Abstract

Detection and quantification of hemoglobin in skeletal or dried remains could potentially have versatile utilization with the forensic field. However, the majority of studies within this area focus on the identification of hemoglobin and its derivatives rather than determining the relative amount of protein present. This research focuses on the use of leuco crystal violet with Ultraviolet/Visible spectroscopy and applying cyclic voltammetry for the determination and quantification of hemoglobin in a tooth and femur sample. Using leuco crystal violet, possible interferences were also investigated, and it was determined that ferrous iron samples appeared to actually mask the overall reaction, while glycine had no obvious effect. Albumin showed no apparent interference, while myoglobin produced a very faint green color that absorbed at a different wavelength, so this species produced minimal interference. Extractions of hemoglobin from a femur bone using 0.5 M hydrochloric acid and from a tooth using 6.0 M urea all yielded the violet product when reacted with leuco crystal violet. Using the measured absorbance of each positive extraction, the corresponding amount of hemoglobin in each sample was determined. Hemoglobin standards and extraction samples from the tooth and femur bone were also analyzed by the electrochemical method, cyclic voltammetry. Hemoglobin levels in the tooth and femur were found to be in the micromolar range in both the spectroscopic and electrochemical analysis, and these levels are consistent with a decrease in hemoglobin concentration post-extraction and post-mortem.

Keywords: Hemoglobin; Spectroscopic detection; Leuco crystal violet; Electrochemical detection; Bone

Introduction

Hemoglobin (Hb) is a major oxygen-carrying molecule composed of four globular protein units and four heme moieties that consist of an iron (II) atom held in the surrounding porphyrin ring, which serves as a binding site for oxygen. The four heme groups duplicate the activity of the enzyme horseradish peroxidase [1], and this reactivity plays a critical role for the determination of Hb in this work. The detection and quantification of Hb from bone samples could have some potentially useful forensic applications such as helping to establish a time since death, identification of certain health and genetic traits, supporting classification in pink teeth cases, and possible determination of wound age in dry bone fractures. Certain blood proteins such as albumin and Hb survive well over time within bone [2]. The feasibility of finding Hb in bone samples of varying age is supported by the ability to detect Hb breakdown products in a fossilized "Tyrannosaurus rex" skeleton [3,4]. Hb is well-preserved as it crystallizes easily which may contribute to the stability of the protein, and the rigid structure of heme could possibly add to the durability of the protein [3]. These findings further support the idea that Hb can be extracted from bone, detected, and quantified.

The majority of studies within this area tend to focus on identifying Hb and its derivatives from bone samples. Methods such as mass spectrometry, immunosassays, and chemiluminescence have been used to characterize Hb and other proteins from skeletal remains with varying results. More specifically, two recent studies have focused on the identification of Hb, its derivatives, and related species in bone and teeth samples. Both Pickworth’s benzidine method and auto-fluorescence were utilized to detect Hb and its derivatives in a case study of shipwrecked remains exhibiting the phenomenon known as pink teeth [5]. Another preliminary study employed certain immunohistological methods to detect hemorrhaging for possible wound age determination in dry bone [6]. Thus far, there has been no quantitative approach to determining Hb levels in bone or teeth samples. This research is directed at determining the feasibility of such analysis by using a combinatorial approach. The two investigative methods of Ultraviolet/Visible spectroscopy and cyclic voltammetry are employed to detect and quantify Hb concentrations in a tooth and femur sample. A comparative evaluation of these results will examine any correlation and the advantages or lack thereof for both methods within this study.

UV/Vis detection is a simple, common method that has been used for the determination of Hb. Hb itself yields an absorption maximum in the wavelength range of 415-476 nm, but its molar absorptivity is fairly low with a value around 3300 M⁻¹cm⁻¹ [7,8]. A more sensitive approach for detecting Hb is to react it as a catalyst with a chromogenic reagent such as leuco crystal violet (LCV). LCV is the colorless, entirely reduced form of the triaryl methane dye, crystal violet (CV⁺) [9]. LCV is oxidized by hydrogen peroxide in the presence of a biocatalyst such as the heme moiety in Hb. The apparent molar absorptivity of crystal violet formed with LCV in the presence of hydrogen peroxide is 75,600 ± 298 M⁻¹cm⁻¹, approximately 20 times greater than the molar absorptivity of Hb itself [9-11]. LCV is investigated for this research.
over the use of other reagents such as Amido Black and Luminol for several reasons. LCV itself has a high affinity for proteins, a large molar absorbivity, is fairly safe and easy to work with, and yields extremely sensitive results in a much faster timeframe [12]. The reaction of heme with LCV should provide a more specific, longer response for Hb in bone than other colorimetric and luminescent methods. Hence, this is the preferred spectroscopic method for determining and quantifying Hb in bone for this research.

Hb does exhibit some electrochemical activity, as it contains ferrous iron at the center of each of its four porphyrin rings. The electro-activity of Hb can be difficult to measure due to its bulky, globular structure and the lengthy pathway electrons must travel from its center to the surface of the working electrode [11]. Most often the slow kinetic redox activity is analyzed through the use of modified electrodes and other enzymatic means. However, if the sample is scanned at a slow enough rate, this enables sufficient electron transfer between the protein and the electrode for a quantitative measurement to be taken. Cyclic voltammetry (CV) is a highly sensitive method that is likely to register any significant electron transfer in to the presence of Hb due to its dual sweep over a specified potential range. This analysis in combination and comparison with UV/Vis detection should result in selective, sensitive identification and quantification of Hb in bone and teeth samples.

Materials and Methods

Solution preparation

A stock Hb solution (Point Scientific, USA) of 7.447 × 10⁻³ M was prepared. From this, Hb standards of 0.25, 0.50, 1.0, 2.0, 4.0, 10.0, and 20.0 µM were prepared. The stock solution was stored at 4°C. The LCV solution (500 mL) was prepared by mixing 150.0 mL of DI water, 50.0 mL of 30% hydrogen peroxide (Fisher Scientific, USA), 10.0 g of 5-sulphosalicylic acid (Sigma-Aldrich, USA), 4.4 g of sodium acetate (Fisher Scientific, USA), and 1.1 g of LCV (99% pure, Acros, USA). The solution was then brought to volume, inverted 20 times to mix, transferred to an amber bottle, and stored at 4°C. The following extraction solutions were prepared daily to maintain a proper working calibration curve. Cyclic voltammetry (CV) is a highly sensitive method that is likely to register any significant electron transfer in to the presence of Hb due to its dual sweep over a specified potential range. This analysis in combination and comparison with UV/Vis detection should result in selective, sensitive identification and quantification of Hb in bone and teeth samples.

Bone and tooth preparation and extraction

A tooth sample (1-2 years in age post-extraction) was sawed into four different sections, and then sanded down to obtain powder from the innermost portion of the tooth. A femur bone (5-10 years in age post-mortem) was sawed in half at the head and the shaft of the bone and powder was collected from the inner cortical bone (Figure 1). Eleven samples were prepared containing ~0.25 grams of powder from the tooth, femur head, or femur shaft and 1.5 mL of one of the 3 extraction solvents. The samples were shaken well every 10 min. for 1 hour. Each sample then had 2 drops of LCV added, and was stored at 4°C overnight. The resulting samples were:

Tooth in (A) 50% 0.5 M HCl /50% DI Water, (B) 0.5 M HCl, (C) 50% 0.2 M Tris Buffer/50% DI Water, (D) 0.2 M Tris Buffer, and (K) 6.0 M Urea.

Femur shaft in (E) 6.0 M Urea, (F) 0.5 M HCl, (G) 0.2 M Tris Buffer

Femur head in (H) 6.0 M Urea, (I) 0.5 M HCl, (J) 0.2 M Tris Buffer

Preliminary tests with leuco crystal violet

As a preliminary test, solutions of varying Hb concentrations were used to determine if LCV would successfully detect the presence of Hb. Upon the addition of two drops of LCV, each standard turned a varying shade of purple, a positive indication for the presence of Hb. The standards with LCV were allowed to sit overnight at 4°C. The violet color remained in these solutions for up to 48 hours, demonstrating the stability of the Hb/LCV reaction product. However, fresh standards were prepared daily to maintain a proper working calibration curve. Hb was initially detected, so corresponding absorption data was then collected.

UV/Vis spectroscopy

Spectroscopic analysis was performed on a midrange Hb standard (1.0 µM) containing two drops of LCV. The absorption spectrum for this Hb standard as shown in Figure 2 yielded a maximum wavelength value of 590 nm. The spectrum was obtained using an Ocean Optics (USA) Red Tide USB650 Spectrometer with a 1 cm path length. Subsequent analysis for all standards and extraction results was performed on a Spectronic (UK) 20 Genesys set to a λmax of 590 nm using a 1 cm path length.

Spectroscopic measurements were taken three times for each standard (average s of ± 0.002 Abs. units) and sample (average s of ± 0.002 Abs. units), and the corresponding average values were utilized for data analysis and interpretation.

Interferences

The effect of interferences such as ferrous iron, glycine, albumin, and myoglobin possibly masking or enhancing the response for Hb in the presence of LCV was investigated. The presence of ferrous iron (Fe²⁺) in skeletal remains is often an indication of dietary behavior,
which could lead to a false response for Hb with LCV [13]. Glycine is an amino acid used for a variety of health reasons including the treatment of iron deficiency anemia [14,15]. Glycine chelates iron, so its effect on the Hb response to LCV was also studied. A few granules of ferrous ammonium sulfate, (NH₄)₂Fe(SO₄)₂·6H₂O, Fisher Scientific, USA), ferrous sulfate, (FeSO₄ Mallinckrodt Baker, USA) and glycine (Fisher Scientific, USA) were tested on the 1.0 µM Hb standard with LCV. The results of these interferences on the reaction of Hb and LCV can be seen in Figure 3. Ferrous ammonium sulfate immediately reacted to reduce the purple color to approximately one-third of the original color. Ferrous sulfate reacted to leave approximately two-thirds of the purple color. Glycine had no obvious effect on the solution. From these observations, these species do not result in a darker product with LCV, so false positives appear to be minimal, if not unlikely.

Albumin is a blood protein most abundant in human plasma, and it has a similar structure and molecular weight to hemoglobin. Myoglobin is also a blood protein that contains a heme moiety, so the LCV could possibly react and produce an interfering purple color if this species is present in a particular sample being analyzed. Solutions of 1.0 µM albumin (Protea Biosciences, USA) and myoglobin (Protea Biosciences, USA) were prepared in deionized water and treated with two drops of LCV. The results of these reactions can also be seen in Figure 3. Albumin remained a clear solution indicating it will not interfere with the reaction of Hb and LCV. Myoglobin reacted to produce a very faint green colored solution, which indicates there may be a slight interference in the presence of this particular protein, but not of a significant nature. This faint green product also absorbed at a wavelength different from the λₘₐₓ of 590 nm for LCV. A simple background correction eliminated the interference response of myoglobin with LCV.

**Electrochemistry**

Cyclic voltammetry measurements were performed using a CH Instruments (USA) 800B Electrochemical Analyzer. The three-electrode system utilized a 2 mm gold disk working electrode (CH Instruments, USA), a Ag/AgCI reference electrode (CH Instruments, USA), and a 0.5 mm Platinum auxiliary wire (Alfa Aesar, USA). Hb standards and samples were prepared in 0.05 M phosphate buffer and the appropriate extraction solution, and subsequently analyzed at a scan rate of 0.005 V/s. Scan rates faster than this value did not yield as high peak currents for each standard and sample, so the aforementioned rate was determined to be optimal.

CV measurements were run a total of three times for all standards (average s of ± 9.3 × 10⁻⁸ A) and samples (average s of ± 8.9 × 10⁻⁸ A), and the corresponding average values were utilized for data analysis.

**Results and Discussion**

The ultimate objective of this research was to detect and quantify Hb in tooth and bone samples by spectrophotometric measurement using LCV and electrochemical analysis using cyclic voltammetry. Three different extraction solutions (0.5 M HCl, 0.2 M Tris, and 6.0 M Urea) were used to extract and detect Hb in tooth, femur head, and femur shaft samples. The use of these solutions was based on previous extraction studies [16,17]. Positive results for the following three samples were obtained: the femur shaft sample in 0.5 M HCl (F), the femur head sample in 0.5 M HCl (I), and the tooth sample in 6.0 M urea (K). Each of these three samples contained extracted Hb that was detectable with the LCV by UV/Vis spectroscopy (Figure 4) and by cyclic voltammetry (Figure 5).

To quantify these positive results spectrophotometrically with LCV, Hb standards were prepared and a calibration curve was obtained. An Hb concentration of 14.9 nM was determined to establish the limit of quantification for this set of measurements. Absorbance measurements for the three positive samples from the tooth and femur were obtained. By using the slope equation from the calibration curve, the Hb concentrations for each of the three positive samples were calculated. The Hb concentrations for each sample are given in Table 1. These three samples are all less concentrated than the normal Hb levels of a human, approximately 2 mM, which supports that Hb levels in bone decrease over time post-mortem and post-extraction.
The results obtained electrochemically were less sensitive than the spectroscopic method using LCV. A higher limit of quantification of 241 nM was established for this particular analysis, which is most likely due to the slow kinetic electron transfer. The bulky nature of the Hb molecule limits the mobility of electrons towards the working electrode surface. Utilizing the data from the voltammetric calibration curve, levels of Hb were determined for the three positive extraction sample as shown in Table 1. These sample concentrations also show a similar decrease in Hb both post-mortem and post-extraction.

These Hb concentration values illustrate that although the spectroscopic method using LCV yields lower limits of quantification for Hb, there is no sample modification necessary for electrochemical analysis, so samples could be further utilized for immunoassay or DNA testing. The combination of these two techniques demonstrates that Hb levels in tooth and bone can be quantified, and there is a significant decrease in Hb concentration post-mortem and post-extraction. While there is some discrepancy among the Hb values determined for both methods, each analysis method shows the feasibility of quantifying the protein within skeletal samples. Although these techniques demonstrate that quantification of Hb in bone and teeth is possible, further research with a variety of samples ranging in age post-mortem would be necessary to establish any specific correlation between Hb levels and a time since death or other forensic phenomena.

The measurement of Hb levels in bone makes this work unique, as the majority of studies focus solely on the identification of Hb and related species in bone and teeth, not on quantification. These methods might be potentially useful in certain cases where the ability to quantify Hb concentrations in tooth and bone samples would be advantageous. The detection method involving LCV is preferable to other colorimetric and luminescent reagents due to its sensitivity, stability, simplicity, and safety. The purple reaction product can last to other colorimetric and luminescent reagents due to its sensitivity, advantage of this method.

The use of LCV is slightly destructive to the extraction sample, so this method does limit the amount of biological and immunological testing possible. This drawback can be overcome by the use of cyclic voltammetry as a quantification method for Hb, because no modification of the extraction sample is required. Another small limitation is the possibility of false positives, but these appear to be minimal for this particular analysis. Environmental factors, age, and contextual variables could play a role in the subsequent sensitivity for the quantitation method, but the specificity and relative stability of the LCV reaction product and the electron transfer process in cyclic voltammetry should not be compromised. Overall, these results demonstrate that UV/Vis spectroscopy using LCV and electrochemical analysis are capable of providing specific and quantitative detection of Hb in bone and teeth. These methods would be mutually advantageous if conjunction with other methods such as auto-fluorescence and immunohistology.

### Conclusion

Quantitative determination of hemoglobin (Hb) in skeletal or dried remains could potentially have a wide range of forensic applications. There have been successful studies illustrating that blood proteins such as certain Hb components can be recovered and identified from fossilized or skeletal remains. This research focused on the use of LCV with UV/Vis spectroscopy and cyclic voltammetry for the detection and quantification of Hb in tooth and femur samples. The reaction of LCV and hydrogen peroxide with the heme group of Hb yields a dark purple color. Using LCV, possible interferences were investigated, and it was determined that ferrous samples appear to reduce the CV+ back to LCV, while glycine did not appear to affect the reaction with Hb. A slight interference by myoglobin was overcome by implementing a background correction, while albumin did not appear to hinder the reaction of Hb with LCV. Hb was successfully extracted from tooth and femur bone samples and was subsequently quantified using the reaction of LCV spectrophotometrically and by electrochemical analysis. The extractions of the femur head and shaft with 0.5 M HCl and the extraction of the tooth with 6.0 M urea yielded measurable amounts of Hb given in Table 1, all of which are consistent with a decrease in concentration post-mortem and post-extraction. Thus, Hb levels in bone and tooth samples were detected and quantified by both methods. While these particular results have not yet established any direct relationship between Hb levels in human remains and a post-mortem interval or other phenomena, the combination of these two analytical techniques illustrates the possibilities for various applications of quantifying Hb in bone and teeth. Future research would focus on the application of these methods to a wider variety of bone samples ranging in morphology, age, and condition, while minimizing some of the possible limitations within this method.

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### References

1. Introna F Jr, Di Vella G, Campobasso CP (1999) Determination of postmortem interval from old skeletal remains by image analysis of luminol test results. J Forensic Sci 44: 535-538.
2. Cattaneo C, Gelstorpe K, Phillips P, Sokol RJ (1992) Reliable identification of human albumin in ancient bone using ELISA and monoclonal antibodies. Am J Phys Anthropol 87: 305-372.
3. Schweitzer MH, Marshall M, Carron K, Bohle DS, Busse SC, et al. (1997) Heme compounds in dinosaur trabecular bone. Proc Natl Acad Sci USA 94: 6291-6296.
4. Schweitzer MH, Suo Z, Avci R, Asara JM, Allen MA, et al. (2007) Analyses of soft tissue from Tyrannosaurus rex suggest the presence of protein. Science 316: 277-280.
5. Campobasso CP, Di Vella G, De Donno A, Santoro V, Favia G, et al. (2006) Heme compounds in dinosaur trabecular bone. Proc Natl Acad Sci USA 94: 6291-6296.
6. Campobasso CP, Di Vella G, De Donno A, Santoro V, Favia G, et al. (2006) Heme compounds in dinosaur trabecular bone. Proc Natl Acad Sci USA 94: 6291-6296.
7. Hempe JM, Craver RD (2000) Separation of hemoglobin variants with similar charge by capillary isoelectric focusing: value of isoelectric point for identification of common and uncommon hemoglobin variants. Electrophoresis 21: 743-748.
8. Zijlstra WG, Buursma A, Meeuwsen-van der Roest WP (1991) Absorption spectra of human fetal and adult oxyhemoglobin, de-oxyhemoglobin, carboxyhemoglobin, and methemoglobin. Clin Chem 37: 1633-1638.
9. Mottola HA, Simpson BE, Gorin G (1970) Absorptiometric Determination of Hydrogen Peroxide in Submicrogram Amounts with Lueco Crystal Violet and Peroxidase as Catalyst. Anal Chem 42: 410-411.
10. Gere EP, Bérczi B, Simándi P, Wittmann G, Dombi A (2002) Simultaneous Determination of Hydrogen Peroxide and Organic Hydroperoxides in Water. Intern J Environ Anal Chem 82: 443-450.
11. Bodziak WJ (1996) Use of leuco crystal violet to enhance shoe prints in blood. Forensic Sci Int 62: 45-52.
12. Dai Z, Liu S, Ju H, Chen H (2004) Direct electron transfer and enzymatic activity of hemoglobin in a hexagonal mesoporous silica matrix. Biosens Bioelectron 19: 861-867.
13. Boyer R (2002) Concepts in Biochemistry. (2nd edn) Brooks/Cole, Pacific Grove, CA, USA.
14. Hopkins S (1979) Controlled iron release tablets. Drugs of Today 15: 404-406.
15. Jin YS, Lai JQ, Zhao XF, Meng J, Yin S (2005) Evaluation on glycine chelated iron (II) in improving nutritional anaemia in rats. Wei Sheng Yan Jiu 34: 344-346.
16. Ascenzi A, Brunori M, Citro G, Zito R (1985) Immunological detection of hemoglobin in bones of ancient Roman times and of Iron and Eneolithic Ages. Proc Natl Acad Sci USA 82: 7170-7172.
17. Jiang X, Ye M, Jiang X, Liu G, Feng S, et al. (2007) Method development of efficient protein extraction in bone tissue for proteome analysis. J Proteome Res 6: 2287-2294.