego se diseccionó la corteza cerebral con un tubo de succión y pinzas para observar e inspeccionar el tejido cerebral subyacente. Se evaluaron el color, la firmeza cortical, la viscosidad y adherencia de la piamadre y la reacción de los participantes ante esta conservación. El color de los cerebros en todas las concentraciones fue ligeramente amarillo. Sin embargo, el área temporal del cerebro, conservado con un 20% de AA, fue significativamente más rosada. La piamadre de la corteza de todas las muestras fue de consistencia firme y difícil de separar; una mayor resistencia se observó en el tejido cerebral preservado con 98.5% de AA. La consistencia del tejido cerebral en todas las muestras era líquida. Los cerebros conservados en AA a una concentración superior al 60% recibieron puntuaciones de satisfacción más elevadas. Concluimos que el ácido acético desempeña un papel en la conservación del cerebro, permitiendo el entrenamiento en cirugía de base de cráneo, por lo que recomendamos concentraciones de AA superiores al 60% para una satisfacción máxima por parte de los participantes.

PALABRAS CLAVE: Ácido acético; Concentración; Conservación; Cerebro; Cadáver fresco.
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