

**Sexual dimorphism and population variation of a selection of vertebrae in a South  
African sample**

by

Shannon Gibbs

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## ABSTRACT

In the discipline of forensic anthropology, numerous methods exist to establish a biological profile. However, many methods rely on established standards created from a limited number of skeletal elements. If these skeletal elements are not recovered or are fragmentary, the established methods cannot be applied, and the biological profile cannot be estimated accurately. As such, further exploration into the variation of numerous alternative skeletal elements and their potential use in forensic analyses is required. The purpose of this study was to explore the osteometric variation of a selection of vertebrae to examine the population affinity and sexual dimorphism in a South African sample. Additionally, this study aimed to establish if a universal discriminant analysis can be used to estimate population affinity and sex without the prior determination of the specific number of vertebrae within the vertebral column.

The current study consisted of two components, centred firstly around the exploration of vertebral variation and secondly on its use in predictive models. The variation exploration component quantified the osteometric variation among vertebrae to determine if there are statistically significant differences among vertebrae to see if non-specific, universal formulae - regardless of the specific number of a vertebra (e.g., T1, T2 etc.) - can be created. Secondly, the classification component aimed to apply the vertebral differences to develop classification standards to estimate population affinity and sex in a sample of black and white South Africans.

For the variation exploration component of the study, a series of 20 measurements were collected to assess the variation within and among the cervical, thoracic, and lumbar vertebrae of a sample of 30 black South African males. Out of the original 20 measurements, 14 were observed to be sufficiently repeatable and were retained for the creation of predictive models. The vertebrae within each type were compared using ANOVA and Tukey's HSD, which revealed that the atypical vertebrae (i.e., C7, T1, T12) were significantly different from the typical vertebrae within each vertebral region. As such, the atypical vertebrae are too different from the remaining vertebrae in each of the relevant subgroups and are not useful for non-specific, universal formulae.

For the classification component, a selection of cervical, thoracic and lumbar vertebrae of 180 black and white South African adult males and females were measured. Both univariate models, as well as multivariate models (combining all of the measurements), were created for

each vertebra to assess how accurately the vertebrae can predict population affinity and sex when using linear discriminant analysis. Finally, combined models were also created using the measurement means for all vertebrae in each subtype (i.e., a universal cervical model, universal upper thoracic model, universal lower thoracic model, and universal lumbar model).

The univariate models using each measurement collected for each sub-type of vertebra separately, resulted in classification accuracies of 50% to 90% for population affinity and 35% to 92% for sex when considering all vertebrae. Overall, the cervical vertebrae (C3-C6) presented with the highest accuracies for sex estimation but yielded poor results for population affinity estimation, where both the thoracic (T2-T11) and lumbar (L1-L4) vertebrae performed comparably well. Among the measurements, the vertebral body, spinous process and transverse process lengths and height measurements performed the best across all vertebrae. The universal univariate models assessing each measurement collected for the combined cervical, thoracic and lumbar vertebrae within each sub-type presented with classification accuracies ranging between 78% and 82% for population affinity and between 73% and 84% for sex estimation, where the upper thoracic vertebrae (T2-T6) and lower thoracic vertebrae (T7-T11) performed the best. Multivariate models using all of the measurements per sub-type vertebrae resulted in classification accuracies between 73% and 94% for population affinity and 70% and 89% for sex estimation. Finally, the universal multivariate model assessed all the vertebrae together (cervical, thoracic and lumbar vertebrae), and demonstrated classification accuracies between 81.7% and 87.2% for population affinity estimation, and 81.7% and 87.2% for sex estimation, where the lower thoracic (T7-T11) vertebrae performed the worst for both biological parameters.

Ultimately the best results were achieved using the multivariate models, as the inclusion of more variables maximises group differences. Combined models that made use of means irrespective of vertebra number yielded good accuracies. Therefore, a universal formula to estimate population affinity and sex can be used without *a priori* knowledge of the vertebral number. Both univariate and multivariate approaches achieve fairly high classification accuracies. The satisfactory performance of the univariate models also allows for the estimation of population affinity and sex from fragmentary or incomplete vertebrae.

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our backs  
tell stories  
no books have  
the spine to  
carry

-Rupi Kaur

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## CHAPTER 1: INTRODUCTION

Forensic anthropologists reconstruct biological profiles from incomplete or skeletonised remains. A biological profile serves to assist medico-legal investigators in possibly identifying an unknown individual and includes parameters such as age, stature, sex and population affinity (Ousley *et al.*, 2009; Krüger *et al.*, 2015; Liebenberg, 2015; Krüger *et al.*, 2017; Liebenberg *et al.*, 2019). Several methods exist to estimate population affinity and sex for South Africa using various skeletal elements (L'Abbé *et al.*, 2013; Liebenberg, 2015; Krüger *et al.*, 2018; Liebenberg *et al.*, 2019). Current methods typically concentrate on larger skeletal elements, specifically, the pelvis, skull and long bones. But skeletal remains are often incomplete or taphonomically altered when recovered in a forensic context. Therefore, other skeletal elements should also be investigated to establish classification standards (Pastor, 2005; Krüger, 2015; Torimitsu *et al.*, 2016; Liebenberg *et al.*, 2019).

To further explore additional skeletal elements, several studies have also assessed smaller elements of the axial skeleton for population affinity and sex estimation. The axial skeleton and its associated muscles provide support, allow movement and disperse load-bearing forces throughout the body (Marino, 1995; Wescott, 2000; Yu *et al.*, 2008; Kibii *et al.*, 2010; Marlow and Pastor, 2011; Bethard and Seet, 2013; Gama *et al.*, 2015; Ostrofsky and Churchill, 2015; Ramadan *et al.*, 2017; Hora and Sladek, 2018; Oura *et al.*, 2018; Jeong *et al.*, 2019; Rozendaal *et al.*, 2020).

Previous studies have implemented a variety of methods to estimate population affinity and sex from specific vertebrae (Berry *et al.*, 1987; Marino, 1995; Cheng *et al.*, 1998; Kibii *et al.*, 2010; Gama *et al.*, 2015; Torimitsu *et al.*, 2016; Montasser *et al.*, 2017; Ramadan *et al.*, 2017; Hora and Sladek, 2018; Jeong *et al.*, 2019; Rozendaal *et al.*, 2020). For example, some authors achieved high classification accuracies (64% - 92.9%) using the first and second cervical, twelfth thoracic or first lumbar vertebrae (Marlow *et al.*, 2011; Ostrofsky *et al.*, 2015; Torimitsu *et al.*, 2016; Hora and Sladek, 2018). While many studies have reported marked sex differences in the specific vertebrae, research on population variation, particularly in the South African population, is limited (Kibii *et al.*, 2010; Ostrofsky *et al.*, 2015; Ünlütürk, 2016).

The purpose of the study was, therefore, to assess the osteometric variation within and among a selection of vertebrae to determine their value for estimating population affinity and sex. More specifically, different vertebrae were compared to see if it would be possible to create

universal standards irrespective of the specific vertebral number and to test how the predictive power of universal standards compares to vertebra-specific standards when employing both univariate and multivariate models.

## CHAPTER 2: LITERATURE REVIEW

### 2.1 Forensic Anthropology and the Biological Profile

Forensic anthropologists frequently receive decomposed remains that require specialised analyses to facilitate personal identification. Several methods that evaluate the size and shape of the human skeleton exist to estimate a biological profile using various skeletal elements, including osteometric and morphoscopic approaches. The osteometric approach involves taking measurements from human remains compared to the morphoscopic approach, which involves visually assessing the morphological variants of human remains (typically the skull and the pelvis). The estimation of population affinity and sexual dimorphism is an essential aspect of the biological profile used to assist in identifying unknown individuals from complete or fragmentary remains (Ousley *et al.*, 2009; Krüger *et al.*, 2015; Liebenberg, 2015; Krüger *et al.*, 2017; Liebenberg *et al.*, 2019).

Population affinity refers to skeletal characteristics that differ among populations and are correlated to several factors, such as genetics, genetic history (species' genetic variation and the relationships between different populations), environment, socioeconomic status, and social boundaries (inequalities in social opportunities and access to resources), all of which can influence skeletal or biological variations (Sauer, 1992; Posel, 2001; Bergström *et al.*, 2020; Ross and Pilloud, 2021; Winburn and Algee-Hewitt, 2021). The observable traits, such as skin colour, facial characteristics, skeletal morphology and behavioural attributes, are interpreted into a social labelling system used by law enforcement to identify unknown individuals (Ousley *et al.*, 2009; Klales *et al.*, 2012; Klales and Kenyhercz, 2015; Krüger *et al.*, 2015; Liebenberg, 2015). For population affinity estimation, the osteometric approach has yielded better accuracies than the morphological or morphoscopic approach (Hefner, 2009; L'Abbé *et al.*, 2011; Ousley and Jantz, 2012; Stull *et al.*, 2014). The cranium has traditionally been recognised as the most accurate skeletal element for population affinity estimation, with classification accuracies ranging between 73% and 89% for South African population groups (L'Abbé *et al.*, 2013; Stull *et al.*, 2014). More recently, the postcranial skeleton has shown to be just as valuable for population affinity estimation, with accuracies of 85% when assessing black, white and coloured South Africans (Liebenberg, 2015).

Sexual dimorphism refers to the size and shape variation between males and females. Bone growth is a continuous process that persists into early adulthood and is influenced by genetics,

environment, and hormones. The presence and concentration of sex hormones (oestrogen and testosterone) have variable effects on bone growth, resulting in general differences in overall body size and proportions between male and female individuals (Cabo *et al.*, 2012; Wells, 2012; Krüger, 2015). In low concentrations, oestrogen can promote the secretion of growth hormones stimulating growth spurts in both sexes. However, at higher concentrations, oestrogen cause epiphyses to fuse and thereby halt the growth of long bones. The stimulation of endosteal and trabecular bone growth, as well as the increase in tensile bone strength, is also regulated by oestrogen (Juil, 2001; Cabo *et al.*, 2012; Krüger, 2015). On the other hand, testosterone promotes cortical bone growth in male adolescents and increases cortical diameter as osteoblastic activity is stimulated and osteoblast apoptosis is inhibited. In males, both oestrogen and testosterone work together to stimulate periosteal bone and cortical bone growth during puberty. As a result, testosterone builds wider and stronger bones in males, with increased bone mass in certain skeletal regions, such as the lumbar vertebrae, resulting in sexual dimorphism (Bonjour *et al.*, 1991; Callewaert *et al.*, 2010; Krüger, 2015).

Sexual dimorphism is population-specific and the degree to which males and females differ from one another can vary among and within groups or populations (Barrier and L'abbé, 2008; Krüger *et al.*, 2017; Liebenberg *et al.*, 2019). Therefore, population-specific standards are important for classifying the sex of unknown individuals (Seeman, 2001; Pastor, 2005; Steyn and Patriquin, 2009; Krüger *et al.*, 2015; Liebenberg *et al.*, 2019). For sex estimation, both morphoscopic and osteometric methods have been shown to be accurate. While measurements from long bones are more reliable and achieve higher accuracies (up to 94%) in estimating sex than the skull, morphological differences of the pelvis and skull are very useful when long bones are not available for analysis. While the pelvis yields accuracies of up to 90%, long bone analysis have yielded accuracies between 86% and 96% (Phenice, 1969; Walker, 2008; Spradley and Jantz, 2011; Klales *et al.*, 2012).

Numerous studies have been conducted on the South African population regarding population affinity and sex estimation, improving the ability of forensic anthropologists to create biological profiles (L'Abbé EN ; Elliott and Collard, 2009; Ousley *et al.*, 2009; Spradley *et al.*, 2011; Klales *et al.*, 2012; Klales *et al.*, 2015; Krüger *et al.*, 2015; Liebenberg *et al.*, 2015; Hefner, 2016; Spradley and Jantz, 2016; Krüger *et al.*, 2017). Unfortunately, the estimation of population affinity and sex commonly relies on fragile elements (such as the pubic bone or craniofacial bones), often lost or destroyed in forensic and archaeological contexts. In the post-

mortem context, remains may present with taphonomic changes, such as faunal damage, thermal alterations and weathering, frequently resulting in fragmentation of the bones. In some cases, only limited or fragmentary elements are recovered, making new techniques necessary to attempt a possible identification (Marlow *et al.*, 2011; Gama *et al.*, 2015; Torimitsu *et al.*, 2016). Therefore, other skeletal elements, such as the vertebrae, should be investigated to establish classification standards for instances when fragmentary or incomplete remains are recovered (Pastor, 2005; Krüger, 2015; Torimitsu *et al.*, 2016; Liebenberg *et al.*, 2019). Vertebrae are often fairly well preserved, with minimal damage and weathering with the exception of the transverse processes and spinous processes. Since the vertebral column is comprised of so many vertebrae, the possibility of at least some vertebrae being available and intact is great, even if the remains have substantial damage and weathering, as thick layers of muscles and ligaments surround the vertebrae.

## 2.2 General Vertebral Morphology and Bone Growth

During embryological development, vertebrae develop through the formation of somites within the axial skeleton. Somites are segmented units that develop along the neural tube and become individualised into different vertebrae types by the action of *Hox* genes (Carapuço *et al.*, 2005). The human vertebral formula (as determined by the complex of *Hox* genes) consists of seven cervical vertebrae, twelve thoracic vertebrae, five lumbar vertebrae, five sacral vertebrae and three to four coccygeal vertebrae. The general components that make up vertebrae include a vertebral body and posterior vertebral arch (Sadler and Langman, 2010; Drake *et al.*, 2016). The vertebral body is the main weight-bearing part of the vertebra, separated in the vertebral column by the intervertebral disc. The body size is dependent on the type of vertebra, where the amount of weight distribution differs in the different regions of the spine. For example, the cervical vertebrae are the smallest and are specialised to support and allow movement of the skull. In contrast, the lumbar vertebrae are the largest and are structurally specialised to support the weight of the upper body (torso) (Waxenbaum and Futterman, 2018).

The vertebral arch forms the lateral and posterior aspects of the vertebral foramen, where the surrounding elements form the vertebral canal. The vertebral canal is firmly anchored to the posterior aspect of the vertebral body by two pedicles. A flat sheet of bone extends from each pedicle to meet at the midpoint, forming the roof of the vertebral arch. Posteriorly and inferiorly, a spinous process extends from the lamina junction, which serves as a ligament and muscle attachment site. Transverse processes extend from the pedicles and lamina and are a

site for articulating the ribs to the thoracic vertebrae. Additionally, superior and inferior articular processes project from the junction between the lamina and pedicles, which are articulation sites for the sequential vertebrae (Drake *et al.*, 2016).

Overall, the seven cervical vertebrae are typically small and square-shaped, presenting with characteristic round transverse foramina (found on the transverse processes), triangular vertebral foramina and short and bifid (split) spinous processes. The first and second cervical vertebrae (atlas and axis, respectively) are considered atypical vertebrae that are specialised to carry and move the head. The twelve thoracic vertebrae are characterised by their articulation facets, more specifically demifacets, located on the lateral aspect of the vertebral bodies for articulation with the ribs. The thoracic vertebrae are somewhat heart-shaped and present with a circular vertebral foramen. The five lumbar vertebrae are distinguishable from the other vertebrae due to their weight-bearing capacity making them considerably larger, with kidney-shaped bodies and large, blunt spinous processes (White and Folkens, 2005; Sadler *et al.*, 2010; Drake *et al.*, 2016).

### **2.3 Extrinsic and Intrinsic Influences on Vertebral Variation**

Various factors affect vertebral morphology, including sex, age, growth and development, pathology, and physical activity. Sexual dimorphism has been noted in the growth and development of vertebrae (Cheng *et al.*, 1998; Wescott, 2000; Pastor, 2005; Soegiharto *et al.*, 2008; Yu *et al.*, 2008; Marlow *et al.*, 2011; Gama *et al.*, 2015; Ostrofsky *et al.*, 2015; Torimitsu *et al.*, 2016; Montasser *et al.*, 2017; Ramadan *et al.*, 2017; Oura *et al.*, 2018; Rozendaal *et al.*, 2020). The vertebrae of females are relatively taller and thinner in the coronal plane than in males. The female vertebral body height is greater than males, while the male vertebral body transverse diameter is greater than females. From the age of eight to about 21 years of age (i.e., when maturity is reached), the thoracic and lumbar vertebrae (specifically from the sixth thoracic to the fifth lumbar vertebrae) increase substantially in size and dimensions in males compared to females (Bellemare *et al.*, 2003).

Furthermore, sexual dimorphism in spinal growth produces a thinner thoracolumbar spine in the coronal plane of females compared to males, indicating sexual dimorphism in the shape of the vertebral body due to differential growth rates between males and females during puberty (Taylor and Twomey, 1984). Given the differences caused by testosterone, male vertebrae have been found to be 25% larger than those of females (Bellemare *et al.*, 2003). Previous studies

have reported that the vertebral column is heavily associated with sex differences in the thoracic cage related to the respiratory system. More specifically, males have larger muscles connected to their transverse processes than females (Bellemare *et al.*, 2006; Bastir *et al.*, 2014; Weaver *et al.*, 2014).

Many changes occur within the skeleton through either growth or degeneration, which can affect the rates at which these processes occur between males and females. Several studies have noted sexually dimorphic patterns in the ageing process of vertebrae (Atkinson, 1967; Mosekilde, 2000; Albert *et al.*, 2010; Aylott *et al.*, 2012; Baidas, 2012; Albert and Maier, 2013; Chiara *et al.*, 2016; Ramadan *et al.*, 2017). Age-related changes of the vertebrae have been observed, especially among lumbar vertebrae, although all vertebrae may be affected. With increased age, the transverse diameter of the endplates and vertebral body increases. Conversely, a decrease in anterior and posterior vertebral height is seen, which produces a wedge-shaped vertebral appearance with age (Snodgrass, 2004a; Garoufi *et al.*, 2020). Essentially, vertebrae become flatter and wider with age. In addition to the dimensional changes of the body, osteophyte development is commonly observed along the articular surfaces. Osteophytes refer to lipping around the articular surfaces, also known as bony spurs. The bony spurs develop due to various internal and external factors, including physical activity (i.e., strenuous labour, sport, and mechanical exertion), age and degenerative diseases. In a normal vertebral column, osteophyte development occurs after the epiphyseal ring fusion, and as the bony spurs appear, osteophytes increase with age in both severity and frequency (Snodgrass, 2004b; Van der Merwe *et al.*, 2006b; Klaassen *et al.*, 2011; Ezra *et al.*, 2019).

Previous studies have also suggested that the lumbar vertebrae show a pattern of sexual dimorphism, which may be due to the different load-bearing capacities between male and female individuals (Berry *et al.*, 1987; Cheng *et al.*, 1998; Ostrofsky *et al.*, 2015; Oura *et al.*, 2018). Load-bearing capacity, such as physical activity, affects bone mass, microarchitecture, and size throughout life, relates to differences in gait between males and females, which effects the pelvic shape, leading to possible sexual dimorphism. An increase in load-bearing capacity of the vertebrae has, in addition, been associated with an increase in vertebral dimensions and bone mass (Junno *et al.*, 2013; Oura *et al.*, 2016). Additionally, studies demonstrated that males in their twenties are more likely to have a higher bone mass and strength peak than females in their twenties, which may translate to osteometric differences (Pastor, 2005; Yu *et al.*, 2008; Torimitsu *et al.*, 2016).

A number of studies have found that osteophytes form in different patterns in females and males, with females showing greater variation (Allbright, 2007; Garoufi *et al.*, 2020). Van der Merwe *et al.* (2006a) focused on osteophyte formation in the South African population, which indicated that cervical and lumbar vertebrae had more significant osteophyte development with greater variation in males. Although age is a factor in osteophyte development, the main reason for the appearance of bony spurs is extensive pressure exerted on the vertebrae, which weakens and progresses due to age (Van der Merwe *et al.*, 2006a; Klaassen *et al.*, 2011). Another factor to consider is premature degeneration, which occurs either from an increase in load-bearing capacity, early degenerative disease onset (e.g., second osteoporosis) or trauma to the vertebrae (Scher, 1990; Kerttula *et al.*, 2000; Junno *et al.*, 2013).

#### **2.4 Practical Application and Previous Studies Assessing Vertebrae**

Research investigating population affinity and sex has assessed various methods, including morphological, metric, and scoring methods, to estimate population affinity and sex using vertebrae (Cheng *et al.*, 1998; Wescott, 2000; Pastor, 2005; Soegiharto *et al.*, 2008; Yu *et al.*, 2008; Marlow *et al.*, 2011; Gama *et al.*, 2015; Ostrofsky *et al.*, 2015; Torimitsu *et al.*, 2016; Montasser *et al.*, 2017; Ramadan *et al.*, 2017; Oura *et al.*, 2018; Rozendaal *et al.*, 2020). However, previous studies did not create standards for vertebral measurements based on the results and typically concentrated on larger skeletal elements (i.e., pelvis, skull and long bones). Even though vertebrae can be distinguished from one another by type, identifying the exact vertebral number (e.g. fourth thoracic vertebrae) may be difficult and time-consuming, especially if the vertebrae are fragmentary and no consecutive vertebrae are present (Cheng *et al.*, 1998; Kibii *et al.*, 2010; Gama *et al.*, 2015; Torimitsu *et al.*, 2016). The lack of anthropological research on vertebrae limits the accuracy and reliability of standards, as previous studies employing vertebrae focus on cadaveric or scan materials rather than skeletal elements. The development of standards for vertebrae may assess variations among and within the different vertebrae exploring whether standards are interchangeable or if vertebrae type-specific standards need to be created (Hora and Sladek, 2018). Therefore, universal, non-specific formulae for vertebrae can be beneficial as standards could be applied interchangeably across vertebrae without first establishing the vertebral type or number.

Numerous studies have evaluated the variation of the first and second cervical vertebrae using multiple measurements, which achieved accuracies between 70.91% and 92.9% for sex estimation (Marino, 1995; Wescott, 2000; Marlow *et al.*, 2011; Gama *et al.*, 2015; Torimitsu

*et al.*, 2016; Rozendaal *et al.*, 2020). A variety of studies (Marlow *et al.*, 2011; Bethard *et al.*, 2013) validated the method published by Wescott (2000) to estimate sex from the first and second cervical vertebrae. Overall, sex differences were observed between males and females, with males having larger vertebral body dimensions than females (Wescott, 2000; Marlow *et al.*, 2011; Bethard *et al.*, 2013).

Similarly, several studies focused on the lower thoracic and upper lumbar vertebrae in various population groups to estimate sex and noted accuracies between 64% and 90% (Yu *et al.*, 2008; Ostrofsky *et al.*, 2015; Hora and Sladek, 2018; Oura *et al.*, 2018; Garoufi *et al.*, 2020). Overall, males were found to present with larger measurement means than females when assessing the lumbar vertebrae. Furthermore, the lumbar vertebrae yielded greater classification accuracies than the studies using thoracic and cervical vertebrae.

Limited research has been done on vertebrae of the South African population to estimate population affinity and sex (Kibii *et al.*, 2010; Ostrofsky *et al.*, 2015; Ünlütürk, 2016). Kibii *et al.* (2010) focused on the seventh cervical vertebrae of South African Zulu (black), white, and coloured individuals to explore morphometric variation. The study found that sexual dimorphism was observable in vertebrae, with males having substantially greater dimensions than females (Kibii *et al.*, 2010). Ostrofsky *et al.* (2015) focused on the morphological differences using the lumbar region to estimate the sex of 98 black South Africans. Their results yielded accuracies ranging from 64.6% to 83.5%, with the vertebral body generating the highest accuracy (Ostrofsky *et al.*, 2015). Ünlütürk (2016) examined the vertebrae of black and white South Africans to estimate population affinity. The sample was composed of complete vertebrae from 144 individuals. The analysis provided a correct classification rate of 85% to 88%. Compared to the classification rates noted with skeletal elements such as the skull and the pelvis, the results reported by Ünlütürk (2016) are comparable. However, as the sample decreased due to missing or damaged vertebrae, the analyses had to counteract a lot of missing data, where more sacral measurements than vertebral measurements were taken with a limited amount of measurements (five) taken from the vertebrae (Ünlütürk, 2016).

In previous studies, authors usually focused on the atypical or transitional vertebrae, so limited evidence of comparable differences in and among the different typical vertebrae exists, and authors typically did not compare the specific types of vertebrae to one another but rather focused on atypical vertebrae (first cervical, second cervical and fifth lumbar vertebrae), and studies were often limited to sex estimation. Furthermore, little to no research is available on

the vertebral variations among the various South African population groups, and limited studies have created established standards for population affinity and sex estimation among typical vertebrae (Wescott, 2000; Pastor, 2005; Yu *et al.*, 2008; Kibii *et al.*, 2010; Marlow *et al.*, 2011; Gama *et al.*, 2015; Ostrofsky *et al.*, 2015; Torimitsu *et al.*, 2016; Ünlütürk, 2016; Jeong *et al.*, 2019; Garoufi *et al.*, 2020; Rozendaal *et al.*, 2020).

## CHAPTER 3: EXPLORATION OF VERTEBRAL VARIATION

To effectively explore sexual dimorphism and population affinity in the vertebrae of modern South Africans, this study consisted of two components. First, the variation exploration component and second, the classification component. The variation exploration component evaluated the size variation among vertebrae, both within and among vertebral types (cervical, thoracic, and lumbar vertebrae), to determine the possibility of creating universal vertebral standards for population affinity and sex estimation. Furthermore, measurement error was assessed to demonstrate the degree of measurement repeatability. Evaluation of measurement error and variability is important when creating standards based on metric data, as measurements need to be repeatable and valid.

### 3.1 South African Population Groups

South Africa is a country with a diverse and vast heritage that consists of various ancestries with varying geographical origins (i.e., indigenous San, Khoe-Khoe African, European, Asian and Indian). The 2021 South African mid-year population estimates reported that black South Africans comprised 80.9% of South Africa's population, while white South Africans and coloured South Africans comprised 7.7% and 8.8% of the population, respectively ([www.statssa.gov.za](http://www.statssa.gov.za)).

South Africans were classified into strict social categories during the segregation era (1948 to 1990). South African social segregation (*Apartheid* era) imposed legal separation, which allowed limited gene flow amongst the South African population groups. South Africans are mainly classified into four groups: white South Africans, black South Africans, coloured South Africans and Indians and other Asian South Africans (Christopher, 1992; Bickford-Smith, 1995; Christopher, 2001; Ribot, 2004).

Black South Africans are said to have originated as a result of the southward migration of Bantu-speaking individuals from western-central Africa into southern Africa (Ribot, 2004; Franklin *et al.*, 2007; Franklin *et al.*, 2010). In South Africa, the descendants of the Bantu-speakers further subdivided with time into different ethnic groups; this includes Nguni (Xhosa, Zulu, Northern Ndebele, Southern Ndebele and Swazi), Sotho (Pedi and Basotho), Tswana, Venda and Tsonga (Hall and Morris, 1983; Savage, 1986; Larson and Ribot, 2004; Ribot, 2004). White South Africans are mainly descendants of European settlers who arrived during

multiple colonisation periods, specifically settlers from the Netherlands, Belgium, Germany, France, Portugal, and Britain (Steyn and İşcan, 1997; İşcan and Steyn, 1999).

### **3.2 Skeletal Sample**

The sample for this component of the study consists of the cervical, thoracic and lumbar vertebrae (except C1, C2, and L5) of 30 black South African adult males. The skeletal remains were obtained from the Pretoria Bone Collection (PBC), which is housed in the Department of Anatomy at the University of Pretoria. Because this component focused only on variation among the different vertebrae, by looking at a single population and sex group (black South African adult males), any variation observed should inherently be due to differences in the vertebral structure and morphology rather than differences attributed to population and sex variation.

Individuals that presented with extensive pathology (e.g., osteophytosis or vertebral compression), vertebral deformity (e.g., scoliosis, spondylolysis, or sacralisation), an indication of spinal surgical procedures, or post-mortem damage were excluded from this study component, however, individuals that presented with minimal pathology were included in the sample. Additionally, individuals that did not present with all 24 vertebrae were excluded.

### **3.3 Methods**

#### **3.3.1 Osteometric Analysis**

A series of 20 measurements (Table 3.1, Appendix I) were chosen to assess the variation of cervical, thoracic, and lumbar vertebrae. Measurements were collected using a digital sliding caliper according to definitions collected from several previous studies (Cheng *et al.*, 1998; Marlow *et al.*, 2011; Gama *et al.*, 2015; Oura *et al.*, 2018). Osteometric data was collected on all cervical, thoracic, and lumbar vertebrae except the first and second cervical vertebrae and the fifth lumbar vertebrae. Refer to Table 3.1 for the measurement abbreviations and Appendix I for detailed measurement definitions.

**Table 3.1:** Measurement abbreviations. Refer to Appendix I and II for more information regarding the measurements.

<b>Maximum Superior Sagittal Length</b>	XSLs	<b>Pedicle Height</b>	PH
<b>Maximum Anterior Height</b>	XAH	<b>Pedicle Width</b>	PW
<b>Maximum Centroid Height</b>	XCH	<b>Maximum Pedicle Length</b>	XPL
<b>Maximum Posterior Height</b>	XPH	<b>Transverse Process Length</b>	TPL
<b>Maximum Superior Articular Length</b>	XALs	<b>Maximum Transverse Process Width</b>	XTPW
<b>Maximum Inferior Articular Length</b>	XALi	<b>Maximum Superior Articular Process Width</b>	XAPWs
<b>Maximum Superior Articular Width</b>	XAWs	<b>Maximum Inferior Articular Width</b>	XAPWi
<b>Maximum Inferior Articular Width</b>	XAWi	<b>Articular Process Height</b>	APH
<b>Foramen Sagittal Length</b>	FSL	<b>Maximum Spinous Process Length</b>	XSPL
<b>Foramen Coronal Width</b>	FCW	<b>Maximum Inferior Sagittal Length</b>	XSLi

### 3.3.2 Statistical Analysis

The intra- and inter-observer error rates were tested to evaluate the repeatability of the 20 measurements. For the intra-observer error rate, the measurements were performed twice by the principal investigator. For the inter-observer error rate, the measurements were performed twice by two different observers, the principal investigator and a second observer. To measure observer error, absolute and relative technical error of measurements (TEM and %TEM, respectively) were calculated, which are useful in comparing sets of repeated measurements (Ulijaszek and Lourie, 2005; Geeta *et al.*, 2009). TEM uses a co-efficient to determine the precision, quality, and accuracy of the measurements and provides an absolute value (in mm) to indicate discrepancies among sets of measurements. Additionally, %TEM offers a relative value of measurement discrepancies by also taking the size of the measurement into account.

Multiple statistical analyses were conducted using R (version 4.0.5) and the RStudio environment (R Core Team, 2020). Pearson's correlations were used to indicate the strength of the linear relationship between the measurements. A linear correlation coefficient greater than zero indicates a positive relationship. In contrast, a value less than zero signifies a negative relationship (Taylor, 1990; Asuero *et al.*, 2006; Torimitsu *et al.*, 2016; Schober *et al.*, 2018). A Holm's adjustment was added to reduce Type I errors that can occur with multiple comparisons (Holm, 1979; Kalnins, 2018). The correlation coefficients were displayed using the *corrplot* package in R to assist in interpreting the relationships among the measurements and search for multicollinearity using shapes and colours as illustrations. The ellipses show both the sign and magnitude of the correlation value using shades of red and blue (Friendly, 2002). The different shades of red and blue indicate the various negative and positive correlations, with the width and shade/spectrum of the ellipse demonstrating the magnitude of the correlation between the measurements. A smaller ellipse indicates a stronger correlation,

and a larger ellipse indicates a weaker correlation. For visualisation of the correlation, variables were ordered to display the patterns of relations using angular order of the eigenvector (AOE). AOE relationships between vectors indicate correlations among variables, and therefore, an order based on the angular relationships naturally places the most similar variables together. Variables are ordered based on the angles formed by the first two or three eigenvectors (Friendly, 2002). As a result, all variables are contained within a 90-degree segment of the p-space, and most correlations are positive or near zero. Additionally, a significance test was added, and any non-significant correlation coefficients were omitted from the plot (Friendly, 2002; Kumari, 2008; Wei *et al.*, 2017).

ANOVA was used to test for significant differences between variables. First, ANOVA was used to test for differences among the different vertebrae within a single type (e.g., comparing the same measurements collected from C3, C4, C5 and C6). Subsequently, ANOVA was also used to test for significant differences among the subtypes (i.e. comparing the same measurements among the cervical, thoracic and lumbar vertebrae) (Fenech, 1979; Madrigal, 2012). ANOVA assumes a normal distribution of the data, as well as homogeneity of variance, which means the variance among the groups or samples should be approximately equal. Additionally, ANOVA assumes that the measurements and variables are independent of each other (Stahle and Wold, 1989; Tabachnick and Fidell, 2019). Additionally, Tukey's HSD test was applied *post hoc* to identify where the specific differences occurred.

### **3.4 Results: Measurement Repeatability**

A series of measurements (Appendix I) were compiled from previous studies and assessed to gauge the repeatability of the measurements. In the current study, both the intra- and inter-observer errors demonstrated widespread variation with many measurements presenting with high error rates. Overall, the intra- and inter-observer error rates ranged between 0.00mm and 3.54mm (TEM); 0.00% and 29.79% (%TEM) and between 0.00mm and 7.29mm (TEM) and 0.00% and 27.03% (%TEM), respectively (Table 3.2 – 3.5). The intra-observer error rate was overall lower than the inter-observer error rate, except for the measurements of the lower thoracic vertebrae, where the upper range of the error for inter-observer error rate was lower than the intra-observer errors. The measurements collected from the lower thoracic and lumbar vertebrae were overall noted to be the most repeatable (Table 3.3 – 3.5).

The measurements with the highest TEM and %TEM for both inter- and intra-observer error rates varied, but the maximum centroid height, maximum articular process height, maximum superior articular length, maximum superior articular width, pedicle height, pedicle width, foramen sagittal length, foramen coronal width consistently presented with large observer errors. A cut-off point of 7% was selected, and any measurement with a %TEM greater than 7% was considered unreliable and was therefore excluded from further classification analyses.

**Table 3.2:** Absolute technical error of measurement (TEM) and relative technical error of measurement (%TEM) assessing the intra- and inter-observer error rates for the cervical vertebrae (C3 – C7). Refer to Appendix I for measurement names and definitions. Bold indicates the highest %TEM.

	Inter										Intra									
	C3		C4		C5		C6		C7		C3		C4		C5		C6		C7	
	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM
<b>XSLs</b>	0.45	2.57	0.45	2.69	0.45	2.66	0.32	1.85	0.32	1.81	0.71	4.14	0.45	2.69	0.45	2.66	0.45	2.60	0.00	0.00
<b>XCH</b>	0.84	<b>7.68</b>	1.00	<b>9.26</b>	0.84	<b>8.28</b>	1.10	<b>10.74</b>	1.45	<b>11.59</b>	1.00	<b>9.26</b>	1.34	<b>12.42</b>	0.95	<b>9.39</b>	1.22	<b>11.66</b>	1.48	<b>11.59</b>
<b>XAH</b>	0.45	3.44	0.63	4.72	1.05	<b>8.67</b>	0.84	6.59	0.84	5.62	0.55	4.12	0.45	3.34	0.77	6.35	0.84	6.49	0.84	5.62
<b>XPH</b>	0.32	2.24	0.32	2.34	0.32	2.41	0.55	4.31	1.14	<b>7.97</b>	0.45	3.11	0.55	3.94	0.32	2.41	1.00	<b>7.58</b>	0.95	6.28
<b>XALs</b>	1.10	<b>7.40</b>	1.61	<b>11.36</b>	1.14	<b>8.32</b>	1.82	<b>12.70</b>	1.00	6.85	1.05	<b>7.55</b>	1.00	<b>7.58</b>	0.45	3.44	1.38	<b>9.92</b>	0.55	3.83
<b>XALi</b>	1.10	<b>7.11</b>	1.22	<b>7.90</b>	1.00	6.41	1.14	<b>7.36</b>	0.89	6.13	0.45	3.06	0.45	3.02	0.63	4.11	0.45	3.02	0.63	4.45
<b>XAWs</b>	1.18	6.10	1.38	<b>7.07</b>	2.14	<b>10.41</b>	2.37	<b>10.96</b>	2.37	<b>9.94</b>	0.77	4.12	0.63	3.36	0.95	4.92	1.22	5.92	1.34	6.10
<b>XAWi</b>	0.77	4.30	1.10	6.02	1.26	6.73	1.73	<b>8.41</b>	1.58	6.45	0.77	4.50	0.77	4.35	0.77	4.26	0.77	3.87	0.84	3.56
<b>FSL</b>	0.95	6.28	1.00	6.67	1.14	<b>7.36</b>	0.55	3.44	1.18	<b>7.58</b>	0.55	3.78	0.63	4.33	1.00	6.58	0.55	3.49	0.95	6.12
<b>FCW</b>	2.17	<b>9.15</b>	2.10	<b>8.53</b>	1.95	<b>7.86</b>	1.67	6.44	1.61	6.35	0.55	2.43	1.00	4.31	0.89	3.73	0.45	1.77	1.00	4.07
<b>PH</b>	0.32	4.45	0.45	6.04	0.84	<b>11.16</b>	0.55	<b>7.94</b>	0.45	5.32	0.71	<b>9.69</b>	0.63	8.32	0.84	<b>11.16</b>	0.84	<b>12.49</b>	1.05	<b>12.34</b>
<b>PW</b>	0.84	<b>15.79</b>	1.00	<b>20.00</b>	0.63	<b>10.90</b>	0.63	<b>10.54</b>	0.71	<b>10.88</b>	0.84	<b>15.21</b>	0.77	<b>14.34</b>	0.63	<b>10.90</b>	0.63	<b>10.54</b>	0.84	<b>12.87</b>
<b>XPL</b>	1.00	<b>20.83</b>	0.84	<b>17.07</b>	1.05	<b>21.40</b>	0.77	<b>14.34</b>	1.26	<b>22.59</b>	0.32	<b>7.35</b>	0.84	<b>19.46</b>	0.77	<b>17.60</b>	0.45	<b>8.94</b>	0.32	6.45
<b>TPL</b>	1.45	4.46	0.71	2.15	0.71	2.03	1.48	4.12	0.71	1.86	1.48	4.58	1.41	4.29	0.55	1.57	1.52	4.25	1.52	4.00
<b>XTPW</b>	1.26	2.40	0.95	1.74	6.32	<b>11.40</b>	0.00	0.00	0.32	0.47	1.26	2.40	1.76	3.30	1.00	1.72	1.26	2.05	2.86	4.04
<b>XAPWs</b>	1.30	2.76	1.34	2.73	0.89	1.77	0.55	1.02	0.55	1.03	1.30	2.76	1.61	3.26	1.34	2.64	0.55	1.03	0.45	0.84
<b>XAPWi</b>	1.18	2.38	1.41	2.76	1.30	2.43	0.77	1.46	2.28	4.51	0.77	1.57	1.55	3.00	1.45	2.72	0.45	0.84	3.45	6.91
<b>APH</b>	1.10	5.02	1.00	4.55	0.55	2.46	1.26	5.55	4.28	<b>15.22</b>	0.71	3.32	1.55	<b>7.17</b>	1.22	5.40	2.57	<b>10.98</b>	1.41	5.24
<b>XSPL</b>	0.71	4.23	0.71	4.75	0.55	3.20	0.32	1.29	0.63	1.96	0.71	4.29	1.05	<b>7.04</b>	2.32	<b>13.06</b>	1.00	4.13	1.92	5.92
<b>XSLi</b>	0.77	4.40	0.84	4.73	0.32	1.77	0.45	2.54	0.00	0.00	0.55	3.09	0.84	4.73	0.32	1.77	0.45	2.54	0.45	2.69
<b>min</b>	0.32	2.24	0.32	1.74	0.32	1.77	0.00	0.00	0.00	0.00	0.32	1.57	0.45	2.69	0.32	1.57	0.45	0.84	0.00	0.00
<b>max</b>	2.17	20.83	2.10	20.00	6.32	21.40	2.37	14.34	4.28	22.59	1.48	15.21	1.76	19.46	2.32	17.60	2.57	12.49	3.45	12.87
<b>mean</b>	0.96	6.15	1.00	6.65	1.22	6.88	0.94	6.08	1.18	6.71	0.77	5.02	0.96	6.21	0.88	5.82	0.90	5.71	1.14	5.65

**Table 3.3:** Absolute technical error of measurement (TEM) and relative technical error of measurement (%TEM) assessing the intra- and inter-observer error rates for the upper thoracic vertebrae (T1 – T6). Refer to Appendix I for measurement names and definitions. Bold indicates the highest %TEM.

	Inter												Intra											
	T1		T2		T3		T4		T5		T6		T1		T2		T3		T4		T5		T6	
	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM
<b>XSLs</b>	0.32	1.89	0.32	1.75	0.95	4.96	0.71	3.29	1.05	4.62	0.55	2.25	0.63	3.79	0.55	2.96	0.89	4.43	1.14	5.21	0.55	2.37	0.45	1.83
<b>XCH</b>	2.19	<b>15.32</b>	3.08	<b>18.24</b>	1.14	7.08	1.26	7.81	1.64	<b>10.33</b>	1.76	<b>10.54</b>	1.52	<b>10.46</b>	1.18	7.40	1.18	7.40	1.52	<b>9.42</b>	1.14	6.99	1.52	<b>8.97</b>
<b>XAH</b>	2.45	<b>14.85</b>	2.26	<b>11.70</b>	0.95	5.30	0.45	2.51	0.55	3.09	0.55	3.03	0.55	3.26	0.55	2.93	0.71	3.95	0.32	1.77	0.55	3.09	0.45	2.46
<b>XPH</b>	1.14	6.71	0.95	5.13	0.55	2.90	0.45	2.38	0.71	3.70	0.45	2.31	0.77	4.45	0.77	4.08	0.55	2.90	0.45	2.38	0.45	2.33	0.45	2.31
<b>XALs</b>	1.18	7.99	1.38	8.89	0.84	4.89	1.45	7.67	2.02	<b>10.18</b>	1.61	7.61	0.84	5.81	0.55	3.78	1.45	8.57	0.45	2.48	0.63	3.36	0.45	2.21
<b>XALi</b>	0.89	5.92	1.05	6.06	0.63	3.44	1.55	7.82	1.61	7.68	1.26	5.80	0.77	5.23	1.30	8.00	0.77	4.21	0.55	2.87	0.45	2.21	0.95	4.54
<b>XAWs</b>	1.26	5.23	1.38	5.74	0.84	3.59	1.38	6.13	1.48	6.18	1.34	5.41	1.48	6.31	1.48	6.31	0.84	3.69	0.71	3.17	0.89	3.86	0.84	3.47
<b>XAWi</b>	2.55	9.96	2.59	10.31	0.55	2.25	1.05	4.18	1.10	4.31	1.58	6.10	1.18	4.77	1.05	4.21	0.84	3.47	1.05	4.25	0.77	3.10	0.63	2.49
<b>FSL</b>	1.22	7.38	1.55	8.95	1.58	9.04	2.32	13.67	1.84	<b>10.98</b>	2.28	<b>13.26</b>	0.63	3.90	0.84	4.89	1.10	6.44	1.00	6.25	1.48	<b>9.16</b>	1.18	7.40
<b>FCW</b>	3.44	<b>16.20</b>	3.73	<b>18.46</b>	1.48	7.97	1.18	6.72	1.14	6.75	1.34	8.08	1.48	7.65	0.89	4.71	1.26	7.11	0.84	4.95	0.71	4.29	0.84	5.33
<b>PH</b>	2.63	<b>25.50</b>	2.63	<b>21.18</b>	0.77	6.15	0.55	4.53	1.05	9.28	1.00	8.20	0.77	7.52	0.71	5.75	0.63	5.02	0.63	5.27	1.34	<b>11.77</b>	0.63	5.27
<b>PW</b>	0.89	11.04	0.95	<b>13.75</b>	0.84	<b>14.18</b>	0.84	<b>17.07</b>	0.55	<b>10.33</b>	1.05	<b>19.79</b>	0.71	8.95	0.77	<b>12.49</b>	0.84	<b>16.41</b>	0.71	<b>15.04</b>	0.89	<b>17.89</b>	0.84	<b>17.80</b>
<b>XPL</b>	1.18	<b>24.15</b>	0.84	<b>19.46</b>	0.77	<b>15.49</b>	0.84	<b>16.41</b>	0.55	<b>10.33</b>	1.26	<b>23.42</b>	0.77	<b>16.14</b>	0.32	7.71	1.14	<b>25.34</b>	0.84	<b>17.07</b>	0.71	<b>13.86</b>	0.45	8.60
<b>TPL</b>	0.63	1.71	0.84	2.41	1.22	3.89	1.41	4.78	0.95	3.05	1.70	5.44	1.05	2.86	2.26	6.62	1.22	3.96	1.58	5.36	0.89	2.94	0.89	2.92
<b>XTPW</b>	1.00	1.32	1.05	1.51	1.45	2.29	0.63	1.05	0.45	0.73	0.55	0.89	2.55	3.41	2.55	3.74	0.45	0.71	1.00	1.67	0.00	0.00	0.55	0.89
<b>XAPW<sub>s</sub></b>	0.45	0.89	0.45	1.14	0.77	2.18	0.45	1.32	0.45	1.41	0.32	1.01	4.54	9.11	1.34	3.42	0.71	2.00	0.55	1.62	1.48	4.72	0.00	0.00
<b>XAPWi</b>	1.22	2.99	1.58	4.24	1.38	3.97	1.73	5.35	1.45	4.49	2.28	7.13	1.67	4.06	0.55	1.44	1.05	2.99	0.77	2.33	0.32	0.96	1.05	3.19
<b>APH</b>	0.55	1.92	2.88	8.70	0.32	1.00	0.77	2.42	1.00	3.16	1.10	3.28	1.64	5.63	0.55	1.69	0.45	1.42	0.77	2.46	0.45	1.42	0.89	2.66
<b>XSPL</b>	0.00	0.00	0.32	0.83	0.77	1.91	0.71	1.70	1.61	3.70	1.58	3.66	0.32	0.86	0.71	1.86	0.84	2.05	0.95	2.27	2.05	4.69	1.58	3.66
<b>XSLi</b>	0.32	1.81	0.32	1.62	1.10	5.22	0.45	2.00	0.32	1.56	0.55	2.22	0.32	1.79	0.45	2.28	0.55	2.55	0.45	1.98	0.45	1.86	0.55	2.20
<b>min</b>	0.00	0.00	0.32	0.83	0.32	1.00	0.45	1.05	0.45	0.73	0.32	0.89	0.32	0.86	0.32	1.44	0.45	0.71	0.32	1.62	0.00	0.00	0.00	0.00
<b>max</b>	3.44	25.50	3.73	21.18	1.58	15.49	2.32	17.07	2.02	10.98	2.28	23.42	4.54	16.14	2.55	12.49	1.45	25.34	1.58	17.07	2.05	17.89	1.58	17.80
<b>mean</b>	1.28	8.14	1.51	8.50	0.95	5.38	1.01	5.94	1.08	5.78	1.24	7.22	1.26	6.01	1.00	4.95	0.89	5.90	0.83	5.04	0.83	5.21	0.77	4.53

**Table 3.4:** Absolute technical error of measurement (TEM) and relative technical error of measurement (%TEM) assessing the intra- and inter-observer error rates for the lower thoracic vertebrae (T7 – T12). Refer to Appendix I for measurement names and definitions. Bold indicates the highest %TEM.

	Inter												Intra											
	T7		T8		T9		T10		T11		T12		T7		T8		T9		T10		T11		T12	
	TEM	% TEM	TE M	% TEM	TEM	% TEM	TEM	% TEM	TE M	% TEM	TEM	% TEM	TEM	% TEM	TE M	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM	TEM	% TEM
XSLs	0.55	2.16	0.55	2.07	0.63	2.33	0.45	1.63	0.32	1.17	0.32	1.12	0.45	1.76	0.32	1.19	0.77	2.85	0.32	1.15	0.32	1.17	0.45	1.57
XCH	1.14	6.52	0.63	3.61	1.64	8.82	1.00	5.19	1.82	8.78	1.92	8.63	1.64	<b>9.19</b>	1.10	6.15	1.61	<b>8.40</b>	0.89	4.52	1.30	6.24	1.95	<b>8.70</b>
XAH	0.63	3.44	0.63	3.29	0.63	3.04	0.45	2.05	0.32	1.42	0.00	0.00	0.45	2.43	0.45	2.35	0.63	3.04	0.45	2.05	0.32	1.42	0.00	0.00
XPH	0.32	1.51	0.32	1.50	1.00	4.59	1.00	4.42	0.45	1.86	0.00	0.00	0.00	0.00	0.32	1.53	1.00	4.59	0.77	3.46	0.45	1.86	0.32	1.22
XALs	1.38	6.18	1.30	5.60	1.34	5.50	1.64	6.65	1.76	7.19	1.87	7.39	0.89	4.14	0.00	0.00	1.18	5.10	0.77	3.34	0.63	2.73	0.71	2.98
XALi	1.64	6.67	1.70	7.07	2.07	<b>8.40</b>	1.73	6.98	3.56	13.76	1.64	6.25	1.34	5.99	0.55	2.37	1.58	6.79	0.71	3.03	0.71	2.93	0.32	1.25
XAWs	1.64	6.25	0.89	3.22	1.34	4.72	1.92	6.27	1.52	4.58	1.61	4.16	1.05	4.08	0.55	2.01	1.00	3.55	1.14	3.81	1.14	3.49	1.38	3.66
XAWi	1.18	4.17	1.22	4.21	0.84	2.71	1.97	5.86	1.45	3.82	1.34	3.30	1.34	4.90	0.77	2.75	0.55	1.80	2.45	<b>7.61</b>	0.32	0.85	1.00	2.50
FSL	1.70	<b>10.45</b>	2.41	<b>14.87</b>	1.73	<b>10.43</b>	2.10	12.79	1.73	9.62	3.27	16.95	1.48	<b>9.39</b>	1.26	<b>8.32</b>	1.22	7.90	1.61	<b>10.21</b>	1.22	<b>7.16</b>	2.39	<b>13.05</b>
FCW	0.77	4.78	0.77	4.67	0.55	3.24	0.55	3.20	0.71	3.78	1.90	8.47	0.32	1.96	0.55	3.44	0.71	4.18	1.41	<b>8.22</b>	0.63	3.44	1.22	5.80
PH	0.45	3.49	1.10	8.43	0.84	5.85	0.84	5.33	0.45	2.63	0.45	2.51	1.61	<b>12.22</b>	0.84	6.59	1.34	<b>9.32</b>	0.71	4.50	0.32	1.87	0.32	1.79
PW	0.84	<b>14.18</b>	0.84	<b>12.13</b>	1.00	<b>13.51</b>	1.34	16.77	1.22	14.41	1.41	14.73	1.55	<b>29.79</b>	1.87	<b>32.82</b>	1.61	<b>24.43</b>	1.00	<b>13.89</b>	0.89	<b>11.47</b>	1.38	<b>15.84</b>
XPL	0.71	<b>11.98</b>	1.30	<b>20.06</b>	1.70	<b>27.03</b>	0.55	9.61	1.10	21.07	0.77	11.74	0.77	<b>13.83</b>	1.14	<b>20.73</b>	0.95	<b>17.25</b>	0.45	<b>8.28</b>	0.71	<b>13.86</b>	0.63	<b>9.88</b>
TPL	1.18	3.87	0.95	3.15	1.18	4.02	0.84	2.90	0.32	1.20	0.55	2.01	0.55	1.84	0.84	2.80	0.95	3.26	1.00	3.52	0.32	1.21	0.84	3.13
XTPW	1.34	2.23	1.41	2.39	1.84	3.19	1.22	2.20	1.22	2.37	0.45	0.83	1.30	2.17	1.38	2.31	2.26	3.91	0.77	1.39	0.77	1.50	0.45	0.83
XAPWs	0.55	1.80	0.55	1.74	1.30	3.82	1.10	3.06	0.45	1.20	0.71	1.88	0.55	1.81	0.77	2.48	0.95	2.78	1.05	2.92	0.55	1.48	0.84	2.23
XAPWi	2.07	6.26	2.49	<b>7.32</b>	1.18	3.22	0.77	2.06	3.18	8.39	0.71	2.27	1.95	5.77	1.64	4.74	0.55	1.48	0.55	1.45	0.32	0.86	0.55	1.76
APH	0.77	2.21	1.14	3.21	0.63	1.79	0.45	1.20	0.32	0.80	0.32	0.69	0.63	1.83	0.63	1.78	0.45	1.26	0.32	0.85	0.00	0.00	0.32	0.69
XSPL	0.45	1.06	1.34	3.21	0.71	1.76	1.14	3.01	0.89	2.71	0.55	1.64	1.00	2.37	1.38	3.29	0.71	1.76	1.00	2.65	0.71	2.14	0.45	1.34
XSLi	0.77	2.90	0.63	2.30	0.32	1.15	0.45	1.64	0.32	1.14	0.32	1.08	0.77	2.90	0.32	1.16	0.00	0.00	0.32	1.16	0.00	0.00	0.55	1.86
min	0.32	1.06	0.32	1.50	0.32	1.15	0.45	1.20	0.32	0.80	0.00	0.00	0.00	0.00	0.00	0.00	0.45	1.26	0.32	0.85	0.00	0.00	0.00	0.00
max	2.07	14.18	2.49	20.06	2.07	27.03	2.10	16.77	3.56	21.07	3.27	16.95	1.95	29.79	1.87	32.82	2.26	24.43	2.45	13.89	1.30	13.86	2.39	15.84
mean	1.00	5.11	1.11	5.70	1.12	5.96	1.08	5.14	1.15	5.59	1.01	4.78	0.99	6.08	0.86	5.67	1.05	5.98	0.91	4.57	0.61	3.46	0.82	4.12

**Table 3.5:** Absolute technical error of measurement (TEM) and relative technical error of measurement (%TEM) assessing the intra- and inter-observer error rates for the lumbar vertebrae (L1 – L4). Refer to Appendix I for measurement names and definitions. Bold indicates the highest %TEM.

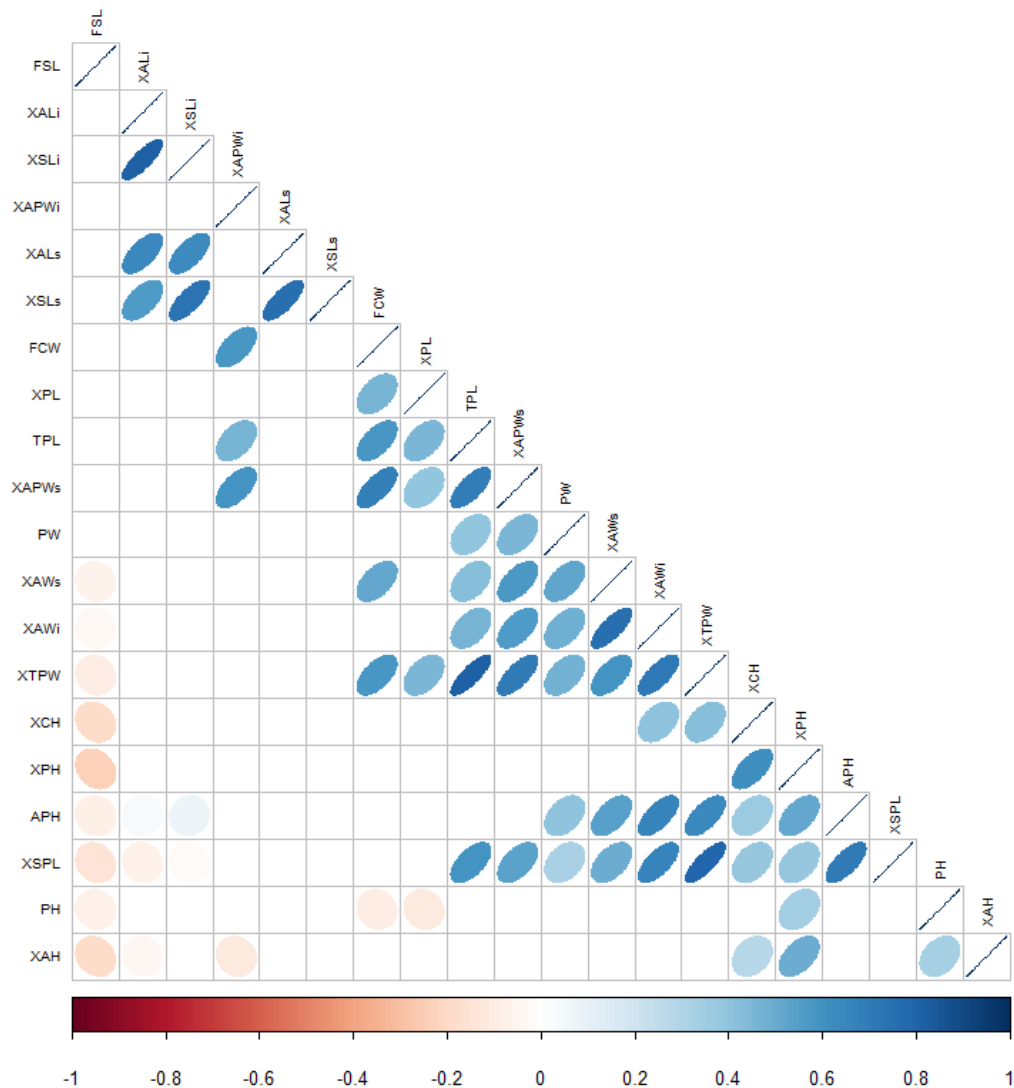
	Inter								Intra							
	L1		L2		L3		L4		L1		L2		L3		L4	
	TEM	%TEM	TEM	%TEM	TEM	%TEM	TEM	%TEM	TEM	%TEM	TEM	%TEM	TEM	%TEM	TEM	%TEM
<b>XSLs</b>	0.32	1.08	0.63	2.07	0.00	0.00	2.26	6.70	0.55	1.88	0.55	1.78	0.00	0.00	0.00	0.00
<b>XCH</b>	1.38	5.82	1.41	5.99	2.43	<b>10.42</b>	1.38	6.18	1.55	6.56	1.22	5.12	1.97	<b>8.33</b>	1.38	6.18
<b>XAH</b>	1.14	4.40	0.63	2.36	0.32	1.18	0.32	1.18	1.05	4.02	0.63	2.36	0.32	1.18	0.71	2.65
<b>XPH</b>	1.00	3.60	0.55	1.96	0.32	1.13	0.55	2.01	0.95	3.42	0.32	1.14	0.32	1.13	0.71	2.61
<b>XALs</b>	2.35	<b>8.92</b>	1.64	5.89	2.10	<b>7.18</b>	1.45	5.01	1.45	5.82	0.55	2.05	1.48	5.26	0.63	2.28
<b>XALi</b>	2.14	<b>7.83</b>	1.14	3.89	1.18	3.97	2.14	6.83	0.63	2.41	1.14	4.09	0.77	2.71	0.95	3.11
<b>XAWs</b>	2.00	5.08	2.19	5.14	1.79	4.01	1.55	3.30	1.34	3.48	0.71	1.70	1.22	2.79	1.00	2.16
<b>XAWi</b>	2.51	5.96	1.00	2.23	1.76	3.71	1.73	3.56	1.45	3.53	1.52	3.45	1.05	2.27	1.18	2.47
<b>FSL</b>	2.93	<b>15.12</b>	3.11	<b>16.84</b>	2.57	<b>14.43</b>	2.93	<b>15.43</b>	1.64	<b>8.88</b>	2.05	<b>11.64</b>	1.34	7.99	1.97	<b>11.03</b>
<b>FCW</b>	1.55	6.68	0.45	1.93	1.22	5.40	0.55	2.33	1.26	5.55	1.10	4.76	1.14	4.98	0.55	2.31
<b>PH</b>	0.32	1.94	0.71	4.29	0.45	2.76	0.32	2.12	0.45	2.73	0.63	3.86	0.45	2.76	0.77	5.10
<b>PW</b>	0.32	2.90	0.32	3.07	1.05	<b>8.13</b>	1.00	6.41	0.00	0.00	0.71	7.00	1.22	<b>9.64</b>	0.45	2.94
<b>XPL</b>	0.32	4.58	0.45	6.78	1.00	<b>15.63</b>	0.89	14.43	0.45	6.58	0.84	13.72	0.45	<b>7.45</b>	0.95	<b>16.64</b>
<b>TPL</b>	0.63	1.99	0.71	1.81	2.26	5.14	1.76	4.22	0.63	2.01	0.71	1.82	1.05	2.41	1.55	3.82
<b>XTPW</b>	0.45	0.71	0.45	0.56	0.45	0.51	0.45	0.55	0.45	0.71	0.32	0.40	0.71	0.81	0.32	0.39
<b>XAPWs</b>	2.93	<b>8.52</b>	2.76	<b>7.41</b>	3.33	<b>8.48</b>	3.07	6.91	2.21	6.34	2.81	7.66	3.54	<b>9.00</b>	2.86	6.42
<b>XAPWi</b>	0.00	0.00	0.32	0.94	0.45	1.17	0.77	1.73	0.77	2.50	0.32	0.94	0.63	1.66	0.32	0.71
<b>APH</b>	0.45	0.98	0.45	0.96	0.71	1.53	0.45	1.01	0.32	0.70	0.63	1.34	0.55	1.19	0.55	1.24
<b>XSPL</b>	0.00	0.00	0.71	1.93	1.22	3.37	1.10	3.28	0.00	0.00	0.95	2.57	0.45	1.22	1.18	3.56
<b>XSLi</b>	0.32	1.04	0.63	1.98	7.23	20.64	0.71	1.99	0.32	1.04	0.55	1.71	0.45	1.36	0.32	0.90
<b>min</b>	0.00	0.00	0.32	0.56	0.00	0.00	0.32	0.55	0.00	0.00	0.32	0.40	0.00	0.00	0.00	0.00
<b>max</b>	2.93	15.12	3.11	16.84	3.33	15.63	3.07	15.43	2.21	8.88	2.81	13.72	3.54	9.64	2.86	16.64
<b>mean</b>	1.20	4.53	1.03	4.00	1.29	5.17	1.30	4.90	0.90	3.53	0.93	4.07	0.98	3.83	0.95	3.98

### 3.5 Results: Exploration of Vertebral Variation

To determine whether universal, non-specific standards were possible for assessing vertebrae or if specific vertebrae had to be identified before further analysis, the dimensions of the vertebrae in the various types were compared.

#### 3.5.1 Cervical Vertebrae

The correlation coefficients for cervical vertebrae demonstrated both positive and negative relationships among the variables (Figure 3.1). Moderate ( $r = 0.33 - 0.68$ ) to strong ( $r = 0.71 - 0.81$ ) positive correlations were noted between maximum inferior sagittal length and maximum inferior articular length ( $r = 0.81$ ), maximum transverse process width and transverse process length ( $r = 0.81$ ), maximum spinous process length and maximum transverse process width ( $r = 0.80$ ), maximum superior sagittal length and maximum superior articular length ( $r = 0.75$ ) and maximum inferior articular width and maximum superior articular width ( $r = 0.76$ ) measurements. Foramen sagittal length was the only variable to show negative correlations ( $r = -0.06 - -0.23$ ).



**Figure 3.1:** Correlation plot depicting the relationship among cervical vertebrae (C3-C7) variables. Red indicates negative correlations, blue indicates positive correlations, the ellipse width indicates the magnitude of the correlation, and the blank boxes indicate non-significant correlations. Refer to Appendix I for measurement names and definitions.

The ANOVA tests detected statistically significant differences among the different vertebral measurements in the cervical vertebrae for all variables, except for maximum superior sagittal, maximum superior articular length, maximum inferior articular length, foramen sagittal length, and maximum inferior sagittal length (Table 3.6). When the cervical vertebrae were further explored with Tukey’s HSD, results revealed significant differences for various measurements in the atypical cervical vertebrae, more often among the third, fifth, sixth, and seventh cervical vertebrae (Appendix III, Figures 1-20). Furthermore, results revealed significant differences in

the height of the pedicle, transverse processes, and articular width measurements of different cervical vertebrae.

**Table 3.6:** ANOVA results evaluating the cervical vertebrae for each measurement. Bold indicates significance. Refer to Appendix I for measurement names and definitions.

Cervical Vertebrae		
	F Value	Pr(>F)
XSLs	1.005	0.38
XCH	5.498	< <b>0.05</b>
XAH	10.84	< <b>0.05</b>
XPH	6.651	< <b>0.05</b>
XALs	0.808	0.52
XALi	0.535	0.71
XAWs	18.52	< <b>0.05</b>
XAWi	48.32	< <b>0.05</b>
FSL	1.066	0.38
FCW	4.848	< <b>0.05</b>
PH	6.929	< <b>0.05</b>
PW	12.82	< <b>0.05</b>
XPL	4.075	< <b>0.05</b>
TPL	12.93	< <b>0.05</b>
XTPW	41.44	< <b>0.05</b>
XAPWs	13.68	< <b>0.05</b>
XAPWi	8.83	< <b>0.05</b>
APH	46.73	< <b>0.05</b>
XSPL	137.2	< <b>0.05</b>
XSLi	0.535	0.71

### 3.5.2 Upper Thoracic Vertebrae

The correlation coefficients in the upper thoracic vertebrae (T1-T6) demonstrated both positive and negative relationships among the variables (Figure 3.2). Substantial weak negative ( $r = -0.14$  -  $-0.62$ ) correlations were shown throughout the measurements. Positive correlations ( $r = 0.65$  –  $0.93$ ) were also noted throughout the upper thoracic vertebral measurements. The strongest positive correlations were noted between maximum transverse process width and transverse process length ( $r = 0.93$ ) maximum inferior and superior articular process width ( $r = 0.86$ ), maximum inferior and superior articular length ( $r = 0.92$ ) and maximum superior and inferior sagittal length ( $r = 0.93$ ).



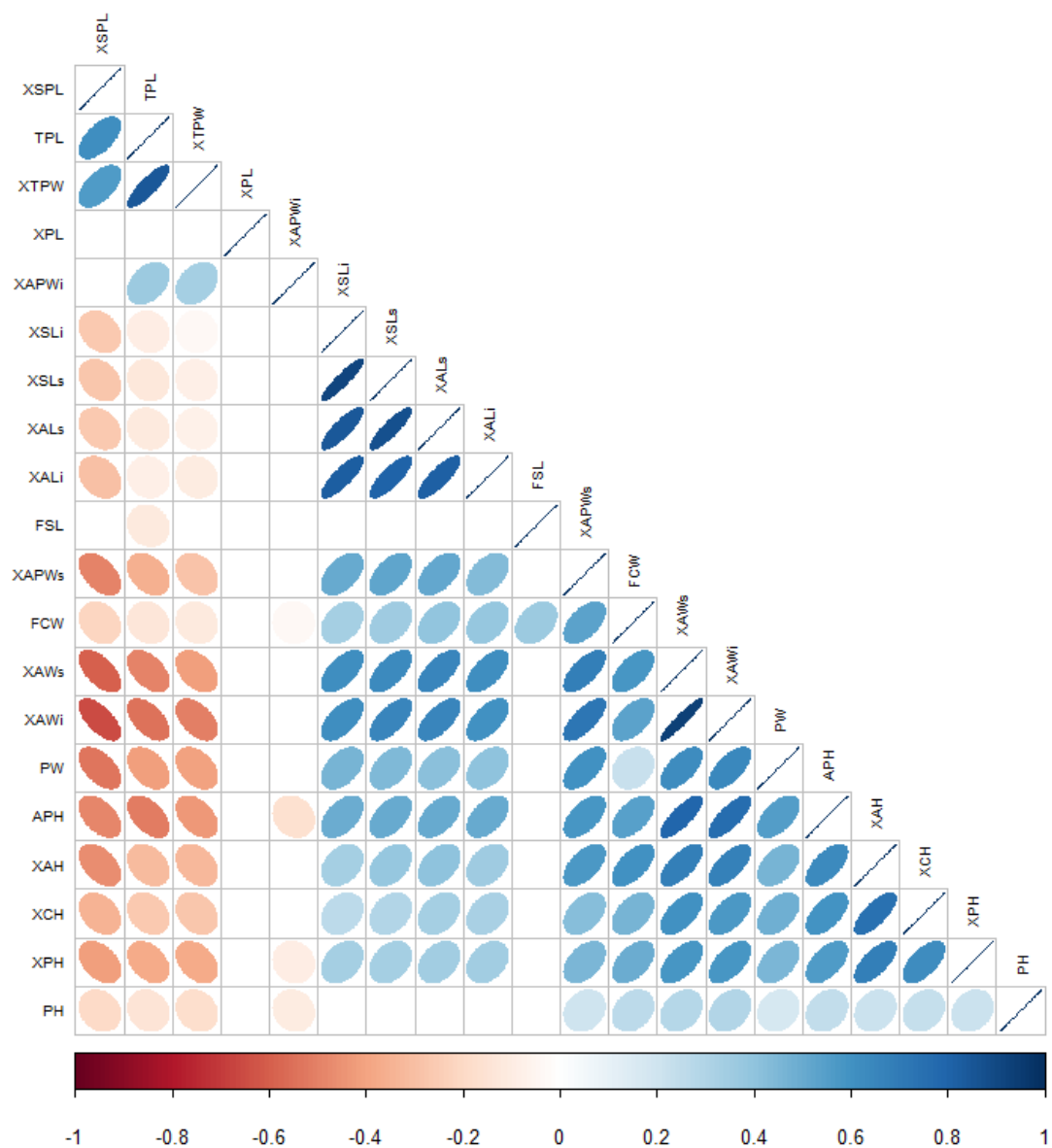
articular widths, foramen sagittal lengths, and pedicle lengths of the different upper thoracic vertebrae did not differ significantly.

**Table 3.7:** ANOVA results evaluating the upper thoracic vertebrae for each measurement. Bold indicates significance. Refer to Appendix I for measurement names and definitions.

<b>Upper Thoracic Vertebrae</b>		
	<b>F Value</b>	<b>Pr(&gt;F)</b>
<b>XSLs</b>	52.62	< <b>0.05</b>
<b>XCH</b>	19.31	< <b>0.05</b>
<b>XAH</b>	9.697	< <b>0.05</b>
<b>XPH</b>	23.94	< <b>0.05</b>
<b>XALs</b>	49.89	< <b>0.05</b>
<b>XALi</b>	48.55	< <b>0.05</b>
<b>XAWs</b>	0.955	0.45
<b>XAWi</b>	1.454	0.21
<b>FSL</b>	0.54	0.746
<b>FCW</b>	23.57	< <b>0.05</b>
<b>PH</b>	23.5	< <b>0.05</b>
<b>PW</b>	56.83	< <b>0.05</b>
<b>XPL</b>	1.853	0.11
<b>TPL</b>	24.8	< <b>0.05</b>
<b>XTPW</b>	40.54	< <b>0.05</b>
<b>XAPWs</b>	86.74	< <b>0.05</b>
<b>XAPWi</b>	36.39	< <b>0.05</b>
<b>APH</b>	13.81	< <b>0.05</b>
<b>XSPL</b>	15.26	< <b>0.05</b>
<b>XSLi</b>	36.13	< <b>0.05</b>

### 3.5.3 Lower Thoracic Vertebrae

Correlation coefficients for lower thoracic vertebrae (T7-T12) demonstrated positive and negative weak relationships among the variables (Figure 3.3). Maximum spinous process length, transverse process length and maximum transverse process width showed weak negative correlations ( $r = -0.09$  -  $-0.66$ ) throughout the measurements. Additionally, weak positive correlations were noted between the lower thoracic vertebral measurements. The strongest positive correlations were noted between maximum transverse process width and transverse process length ( $r = 0.85$ ), maximum inferior and superior articular process length ( $r = 0.82$ ), maximum inferior and superior articular widths ( $r = 0.94$ ), maximum superior and inferior sagittal length ( $r = 0.92$ ).



**Figure 3.3:** Correlation plot depicting the relationship among lower thoracic vertebrae (T7-T12) variables. Red indicates negative correlations, blue indicates positive correlations, the ellipse's size indicates the magnitude of the correlation, and the blank boxes indicate non-significant correlations. Refer to Appendix I for measurement names and definitions.

In the lower thoracic vertebrae (T7-T12), ANOVA detected significant differences for all variables, except the foramen sagittal length (Table 3.8). When further explored with Tukey's HSD, significant differences were found among the different lower thoracic vertebra types (i.e., T7-T12). When compared to the typical vertebrae, such as the seventh, ninth, and tenth thoracic

vertebrae, the atypical lower thoracic vertebrae (T12) showed no significant differences using Tukey's HSD (Appendix III, Figures 31-50).

**Table 3.8:** ANOVA results evaluating the lower thoracic vertebrae for each measurement. Bold indicates significance. Refer to Appendix I for measurement names and definitions.

<b>Lower Thoracic Vertebrae</b>		
	<b>F Value</b>	<b>Pr(&gt;F)</b>
<b>XSLs</b>	9.163	<b>&lt; 0.05</b>
<b>XCH</b>	17.39	<b>&lt; 0.05</b>
<b>XAH</b>	37.15	<b>&lt; 0.05</b>
<b>XPH</b>	23.06	<b>&lt; 0.05</b>
<b>XALs</b>	9.94	<b>&lt; 0.05</b>
<b>XALi</b>	6.497	<b>&lt; 0.05</b>
<b>XAWs</b>	59.83	<b>&lt; 0.05</b>
<b>XAWi</b>	76.97	<b>&lt; 0.05</b>
<b>FSL</b>	2.65	<b>&lt; 0.05</b>
<b>FCW</b>	16.72	<b>&lt; 0.05</b>
<b>PH</b>	3.136	<b>&lt; 0.05</b>
<b>PW</b>	18.42	<b>&lt; 0.05</b>
<b>XPL</b>	0.845	0.52
<b>TPL</b>	32.19	<b>&lt; 0.05</b>
<b>XTPW</b>	28.46	<b>&lt; 0.05</b>
<b>XAPWs</b>	26.57	<b>&lt; 0.05</b>
<b>XAPWi</b>	19.23	<b>&lt; 0.05</b>
<b>APH</b>	59.06	<b>&lt; 0.05</b>
<b>XSPL</b>	33.84	<b>&lt; 0.05</b>
<b>XSLi</b>	7.716	<b>&lt; 0.05</b>

### 3.5.4 Lumbar Vertebrae

The lumbar vertebral correlation coefficients demonstrated both positive and negative correlations between measurements (Figure 3.4). The moderate ( $r = 0.29 - 0.64$ ) to strongest ( $r = 0.71 - 0.81$ ) positive correlations were noted between maximum transverse process width and transverse process length ( $r = 0.94$ ), maximum inferior and superior articular process width ( $r = 0.88$ ), maximum inferior and superior articular length ( $r = 0.83$ ), maximum superior sagittal length and maximum superior articular length ( $r = 0.90$ ) and maximum inferior sagittal length and maximum inferior articular width ( $r = 0.90$ ).



**Table 3.9:** ANOVA results evaluating the lumbar vertebrae for each measurement. Bold indicates significance. Refer to Appendix I for measurement names and definitions.

<b>Lumbar Vertebrae</b>		
	<b>F Value</b>	<b>Pr(&gt;F)</b>
<b>XSLs</b>	20.79	<b>&lt; 0.05</b>
<b>XCH</b>	7.496	<b>&lt; 0.05</b>
<b>XAH</b>	3.178	<b>&lt; 0.05</b>
<b>XPH</b>	5.724	<b>&lt; 0.05</b>
<b>XALs</b>	10.12	<b>&lt; 0.05</b>
<b>XALi</b>	13.55	<b>&lt; 0.05</b>
<b>XAWs</b>	11.84	<b>&lt; 0.05</b>
<b>XAWi</b>	10.65	<b>&lt; 0.05</b>
<b>FSL</b>	2.099	0.11
<b>FCW</b>	1.677	0.18
<b>PH</b>	8.563	<b>&lt; 0.05</b>
<b>PW</b>	22.09	<b>&lt; 0.05</b>
<b>XPL</b>	1.976	0.12
<b>TPL</b>	25.02	<b>&lt; 0.05</b>
<b>XTPW</b>	32.45	<b>&lt; 0.05</b>
<b>XAPWs</b>	14	<b>&lt; 0.05</b>
<b>XAPWi</b>	40.84	<b>&lt; 0.05</b>
<b>APH</b>	1.059	0.37
<b>XSPL</b>	5.189	<b>&lt; 0.05</b>
<b>XSLi</b>	19.03	<b>&lt; 0.05</b>

### 3.6 Summary

- Maximum centroid height, foramen sagittal length, foramen coronal width, pedicle height, pedicle width, and maximum pedicle length measurements were significantly different between the vertebrae and thus will not be useful for the creation of universal formulae.
- Several measurements were shown to have repeatability issues; thus, maximum centroid height, foramen sagittal length, foramen coronal width, and pedicle height, width, and maximum pedicle length measurements were removed from further analyses.
- The atypical vertebrae (C7, T1 and T12) showed vast differences, along with the vertebrae closest to the atypical vertebrae (C6, T2, T11, L1 and L4).
- The sixth cervical, second thoracic, eleventh thoracic and fourth lumbar vertebrae have been excluded from the study because they differ too much from the rest of the vertebrae and will not be useful for the creation of universal formulae.

## CHAPTER 4: POPULATION AFFINITY AND SEX ESTIMATION FROM THE VERTEBRAE

In chapter 3, the variation exploration component explored the size variation among sequential vertebrae within and among the different subtypes (i.e., cervical, thoracic, and lumbar vertebrae) to determine possible single vertebral or combined vertebral standards for population affinity and sex estimation. Using the information from the variation exploration component, classification models were created to gauge the predictive accuracy of vertebral measurements when estimating population affinity and sex. The classification component focused on the quantifiable differences in the vertebrae in order to develop standards that can be used to estimate population affinity and sex in South Africa. Furthermore, this component assessed whether universal, non-specific standards can be established or if a specific vertebra needs to be identified first before classification can occur.

### 4.1 Skeletal Sample

For the classification component, the study sample consisted of the selection of vertebrae suggested to be useful for a universal formula in the variation exploration component; this includes cervical (C3 to C6), thoracic (T2 to T11), and lumbar (L1 to L4) vertebrae of 120 adult South African individuals (60 black and 60 white) with equal sex distribution (30 males and 30 females per population group). The morphology of the thoracic vertebrae is variable, as the upper and lower thoracic play different roles in the vertebral column, for example, the upper thoracic (T1-T6) has less load-bearing capacity compared to the lower thoracic (T7-T12) vertebrae. Therefore, the thoracic vertebrae were separated into upper thoracic (T1-T6) and lower thoracic (T7-T12) vertebrae. The skeletal remains were obtained from the Pretoria Bone Collection (PBC), housed in the Department of Anatomy at the University of Pretoria.

All personal information was anonymised for each individual, with only the sex, age, and population affinity recorded for the study. The sample included individuals between 27 and 93 years of age. However, it should be acknowledged that the dimensions of the vertebral body may be affected in older individuals due to vertebral compression and osteophyte growth. Table 4.1 provides the mean age distribution for the classification component of the skeletal sample for each population group and sex.

For bilateral measurements (i.e., pedicle and transverse process measurements), only the left side was measured. Any measurements that were affected by the extensive presence of pathology (e.g., osteophytosis or vertebral compression), vertebral deformity (e.g., scoliosis, spondylolysis, or sacralisation), an indication of spinal surgical procedures, or post-mortem damage were excluded from the study, however, individuals with minimal presence of pathology were included in the sample. Additionally, the exclusion of one vertebra or measurement did not exclude the individual from the sample.

**Table 4.1:** Mean age distribution of the skeletal sample.

	Black			White		
	Males (n=30)	Females (n=30)	Pooled (n=60)	Males (n=30)	Females (n=30)	Pooled (n=60)
Age (years)	35	38	37	55	59	57

## 4.2 Methods

After exploring the variation of the various vertebrae and their measurements in chapter 3, measurements from a range of typical and non-transitional vertebrae were selected for the creation of single vertebral univariate and multivariate models and combined universal univariate and multivariate models for population affinity and sex estimation. In the current chapter, the vertebral measurements were explored for their potential to estimate population affinity and sex.

Univariate models were created to assess the accuracy of each individual measurement for each vertebra in identifying population affinity and sex, which could be useful when only fragmentary vertebrae are available. Universal univariate models were created for each individual measurement per population affinity and sex for each vertebral sub-type (i.e., combined models for cervical (C3-C6), upper thoracic (T2-T6), lower thoracic (T7-T11) and lumbar vertebrae (L1-L4)).

Multivariate models were also created to estimate population affinity and sex. The multivariate models were created for each individual vertebra and included the various measurements retained after stepwise analyses for those vertebrae. Additionally, combined universal multivariate vertebral models were created that included all the measurements from the various vertebrae sub-types (i.e., models for cervical (C3-C6), upper thoracic (T2-T6), lower thoracic (T7-T11) and lumbar vertebrae (L1-L4)). The individual multivariate models could be useful

for population affinity and sex estimation if intact vertebrae of known vertebra sub-type and number were available for analysis. However, in cases where the specific vertebra number is not identifiable, the universal multivariate models could be useful, as only the sub-type needs to be identified to estimate population and sex.

#### 4.2.1 Osteometric analysis

A series of 14 measurements were retained from the variation exploration component to assess the cervical, thoracic, and lumbar vertebrae in the classification component. The principal investigator collected osteometric data on a selection of cervical, thoracic, and lumbar vertebrae, including the third to sixth cervical vertebrae (C3 - C6), the second to eleventh thoracic (T2 - T11) and first to fourth lumbar (L1 - L4) vertebrae. The first, second and seventh cervical vertebrae, first and twelfth thoracic and fifth lumbar vertebrae were excluded due to morphological variability, as established in the variation exploration component. Refer to Table 4.2 for the retained measurements and their associated measurement abbreviations.

**Table 4.2:** Measurement abbreviations for the variables retained for the classification component of the study. Refer to Appendix I and II for more measurement information.

<b>Maximum Superior Sagittal Length</b>	XSLs	<b>Transverse Process Length</b>	TPL
<b>Maximum Anterior Height</b>	XAH	<b>Maximum Transverse Process Width</b>	XTPW
<b>Maximum Posterior Height</b>	XPH	<b>Maximum Inferior Articular Width</b>	XAPWi
<b>Maximum Superior Articular Length</b>	XALs	<b>Articular Process Height</b>	APH
<b>Maximum Inferior Articular Length</b>	XALi	<b>Maximum Spinous Process Length</b>	XSPL
<b>Maximum Superior Articular Width</b>	XAWs	<b>Maximum Inferior Sagittal Length</b>	XSLi
<b>Maximum Inferior Articular Width</b>	XAWi	<b>Maximum Pedicle Length</b>	XPL

#### 4.2.2 Statistical Analysis

All statistical analyses were performed using R (version 4.0.5) and the RStudio environment (R Core Team, 2020). Outliers were detected and removed before statistical analysis using univariate boxplots. Descriptive statistics, including the mean and standard deviations, were assessed to analyse patterns in the vertebral measurements of the various population and sex groups. ANOVA was used to test for significant differences in the vertebral measurements between the population and sexes groups. A multitude of univariate and multivariate models was employed to identify the most accurate combination of predictor variables. After taking the mean of each vertebra separately, the mean of each measurement was combined per subtype of vertebra. In this manner, the method not only developed a univariate or specific formula but also determined if universal formulas could be useful. Finally, linear discriminant analysis

(LDA) was run with univariate and multivariate classification models in order to assess the potential of the measurements in estimating population affinity and sex. In biological anthropology, LDA is a classification method that incorporates measurements to evaluate biological parameters (sex, population affinity) using mathematical formulae (Ousley *et al.*, 2012; Tabachnick *et al.*, 2019). LDA calculates factors or weights for each variable (i.e., measurements), which maximises the mean differences among groups. The factors can then be used to classify unknown individuals. An unknown individual classification is based on the similarities and discriminant function scores produced with LDA, allowing comparison to each reference group centroid, thereby classifying the unknown individual into the group with the smallest distance from the centroid (Ousley *et al.*, 2012).

There are several requirements for the LDA to produce functions with the highest possible classification accuracy, including adequate sample size, multivariate normality, no outliers and equal group variation. Firstly, the sample size is extremely important, as accurate calculations are related to the number of measurements analysed. The basic criterion needed is that the smallest sample must be at least one individual larger than the number of variables used in the model (Ousley *et al.*, 2012; Tabachnick *et al.*, 2019). Another requirement is multivariate normality of data, both univariate and multivariate normality, to accurately estimate the parameters and probabilities of LDA (Ousley *et al.*, 2012; Tabachnick *et al.*, 2019). Outliers are typically measurements or individuals that are outside the normal variance of the sample. Outliers can be due to measurement error, recording error, an encoding error, human variation, pathology, or a combination of these reasons. Therefore, removing outliers, especially in multivariate analyses, is required and recommended, especially if the sample size is large enough (Ousley *et al.*, 2012). Lastly, equal group variation (i.e., homogeneity of variance), which is dependent on the standard deviations, is a measure of variation within a group that quantifies differences among the classified groups. Therefore, as in univariate analyses, the within-group variation must be relatively similar to the centroid to make a comparison using a common measure of variability (Ousley *et al.*, 2012; Tabachnick *et al.*, 2019).

Multivariate models were created using all the measurements taken per vertebra to test the classification accuracies of each vertebral sub-type (i.e., L1, L2, L3, L4, respectively) and universal models were created using all the measurements combining each vertebral sub-type (i.e., combining all lumbar vertebrae simultaneously (L1-L4)). Table 4.3 provides the variables included in each univariate and multivariate classification model and associated measurements.

**Table 4.3:** Variables included in each univariate and multivariate classification models.

Vertebrae sub-type	Vertebrae	<i>n</i>
Individual cervical models	C3	74
	C4	80
	C5	69
	C6	63
Universal cervical model	Combining all cervical vertebrae (C3-C7)	286
Individual upper thoracic models	T2	96
	T3	107
	T4	97
	T5	88
	T6	89
Universal upper thoracic model	Combining all upper thoracic vertebrae (T2-T6)	478
Lower thoracic models	T7	89
	T8	96
	T9	91
	T10	96
	T11	99
Universal lower thoracic model	Combining all lower thoracic vertebrae (T7-T11)	475
Lumbar models	L1	73
	L2	59
	L3	51
	L4	43
Universal lumbar model	Combining all lumbar vertebrae (L1-L4)	226

## 4.3 Results

### 4.3.1 Cervical Vertebrae

The descriptive statistics demonstrated variability among the cervical vertebrae of black and white South African males and females. Tables A1 to A4 (see Appendix IV) present the descriptive statistics of the cervical vertebrae, including the sample size, mean and standard deviation for each measurement. Overall, white South Africans ( $\mu = 5.31 - 55.51$ ) had larger measurement means than black South Africans ( $\mu = 4.8 - 52.96$ ), except for the maximum superior sagittal length and the maximum transverse process width which were smaller. Compared to females ( $\mu = 4.97 - 51.89$ ), males ( $\mu = 5.13 - 56.11$ ) had greater measurement means. From the third cervical to the sixth cervical vertebra, the transverse and spinous processes increased in size, while the articular widths became larger to accommodate the articulation of larger sequential vertebra.

ANOVA demonstrated significant differences between black and white South Africans for most of the measurements across all cervical vertebrae, with a few exceptions. Specifically, maximum superior sagittal length was only significantly different between the population groups for the sixth cervical vertebra, while transverse process length, maximum transverse

process width and maximum spinous process length were significantly different for all cervical vertebrae, except for the sixth cervical vertebra. When comparing the sexes, maximum pedicle length was noted as not significantly different between the sexes for any of the cervical vertebrae. Additionally, articular process height was not significantly different when assessing the third and fourth cervical vertebrae for sex differences (Table 4.4).

**Table 4.4:** ANOVA results evaluating population groups and sexes for each measurement for the cervical vertebra (C3-C6). Bold indicates significant variables. Refer to Appendix I for measurement names and definitions.

	Population				Sex			
	C3	C4	C5	C6	C3	C4	C5	C6
<b>XSLs</b>	0.466	0.022	0.047	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XAH</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XPH</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XALs</b>	0.014	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XALi</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XAWs</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.024	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XAWi</b>	0.186	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XPL</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.120	0.369	0.708	0.328
<b>TPL</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.144	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XTPW</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.584	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XAPWi</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>APH</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.038	0.020	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XSPL</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.473	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XSLi</b>	0.877	0.160	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>

With the universal cervical model (combined means of C3 – C6), ANOVA detected statistically significant differences among the vertebral body lengths and articular process, spinous process and articular widths of white South Africans. Among black South Africans, the only measurements that did not differ significantly included the maximum superior sagittal length, maximum posterior height, maximum superior articular length, maximum inferior articular length, and maximum inferior sagittal length. Similar results were seen when comparing the sexes. Among males, the only measurements that were not significantly different included the maximum superior sagittal length, maximum posterior height, maximum superior articular length, maximum inferior articular length, and maximum inferior sagittal length. In females, the vertebral body lengths, articular process, spinous process, and articular width did not differ significantly (Table 4.5).

**Table 4.5:** ANOVA results evaluating population groups and sexes for each measurement using the universal cervical model (C3-C6). Bold indicates significant variables. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Blacks	Whites	Males	Females
XSLs	0.427	<0.01	0.026	<0.01
XAH	<0.01	<0.01	<0.01	0.023
XPH	0.155	0.427	0.199	0.565
XALs	0.236	<0.01	0.163	<0.01
XALi	0.073	<0.01	0.010	<0.01
XAWs	<0.01	<0.01	<0.01	<0.01
XAWi	<0.01	<0.01	<0.01	<0.01
XPL	<0.01	<0.01	<0.01	<0.01
TPL	<0.01	0.158	<0.01	0.074
XTPW	<0.01	<0.01	<0.01	<0.01
XAPWi	<0.01	<0.01	<0.01	<0.01
APH	<0.01	<0.01	<0.01	<0.01
XSPL	<0.01	<0.01	<0.01	<0.01
XSLi	0.672	<0.01	0.208	<0.01

When attempting population affinity estimation using univariate models, the classification accuracies ranged between 0% and 89% for the third cervical vertebra, between 44% and 81% for the fourth cervical vertebra, between 35% and 86% for the fifth cervical vertebra, and between 42% and 84% for the sixth cervical vertebra (Table 4.6). For sex estimation, the univariate models yielded accuracies between 56% and 89% for the third cervical vertebra, between 38% and 86% for the fourth cervical vertebra, between 35% and 81% for the fifth cervical vertebra, and between 22% and 98% for the sixth cervical vertebrae (Table 4.7).

**Tables 4.6:** Univariate LDA correct classification accuracy (%) for population affinity using the cervical vertebrae (C3-C6). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	C3						C4						C5						C6					
	Black		White		Total		Black		White		Total		Black		White		Total		Black		White		Total	
	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
XSLs	19	41	18	64	37	50	29	60	16	69	45	<b>80</b>	32	65	11	55	43	62	31	82	14	56	45	71
XAH	33	72	18	64	51	69	36	75	20	56	56	64	35	74	13	65	48	70	23	61	13	52	36	57
XPH	31	67	19	68	50	68	40	83	22	64	62	68	42	86	12	60	54	<b>78</b>	28	74	13	52	41	65
XALs	23	50	15	54	38	51	35	73	14	78	49	76	35	71	7	35	42	61	24	63	16	64	40	63
XALi	25	54	19	68	44	59	26	54	23	83	49	78	35	71	10	50	45	65	27	71	18	72	45	71
XAWs	25	54	19	68	44	59	29	60	24	75	53	70	28	57	15	75	43	62	31	82	17	68	48	<b>76</b>
XAWi	23	50	13	46	36	49	31	65	25	75	56	70	29	59	13	65	42	61	25	66	12	48	37	59
XPL	38	59	18	43	56	53	24	50	22	47	46	56	38	78	8	40	46	67	25	66	15	60	40	63
TPL	27	83	12	64	39	76	36	75	24	67	60	64	36	73	14	70	50	72	17	45	9	36	26	41
XTPW	38	72	24	86	62	78	33	69	24	75	57	74	30	61	10	50	40	58	16	42	15	60	31	49
XAPWi	35	76	17	89	52	<b>82</b>	42	88	25	67	67	68	35	71	14	70	49	71	21	55	18	72	39	62
APH	36	78	17	61	53	72	34	71	26	61	60	59	31	63	11	55	42	61	32	84	13	52	45	71
XSPL	32	70	19	68	51	69	37	77	22	58	59	64	30	61	13	65	43	62	24	63	15	60	39	62
XSLi	0	0	5	18	5	1	30	63	18	64	48	73	25	51	12	60	37	54	28	74	13	52	41	65

**Tables 4.7:** Univariate LDA correct classification accuracy (%) for sex using the cervical vertebrae (C3-C6). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	C3						C4						C5						C6					
	Females		Males		Total		Females		Males		Total		Females		Males		Total		Females		Males		Total	
	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
XSLs	34	89	25	69	59	<b>80</b>	34	81	27	71	61	<b>76</b>	28	78	20	61	48	70	23	74	13	41	36	57
XAH	27	71	20	56	47	64	33	79	23	61	56	70	23	64	14	42	37	54	21	68	18	56	39	62
XPH	27	71	23	64	50	68	33	79	21	55	54	68	16	44	20	61	36	52	26	84	18	56	44	70
XALs	28	74	28	78	56	76	36	86	21	55	47	71	20	56	23	70	43	62	19	61	18	56	37	59
XALi	28	74	30	83	58	78	26	62	29	76	55	69	31	86	19	58	50	72	19	61	18	56	37	59
XAWs	25	66	27	75	52	70	23	55	23	61	46	56	21	58	21	64	42	61	15	48	22	69	37	59
XAWi	25	66	27	75	52	70	23	55	23	61	46	58	24	67	21	64	45	65	25	74	18	53	43	63
XPL	24	63	17	47	41	56	16	38	15	39	31	39	11	31	25	76	36	52	14	45	7	22	21	33
TPL	23	61	24	67	47	64	28	67	30	79	58	73	28	78	19	58	47	68	22	71	23	72	45	71
XTPW	28	74	27	75	55	74	30	71	27	71	57	71	29	81	22	67	51	<b>74</b>	26	84	22	89	48	<b>76</b>
XAPWi	26	68	24	67	53	68	29	69	18	47	47	59	27	75	21	64	48	70	22	71	26	81	48	<b>76</b>
APH	22	58	22	61	44	59	23	55	21	55	44	55	27	75	20	61	47	68	19	61	19	59	38	60
XSPL	26	68	21	58	47	64	28	67	22	58	50	63	23	64	19	58	42	61	21	98	19	59	40	63
XSLi	31	82	23	64	54	73	31	74	25	66	56	70	22	61	22	67	44	64	19	61	20	63	39	62

The accuracies of the univariate universal cervical (measurement means of C3 to C6) models were slightly lower than those of the individual univariate models, ranging between 40% and 78% for population affinity and between 37% and 84% for sex estimation (Table 4.8).

**Table 4.8:** Univariate LDA correct classification accuracies (%) for population affinity and sex estimation using the universal cervical model (C3-C6). Bold indicates the highest total accuracy. Refer to Appendix I for measurement names and definitions.

variable	Population						Sex					
	Black		White		Total		Females		Males		Total	
	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
<b>XSLs</b>	109	60	58	55	167	58	112	76	95	68	207	<b>72</b>
<b>XAH</b>	102	56	73	70	175	61	84	57	89	64	173	60
<b>XPH</b>	141	78	66	63	207	72	115	78	74	53	189	66
<b>XALs</b>	131	72	46	44	177	62	124	84	73	53	197	69
<b>XALi</b>	138	76	52	50	190	66	87	59	110	79	197	69
<b>XAWs</b>	121	67	63	60	184	64	95	65	71	51	166	58
<b>XAWi</b>	124	69	63	60	187	65	101	69	74	53	175	61
<b>XPL</b>	72	40	71	68	143	50	54	37	77	55	131	46
<b>TPL</b>	125	69	73	70	198	69	103	70	85	61	188	66
<b>XTPW</b>	117	65	56	53	173	60	97	66	99	71	196	68
<b>XAPWi</b>	136	75	76	72	212	<b>74</b>	96	65	97	70	193	67
<b>APH</b>	114	63	74	70	188	66	91	62	85	61	176	62
<b>XSPL</b>	128	71	57	54	185	65	93	63	77	55	170	59
<b>XSLi</b>	107	59	60	57	167	58	103	70	90	64	193	67

Multivariate models were created by incorporating all of the measurements collected per vertebra. The accuracies ranged between 77.5% and 83.8% for population affinity and between 73% and 82.4% for sex. The multivariate classification accuracies of the universal cervical model were 80.1% and 76.2% for population affinity and sex, respectively. Stepwise variable selection in multivariate models resulted in variable model performances, where different combinations of variables were included in the models for each vertebra. However, maximum pedicle length, maximum inferior articular length, maximum superior articular width, maximum spinous process length, maximum pedicle length, and maximum articular height were frequently removed from the analyses for both population affinity and sex estimation (Table 4.9).

**Table 4.9:** Classification accuracies (%) for the multivariate cervical models and multivariate universal cervical model (C3 – C6) when using all variables and with stepwise variable selection. Refer to Appendix I for measurement names and definitions.

Model	Population				Sex			
	<i>n</i>	LDA (%)	Stepwise (%)	Variables removed	<i>n</i>	LDA (%)	Stepwise (%)	Variables removed
<b>C3</b>	74	82.4	87.9	XPL, XALs, XAWs	74	83.8	86.2	XAWi, XTPW
<b>C4</b>	80	76.3	86.3	XALi, XAWs, XALs, XSPL, XAH	80	77.5	81.3	XPL, XAWi
<b>C5</b>	69	78.2	87.1	APH, XSLi, XPH, XAWs, XALi	69	79.7	85.7	XALs, XAH
<b>C6</b>	63	73	82.9	XALi, XALs, XAH, XPH, XSLi, XSPL	63	77.8	81.2	XAPWi, XALi, XALs, XSLi
<b>Universal cervical model</b>	286	80.1	81.8	XPL, XSLi	286	76.2	81.9	XAH, XPH, XAPi, APH

### 4.3.2 Upper Thoracic Vertebrae

Tables A6 to A9 (see Appendix IV) present the descriptive statistics of the cervical vertebrae, including the sample size, mean and standard deviation for each measurement. White South Africans ( $\mu = 5.26 - 64.93$ ) had overall larger measurement means than black South Africans, ( $\mu = 4.48 - 67.86$ ) and males ( $\mu = 5.03 - 64.46$ ) had larger measurement means than females ( $\mu = 4.69 - 57.88$ ). From the second to sixth thoracic vertebrae, the vertebral body dimensions, specifically the maximum superior sagittal length ( $\mu = 16.56 - 26.40$ ), the maximum inferior sagittal length ( $\mu = 17.83 - 27.35$ ), and the spinous process length ( $\mu = 36.32 - 49.58$ ), increased in size.

ANOVA demonstrated significant differences between black and white South Africans for most of the measurements across all upper thoracic vertebrae, with a few exceptions. Specifically, when assessing the fourth thoracic vertebrae, maximum pedicle length was noted as not significantly different between black and white South Africans. When comparing the sexes, ANOVA demonstrated significant differences for most measurements, except pedicle length, which did not differ significantly when assessing the second and third thoracic vertebrae (Table 4.10).

**Table 4.10:** ANOVA results evaluating population groups and sexes for each measurement for the upper thoracic vertebrae (T2-T6). Bold indicates the significant variables. Refer to Appendix I for measurement names and definitions.

	Population					Sex				
	T2	T3	T4	T5	T6	T2	T3	T4	T5	T6
XSLs	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAH	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XPH	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XALs	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XALi	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAWs	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAWi	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XPL	<0.01	<0.01	0.276	<0.01	<0.01	0.439	0.138	<0.01	<0.01	<0.01
TPL	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XTPW	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAPWi	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
APH	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XSPL	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XSLi	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01

The universal upper thoracic model combined the means of T2 – T6. ANOVA detected significant differences among the measurements of the upper thoracic vertebrae, which demonstrated significant differences between black and white South Africans for most measurements, except for the maximum anterior height, maximum superior articular width, maximum pedicle length and articular process height when assessing black South Africans, and maximum anterior height for white South Africans. When comparing the sexes, maximum anterior height was noted as not significantly different (Table 4.11).

**Table 4.11:** ANOVA results evaluating population groups and sexes for each measurement for all the upper thoracic vertebrae (T2-T6). Bold indicates the significant variables. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Blacks	Whites	Males	Females
XSLs	<0.01	<0.01	<0.01	<0.01
XAH	0.224	0.015	0.142	0.169
XPH	<0.01	<0.01	<0.01	<0.01
XALs	<0.01	<0.01	<0.01	<0.01
XALi	<0.01	<0.01	<0.01	<0.01
XAWs	0.082	<0.01	<0.01	<0.01
XAWi	<0.01	<0.01	<0.01	<0.01
XPL	0.014	<0.01	<0.01	<0.01
TPL	<0.01	<0.01	<0.01	<0.01
XTPW	<0.01	<0.01	<0.01	<0.01
XAPWi	<0.01	<0.01	<0.01	<0.01
APH	0.026	<0.01	<0.01	<0.01
XSPL	<0.01	<0.01	<0.01	<0.01
XSLi	<0.01	<0.01	<0.01	<0.01

When attempting univariate population affinity estimation using LDA in the upper thoracic vertebrae, the classification accuracies ranged between 42% and 84% for the second thoracic vertebra, between 52% and 89% for the third thoracic vertebra, between 51% and 92% for the fourth thoracic vertebra, between 48% and 89% for the fifth thoracic vertebra, and between 48% and 88% for the sixth thoracic vertebra (Table 4.12). The classification accuracies for the univariate models for sex estimation from 22% to 98% for the second thoracic vertebra, from 48% to 89% for the third thoracic vertebra, from 47% to 79% for the fourth thoracic vertebra, from 52% to 82% for the fifth thoracic vertebra, and from 50% to 82% for the sixth thoracic vertebra (Table 4.13).

**Table 4.12:** Univariate LDA correct classification accuracy (%) for population affinity using the upper thoracic vertebrae (T2-T6). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	T2						T3						T4						T5						T6					
	Black		White		Total		Black		White		Total		Black		White		Total		Black		White		Total		Black		White		Total	
	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%
XSLs	39	75	22	50	61	64	42	74	31	62	73	68	44	88	37	79	81	84	40	87	30	71	70	80	42	88	32	78	74	<b>83</b>
XAH	29	56	28	64	57	59	33	58	36	72	69	64	27	54	41	87	68	70	40	87	23	55	63	72	38	79	23	56	61	69
XPH	45	87	23	52	68	71	42	74	28	56	70	65	32	64	41	87	73	75	36	78	32	76	68	77	33	69	36	88	69	78
XALs	39	75	27	61	66	69	34	60	27	54	61	57	41	82	39	83	80	82	41	89	33	79	74	84	37	77	33	80	70	79
XALi	40	77	22	50	62	65	51	89	37	74	88	<b>82</b>	45	90	33	70	78	80	38	83	34	81	72	82	39	81	30	73	69	78
XAWs	36	69	34	77	70	73	34	60	36	70	70	65	36	72	35	74	71	73	35	76	27	64	62	70	39	81	28	68	67	75
XAWi	40	77	27	61	67	70	44	77	27	54	71	66	36	72	30	64	66	68	37	80	28	67	65	74	40	83	87	66	127	75
XPL	39	75	36	82	75	<b>78</b>	38	67	35	70	73	68	27	54	24	51	51	53	25	54	20	48	45	51	23	48	24	59	47	53
TPL	38	73	27	61	65	68	34	60	37	74	71	66	34	68	36	77	70	72	37	80	34	81	71	81	36	75	30	73	66	74
XTPW	35	67	28	64	63	66	39	68	35	70	74	69	41	82	37	79	78	80	30	65	33	79	63	72	36	75	33	80	69	78
XAPWi	39	75	33	75	72	75	41	72	35	70	76	71	38	76	36	77	74	76	31	67	37	88	68	77	37	77	33	80	70	79
APH	28	54	28	64	56	58	43	75	26	52	69	64	39	78	35	74	74	76	30	65	28	67	58	66	36	75	27	66	63	71
XSPL	31	60	66	66	97	63	36	63	30	60	66	62	36	72	34	72	70	72	37	80	30	71	67	76	37	77	32	78	69	78
XSLi	36	69	28	64	64	66	44	77	37	74	81	76	46	92	38	81	84	<b>87</b>	39	85	37	88	76	<b>86</b>	42	88	30	73	72	81

**Table 4.13:** Univariate LDA correct classification accuracy (%) for sex using the upper thoracic vertebrae (T2-T6). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	T2						T3						T4						T5						T6					
	Female		Male		Total		Female		Male		Total		Female		Male		Total		Female		Male		Total		Female		Male		Total	
	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%
XSLs	38	83	27	54	65	68	41	76	33	62	74	69	32	68	28	56	60	62	26	57	30	61	56	64	26	67	25	50	51	57
XAH	33	72	38	76	71	74	32	59	38	72	70	65	22	47	39	78	61	63	20	51	36	73	56	64	32	82	26	52	58	65
XPH	29	63	43	86	72	<b>75</b>	41	76	30	57	71	66	27	57	39	78	66	68	28	72	31	63	59	67	23	59	35	70	58	65
XALs	38	83	32	64	70	73	39	72	40	75	79	74	32	68	33	66	65	67	29	74	28	57	57	64	25	64	30	60	55	62
XALi	39	85	27	54	66	69	40	74	29	55	69	64	37	79	28	56	65	67	29	74	32	65	61	69	29	74	29	58	58	65
XAWs	30	65	34	68	64	67	36	67	41	77	77	72	35	74	37	74	72	<b>74</b>	32	82	31	63	63	72	30	77	28	56	58	65
XAWi	37	80	30	60	67	70	48	89	34	64	82	<b>77</b>	36	77	33	66	69	71	28	72	34	69	62	70	29	74	25	50	54	61
XPL	30	65	31	62	61	64	26	48	26	49	52	49	31	66	31	62	62	64	26	67	29	59	55	63	24	62	36	72	60	<b>67</b>
TPL	28	61	39	78	67	70	37	69	43	81	80	75	31	70	36	72	67	69	26	67	38	78	64	73	28	72	31	62	59	62
XTPW	29	63	36	72	65	68	40	74	39	74	79	74	35	74	34	68	69	71	26	67	39	80	65	74	24	62	34	68	58	65
XAPWi	34	74	34	68	68	71	38	70	35	66	73	68	34	72	35	70	69	71	23	59	36	73	59	67	22	56	35	70	57	64
APH	31	67	37	74	68	71	43	80	29	55	72	67	34	72	33	66	67	69	29	74	38	78	67	<b>76</b>	22	56	34	68	56	62
XSPL	31	67	35	70	66	69	37	69	34	54	71	66	32	68	33	66	65	67	25	64	33	67	58	70	25	64	29	58	54	61
XSLi	34	74	32	64	66	69	38	70	34	67	72	67	32	68	27	54	59	61	26	67	31	63	57	65	21	54	27	54	48	54

Compared to the individual univariate upper thoracic models, the accuracy of the universal upper thoracic model was slightly lower, with correct classification ranging between 54% and 79% for population affinity and between 57% and 79% for sex estimation (Table 4.14).

**Table 4.14:** Univariate LDA correct classification accuracy (%) for population affinity and sex using the upper thoracic vertebrae (T2-T6). Bold indicates the highest total accuracy. Refer to Appendix I for measurement names and definitions.

variable	Population						Sex					
	Black		White		Total		Females		Males		Total	
	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
<b>XSLs</b>	192	76	135	60	327	68	135	60	159	63	294	62
<b>XAH</b>	141	56	178	79	319	67	128	57	193	77	321	67
<b>XPH</b>	163	64	169	75	332	69	143	63	177	70	320	67
<b>XALs</b>	183	72	144	64	327	68	157	69	146	58	303	63
<b>XALi</b>	206	81	135	60	341	71	146	65	156	62	302	63
<b>XAWs</b>	191	75	142	63	333	70	179	79	158	63	337	<b>71</b>
<b>XAWi</b>	167	66	162	72	329	69	159	69	179	71	338	70
<b>XPL</b>	136	54	150	67	286	60	111	49	153	61	264	55
<b>TPL</b>	172	68	152	68	324	68	152	67	179	71	331	69
<b>XTPW</b>	183	72	172	77	355	<b>74</b>	155	69	172	68	327	68
<b>XAPWi</b>	188	74	158	70	346	72	139	62	185	73	324	68
<b>APH</b>	167	66	152	68	319	67	158	70	171	68	329	69
<b>XSPL</b>	176	69	134	60	310	65	163	72	149	59	312	65
<b>XSLi</b>	203	80	141	62	344	72	145	64	152	60	297	62

Multivariate models were created incorporating all the measurements collected per vertebra for the upper thoracic vertebrae. Classification accuracies for the multivariate models ranged between 67% and 81.3% for population affinity and between 78.5% and 91.1% for sex estimation. The multivariate classification accuracies of the universal upper thoracic model were 80.5% and 67.6% for population affinity and sex, respectively. When employing stepwise variable selection for the individual multivariate models and universal models, different combinations of variables were included in the models for each vertebra; however, maximum superior sagittal length, maximum inferior articular length, maximum superior articular width, maximum inferior articular width, maximum transverse process width, maximum spinous process length, were frequently removed from the analyses for both population affinity and sex estimation (Table 4.15).

**Table 4.15:** Classification accuracies (%) for each multivariate upper thoracic models and multivariate universal upper thoracic model (T2-T6) for population affinity and sex. Refer to Appendix I for measurement names and definitions.

Model	Population				Sex			
	<i>n</i>	LDA (%)	Stepwise (%)	Variables removed	<i>n</i>	LDA (%)	Stepwise (%)	Variables removed
<b>T2</b>	96	86.4	86.7	XSLs, XALi	96	74	81.4	APH, XSPL, XAWs, XAPWi
<b>T3</b>	107	78.5	83.5	XSLs, XALi	107	81.3	85.3	XAH, APH, XALi, XAPi
<b>T4</b>	97	81.4	88.7	XAWs, XSLi, XSLs, XALi	97	70.1	81.4	XAWi, XAH, APH, XAPWi, XALi, XTPW
<b>T5</b>	88	81.8	88.8	TPL, XAWi, XTPW	88	67	75.1	XAWs, XSLs
<b>T6</b>	89	85.3	91.1	XALi	89	76.4	83.2	XSLs, XAPi, XSPL
<b>Universal upper thoracic model</b>	478	80.5	82.6	XSPL, XAH, TPL	478	67.6	75.5	XAWi, APH

### 4.3.3 Lower Thoracic Vertebrae

In the lower thoracic vertebrae, the mean and standard deviations differed between white and black South Africans and between males and females. Table A11 to A14 (see Appendix IV) presents the descriptive statistics of the lower thoracic vertebrae, including the sample size, mean and standard deviation for each measurement. The results demonstrated that white South Africans ( $\mu = 5.41 - 64.69$ ) had larger measurements in comparison to black South Africans ( $\mu = 4.88 - 57.66$ ). In addition, when comparing sexes, males ( $\mu = 5.17 - 64.23$ ) had larger measurements than females ( $\mu = 5.10 - 57.70$ ). From the seventh thoracic vertebrae to the eleventh thoracic vertebrae, the dimensions of the vertebral bodies, specifically the maximum superior sagittal length ( $\mu = 23.30 - 31.27$ ), maximum anterior height ( $\mu = 18.07 - 22.44$ ), maximum posterior height ( $\mu = 19.23 - 25.12$ ), maximum inferior articular width ( $\mu = 25.73 - 38.33$ ), articular process height ( $\mu = 32.76 - 40.98$ ), and maximum inferior sagittal length ( $\mu = 23.98 - 31.96$ ), increased from the seventh thoracic vertebrae to the eleventh thoracic vertebrae. Transverse process length ( $\mu = 31.93 - 24.17$ ), maximum transverse process width ( $\mu = 62.83 - 48.65$ ) and spinous process length ( $\mu = 49.47 - 32.93$ ) decreased from the seventh thoracic vertebrae to the eleventh thoracic vertebrae.

ANOVA demonstrated significant differences between black and white South Africans for the majority of the measurements across all of the lower thoracic vertebrae, with a few exceptions.

Specifically, maximum pedicle length was noted as not significantly different between the population groups when assessing the seventh and eighth thoracic vertebrae. Additionally, maximum inferior articular process width was noted as not significantly different between black and white South Africans for the eleventh thoracic vertebra. When comparing the sexes, ANOVA demonstrated significant differences between males and females for most of the measurements across the upper thoracic vertebrae (T7-T11). Maximum pedicle length was noted as not significantly different when assessing the tenth thoracic vertebrae. Furthermore, the maximum anterior height and articular process height of the eleventh thoracic vertebrae did not differ significantly between the sexes (Table 4.16).

**Table 4.16:** ANOVA results evaluating population groups and sexes for each measurement for lower thoracic vertebrae (T7-T11). Bold indicates the significant variables. Refer to Appendix I for measurement names and definitions.

	Population					Sex				
	T7	T8	T9	T10	T11	T7	T8	T9	T10	T11
XSLs	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAH	<0.01	<0.01	<0.01	0.011	<0.01	<0.01	<0.01	<0.01	<0.01	0.109
XPH	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XALs	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XALi	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAWs	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAWi	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XPL	0.554	0.448	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.648	<0.01
TPL	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XTPW	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XAPWi	<0.01	<0.01	<0.01	<0.01	0.054	<0.01	<0.01	<0.01	<0.01	<0.01
APH	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.026
XSPL	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
XSLi	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01

With the universal lower thoracic model, ANOVA demonstrated significant differences between black and white South Africans for the majority of the measurements, except for maximum anterior height, maximum superior articular width, maximum pedicle length and articular process height for black South Africans. When comparing the sexes, maximum anterior height and maximum pedicle length were noted as not significantly different between the sexes (Table 4.17).

**Table 4.17:** ANOVA results evaluating population affinity and sex for each measurement for all the lower thoracic vertebrae (T7-T11). Bold indicates the significant variables. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Blacks	Whites	Males	Females
XSLs	<0.01	<0.01	<0.01	<0.01
XAH	0.281	<0.01	0.113	0.152
XPH	<0.01	<0.01	<0.01	<0.01
XALs	<0.01	<0.01	<0.01	<0.01
XALi	<0.01	<0.01	<0.01	<0.01
XAWs	0.082	<0.01	<0.01	<0.01
XAWi	<0.01	<0.01	<0.01	<0.01
XPL	0.016	<0.01	0.081	0.075
TPL	<0.01	<0.01	<0.01	<0.01
XTPW	<0.01	<0.01	<0.01	<0.01
XAPWi	<0.01	<0.01	<0.01	<0.01
APH	0.026	<0.01	<0.01	<0.01
XSPL	<0.01	<0.01	<0.01	<0.01
XSLi	<0.01	<0.01	<0.01	<0.01

When attempting to classify the individuals using univariate models for the lower thoracic vertebrae, the population affinity estimation accuracy ranged from 25% to 94% for the seventh thoracic vertebrae, from 44% to 89% for the eighth thoracic vertebrae, from 54% to 92% for the ninth thoracic vertebrae, from 48% to 93% for the tenth thoracic vertebrae, and from 50% to 92% for the eleventh thoracic vertebrae (Table 4.18). For sex estimation, the univariate models yielded accuracies between 51% and 87% for the seventh thoracic vertebrae, between 44% and 85% for the eighth thoracic vertebrae, between 45% and 80% for the ninth thoracic vertebrae, between 34% and 81% for the tenth thoracic vertebrae, and between 38% and 85% for the eleventh thoracic vertebrae, respectively (Table 4.19).

**Tables 4.18:** Univariate LDA correct classification accuracy (%) for population affinity using the lower thoracic vertebrae (T7-T11). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	T7						T8						T9						T10						T11					
	Black		White		Total		Black		White		Total		Black		White		Total		Black		White		Total		Black		White		Total	
	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%
XSLs	47	92	24	63	71	<b>80</b>	41	77	33	77	74	77	42	86	32	76	74	81	50	93	30	71	80	83	44	85	39	83	83	<b>84</b>
XAH	36	71	24	63	60	67	35	66	20	47	55	57	36	73	20	78	56	52	36	67	24	57	60	63	28	84	33	70	61	62
XPH	41	80	25	66	66	74	45	85	27	63	72	75	35	71	33	79	68	75	39	72	30	71	69	72	43	83	29	62	72	73
XALs	40	78	30	79	70	79	46	87	32	74	78	<b>81</b>	44	90	28	67	72	79	49	91	32	76	81	84	45	87	35	74	80	81
XALi	42	82	27	71	69	78	47	89	25	58	72	75	44	90	30	71	74	81	48	89	33	79	81	84	47	90	34	72	81	82
XAWs	40	78	28	74	68	76	44	83	29	67	73	76	35	71	32	76	67	74	42	78	30	71	72	75	38	73	29	62	67	68
XAWi	42	82	24	63	66	74	45	85	28	65	73	76	39	80	26	62	65	71	42	78	28	67	70	73	36	69	34	72	70	71
XPL	18	35	25	66	43	48	37	70	16	37	53	55	36	73	23	55	59	65	44	81	20	48	64	67	32	62	29	62	61	62
TPL	39	76	26	68	65	73	34	64	25	58	59	61	33	67	31	74	64	70	36	67	25	60	61	64	33	63	29	62	62	63
XTPW	38	75	27	71	65	73	36	68	30	70	66	69	34	69	30	71	64	70	35	65	28	67	63	66	30	58	30	64	60	61
XAPWi	40	78	31	82	71	<b>80</b>	40	75	34	79	74	77	35	71	33	79	68	75	35	65	26	62	61	64	26	50	29	62	55	56
APH	30	59	30	79	60	67	32	60	30	70	62	65	35	71	29	69	64	70	39	72	21	50	60	63	36	69	30	64	66	67
XSPL	41	80	27	71	68	76	44	83	28	65	72	75	32	65	26	62	58	64	36	67	25	60	61	64	34	65	26	55	60	61
XSLi	45	88	26	68	71	<b>80</b>	45	85	32	85	77	80	45	92	30	71	75	<b>82</b>	49	91	33	79	82	<b>85</b>	48	92	33	70	81	82

**Tables 4.19:** Univariate LDA correct classification accuracy (%) for sex using the lower thoracic vertebrae (T7-T11). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	T7						T8						T9						T10						T11					
	Female		Male		Total		Female		Male		Total		Female		Male		Total		Female		Male		Total		Female		Male		Total	
	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%
XSLs	26	59	29	64	55	62	29	69	32	59	61	64	32	73	37	57	69	65	33	70	31	63	64	67	30	64	30	58	60	61
XAH	31	70	26	58	57	64	33	79	29	54	62	65	24	55	21	45	45	49	34	72	29	59	63	66	22	47	32	62	54	55
XPH	34	77	25	56	59	66	23	55	41	76	64	67	25	57	28	60	53	58	28	60	26	53	54	56	29	62	20	38	49	49
XALs	30	68	27	60	57	64	32	76	29	54	61	64	34	77	30	64	64	70	35	74	25	51	60	63	34	72	29	56	63	64
XALi	33	75	25	56	58	65	30	71	33	61	63	66	35	80	26	55	61	67	35	74	27	55	62	65	35	74	27	52	62	63
XAWs	32	73	27	60	59	66	33	79	29	54	62	65	31	70	33	70	64	70	33	70	28	57	61	64	36	77	32	62	68	69
XAWi	26	59	32	71	58	65	32	76	35	65	67	70	30	68	30	64	60	66	38	81	31	63	69	72	32	68	35	67	67	68
XPL	25	57	39	87	64	72	34	81	24	44	58	60	32	73	24	51	56	62	16	34	0	0	16	16	32	68	35	67	67	68
TPL	35	80	29	64	64	72	32	76	46	85	78	81	33	75	36	77	69	76	38	81	34	69	72	75	33	70	44	85	77	78
XTPW	34	77	30	67	64	72	31	74	41	76	72	75	34	77	35	75	69	76	36	77	36	73	72	75	33	70	38	73	71	72
XAPWi	29	66	27	60	56	63	26	62	37	69	63	66	30	68	33	70	63	69	29	62	34	69	63	66	29	62	37	71	66	67
APH	27	61	34	76	61	69	27	64	36	67	63	66	29	66	28	60	57	63	28	60	29	59	57	59	26	55	25	48	51	52
XSPL	32	73	29	64	61	69	29	69	36	67	65	68	29	66	29	62	58	64	35	74	31	63	66	69	33	70	30	58	63	64
XSLi	26	59	23	51	49	55	29	69	27	50	56	58	32	73	24	51	56	62	35	74	26	53	61	64	35	74	25	48	60	61

The classification accuracy of the universal lower thoracic model was slightly lower than the univariate lower thoracic models, with population affinity estimation ranging from 53% to 81% and sex estimation ranging from 49% to 77% (Table 4.20).

**Table 4.20:** Univariate LDA correct classification accuracy (%) for population affinity and sex using the universal lower thoracic model (T7-T11). Bold indicates the highest total accuracy. Refer to Appendix I for measurement names and definitions.

variable	Population						Sex					
	Black		White		Total		Females		Males		Total	
	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
<b>XSLs</b>	190	76	135	60	325	68	133	59	158	63	291	61
<b>XAH</b>	139	55	178	79	283	67	127	57	193	77	320	67
<b>XPH</b>	162	65	168	75	330	69	142	63	175	70	317	67
<b>XALs</b>	181	72	144	64	325	68	155	69	145	58	300	63
<b>XALi</b>	204	81	134	60	338	71	144	64	155	62	299	63
<b>XAWs</b>	188	75	142	63	330	69	177	79	158	63	335	<b>71</b>
<b>XAWi</b>	164	65	161	72	325	68	154	69	178	71	332	70
<b>XPL</b>	134	53	150	67	284	60	109	49	152	61	261	55
<b>TPL</b>	170	68	170	76	340	72	151	67	178	71	329	69
<b>XTPW</b>	180	72	171	76	351	<b>74</b>	153	68	171	68	324	68
<b>XAPWi</b>	186	74	158	71	344	72	157	70	156	62	313	66
<b>APH</b>	166	66	152	58	318	67	158	71	171	68	329	69
<b>XSPL</b>	173	69	133	59	306	64	161	72	148	59	309	65
<b>XSLi</b>	200	80	141	63	341	72	143	64	151	60	294	62

The measurements per vertebra for the lower thoracic vertebrae were combined into multivariate models, where the classification accuracies ranged between 69.2% and 80% for population affinity estimation and between 78.8% and 89.6% for sex estimation. A variety of combinations of variables were used in stepwise variable selection for each vertebra; however, maximum superior sagittal length, maximum anterior height, maximum anterior height, maximum inferior articular width, maximum transverse process width, maximum spinous process length, and articular process height were frequently removed from the analyses for both population affinity and sex estimation in the multivariate models (Table 4.21).

With the universal multivariate lower thoracic model, the classification accuracies were slightly lower compared to the multivariate models. The universal multivariate classification accuracies were 79.2% and 70.9% for population affinity and sex, respectively. In the universal lower thoracic model, stepwise variables were selected for each vertebra. However, maximum posterior height, maximum superior articular length, and maximum spinous process length were frequently removed from the analyses for sex, while maximum posterior height, maximum inferior articular length, maximum inferior articular process width and articular process height were frequently removed from the analyses for population affinity (Table 4.21).

**Table 4.21:** Classification accuracies (%) for each multivariate lower thoracic models and multivariate universal lower thoracic model (T7-T11) for population affinity and sex. Refer to Appendix I for measurement names and definitions.

Model	n	Population			n	Sex		
		LDA (%)	Stepwise (%)	Variables removed		LDA (%)	Stepwise (%)	Variables removed
<b>T7</b>	89	82	87.4	XSPL, XAH, XPL, XAWi, XAWs, XSLs	89	71.9	81.1	XPH, XSPL, APH, XALs
<b>T8</b>	96	89.6	90.8	XSPL	96	80	81.6	XAH, XALs, XAWi, XALi
<b>T9</b>	91	80.2	83.7	XALi, XAWi	91	69.2	79.2	XAPWi, XPH, XSLs
<b>T10</b>	96	83.3	87.6	XTPW, XAWi	96	71.9	79.1	XAWs, XTPW, XSPL
<b>T11</b>	99	78.8	85	APH, XSPL, XAH	99	72.7	81.8	XTPW, XSLs, TPL, XAH
<b>All lower thoracic</b>	475	79.2	81.7	XPH, XALs, XSPL	475	70.9	74.3	XPH, XALi, XAPWi, APH

#### 4.3.4 Lumbar Vertebrae

A comparison of the means and standard deviations of the lumbar vertebrae in white and black South African males and females revealed considerable variation. Table A11 to A14 (see Appendix IV) presents the descriptive statistics of the cervical vertebra, including the sample size, mean and standard deviation for each measurement. Overall, white South Africans ( $\mu = 8.97 - 77.01$ ) have the larger measurement means compared to black South Africans ( $\mu = 7.72 - 76.68$ ). Overall, males ( $\mu = 7.87 - 80.42$ ) have greater variability and greater measurement means compared to females ( $\mu = 8.83 - 72.84$ ). Several dimensions of the vertebral body, namely the maximum superior sagittal length ( $\mu = 27.26 - 35.23$ ), maximum superior articular length ( $\mu = 23.05 - 31.19$ ), maximum inferior articular length ( $\mu = 24.66 - 31.65$ ), maximum superior articular width ( $\mu = 37.23 - 46.92$ ), maximum inferior articular width ( $\mu = 39.41 - 49.45$ ), and maximum inferior sagittal length ( $\mu = 28.33 - 35.42$ ) increased the lower the vertebrae were positioned in the spine, from the first lumbar vertebrae to the fourth lumbar vertebrae. Additionally, maximum transverse process width ( $\mu = 31.38 - 44.87$ ), maximum pedicle length ( $\mu = 6.96 - 11.38$ ), and maximum inferior articular process width ( $\mu = 26.81 - 44.05$ ) increased from the first lumbar vertebrae to the fourth lumbar vertebrae.

In most of the measurements across the lumbar vertebrae, ANOVA showed significant differences between black and white South Africans. However, the maximum pedicle length of the third and fourth lumbar vertebrae did not differ significantly for population affinity.

Additionally, maximum inferior articular process width was noted as not significantly different between black and white South Africans, when assessing the third lumbar vertebrae and maximum transverse process width when assessing all the lumbar vertebrae. When comparing the sexes, ANOVA demonstrated significant differences between males and females for most of the measurements across all of the lumbar vertebrae, except for maximum anterior height, which was noted as not significantly different when assessing all the lumbar vertebrae. In addition, multiple measurements did not differ significantly when assessing the lumbar vertebrae, specifically the maximum pedicle length and maximum transverse process width when assessing the first lumbar vertebrae, maximum pedicle length when assessing the second lumbar vertebrae, maximum inferior articular process width when assessing the third lumbar vertebrae and maximum transverse process width and articular process height when assessing the fourth lumbar vertebrae (Table 4.22).

**Table 4.22:** ANOVA results evaluating population affinity and sex for each measurement for lumbar vertebrae (L1-L4). Bold indicates the significant variables. Refer to Appendix I for measurement names and definitions.

	Population				Sex			
	L1	L2	L3	L4	L1	L2	L3	L4
<b>XSLs</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XAH</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.469	0.948	0.462	0.202
<b>XPH</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XALs</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XALi</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XAWs</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XAWi</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XPL</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.312	0.946	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>TPL</b>	<b>&lt;0.01</b>	0.085	0.783	0.710	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XTPW</b>	0.074	0.351	0.273	0.754	0.031	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.042
<b>XAPWi</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.256	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.219	<b>&lt;0.01</b>
<b>APH</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	0.041
<b>XSPL</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>
<b>XSLi</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>

For the universal lumbar model, ANOVA showed significant differences between black and white South Africans for most of the measurements, except for maximum anterior height, maximum superior articular width, maximum pedicle length, and articular process height when assessing black South Africans. When comparing the sexes, maximum anterior height and maximum pedicle length were noted as not significantly different for either of the sexes (Table 4.23).

**Table 4.23:** ANOVA results evaluating population affinity and sex for each measurement for the universal lumbar model (L1-L4). Bold indicates the significant variables. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Blacks	Whites	Males	Females
XSLs	<0.01	<0.01	<0.01	<0.01
XAH	<0.01	<0.01	<0.01	<0.01
XPH	<0.01	0.091	0.070	<0.01
XALs	<0.01	<0.01	<0.01	<0.01
XALi	<0.01	<0.01	<0.01	<0.01
XAWs	<0.01	<0.01	<0.01	<0.01
XAWi	<0.01	<0.01	<0.01	<0.01
XPL	0.036	<0.01	<0.01	<0.01
TPL	<0.01	<0.01	<0.01	<0.01
XTPW	<0.01	<0.01	<0.01	<0.01
XAPWi	<0.01	<0.01	<0.01	<0.01
APH	<0.01	<0.01	<0.01	<0.01
XSPL	<0.01	<0.01	<0.01	<0.01
XSLi	<0.01	<0.01	<0.01	<0.01

When assessing the lumbar vertebrae, univariate population affinity estimation revealed classification accuracies between 48% and 93% for the first lumbar vertebrae, between 52% and 81% for the second lumbar vertebrae, between 47% and 82% for the third lumbar vertebrae, and between 14% and 87% for the fourth lumbar vertebrae (Table 4.24). When comparing the sexes, the univariate lumbar models demonstrated sex classification accuracies between 0% and 85% for the first lumbar vertebrae, between 13% and 77% for the second lumbar vertebrae, between 35% and 79% for the third lumbar vertebrae, and between 48% and 81% for the fourth lumbar vertebrae (Table 4.25).

**Tables 4.24:** Univariate LDA correct classification accuracy (%) for population affinity using the lumbar vertebrae (L1-L4). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	L1						L2						L3						L4					
	Black		White		Total		Black		White		Total		Black		White		Total		Black		White		Total	
	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%
XSLs	41	93	19	66	60	82	28	78	15	65	43	73	27	79	12	71	39	76	20	71	11	73	31	72
XAH	33	75	19	66	52	71	29	81	17	74	46	78	26	76	11	65	37	73	22	79	9	60	31	72
XPH	31	71	16	55	47	64	23	64	18	78	41	65	21	62	12	71	33	65	17	61	9	60	26	61
XALs	38	86	21	72	59	81	28	78	17	74	45	76	28	82	12	71	40	78	23	82	13	87	36	84
XALi	34	77	20	69	54	74	25	69	16	70	41	69	22	65	12	71	34	67	20	71	9	60	29	67
XAWs	29	66	18	62	47	64	22	61	13	57	35	59	18	53	8	47	26	51	20	71	9	60	29	67
XAWi	30	68	16	55	46	63	24	67	14	61	38	64	24	71	9	53	33	65	23	82	11	73	34	79
XPL	32	73	16	55	48	66	26	72	14	61	40	68	27	79	9	53	36	71	19	68	13	87	32	74
TPL	24	55	19	66	43	59	20	56	13	57	33	56	19	56	10	59	29	57	14	50	11	73	25	58
XTPW	22	50	20	69	42	58	19	53	12	52	31	53	23	68	10	59	33	65	4	14	4	27	8	19
XAPWi	27	61	14	48	41	56	20	56	13	57	33	56	17	50	8	47	25	49	18	64	9	60	27	63
APH	28	64	17	59	45	62	21	58	12	52	33	56	19	56	11	65	30	59	18	64	10	67	28	65
XSPL	29	66	14	48	43	59	24	57	13	57	37	63	24	71	13	77	37	73	21	75	11	73	32	74
XSLi	38	86	19	66	57	78	28	78	14	61	42	71	23	67	11	65	34	67	17	61	10	67	27	63

**Tables 4.25:** Univariate LDA correct classification accuracy (%) for sex using the lumbar vertebrae (L1-L4). Bold indicates the highest total accuracy per vertebra. Refer to Appendix I for measurement names and definitions.

variable	L1						L2						L3						L4					
	Females		Males		Total		Females		Males		Total		Females		Males		Total		Females		Males		Total	
	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%	n	%
XSLs	25	64	22	65	47	64	22	71	14	50	36	61	14	61	17	61	31	61	10	63	13	48	23	54
XAH	22	56	13	38	35	48	16	52	20	71	36	61	11	48	20	71	31	61	8	50	18	67	26	61
XPH	27	69	17	50	44	60	4	13	0	0	4	7	12	52	14	50	26	51	12	75	16	59	28	65
XALs	30	77	18	53	48	66	22	71	16	57	38	64	16	70	13	46	29	57	10	63	16	59	26	61
XALi	30	77	21	62	51	70	21	68	17	61	38	64	17	74	18	64	35	69	10	63	16	59	26	61
XAWs	30	77	24	71	54	74	23	74	19	68	42	71	15	65	21	75	36	71	12	75	19	70	31	72
XAWi	33	85	24	71	57	78	24	77	19	68	43	73	16	70	18	64	34	67	13	81	16	59	29	67
XPL	0	0	13	38	13	18	15	48	19	68	34	58	17	74	20	71	37	73	13	81	18	67	31	72
TPL	21	54	21	62	42	58	20	65	18	64	38	64	17	74	20	71	37	73	9	56	18	67	27	63
XTPW	18	46	22	65	40	55	20	65	18	64	38	64	15	65	22	79	37	73	9	56	18	67	27	63
XAPWi	28	72	20	59	48	66	19	61	17	61	36	61	13	57	15	54	28	55	11	69	20	74	31	72
APH	27	69	21	62	48	66	17	55	13	46	30	51	14	61	17	61	31	61	9	56	13	48	22	51
XSPL	29	74	19	56	48	66	16	52	15	54	31	53	8	35	17	61	25	49	9	56	16	59	25	58
XSLi	26	67	25	74	51	70	21	68	12	43	33	56	16	70	15	54	31	61	11	69	16	59	27	63

In the universal univariate lumbar model, classification accuracies were slightly lower than those from the individual univariate models, with estimates ranging from 40% to 82% for the population affinity and 45% to 75% for the sexes (Table 4.26).

**Table 4.26:** Univariate LDA correct classification accuracy (%) for population affinity and sex using the universal lumbar model (L1-L4). Bold indicates the highest total accuracy. Refer to Appendix I for measurement names and definitions.

variable	Population						Sex					
	Black		White		Total		Females		Males		Total	
	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
XSLs	108	76	53	63	161	71	70	64	73	62	143	63
XAH	101	71	61	72	162	72	49	45	64	55	113	50
XPH	98	69	55	65	153	68	65	60	55	47	120	53
XALs	117	82	58	69	175	<b>77</b>	72	66	72	62	144	64
XALi	106	75	50	60	156	69	69	63	77	66	146	65
XAWs	87	61	48	57	135	60	78	72	88	75	166	<b>73</b>
XAWi	83	58	57	68	140	62	76	70	83	71	159	70
XPL	98	69	59	70	157	69	64	59	78	57	142	63
TPL	70	49	40	48	110	49	71	65	76	65	147	65
XTPW	73	51	36	43	109	48	62	57	73	62	135	60
XAPWi	86	61	34	40	120	53	77	71	58	50	135	60
APH	77	54	56	67	133	59	62	57	74	63	136	60
XSPL	96	68	60	71	156	69	89	63	66	56	155	60
XSLi	95	67	55	65	150	66	76	70	69	60	145	64

Based on all the measurements collected for each lumbar vertebra, multivariate lumbar models were created. For population affinity estimation the classification accuracies ranged between 69.2% and 80%, and between 78.8% and 89.6% for sex estimation. Stepwise variable selection in the multivariate lumbar models was employed and removed different combinations of variables included for each vertebra, except for the third lumbar vertebra which removed no variables. However, maximum inferior articular length, maximum transverse process width, maximum superior sagittal length, transverse process length and maximum pedicle length were frequently removed from the analyses for both population affinity and sex estimation.

The classification accuracy was slightly lower when comparing the multivariate universal lumbar model with the multivariate lumbar models. The universal multivariate classification accuracies were 79.2% and 70.9% for population affinity and sex, respectively. When employing stepwise variable selection, different combinations of variables were included in the multivariate universal lumbar model; however, maximum spinous process length, maximum inferior sagittal length and maximum inferior articular process width were frequently removed from the analyses for population affinity, while articular process height,

transverse process length and maximum inferior articular length were frequently removed for sex estimation (Table 4.27).

**Table 4.27:** Classification accuracies (%) for each multivariate lumbar models and multivariate universal lumbar model (L1-L4) for population affinity and sex. Refer to Appendix I for measurement names and definitions.

Model	Population				Sex			
	<i>n</i>	LDA (%)	Stepwise (%)	Variables removed	<i>n</i>	LDA (%)	Stepwise (%)	Variables removed
L1	73	79.5	88	XALi, XTPW	73	64.4	77.1	XAPWi, XAWi, APH, XPL, XALs, XTPW
L2	59	84.7	89.7	XSLs, TPL, XPL	59	76.3	84.3	TPL, APH, XSPL, XAPi
L3	51	88.2	90.7	None	51	76.5	86.3	APH, XPH, XSLi
L4	43	88.4	93.5	XAH, XAWi	43	74.4	88.5	XTPW
Universal lumbar model	226	84.1	87.2	XSPL, XSLi, XAPWi	226	80	82.4	APH, TPL, XALi

#### 4.4 Summary

- The vertebral body and spinous process dimensions were noted to increase in size, except for the lower thoracic region, where spinous process length and transverse process dimensions decreased in size.
- Population affinity and sex classification accuracies were high, with populations classifying better when using the thoracic and lumbar vertebrae and sex classifying better when using the cervical vertebrae.
- Univariate models demonstrated the highest classification accuracies compared to multivariate models, with cervical (98% for C6) and thoracic (92% for T4 and T9) vertebrae performing the best.
- Individual multivariate vertebrae models (i.e., L1, L2, L3, L4), particularly for the lumbar vertebrae, classified the best, specifically the third (90.7% and 86.3%) and fourth (93.5% and 88.5%) lumbar vertebrae for both population affinity and sex estimation.
- Multivariate universal models (for each vertebral sub-type) indicated that the upper thoracic (82.6% and 75.5%), lower thoracic (81.7% and 74.3%) and lumbar (87.2% and 82.4%) vertebrae performed the best when classifying according to population affinity and sex estimation.

## CHAPTER 5: DISCUSSION

Several researchers have studied the vertebrae with regard to sexual dimorphism and population variation (Cheng *et al.*, 1998; Wescott, 2000; Pastor, 2005; Soegiharto *et al.*, 2008; Yu *et al.*, 2008; Marlow *et al.*, 2011; Gama *et al.*, 2015; Ostrofsky *et al.*, 2015; Torimitsu *et al.*, 2016; Montasser *et al.*, 2017; Ramadan *et al.*, 2017; Oura *et al.*, 2018; Rozendaal *et al.*, 2020). However, to date, a very limited number of vertebrae have been investigated (i.e., first and second cervical vertebrae, first and twelfth thoracic vertebrae and the fifth lumbar vertebrae) with the focus mainly placed on sexual dimorphism (Cheng *et al.*, 1998; Wescott, 2000; Pastor, 2005; Soegiharto *et al.*, 2008; Yu *et al.*, 2008; Marlow *et al.*, 2011; Gama *et al.*, 2015; Montasser *et al.*, 2017; Oura *et al.*, 2018). The current study aimed to expand on previous research, as previous studies lack universal standards for estimating population affinity and sex using vertebral measurements. Two components were included in the study to effectively investigate sexual dimorphism and population variation among modern South Africans: (1) an exploration component, and (2) a classification component. To determine whether universal vertebral standards for population affinity and sex estimation can be created, the variation exploration component evaluated the size variation among vertebrae both within and among vertebral types (cervical, thoracic, and lumbar vertebrae). Classification models were created based on the information provided by the variation exploration component to estimate population affinity and sex.

In exploring the vertebral variation, the results demonstrated that both the atypical vertebrae (C7, T1 and T12) as well as the vertebrae closest to the atypical vertebrae (i.e., C6, T2, T11, L1 and L4) showed vast differences among the vertebrae and measurements. Because these vertebrae are so different to the rest of the vertebrae, the specific vertebrae will have to be correctly identified prior to further analyses and they will not be useful to establish universal standards. Atypical vertebrae (i.e., C6, T2, T11, L1 and L4) retain characteristics of the preceding vertebrae in the vertebral column, for example, the first thoracic vertebra retains characteristics of the cervical vertebrae such as the spinous process that projects more horizontally and vertebral body that presents more saddle-shaped and interlocking than thoracic vertebrae (White *et al.*, 2005). However, the first lumbar vertebra was included in the sample to explore the lumbar variation attributable to sexual dimorphism and population affinity. Previous research indicated that the first lumbar vertebra produced high accuracies for sex; therefore, the first lumbar vertebra was included to establish whether the first lumbar vertebra

would produce similar accuracies for estimating population affinity as well as to compare accuracies to the remaining lumbar vertebrae.

Multiple measurements showed repeatability issues, with errors greater than 7%. Most of the measurements that presented with large errors were small measurements, where even a small margin of error results in very high %TEM levels. Additionally, the high error rates can be ascribed to difficulty in locating the correct landmark to place the caliper, as the area on the bone (such as the pedicle) tends to be highly variable among and within individuals. To minimize measurement error, it is necessary to use the appropriate instrumentation, understand the measurement definition, and use highly reliable and repeatable measurements. It was noted that minimal experience was required in these measurements. However, knowledge of vertebral landmarks is required as landmarks were difficult to locate on some measurements due to the measurements, additionally, the measurements are not typically used by forensic anthropologists. After the variation of the vertebrae was assessed, a total of 14 measurements were retained for the creation of multivariate and univariate models to estimate population affinity and sex. Multiple multivariate and univariate models were created using all variables for a single vertebra as well as a universal model using all variables for combined sub-type vertebrae (i.e., cervical (C3-C6), upper thoracic (T2-T6), lower thoracic (T7-T11) and lumbar vertebrae (L1-L4)). The universal models were created for vertebrae to be able to estimate population affinity and sex without defining the specific vertebral number.

A series of different models were created to thoroughly capture the variation and explore the potential of the vertebrae when estimating population affinity and sex. While the literature has shown that multivariate models typically produce the best results (Liebenberg *et al.*, 2015), the availability of univariate standards can also be beneficial to biological anthropologists in instances where bones are taphonomically altered or fragmentary. Two iterations of all models were analysed. First, classification models (both univariate and multivariate) were created for each vertebra on its own to determine whether the classification accuracies would be higher if the specific number of the vertebra within the sequence of the vertebral column is known. Second, classification models (both univariate and multivariate) were created for each category of the vertebra to determine if universal discriminant functions can be used to estimate population affinity and sex without identifying the specific number of a vertebra.

When comparing the univariate models among the vertebrae, there were no clear trends regarding which variables performed the best. In other words, the classification accuracy for a

specific variable differed substantially among vertebrae and was not consistently the best indicator of either population affinity or sex. For example, when estimating population affinity, maximum superior sagittal length was one of the poorest classifiers for the third cervical vertebrae; however, it was observed to be the greatest classifier of all the variables when estimating population affinity for the fourth cervical vertebrae. The combination of all of the measurement means per vertebral subtype for the universal models demonstrated overall more intermediate classification accuracies. Thus, there are small variations in the size and shape of connecting vertebrae that variably influence the discriminatory power of measurements from one vertebra to the next. However, by combining the means, the individual vertebra would not need to be identified in order to make use of the method for estimating the parameters of the biological profile.

Results for the univariate models demonstrate higher classification accuracies than the multivariate models, especially for population affinity estimation. The high classification accuracies for most of the univariate models demonstrate that if fragmentary vertebrae are present, single variable analyses can be used to estimate both population affinity and sex. While the current study demonstrated that population affinity and sex estimation is possible using univariate methods, single variable classification has limitations as the use of a single dimension may also result in higher misclassification rates of both population affinity and sex. The high rates of misclassification are due to univariate models only being able to capture the limited amount of human variation expressed in that single variable. More variables allow for more information to be assembled, which might increase classification accuracy (Ousley *et al.*, 2012). While more variables from multiple bones can assess more human variation, even just using multiple measurements from a single vertebra already provides a greater number of measurements than a single variable examined by univariate analysis, thus providing more information about the individual.

While univariate models can be employed with fairly high accuracy, the approach is not necessarily the most effective for properly distinguishing between the population and sex groups. As such, multivariate models are better able to capture the variation and give overall, more robust results. The greater performance of multivariate models compared to univariate models is evident in the classification accuracies, where the multivariate models consistently achieved accuracies above 80%. This result is not unexpected, as many recent publications have confirmed the advantages of employing a combination of variables (Ostrowsky *et al.*,

2015; Krüger *et al.*, 2017; Ramadan *et al.*, 2017; Hora and Sladek, 2018; Oura *et al.*, 2018; Liebenberg *et al.*, 2019; Rozendaal *et al.*, 2020). In other words, multivariate models should be used where possible when estimating population affinity and sex from the vertebrae.

The study also aimed to determine if the specific number of the vertebra needs to be identified prior to further analysis. While some vertebrae can easily be identified due to unique morphological features, others may be much more difficult to identify correctly, particularly some of the mid-cervical and mid-thoracic vertebrae (White *et al.*, 2005). A comparison of the single multivariate vertebra models to the corresponding universal multivariate models revealed that the more specific, single vertebra models achieved slightly higher classification accuracies (about 1% to 9% higher, depending on the vertebra and vertebral sub-type). However, the classification accuracy should not be the only factor taken into consideration when selecting models for analyses. Other important factors include the sample size, as well as the overall generalisability of the model. The sample size of the single vertebra models is fairly small, largely due to missing variables resulting from postmortem damage, osteophytosis or vertebral compression. In turn, the sample size affects how well a model can generalise information to a test sample that was not used to train the model, and of course the overall population. Models made with smaller samples have been shown to overfit data, thereby producing overly optimistic results (Reunanen, 2003; Babyak, 2004; Roberts *et al.*, 2017). But when tested on independent samples, these models often demonstrate a substantial decrease in classification accuracy as the models are too specific and generalise information poorly when applied to the general population (Ousley *et al.*, 2012). Another problem that may arise with the single vertebra models, is if the practitioner identifies a specific vertebra incorrectly and attempts to use the incorrect model (for example using the C3 model on a C4 vertebra). As was noted with the univariate models, each vertebra has a unique combination of dimensions with small albeit significant differences in the size and shape that can affect the accuracy of measurements. Further validation studies are required to establish how the incorrect identification of vertebrae will affect the discriminatory power of the single vertebra models. Based on the results, universal models made by combining all of the vertebrae per sub-type can be used to classify population affinity and sex with fair accuracy. Moreover, the combination of data allows for larger samples and greater generalisability and will not be adversely affected in instances where specific vertebrae are identified incorrectly. Overall, the vertebral sub-types achieved similar accuracies for population affinity estimation, indicating that cervical, thoracic and lumbar models can all be useful for assessing population groups.

A variety of different skeletal elements have been studied in the South African population regarding population affinity and sex estimation (Pastor, 2005; Krüger, 2015; Torimitsu *et al.*, 2016; Liebenberg *et al.*, 2019). The vertebrae are usually preserved relatively well, with minor damage and weathering except for transverse processes and spinous processes. Research exploring population affinity and sex have used various methods and skeletal elements to achieve high classification accuracies. Based on craniometric and postcranial studies on population affinity and sex estimation, the accuracy of this study is comparable to published methods frequently used and recognized as best practices; however, most studies have examined more than two groups (Stull *et al.*, 2014; Krüger *et al.*, 2015; Krüger *et al.*, 2017; Liebenberg *et al.*, 2019). The use of two populations means that group affinity probabilities and classification accuracies are high, especially when comparing white and black South Africans due to their substantial population differences. When a third population, such as Coloured South Africans, is entered into the comparison, typically a decrease in accuracies is seen, especially in the South African population as black and coloured South Africans may present with large amounts of overlap (Stull *et al.*, 2014; Krüger *et al.*, 2017; Liebenberg *et al.*, 2019).

Other than variations among population groups, ageing and bone pathology should also be considered in population and sex estimation using vertebrae. Alterations in vertebrae, especially those of the lumbar spine, have been observed with ageing. The differences between the two populations could result from pathological conditions associated with advanced age (i.e., osteophytes, wedging, compression). In the current sample, the mean age of white South Africans was substantially higher than that of the black South Africans, which is a result of the overall higher mean age of white South Africans in the skeletal collection (L'Abbé *et al.*, 2005). The presence of age-related pathological conditions could affect measurements, particularly vertebral body measurements, used to estimate population and sex. For example, osteophytes can affect the vertebral body height and length, as spurs form on the borders of the vertebral body. With age, the anterior and posterior heights of the vertebrae can also decrease, giving the appearance of a wedge-shaped vertebra, which is more frequent in men than in women (Snodgrass, 2004a; Garoufi *et al.*, 2020). The difference between males and females is partially due to load-bearing capacity differences, brought on by differences in physical activity, which may impact bone mass, microarchitecture, and size throughout life. Additionally, anatomical differences in the transfer of weight between the vertebral column and pelvis between sexes and populations could affect variation in estimations.

The universal multivariate models (for all vertebral subtypes: cervical, thoracic and lumbar vertebrae) were found to have fairly similar, although slightly higher, accuracies to the individual multivariate models, which is most likely the result of the larger sample size obtained by combining the relevant vertebrae measurements per subtype. The increased sample size provided more information for the discriminant functions to create group boundaries, making the models more proficient in quantifying the variation and thereby maximising group differences to classify either population affinity or sex. Both universal univariate and multivariate models performed similarly and yielded similar classification accuracies. Depending on the condition of the remains, practitioners are free to utilise any combination of variables, individual variables or vertebrae on their own while still yielding reliable results.

Interestingly, the thoracic vertebrae demonstrated lower accuracies (around 74% to 75% compared to the 82% obtained with both cervical and lumbar stepwise-selected models) when attempting to estimate sex. Previous studies on sex estimation have shown that cervical and lumbar vertebrae variables are higher than thoracic vertebrae variables (Allbright, 2007). As such the thoracic vertebrae are less useful for sex estimation than the lumbar and cervical models but can still be used in instances where only thoracic vertebrae are recovered. Additionally, similar and higher classification accuracies were obtained in the current study when compared to previous South African studies that used other bone models exploring population affinity and sex estimation (Kibii *et al.*, 2010; Ostrofsky *et al.*, 2015; Ünlütürk, 2016). Several studies have evaluated the variation of the first and second cervical vertebrae using multiple measurements, achieving accuracy ranging between 70.9% and 92.9% for sex estimation, similar to the current research (Marino, 1995; Wescott, 2000; Marlow *et al.*, 2011; Gama *et al.*, 2015; Torimitsu *et al.*, 2016; Rozendaal *et al.*, 2020). In addition, previous studies involving lumbar vertebrae have shown greater classification accuracy than studies of thoracic and cervical vertebrae, as seen in the current study. (Taylor *et al.*, 1984; Ünlütürk, 2016; Hora and Sladek, 2018; Oura *et al.*, 2018).

Based on the results of the current study, white South Africans had higher classification accuracy in most vertebrae than black South Africans, while males had higher accuracy than females. The measurements that demonstrated the most sexual dimorphism and population variation were the vertebral body, spinous process and transverse process length and height measurements. As mentioned, certain measurements proved to be better at estimating population affinity and sex than others. The transverse process length and maximum posterior

height measurements were the most discriminating variables for both population affinity and sex estimation. A possible reason for the increased transverse process and vertebral body dimension measurements observed may be due to the differences in growth patterns between the sexes. In the coronal plane, female vertebrae are relatively taller and thinner than those of males. In comparison with males, the height of the female vertebral body is higher, while the transverse diameter of the male vertebral body is larger. Approximately between the ages of eight and 21 years old, the thoracic and lumbar vertebrae (specifically from the sixth thoracic vertebra to the fifth lumbar vertebra) increase in size and dimensions significantly in males compared to females (Bellemare *et al.*, 2003). Additionally, sexual dimorphism in spinal development results in females having a thinner thoracolumbar spine than males because of differences in growth rates during puberty (Taylor and Twomey, 1984). Male vertebrae have been found to be 25% larger than females because of delayed apoptosis in males associated with increased testosterone levels during puberty (Bellemare *et al.*, 2003). Furthermore, the spinous process is an attachment site for the back muscles, and as males tend to be more muscular than females, the spinous process of males tends to be larger than females, thereby adding to the sexual dimorphism (Atkinson, 1967; Taylor *et al.*, 1984; Bellemare *et al.*, 2003; Allbright, 2007; Masharawi and Salame, 2011; Aylott *et al.*, 2012; Waxenbaum *et al.*, 2018). The larger muscle mass noted in males may also explain why the vertebral body measurements are good sex estimators, especially in the lower thoracic and lumbar vertebrae, as they act as the major load-bearing aspects for the vertebrae and support the upper body weight. The weight-bearing function of the vertebral bodies may also result in larger vertebral bodies in males, which would explain why the measurements associated with the body are more sexually dimorphic.

### **5.1 Practical Application in Forensic Anthropology and Future Recommendations**

Although estimation of population affinity and sex from the vertebrae is not necessary for situations involving complete skeletal remains, the universal formula could be useful in instances where remains are incomplete or fragmentary. The current study demonstrated that even without prior knowledge of vertebral number, population affinity and sex could be estimated successfully from single vertebrae with comparable accuracies to some other common bone models (Stull *et al.*, 2014; Krüger *et al.*, 2015; Krüger *et al.*, 2017; Liebenberg *et al.*, 2019). The vertebrae demonstrated high classifications of 78% to 93% for population affinity and 73% to 92% for sex estimation. Therefore, the vertebrae bone models can be used

as an additional technique when crania and postcranial elements are not available to estimate population affinity and sex. With specialised software, such as FORDISC, a custom database can be imported, as the software creates discriminant functions applying numerous combinations of variables for individual analysis (L'Abbé *et al.*, 2013; Ousley and Jantz, 2013).

In order to meet the inclusion criteria, a large number of older individuals were included in the sample, with the mean age for the white South Africans being higher than the black South African sample. As a result of the composition of the Pretoria Bone Collection, demographics include primarily black and white South Africans, with black South Africans, regardless of age, making up the majority of the collection. On the other hand, white South Africans make up the minority of the collection, with more older individuals represented in the collection (L'Abbé *et al.*, 2005). Because the white South Africans in the skeletal collection have older mean ages, particularly the females, data analyses on sexual dimorphism and population variation may be affected. However, despite the white South African sample having an older mean age, efforts were made to exclude any individuals, vertebrae, or measurements that were clearly influenced by age. Even though age influences the vertebrae and may potentially affect classification accuracy, these influences can typically be seen in the development of osteophytes, in the compression and wedging of vertebral bodies, or in pathology. Until these influences are tested, the age-related effects on classification remain unknown and, in turn, should be further investigated. If there are obvious changes in the vertebral body because of ageing or spinal disease, these measurements of the vertebral body should be excluded from analyses. The age difference between sexes may also affect the analyses, as ageing may alter vertebrae morphology and lead to incorrect classification. Research needs to be conducted on influences on vertebral classification accuracy, considering age cohorts and the interaction between age, sexes, and population groups. Additionally, future research on a larger sample size that includes individuals housed throughout south African collections as well as creating models that account age related changes.

Future research assessing further South African populations, including coloured and Indian South Africans, will improve understanding of the vertebral sexual dimorphism and population variation of modern South Africans. Further explorations, such as assessing the effects of using thermally altered fragmentary vertebrae, using vertebrae with major pathology present and using heavily fragmentary vertebrae could improve the universal formulae and models, allowing further univariate and multivariate analyses to be used in fragmentary or partial

remains. Lastly, to improve the validity and reliability of the universal formula, a validation study should be conducted on an unrelated, independent test sample.

## CHAPTER 6: CONCLUSION

The current study assessed osteometric variation of a selection of vertebrae to examine population variation and sexual dimorphism, and to determine whether a universal discriminant function can effectively predict population affinity and sex without prior identification of vertebral number in a South African sample. Through the use of univariate and multivariate analyses, the results demonstrated that differences exist among the vertebrae of black and white South African males and females. The majority of sexual dimorphism and population variation is attributed to differences in the size of the vertebral body and spinous processes, with males and white South Africans presenting with greater means than females and black South Africans, respectively.

Multivariate models yielded the best results, as the inclusion of multiple variables maximised group differences. With the use of single vertebrae and combined vertebrae sub-type multivariate models, population affinity and sex can be estimated with comparable or even higher accuracies than commonly used methods making use of other postcranial bone models for estimating population affinity and sex. However, only two populations were examined in the current study, while commonly used methods that assess bone models typically include at least three populations (black, white and Coloured South Africans). Furthermore, multivariate universal models are recommended even though a variety of different standards are available for use, such as single vertebrae univariate models and universal univariate models. While not the best tool for exploring human variation, univariate models still allow for accurate estimation of population affinity and sex from single vertebrae, which are particularly useful in the analysis of taphonomically altered remains. It is also recommended that the standards be used with *FORDISC*, after they have been validated, which allows for case-specific analyses where the practitioner has more flexibility in choosing which models to use. The current study has shown the potential of using vertebral models, both univariate and multivariate, to estimate population affinity and sex in a forensic setting.

## REFERENCES

- Albert, ., Mulhern, D., Torpey, M.A., Boone, E. 2010. Age Estimation Using Thoracic and First Two Lumbar Vertebral Ring Epiphyseal Union. *Forensic Science International*.
- Albert, M.A., Maier, C.A. 2013. Epiphyseal Union of the Cervical Vertebral Centra: Its Relationship to Skeletal Age and Maturation of Thoracic Vertebral Centra. *Journal of Forensic Science*. 58(6):1568-1574.
- Allbright, A.S. 2007. Sexual Dimorphism in the Vertebral Column. Masters Masters, The University of Tennessee, Knoxville.
- Asuero, A.G., Sayago, A., Gonzalez, A. 2006. The Correlation Coefficient: An Overview. *Critical reviews in analytical chemistry*. 36(1):41-59.
- Atkinson, P. 1967. Variation in Trabecular Structure of Vertebrae with Age. *Calcified Tissue Research*. 1(1):24-32.
- Aylott, C.E.W., Puna, R., Robertson, P.A., Walker, C. 2012. Spinous Process Morphology: The Effect of Ageing through Adulthood on Spinous Process Size and Relationship to Sagittal Alignment. *European Spine Journal*. 21(5):1007-1012.
- Babyak, M.A. 2004. What You See May Not Be What You Get: A Brief, Nontechnical Introduction to Overfitting in Regression-Type Models. *Psychosomatic medicine*. 66(3):411-421.
- Baidas, L. 2012. Correlation between Cervical Vertebrae Morphology and Chronological Age in Saudi Adolescents. *King Saud University Journal of Dental Sciences*. 3(1):21-26.
- Barrier, I., L'abbé, E. 2008. Sex Determination from the Radius and Ulna in a Modern South African Sample. *Forensic science international*. 179(1):85. e81-85. e87.
- Bastir, M., Higuero, A., Rios, L., Garcia Martinez, D. 2014. Three-Dimensional Analysis of Sexual Dimorphism in Human Thoracic Vertebrae: Implications for the Respiratory System and Spine Morphology. *American Journal of Physical Anthropology*. 155(4):513-521.
- Bellemare, F., Jeanneret, A., Couture, J. 2003. Sex Differences in Thoracic Dimensions and Configuration. *American Journal of Respiratory and Critical Care Medicine*. 168(3):305-312.

- Bellemare, F., Fuamba, T., Bourgeault, A. 2006. Sexual Dimorphism of Human Ribs. *Respiratory Physiology and Neurobiology*. 150(2-3):233-239.
- Bergström, A., McCarthy, S.A., Hui, R., Almarri, M.A., Ayub, Q., Danecek, P., *et al.* 2020. Insights into Human Genetic Variation and Population History from 929 Diverse Genomes. *Science*. 367(6484):eaay5012.
- Berry, J.L., Moran, J.M., Berg, W.S., Steffee, A.D. 1987. A Morphometric Study of Human Lumbar and Selected Thoracic Vertebrae. *Spine*. 12(4):362-367.
- Bethard, J.D., Seet, B.L. 2013. Sex Determination from the Second Cervical Vertebra: A Test of Wescott's Method on a Modern American Sample. *Journal of Forensic Science*. 58(1):101-103.
- Bickford-Smith, V. 1995. South African Urban History, Racial Segregation and the Unique Case of Cape Town? *Journal of Southern African Studies*. 21(1):63-78.
- Bonjour, J.-P., Theintz, G., Buchs, B., Slosman, D., Rizzoli, R. 1991. Critical Years and Stages of Puberty for Spinal and Femoral Bone Mass Accumulation During Adolescence. *The Journal of Clinical Endocrinology & Metabolism*. 73(3):555-563.
- Cabo, L.L., Brewster, C.P., Luengo Azpiazu, J. 2012. Sexual Dimorphism: Interpreting Sex Markers. In: Dirkmaat, D. C. (eds.) *A Companion to Forensic Anthropology*. 1 edn.: Blackwell Publishing Ltd. 248.
- Callewaert, F., Sinnesael, M., Gielen, E., Boonen, S., Vanderschueren, D. 2010. Skeletal Sexual Dimorphism: Relative Contribution of Sex Steroids, Growth Hormone-Insulin-Like Growth Factor-I (Gh-Igf-I) and Mechanical Loading. *Journal of Endocrinology*. 207(2):127-134.
- Carapuço, M., Nóvoa, A., Bobola, N., Mallo, M. 2005. Hox Genes Specify Vertebral Types in the Presomitic Mesoderm. *Genes & Development*. 19(18):2116-2121.
- Cheng, X., Sun, Y., Boonen, S., Nicholson, P., Brys, P., Dequeker, J., *et al.* 1998. Measurements of Vertebral Shape by Radiographic Morphometry: Sex Differences and Relationships with Vertebral Level and Lumbar Lordosis. *Skeletal radiology*. 27(7):380-384.

Chiara, V., Jo, B., Lynnerup, N. 2016. Evaluating Osteological Ageing from Digital Data. *Journal of Anatomy*. 235(2):386-395.

Christopher, A.J. 1992. Segregation Levels in South African Cities, 1911-1985. *The International Journal of African Historical Studies*. 25(3):561-582.

Christopher, A.J. 2001. Urban Segregation in Post-Apartheid South Africa. *Urban studies*. 38(3):449-466.

Drake, R.L., Vogl, W., Mitchell, A.W.M., Gray, H., 2016. *Gray's Anatomy for Students*[Internet]. Philadelphia, PA: Elsevier. [cited Available from:

Elliott, M., Collard, M. 2009. Fordisc and the Determination of Ancestry from Cranial Measurements. *Biology Letters*. 5(6):849-852.

Ezra, D., Hershkovitz, I., Salame, K., Alperovitch-Najenson, D., Slon, V. 2019. Osteophytes in the Cervical Vertebral Bodies (C3–C7)—Demographical Perspectives. *The Anatomical Record*. 302(2):226-231.

Franklin, D., Cardini, A., Oxnard, C.E. 2010. A Geometric Morphometric Approach to the Quantification of Population Variation in Sub-Saharan African Crania. *American Journal of Human Biology*. 22(1):23-35.

Franklin, D., Freedman, L., Milne, N., Oxnard, C.E. 2007. Geometric Morphometric Study of Population Variation in Indigenous Southern African Crania. *American Journal of Human Biology*. 19(1):20-33.

Friendly, M. 2002. Corrgrams: Exploratory Displays for Correlation Matrices. *The American Statistician*. 56(4):316-324.

Gama, I., Navega, D., Cunha, E. 2015. Sex Estimation Using the Second Cervical Vertebra: A Morphometric Analysis in a Documented Portuguese Skeletal Sample. *International Journal of Legal Medicine*. 129(2):365-372.

Garoufi, N., Bertsatos, A., Chovalopoulou, M.-E., Villa, C. 2020. Forensic Sex Estimation Using the Vertebrae: An Evaluation on Two European Populations. *International Journal of Legal Medicine*. 134(6):2307-2318.

- Geeta, A., Jamaiyah, H., Safiza, M., Khor, G., Kee, C., Ahmad, A., *et al.* 2009. Reliability, Technical Error of Measurements and Validity of Instruments for Nutritional Status Assessment of Adults in Malaysia. *Singapore Medical Journal*. 50(10):1013.
- Hall, M., Morris, A. 1983. Race and Iron Age Human Skeletal Remains from Southern Africa: An Assessment. *Social Dynamics*. 9(2):29-36.
- Hefner, J.T. 2009. Cranial Nonmetric Variation and Estimating Ancestry\*. *Journal of Forensic Science*. 54(5):985-995.
- Hefner, J.T. 2016. Biological Distance Analysis, Cranial Morphoscopic Traits, and Ancestry Assessment in Forensic Anthropology. *Biological Distance Analysis*. 301-315.
- Holm, S. 1979. A Simple Sequentially Rejective Multiple Test Procedure. *Scandinavian Journal of Statistics*. 6(2):65-70.
- Hora, M., Sladek, V. 2018. Population Specificity of Sex Estimation from Vertebrae. *Forensic Science International*. 291279 e271-279 e212.
- İşcan, M.Y., Steyn, M. 1999. Craniometric Determination of Population Affinity in South Africans. *International Journal of legal medicine*. 112(2):91-97.
- Jeong, Y., Jeong, G., Pergande, S., Lee, K. 2019. Generating a Vertebrae-Based Method to Discriminate between Korean and U.S. White Male Casualties from the Korean War. *Journal of Forensic Science*. 64(6):1776-1781.
- Junno, J.-A., Paananen, M., Karppinen, J., Tammelin, T., Niinimäki, J., Lammentausta, E., *et al.* 2013. Influence of Physical Activity on Vertebral Strength During Late Adolescence. *The Spine Journal*. 13(2):184-189.
- Juul, A. 2001. The Effects of Oestrogens on Linear Bone Growth. *Apmis*. 7(3):303 - 313.
- Kalnins, A. 2018. Multicollinearity: How Common Factors Cause Type 1 Errors in Multivariate Regression. *Strategic Management Journal*. 39(8):2362-2385.
- Kerttula, L.I., Serlo, W.S., Tervonen, O.A., Pääkkö, E.L., Vanharanta, H.V. 2000. Post-Traumatic Findings of the Spine after Earlier Vertebral Fracture in Young Patients: Clinical and Mri Study. *Spine*. 25(9):1104-1108.

- Kibii, J.M., Pan, R., Tobias, P.V. 2010. Morphometric Variations of the 7th Cervical Vertebrae of Zulu, White, and Colored South Africans. *Clinical Anatomy* 23(4):399-406.
- Klaassen, Z., Tubbs, R.S., Apaydin, N., Hage, R., Jordan, R., Loukas, M. 2011. Vertebral Spinal Osteophytes. *Anatomical science international*. 86(1):1-9.
- Klales, A.R., Kenyhercz, M.W. 2015. Morphological Assessment of Ancestry Using Cranial Macromorphoscopies. *Journal of forensic science*. 60(1):13-20.
- Klales, A.R., Ousley, S.D., Vollner, J.M. 2012. A Revised Method of Sexing the Human Innominate Using Phenice's Nonmetric Traits and Statistical Methods. *American journal of physical anthropology*. 149(1):104-114.
- Krüger, G.C. 2015. Comparison of Sexually Dimorphic Patterns in the Postcrania of South Africans and North Americans. University of Pretoria.
- Krüger, G.C., L'Abbé, E.N., Stull, K.E. 2017. Sex Estimation from the Long Bones of Modern South Africans. *International Journal of Legal Medicine*. 131(1):275-285.
- Krüger, G.C., L'Abbe, E.N., Stull, K.E., Kenyhercz, M.W. 2015. Sexual Dimorphism in Cranial Morphology among Modern South Africans. *International Journal of Legal Medicine*. 129(4):869-875.
- Krüger, G.C., Liebenberg, L., Myburgh, J., Meyer, A., Oettlé, A.C., Botha, D., *et al.* 2018. Forensic Anthropology and the Biological Profile in South Africa. *New Perspectives in Forensic Human Skeletal Identification*. 313-321.
- Kumari, S. 2008. Multicollinearity: Estimation and Elimination. *Journal of Contemporary research in Management*. 3(1):87-95.
- L'Abbé, E.N., Kenyhercz, M., Stull, K.E., Keough, N., Nawrocki, S. 2013. Application of Fordisc 3.0 to Explore Differences among Crania of North American and South African Blacks and Whites. *Journal of Forensic Sciences*. 58(6):1579-1583.
- L'Abbé EN , K.M., Stull KE , Ousley SD Craniometric Assessment of Modern 20 Century Black, White and Coloured\* South Africans. In: Pretoria, U. O. (ed.).

L'Abbé, E.N., Loots, M., Meiring, J.H. 2005. The Pretoria Bone Collection: A Modern South African Skeletal Sample. *Homo: Journal of Comparative Human Biology* 56(2):197-205.

L'Abbé, E.N., Van Rooyen, C., Nawrocki, S.P., Becker, P.J. 2011. An Evaluation of Non-Metric Cranial Traits Used to Estimate Ancestry in a South African Sample. *Forensic Science International*. 209(1-3):195 e191-197.

Larson, A.M., Ribot, J.C. 2004. Democratic Decentralisation through a Natural Resource Lens: An Introduction. *The European Journal of Development Research*. 16(1):1-25.

Liebenberg, L. 2015. *Postcraniometric Analysis of Ancestry among Modern South Africans*. University of Pretoria.

Liebenberg, L., Stull, K.E., L'Abbé, E.N., Botha, D. 2015. Evaluating the Accuracy of Cranial Indices in Ancestry Estimation among South African Groups. *Journal of forensic science*. 60(5):1277-1282.

Liebenberg, L., Krüger, G.C., L'Abbé, E.N., Stull, K.E. 2019. Postcraniometric Sex and Ancestry Estimation in South Africa: A Validation Study. *International Journal of Legal Medicine*. 133(1):289-296.

Marino, E.A. 1995. Sex Estimation Using the First Cervical Vertebra. *American Journal of Physical Anthropology*. 97(2):127-133.

Marlow, E.J., Pastor, R.F. 2011. Sex Determination Using the Second Cervical Vertebra--a Test of the Method. *Journal of Forensic Science*. 56(1):165-169.

Masharawi, Y., Salame, K. 2011. Shape Variation of the Neural Arch in the Thoracic and Lumbar Spine: Characterization and Relationship with the Vertebral Body Shape. *Clinical Anatomy*. 24(7):858-867.

Montasser, M.A., Viana, G., Evans, C.A. 2017. Secular Trends in the Timing of Skeletal Maturation as Assessed by the Cervical Vertebrae Maturation Method. *European Journal of Orthodontics*. 39(2):188-193.

Mosekilde, L. 2000. Age-Related Changes in Bone Mass, Structure, and Strength--Effects of Loading. *Zeitschrift für Rheumatologie*. 59(1):I1-I9.

Ostrofsky, K.R., Churchill, S.E. 2015. Sex Determination by Discriminant Function Analysis of Lumbar Vertebrae. *Journal of Forensic Science*. 60(1):21-28.

Oura, P., Karppinen, J., Niinimäki, J., Junno, J.A. 2018. Sex Estimation from Dimensions of the Fourth Lumbar Vertebra in Northern Finns of 20, 30, and 46 Years of Age. *Forensic Science International*. 290350.e351-350.e356.

Oura, P., Paananen, M., Niinimäki, J., Tammelin, T., Herrala, S., Auvinen, J., *et al.* 2016. Effects of Leisure-Time Physical Activity on Vertebral Dimensions in the Northern Finland Birth Cohort 1966. *Scientific Reports*. 6(1):1-10.

Ousley, S., Jantz, R. 2013. Fordisc 3. *Rechtsmedizin*. 23(2):97-99.

Ousley, S., Jantz, R., Freid, D. 2009. Understanding Race and Human Variation: Why Forensic Anthropologists Are Good at Identifying Race. *American Journal of Physical Anthropology*. 139(1):68-76.

Ousley, S.D., Jantz, R.L. 2012. Fordisc 3 and Statistical Methods for Estimating Sex and Ancestry. In: Dirkmaat, D. C. (eds.) *A Companion to Forensic Anthropology*. 1 edn.: Blackwell Publishing Ltd. 29.

Pastor, R. Year. Published. Sexual Dimorphism in Vertebral Dimensions at the T12/L1 Junction. *Proceedings of the 57th annual meeting of the American Academy of Forensic Sciences*, 2005. 302-303.

Phenice, T.W. 1969. A Newly Developed Visual Method of Sexing the Os Pubis. *Journal of Physical Anthropology* 30(2):297-301.

Posel, D. 2001. Race as Common Sense: Racial Classification in Twentieth-Century South Africa. *African Studies Review*. 44(2):87-114.

Ramadan, N., Abd El-Salam, M.H., Hanon, A.F., El-Sayed, N.F., Al-Amir, A.Y. 2017. Identification of Sex and Age for Egyptians Using Computed Tomography of the First Lumbar Vertebra. *Egyptian Journal of Forensic Sciences*. 7(1):1-8.

Reunanen, J. 2003. Overfitting in Making Comparisons between Variable Selection Methods. *Journal of Machine Learning Research*. 3(Mar):1371-1382.

- Ribot, J.C. 2004. *Waiting for Democracy*. World Resources Institute, Washington, DC, USA.
- Roberts, D.R., Bahn, V., Ciuti, S., Boyce, M.S., Elith, J., Guillerá-Arroita, G., *et al.* 2017. Cross-Validation Strategies for Data with Temporal, Spatial, Hierarchical, or Phylogenetic Structure. *Ecography*. 40(8):913-929.
- Ross, A.H., Pilloud, M. 2021. The Need to Incorporate Human Variation and Evolutionary Theory in Forensic Anthropology: A Call for Reform. *American Journal of Physical Anthropology*. 176(4):672-683.
- Rozendaal, A.S., Scott, S., Peckmann, T.R., Meek, S. 2020. Estimating Sex from the Seven Cervical Vertebrae: An Analysis of Two European Skeletal Populations. *Forensic Science International*. 3061-10.
- Sadler, T.W., Langman, J. 2010. *Langman's Medical Embryology*, Philadelphia, Pa., Lippincott Williams & Wilkins.
- Sauer, N.J. 1992. Forensic Anthropology and the Concept of Race: If Races Don't Exist, Why Are Forensic Anthropologists So Good at Identifying Them? *Social Science & Medicine*. 34(2):107-111.
- Savage, M. 1986. The Imposition of Pass Laws on the African Population in South Africa 1916-1984. *African Affairs*. 85(339):181-205.
- Scher, A. 1990. Premature Onset of Degenerative Disease of the Cervical Spine in Rugby Players. *South African Medical Journal*. 77:557-558.
- Schober, P., Boer, C., Schwarte, L.A. 2018. Correlation Coefficients: Appropriate Use and Interpretation. *Anesthesia and Analgesia*. 126(5):1763-1768.
- Seeman, E. 2001. Sexual Dimorphism in Skeletal Size, Density, and Strength. *The Journal of Clinical Endocrinology and Metabolism*. 86(10):4576-4584.
- Snodgrass, J.J. 2004a. Sex Differences and Aging of the Vertebral Column. *Journal of Forensic Science*. 49(3):198-206.
- Snodgrass, J.J. 2004b. Sex Differences and Aging of the Vertebral Column. *Journal of Forensic Science*. 49(3):JFS2003198-2003196.

Soegiharto, B.M., Cunningham, S.J., Moles, D.R. 2008. Skeletal Maturation in Indonesian and White Children Assessed with Hand-Wrist and Cervical Vertebrae Methods. *American Journal of Orthodontics and Dentofacial Orthopedics*. 134(2):217-226.

Spradley, K.M., Jantz, R.L. 2016. Ancestry Estimation in Forensic Anthropology: Geometric Morphometric Versus Standard and Nonstandard Interlandmark Distances. *Journal of forensic science*. 61(4):892-897.

Spradley, M.K., Jantz, R.L. 2011. Sex Estimation in Forensic Anthropology: Skull Versus Postcranial Elements. *Journal of Forensic Science*. 56(2):289-296.

Stahle, L., Wold, S. 1989. Analysis of Variance (Anova). *Chemometrics and Intelligent Laboratory Systems*. 6(4):259-272.

Steyn, M., İşcan, M.Y. 1997. Sex Determination from the Femur and Tibia in South African Whites. *Forensic science international*. 90(1-2):111-119.

Steyn, M., Patriquin, M. 2009. Osteometric Sex Determination from the Pelvis—Does Population Specificity Matter? *Forensic science international*. 191(1-3):113. e111-113. e115.

Stull, K.E., Kenyhercz, M.W., L'Abbe, E.N. 2014. Ancestry Estimation in South Africa Using Craniometrics and Geometric Morphometrics. *Forensic Science International*. 245206 e201-207.

Tabachnick, B.G., Fidell, L.S. 2019. *Using Multivariate Statistics*, Pearson Education. xiv-815.

Taylor, J.R., Twomey, L.T. 1984. Sexual Dimorphism in Human Vertebral Body Shape. *Journal of Anatomy*. 138(2):281 - 286.

Taylor, R. 1990. Interpretation of the Correlation Coefficient: A Basic Review. *Journal of Diagnostic Medical Sonography*. 6(1):35-39.

Team, R.C. 2020. *R: A Language and Environment for Statistical Computing*. . R Foundation for Statistical Computing.

Torimitsu, S., Makino, Y., Saitoh, H., Sakuma, A., Ishii, N., Yajima, D., *et al.* 2016. Sexual Determination Based on Multidetector Computed Tomographic Measurements of the Second

Cervical Vertebra in a Contemporary Japanese Population. *Forensic Science International*. 266588 e581-588 e586.

Ulijaszek, S.J., Lourie, J.A. 2005. Anthropometric Measurement. *Anthropometry*. 1430.

Ünlütürk, Ö. 2016. Metric Assessment of Ancestry from the Vertebrae in South Africans. *International Journal of Legal Medicine*. 131(4):1123-1131.

Van der Merwe, A., Işcan, M., L'abbé, E. 2006a. The Pattern of Vertebral Osteophyte Development in a South African Population. *International Journal of Osteoarchaeology*. 16(5):459-464.

Van der Merwe, A.E., Işcan, M., L'Abbè, E.N. 2006b. The Pattern of Vertebral Osteophyte Development in a South African Population. *International Journal of Osteoarchaeology*. 16(5):459-464.

Walker, P.L. 2008. Sexing Skulls Using Discriminant Function Analysis of Visually Assessed Traits. *American Journal of Physical Anthropology*. 136(1):39-50.

Waxenbaum, J.A., Futterman, B. 2018. *Anatomy, Back, Lumbar Vertebrae*. Statpearls. StatPearls Publishing.

Weaver, A.A., Schoell, S.L., Stitzel, J.D. 2014. Morphometric Analysis of Variation in the Ribs with Age and Sex. *Journal of anatomy*. 225(2):246-261.

Wei, T., Simko, V., Levy, M., Xie, Y., Jin, Y., Zemla, J. 2017. Package 'Corrplot'. *Statistician*. 56(316):e24.

Wells, J.C. 2012. Sexual Dimorphism in Body Composition across Human Populations: Associations with Climate and Proxies for Short- and Long-Term Energy Supply. *Am J Hum Biol*. 24(4):411-419.

Wescott, D.J. 2000. Sex Variation in the Second Cervical Vertebra. *Journal of Forensic Science*. 45(2):462-466.

White, T.D., Folkens, P.A. 2005. *The Human Bone Manual*, Elsevier.

Winburn, A.P., Algee-Hewitt, B. 2021. Evaluating Population Affinity Estimates in Forensic Anthropology: Insights from the Forensic Anthropology Database for Assessing Methods Accuracy (Fadama). *Journal of Forensic Sciences*. 66(4):1210-1219.

Yu, S.B., Lee, U.Y., Kwak, D.S., Ahn, Y.W., Jin, C.Z., Zhao, J., *et al.* 2008. Determination of Sex for the 12th Thoracic Vertebra by Morphometry of Three-Dimensional Reconstructed Vertebral Models. *Journal of Forensic Science*. 53(3):620-625.

## APPENDICES

### APPENDIX I – MEASUREMENT DEFINITIONS

\*The various measurements included in the study were modified from Cheng (1998); Gama (2015); Marlow and Pastor (2011); Oura *et al.* (2018); and Torimitsu *et al.* (2016).

\*\*Refer to Appendix II for diagrams illustrating the measurements.

**Maximum Superior Sagittal Length (XSLs):** Maximum superior length from the most anterior to the posterior aspect of the vertebral body along the sagittal plane.

**Maximum Inferior Sagittal Length (XSLi):** Maximum inferior length from the most anterior to the posterior aspect of the vertebral body along the sagittal plane.

**Maximum Centroid Height (XCH):** Maximum height at the mid-point of the vertebral body.

**Maximum Anterior Height (XAH):** Maximum height taken at the most anterior aspect of the vertebral body, from the most superior to the most inferior point.

**Maximum Posterior Height (XPH):** Maximum height taken at the midpoint of the posterior aspect of the vertebral body, from the most superior to the most inferior point.

**Maximum Superior Articular Length (XALs):** Maximum length of the superior articular aspect of the vertebral body. The measurement is taken on the inner edge of the ring apophysis from the anterior to posterior aspect.

**Maximum Inferior Articular Length (XALi):** Maximum length of the inferior articular aspect of the vertebral body. The measurement is taken on the inner edge of the ring apophysis from the anterior to posterior aspect.

**Maximum Superior Articular Width (XAWs):** Maximum length of the superior articular aspect of the vertebral body. The measurement is taken on the inner edge of the ring apophysis from the most lateral aspects.

**Maximum Inferior Articular Width (XAWi):** Maximum width of the inferior articular aspect of the vertebral body. The measurement is taken on the inner edge of the ring apophysis from the most lateral aspects.

**Foramen Sagittal Length (FSL):** Length of the vertebral foramen taken at the sagittal plane.

**Foramen Coronal Width (FCW):** Maximum width of the vertebral foramen taken at the coronal plane.

**Pedicle Height (PH):** Height of the pedicle (midpoint) from the most inferior to superior aspect at the coronal plane. The measurement is the most minimum measurement of the pedicle height.

**Pedicle Width (PW):** Width of the pedicle at the midpoint of the coronal plane. For the cervical and lumbar vertebrae, the caliper is placed diagonally at the midpoint of the pedicle. Additionally, the measurement is taken at an angle on the foramen transversarium for the cervical vertebrae and the vertebral foramen. For the thoracic and lumbar vertebrae, place the caliper in the middle of the pedicle.

**Maximum Pedicle Length (XPL):** Maximum pedicle length from the posterior aspect of the vertebral body to the most anterior aspect of the articular process.

**Transverse Process Length (TPL):** Length of the transverse process from where the lamina meets the spinous process to the most lateral aspect of the process. For the cervical vertebrae, the caliper is placed on the posterior tubercle of the transverse process to the midpoint, where the lamina and spinous process meet. The caliper is placed on the most lateral aspect of one transverse process to the midpoint, where the lamina and spinous process meet for the thoracic and lumbar vertebrae.

**Maximum Transverse Process Width (XTPW):** Maximum width of the transverse processes, taken from the most lateral (left end) to the most lateral (right end). For the cervical vertebrae, the caliper is placed on the posterior tubercle of both transverse processes. The caliper is placed on the most lateral aspect of both the transverse processes for the thoracic and lumbar vertebrae.

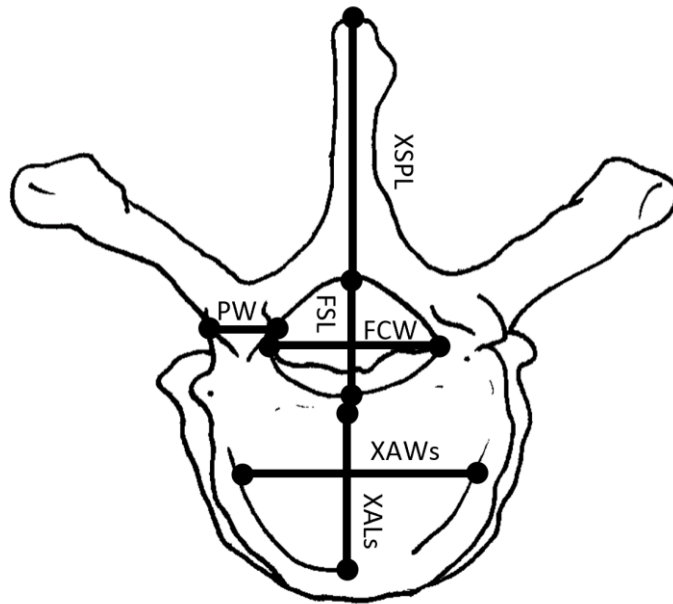
**Maximum Superior Articular Process Width (XAPWs):** Maximum distance between the left and right superior articular processes and the lateral articular borders.

**Maximum Inferior Articular Process Width (XAPWi):** Maximum distance between the left and right inferior articular processes and the lateral articular borders.

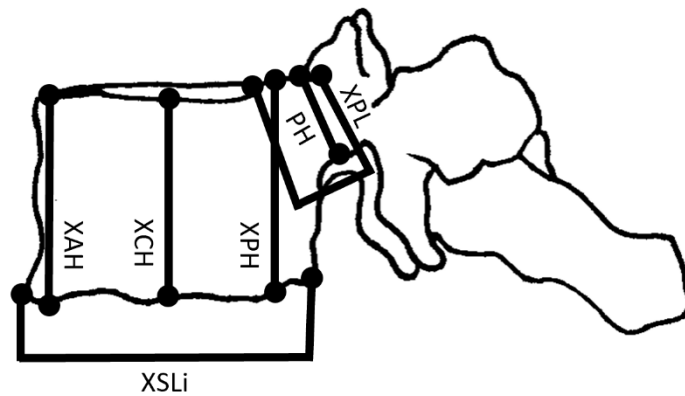
**Articular Process Height (APH):** Maximum height of both articular processes from the most superior aspect to the most inferior of the articular processes.

**Maximum Spinous Process Length (XSPL):** Maximum length of the spinous process from the articular process to the most posterior aspect of the process. For the cervical vertebrae, the measurement is taken from the point where the laminae meet the spinous process to the most posterior aspect of the spinous process, including the bifid vertebrae. The thoracic and lumbar vertebrae measurement is taken from the point where the laminae meet the spinous process to the most inferior-posterior aspect of the spinous process.

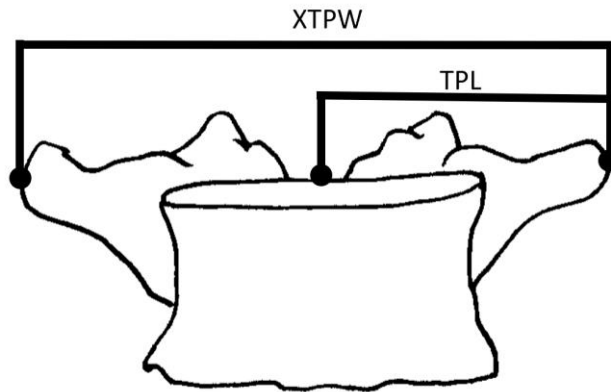
## APPENDIX II – MEASUREMENT ILLUSTRATIONS



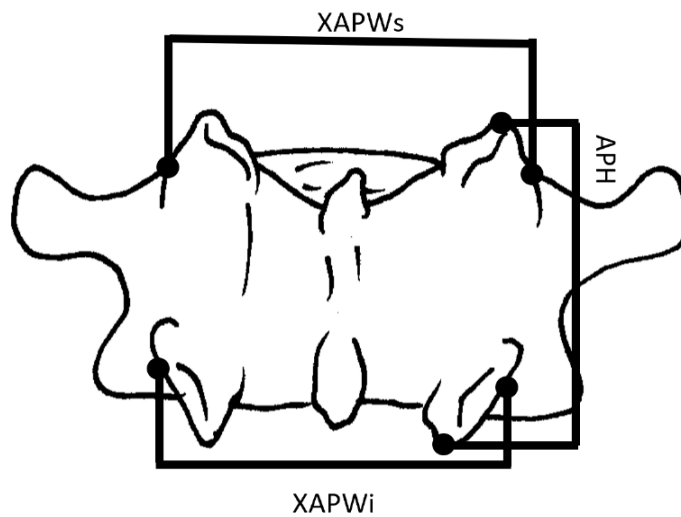
**Figure A:** Superior view of a typical vertebra showing measurements of the vertebral body, foramen, pedicle, and spinous process. Refer to Appendix I for measurement names and definitions.



**Figure B:** Lateral view of a typical vertebra showing measurements of the vertebral body and pedicle. Refer to Appendix I for measurement names and definitions.

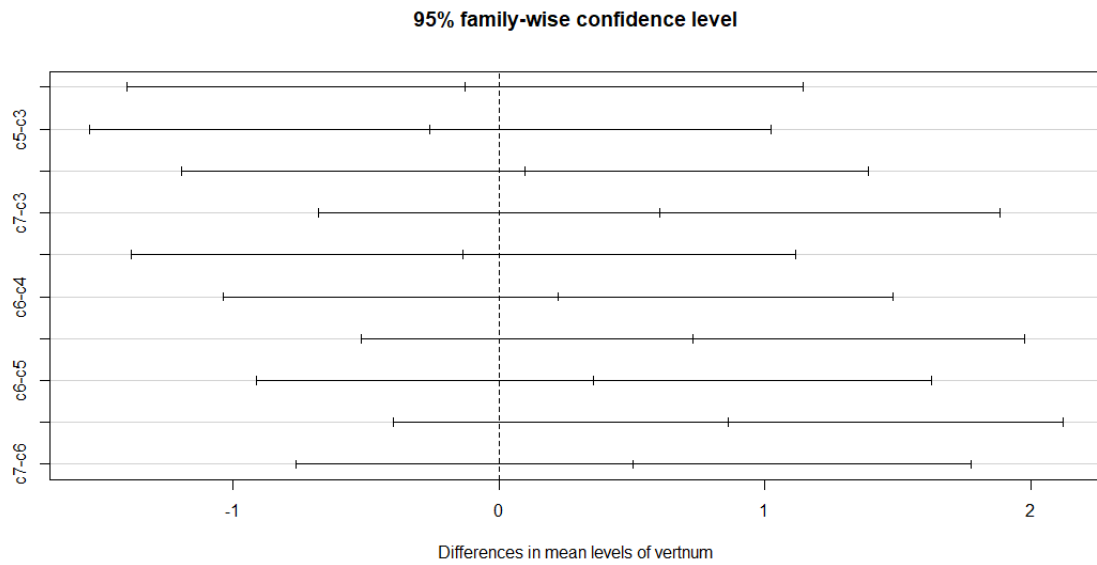


**Figure C:** Anterior view of a typical vertebra showing measurements of the articular process. Refer to Appendix I for measurement names and definitions.

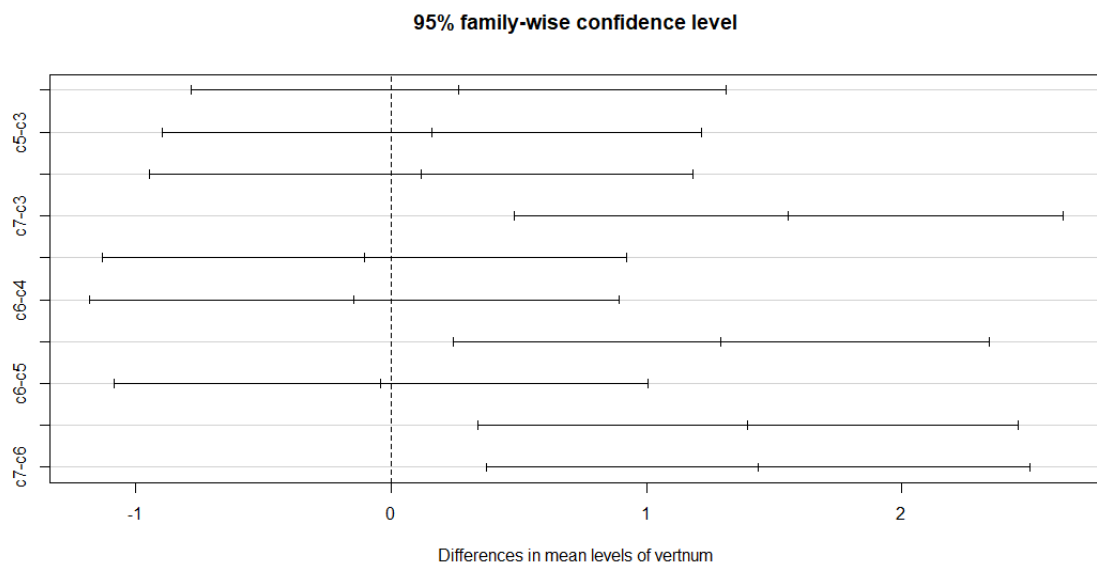


**Figure D:** Posterior view of a typical vertebra showing measurements of the superior and inferior articular processes. Refer to Appendix I for measurement names and definitions.

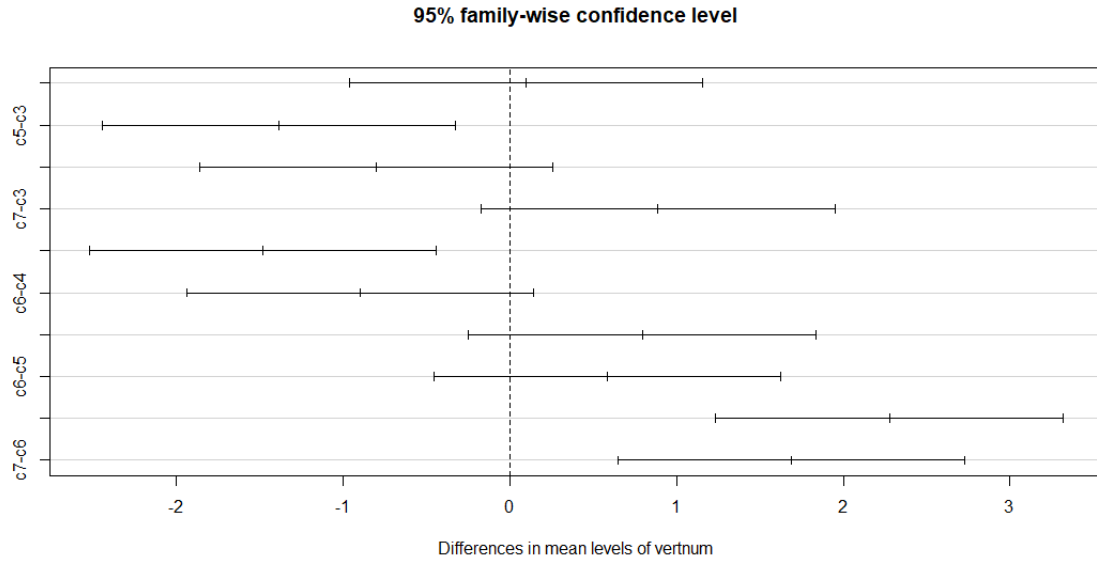
### APPENDIX III – TUKEY'S HSD RESULTS



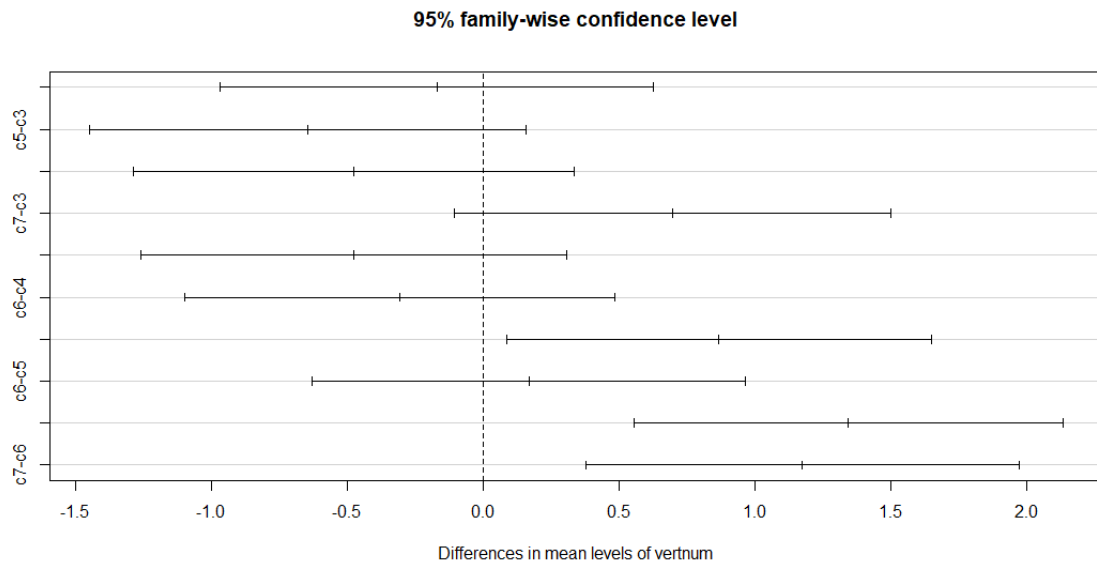
**Figure 1:** Results for the Tukey’s HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Superior Sagittal Length measurement.



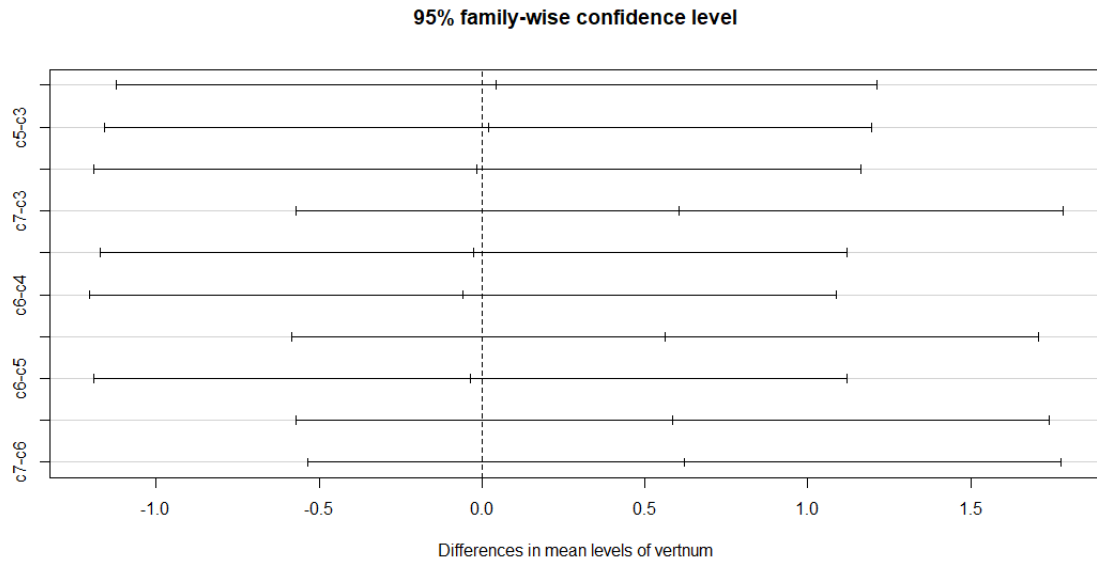
**Figure 2:** Results for the Tukey’s HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Centroid Height measurement.



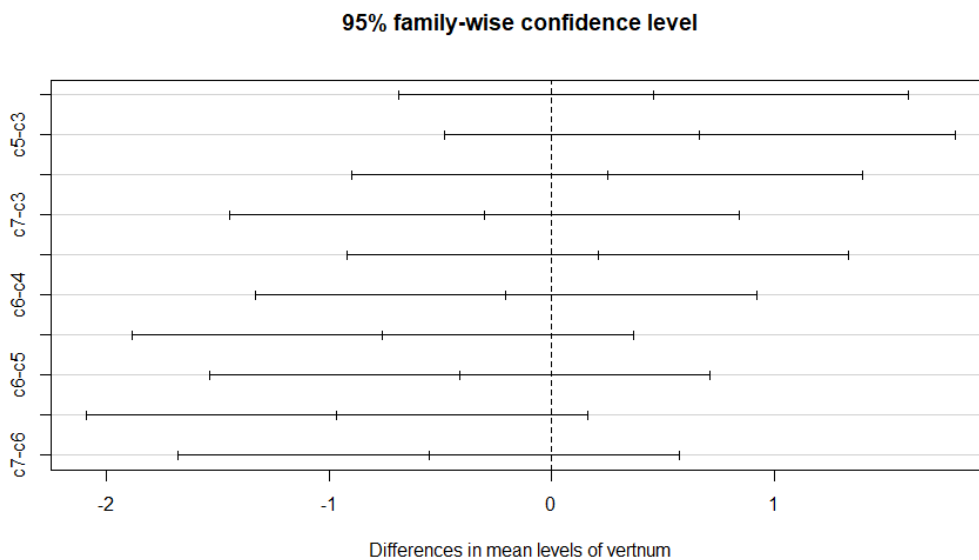
**Figure 3:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Anterior Height measurement.



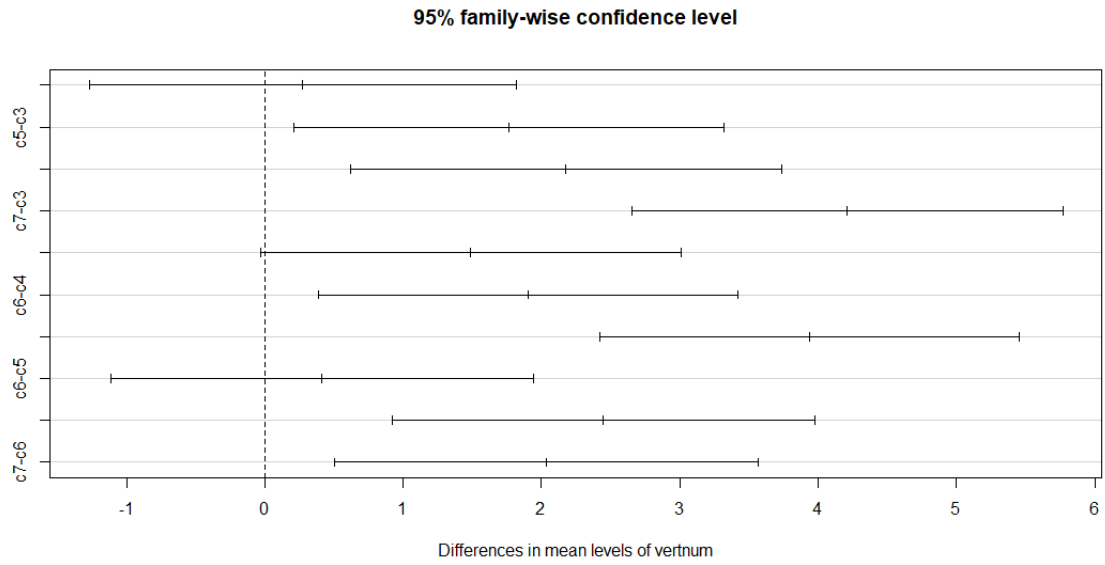
**Figure 4:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Posterior Height measurement.



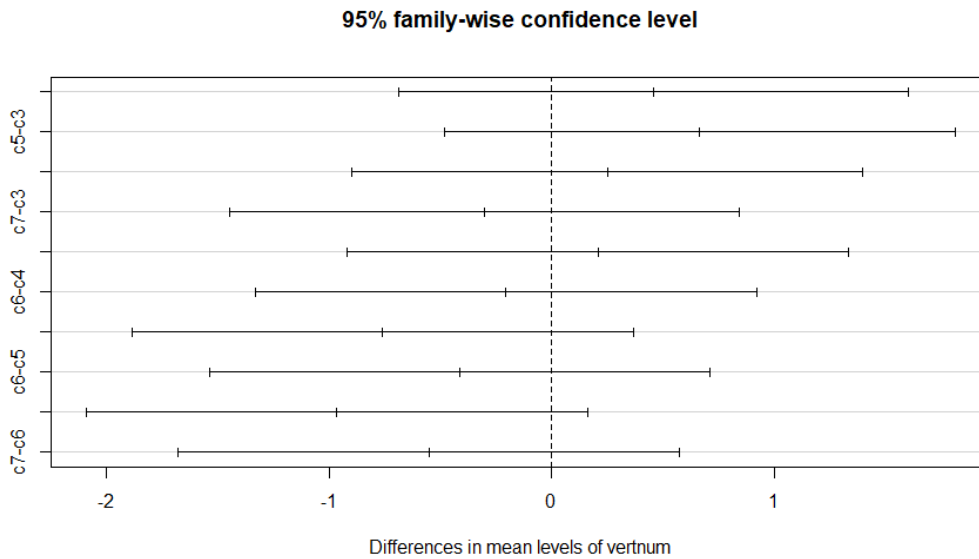
**Figure 5:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Superior Articular Length measurement.



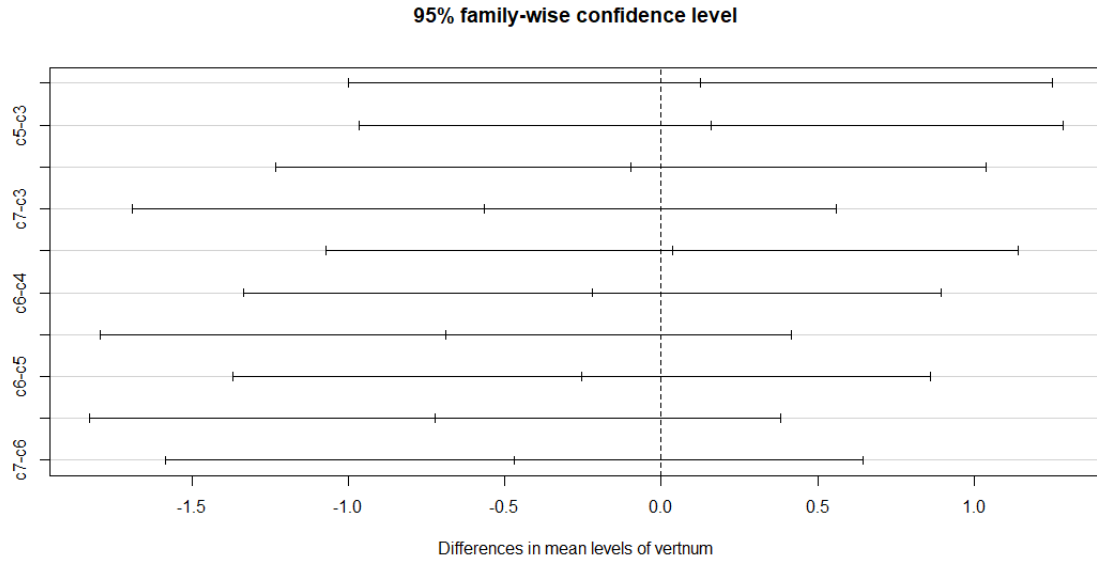
**Figure 6:** Results for the Tukey's HSD differences among the cervical vertebrae (C3-C7) Maximum Superior Articular Length measurement.



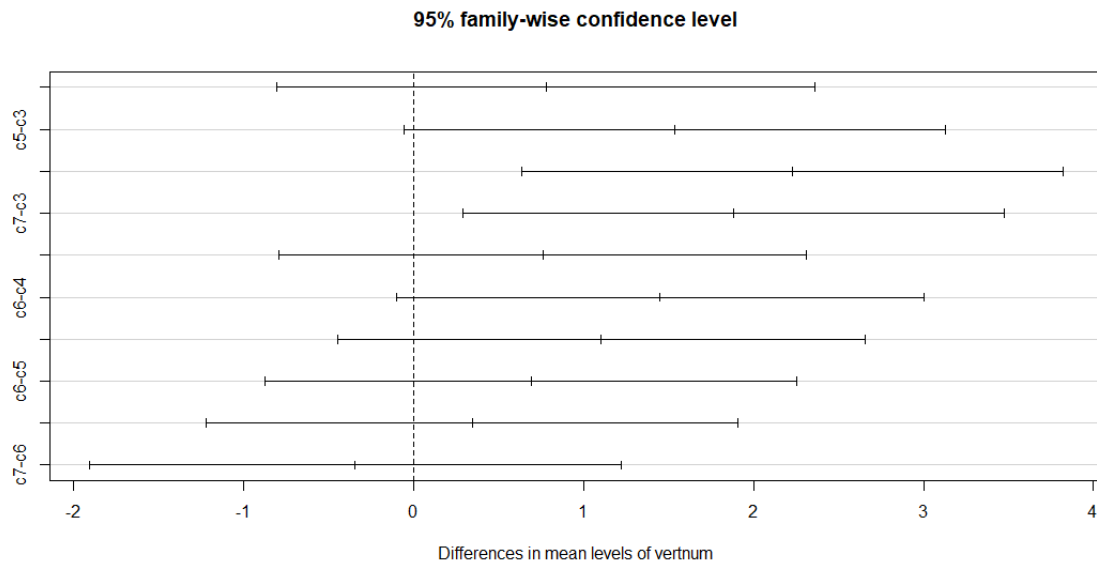
**Figure 7:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Superior Articular Width measurement.



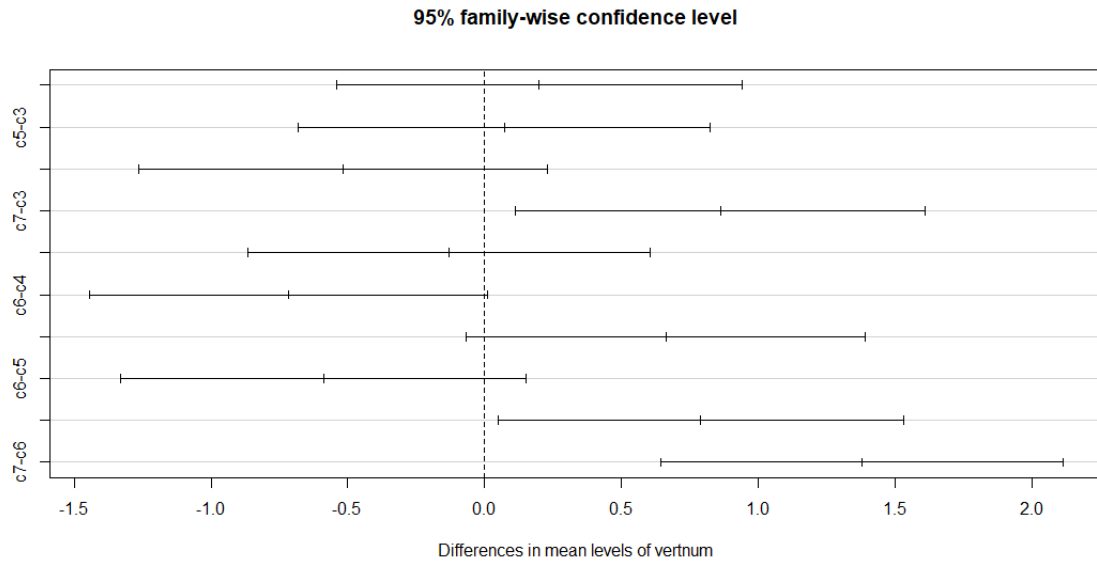
**Figure 8:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Inferior Articular Width measurement.



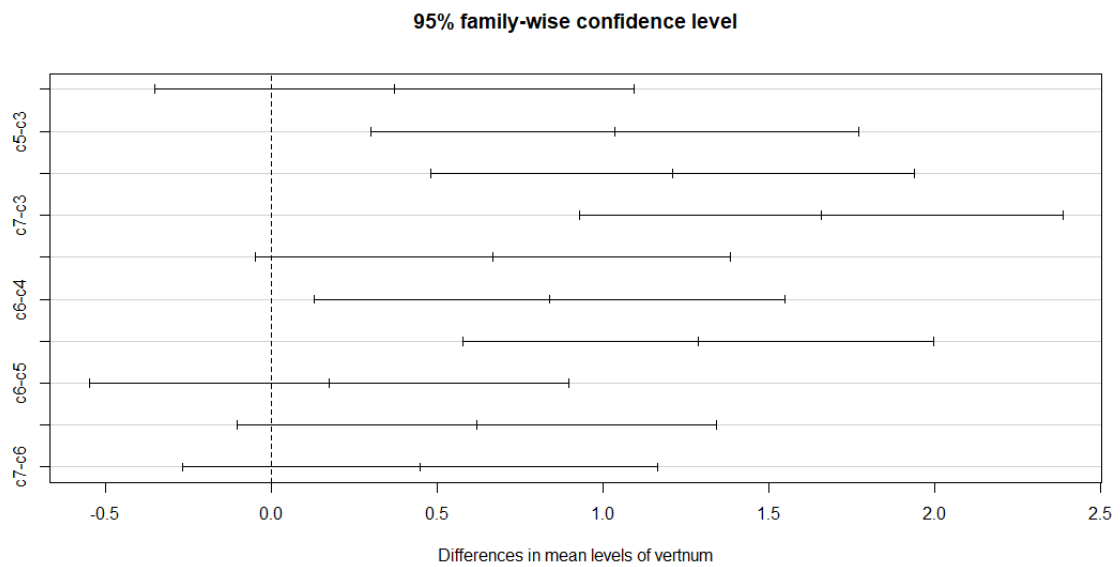
**Figure 9:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Foramen Sagittal Length measurement.



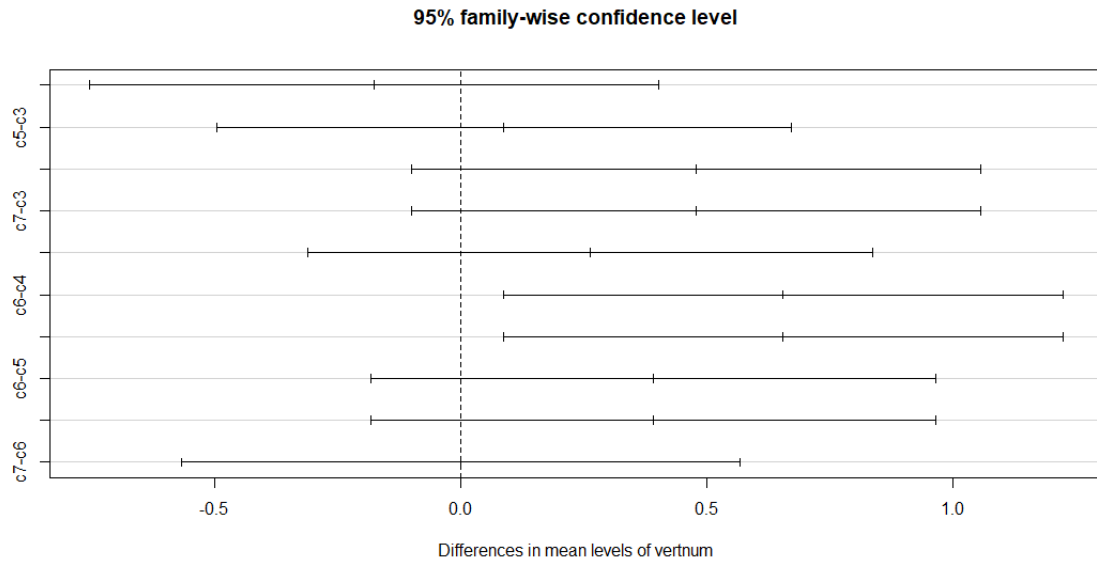
**Figure 10:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Foramen Coronal Width measurement.



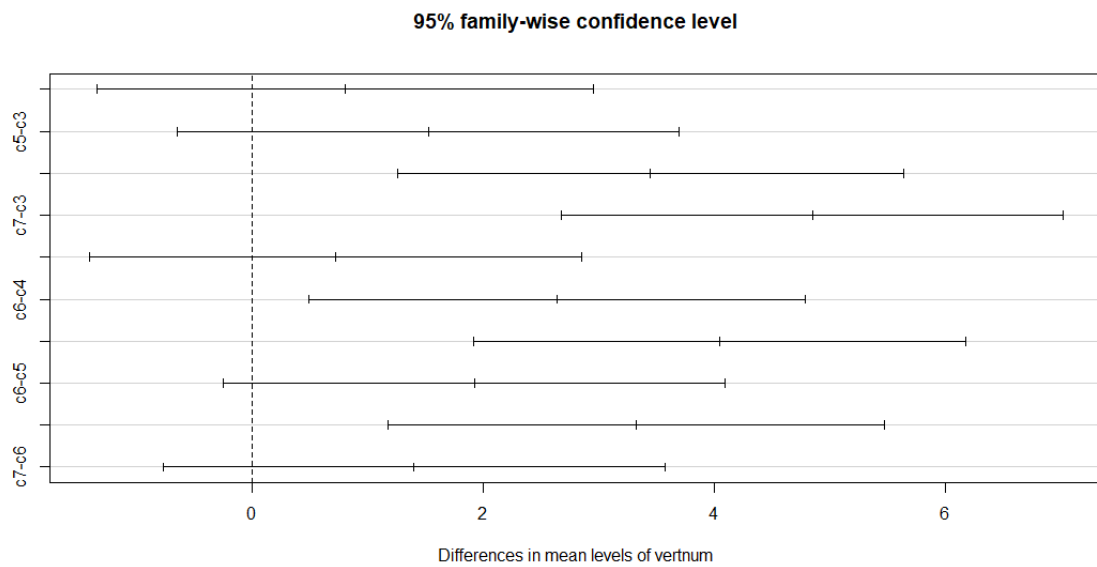
**Figure 11:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Pedicle Height measurement.



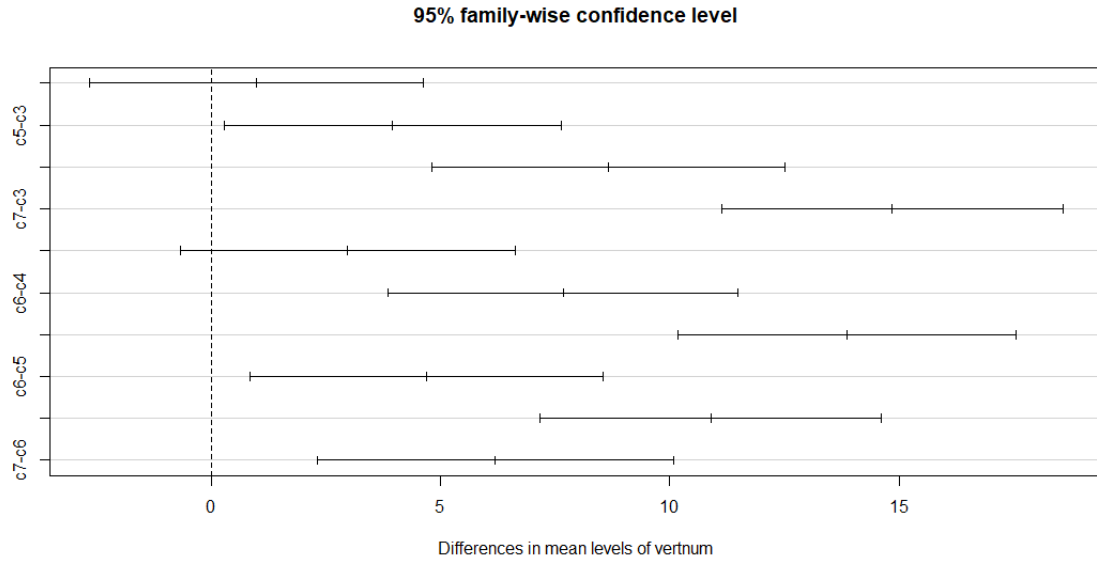
**Figure 12:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Pedicle Width measurement.



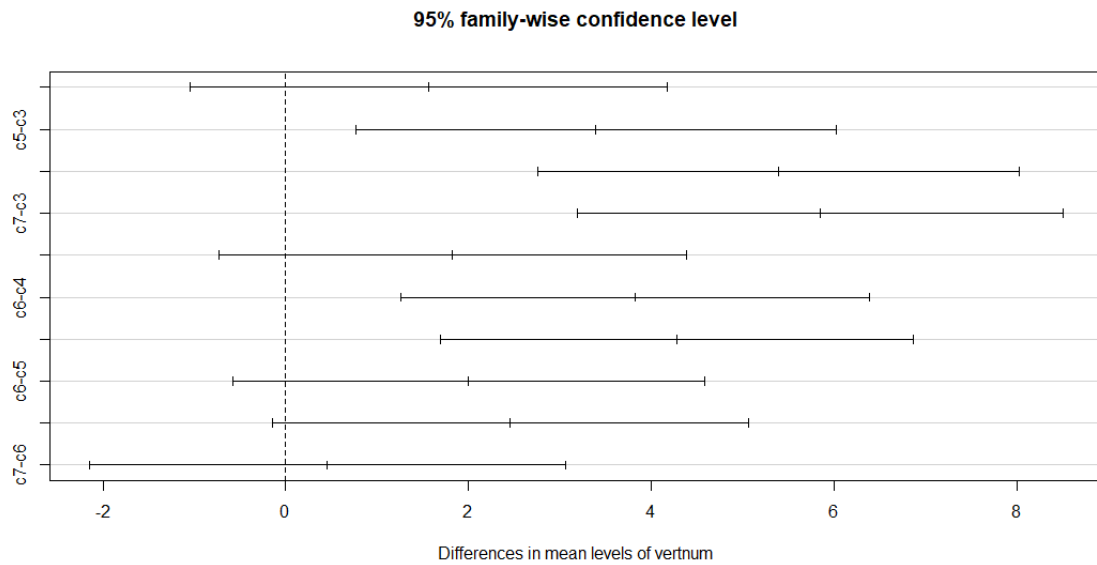
**Figure 13:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Pedicle Length measurement.



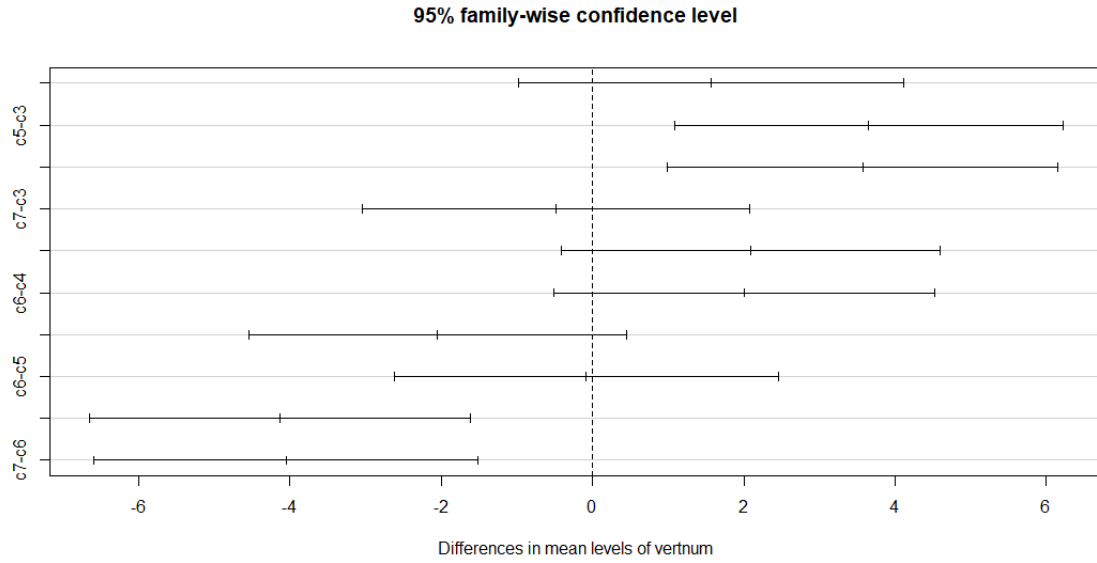
**Figure 14:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Transverse Process Length measurement.



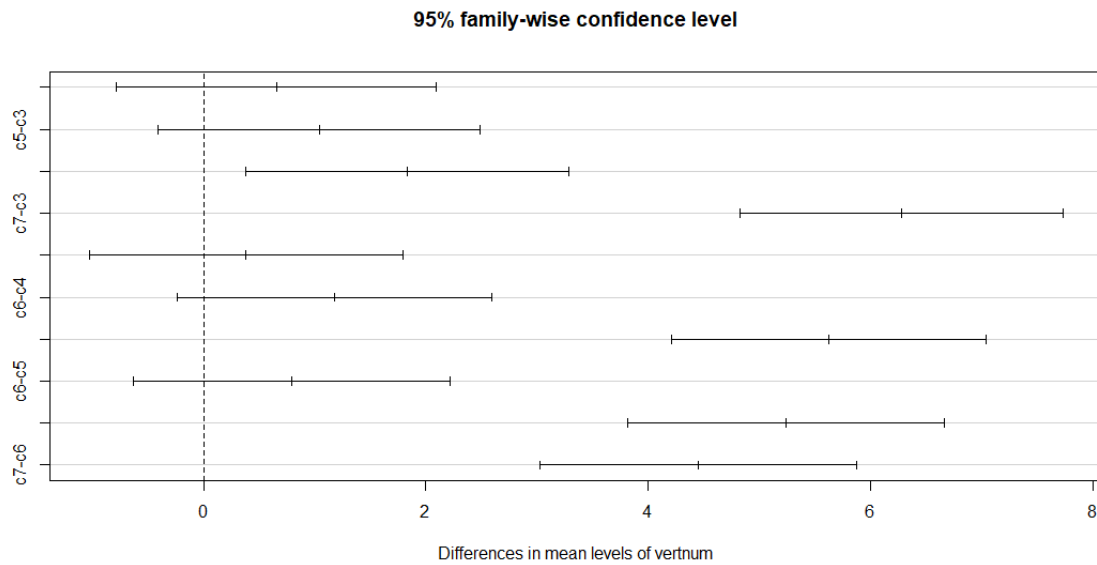
**Figure 15:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Transverse Process Width measurement.



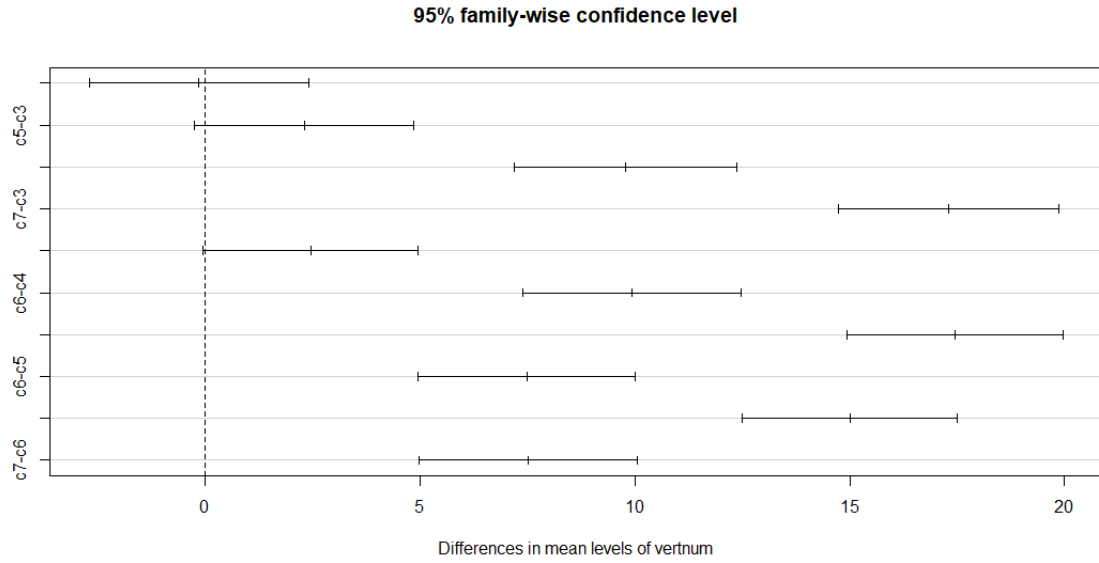
**Figure 16:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Superior Articular Process Width measurement.



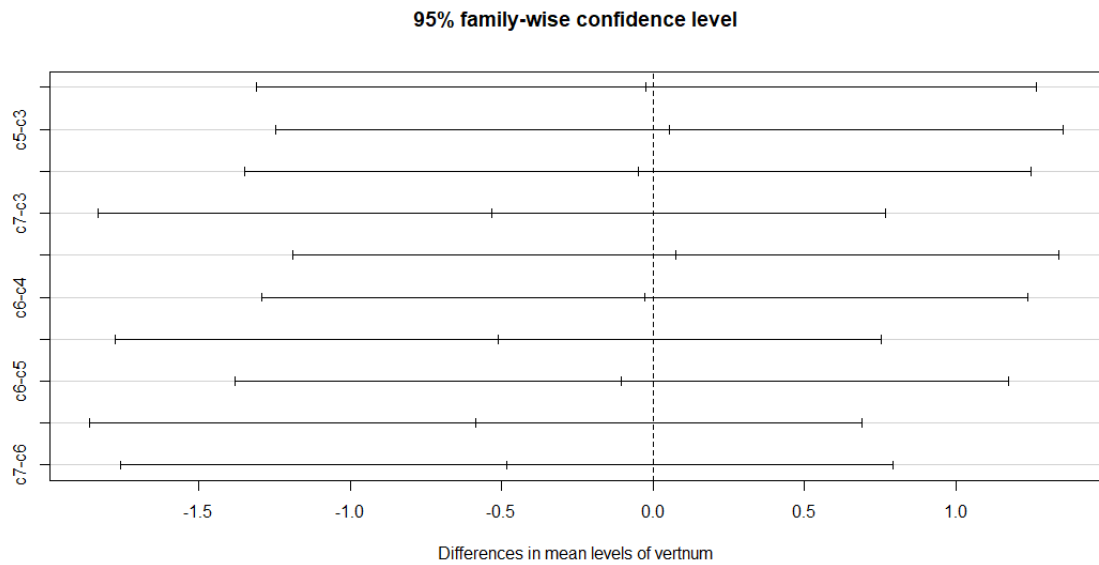
**Figure 17:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Inferior Articular Process Width measurement.



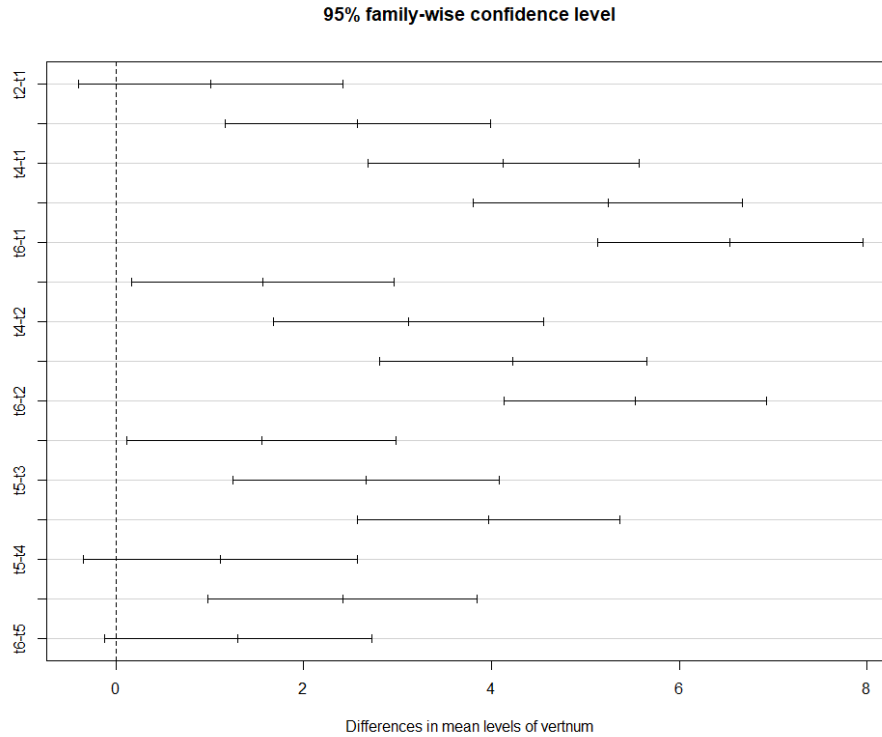
**Figure 18:** Results for the Tukey's HSD illustrating differences among for the cervical vertebrae (C3-C7) Articular Process Height measurement.



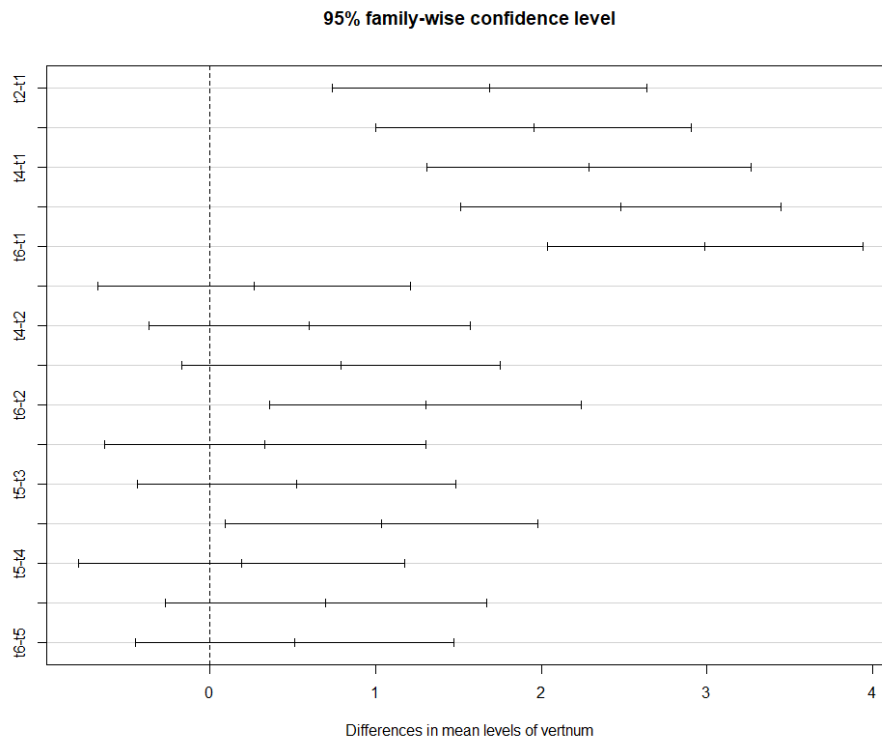
**Figure 19:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Spinous Process Length measurement.



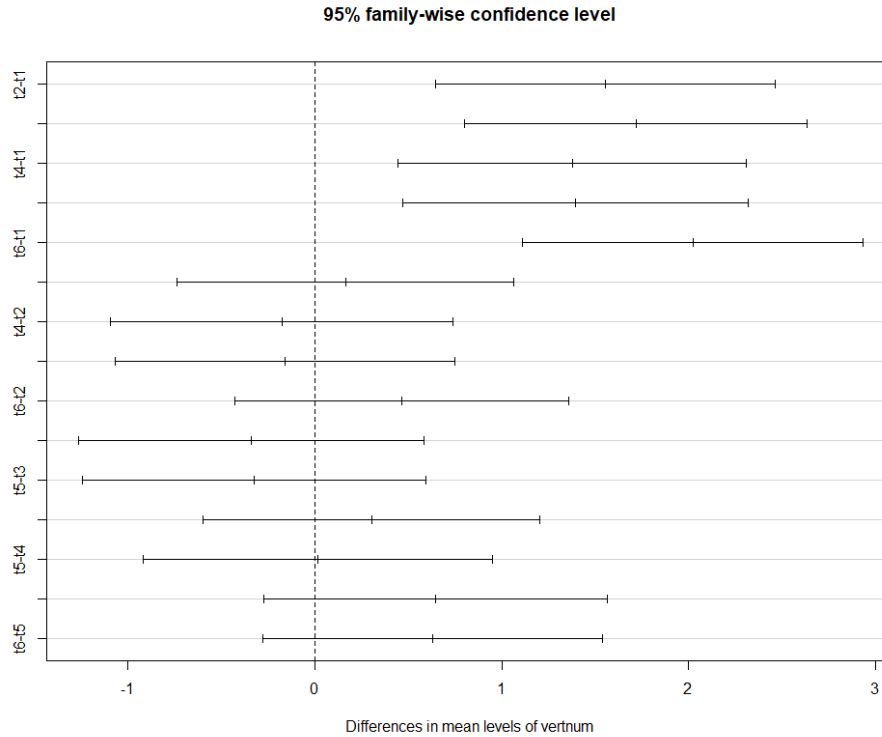
**Figure 20:** Results for the Tukey's HSD illustrating differences among the cervical vertebrae (C3-C7) Maximum Inferior Sagittal Length measurement.



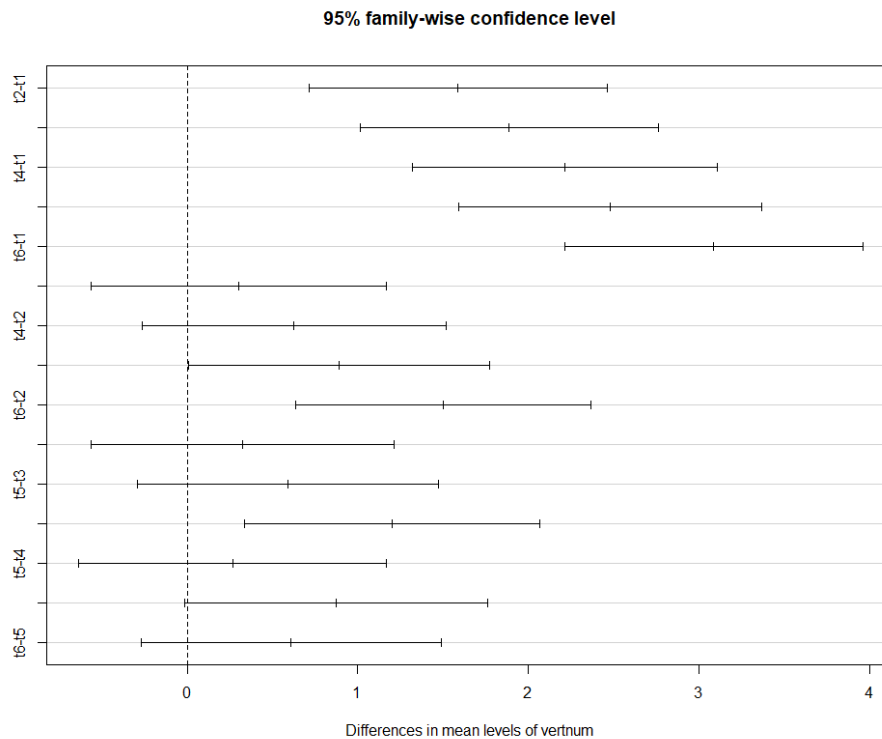
**Figure 21:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Superior Sagittal Length measurement.



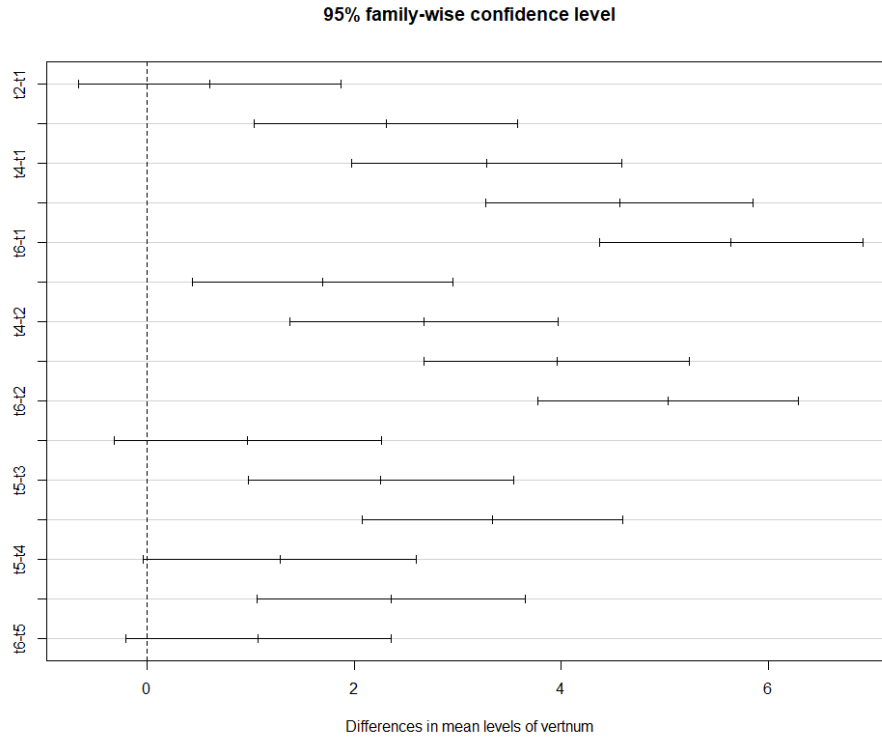
**Figure 22:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Centroid Height measurement.



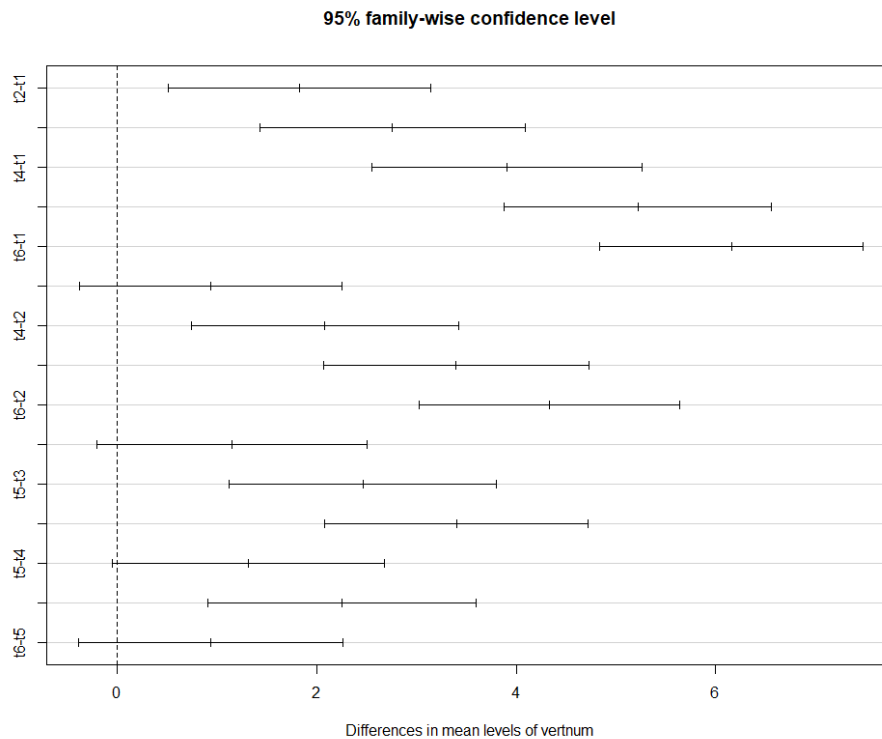
**Figure 23:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Anterior Height measurement.



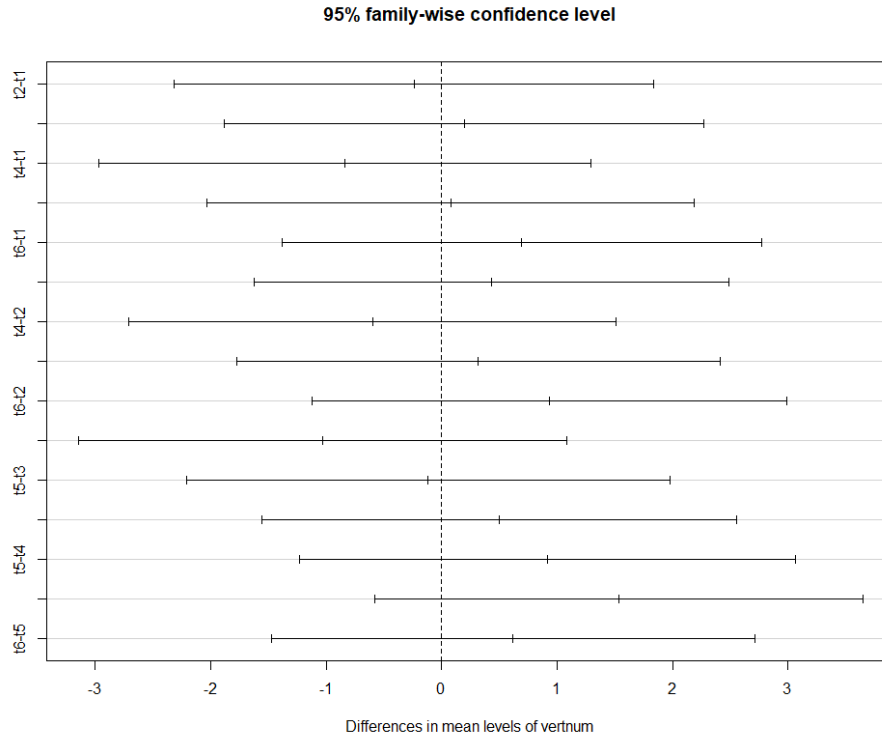
**Figure 24:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Posterior Height measurement.



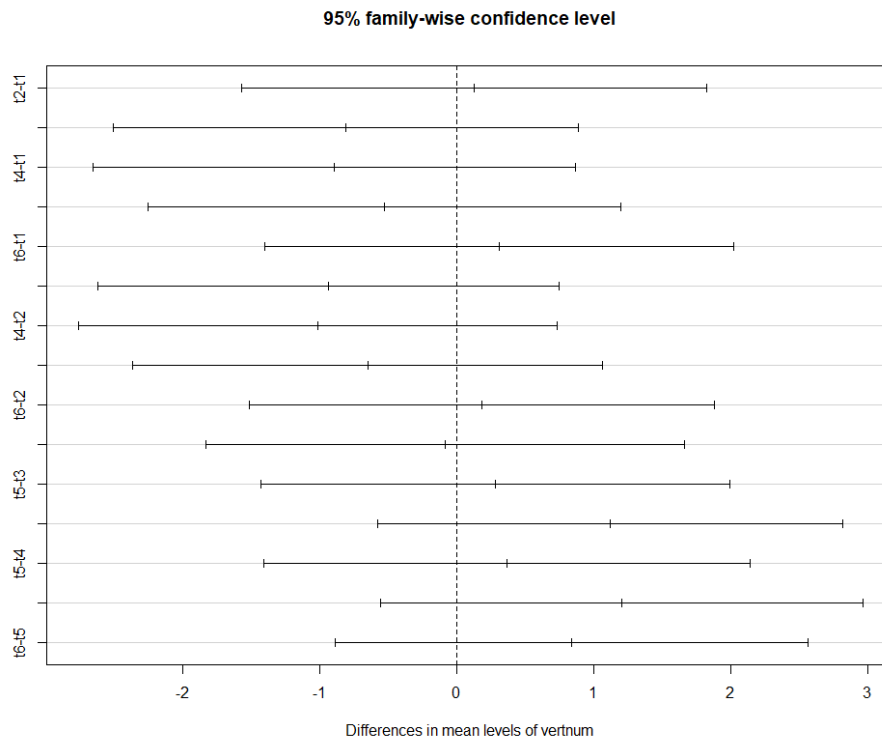
**Figure 25:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Superior Articular Length measurement.



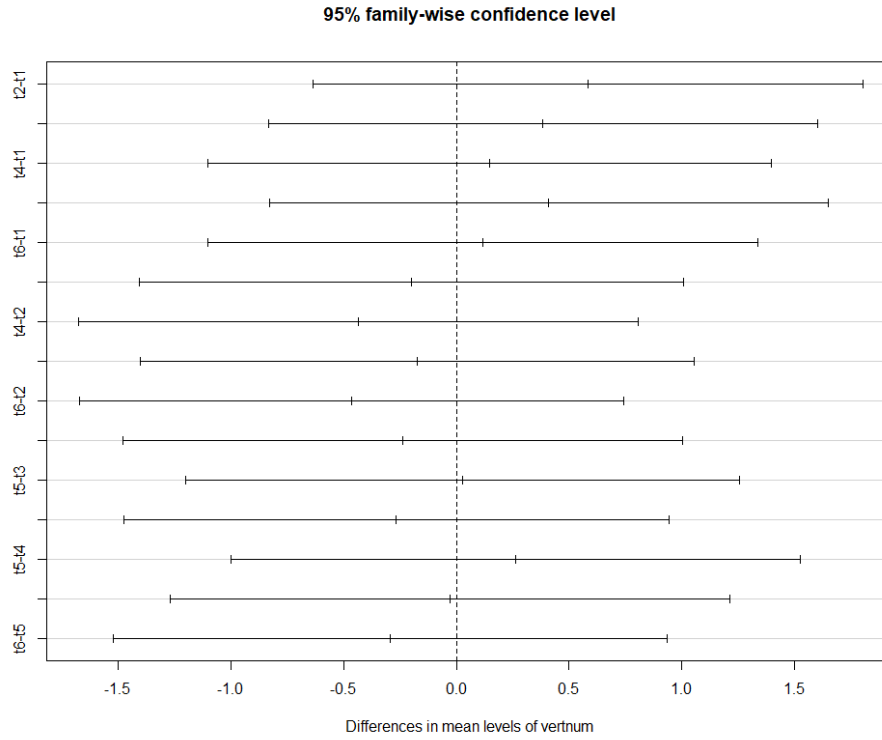
**Figure 26:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Superior Articular Length measurement.



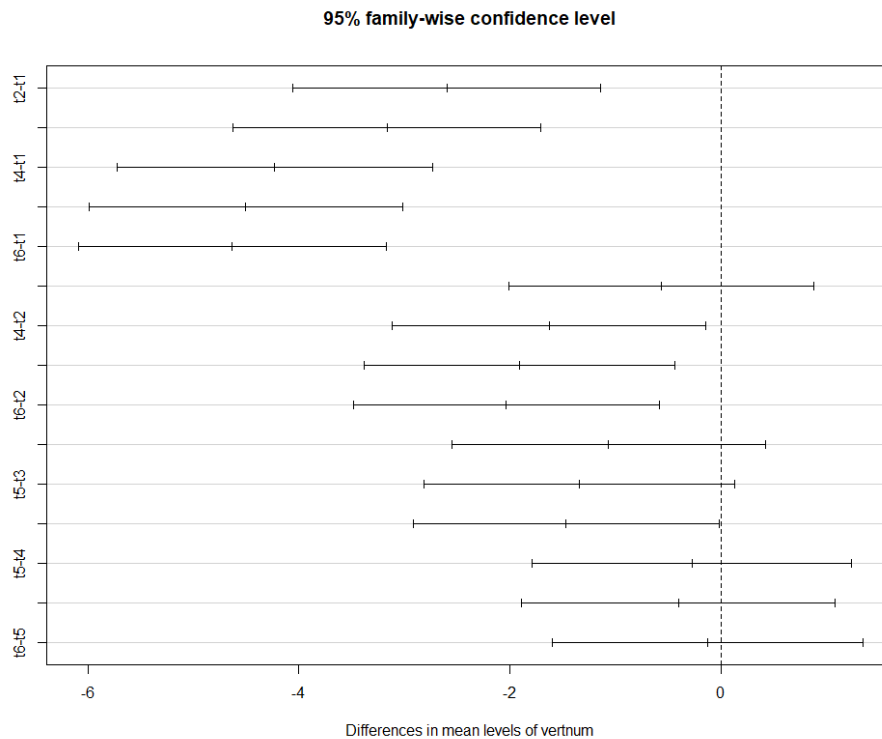
**Figure 27:** Results for the Tukey's HSD differences among the upper thoracic vertebrae (T1-T6) Maximum Superior Articular Width measurement.



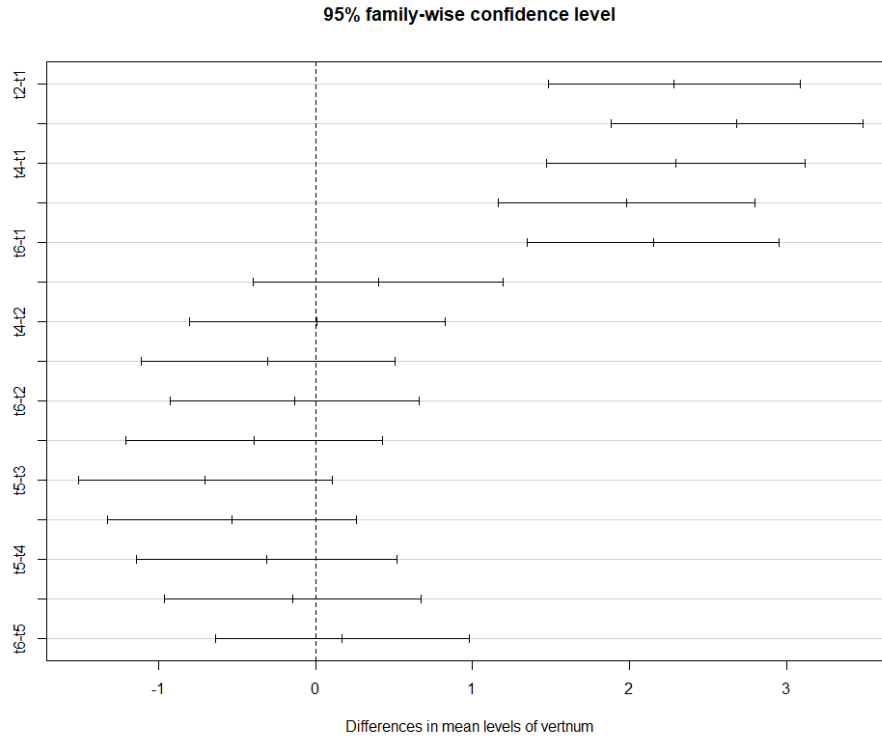
**Figure 28:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Inferior Articular Width measurement.



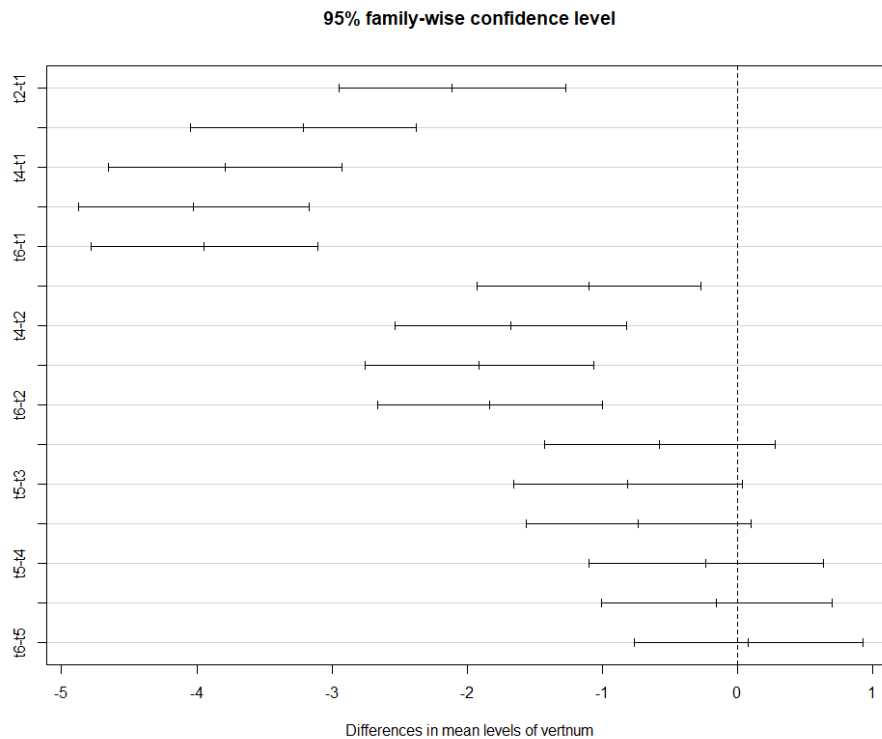
**Figure 29:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Foramen Sagittal Length measurement.



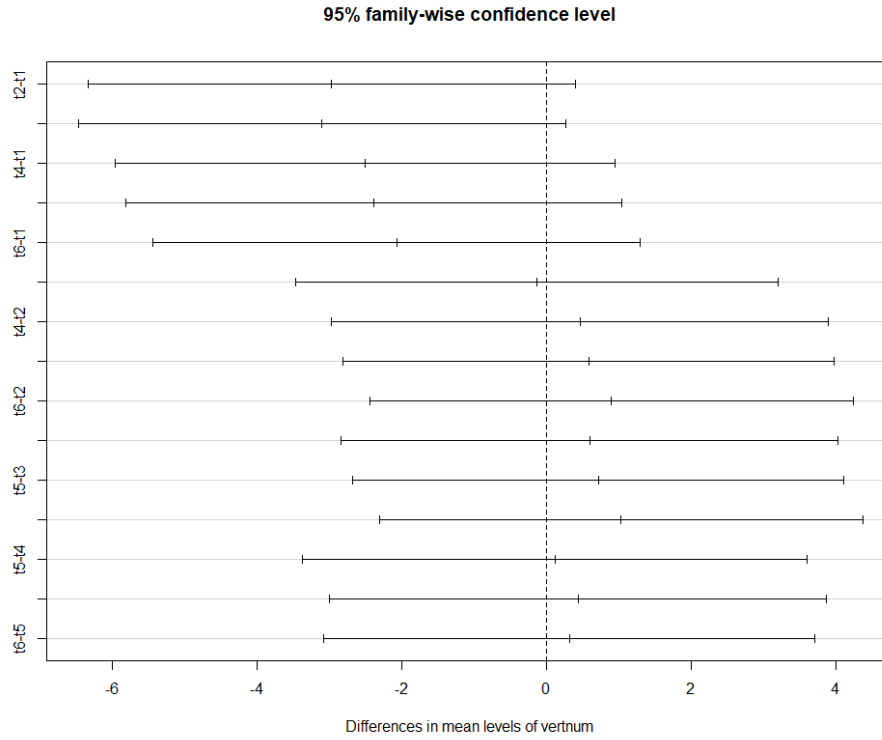
**Figure 30:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Foramen Coronal Width measurement.



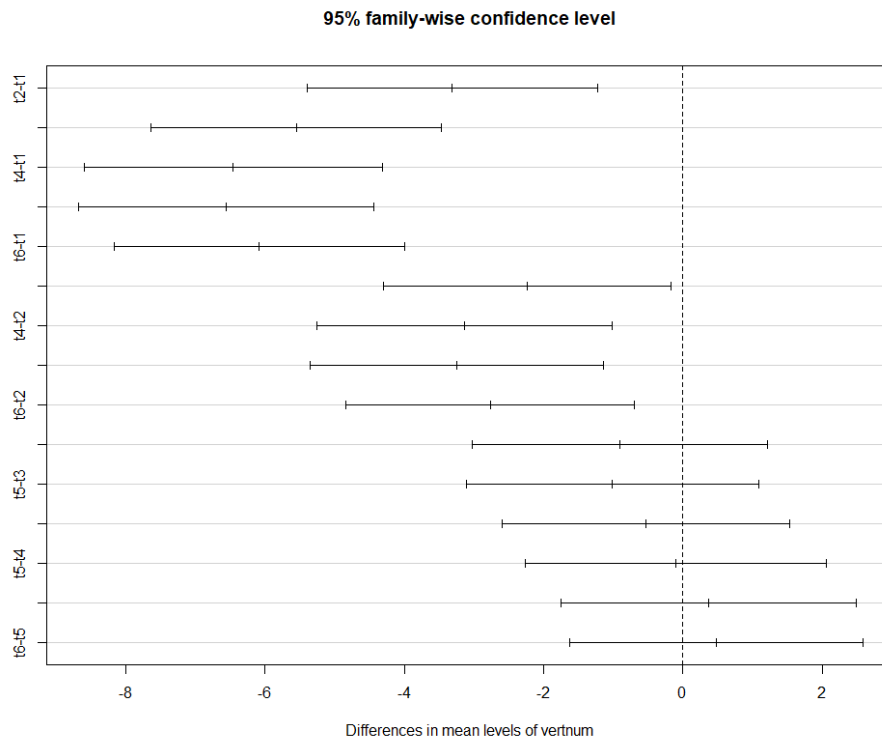
**Figure 31:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Pedicle Height measurement.



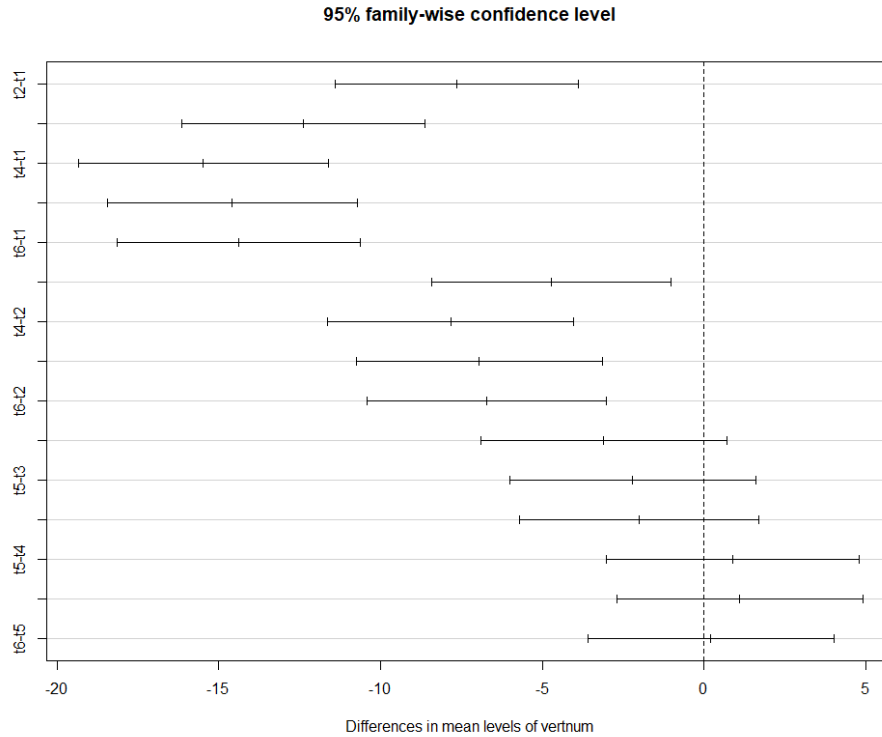
**Figure 32:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Pedicle Width measurement.



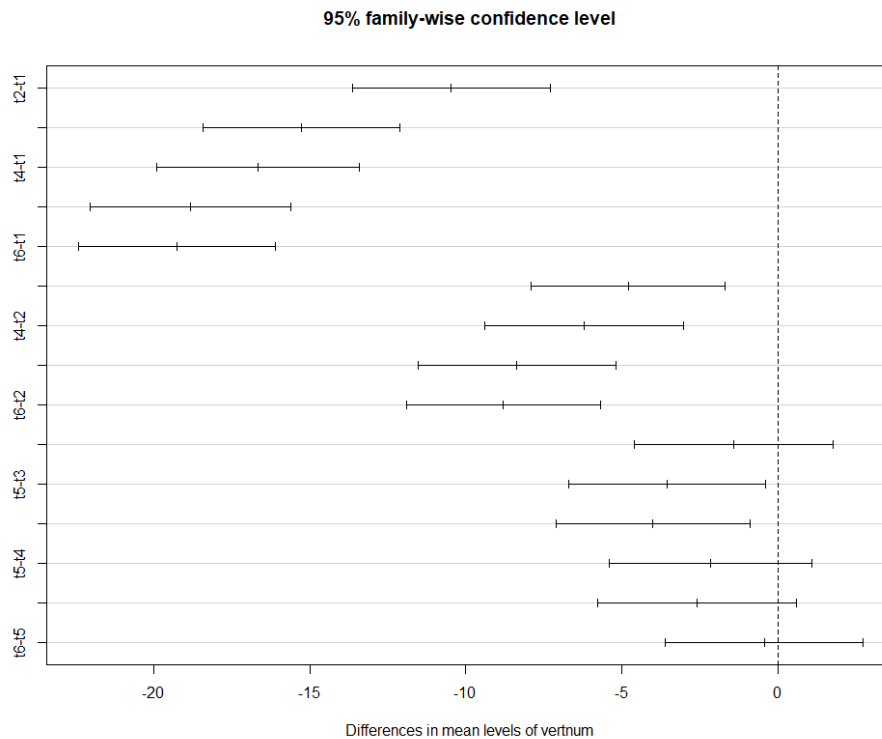
**Figure 33:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Pedicle Length measurement.



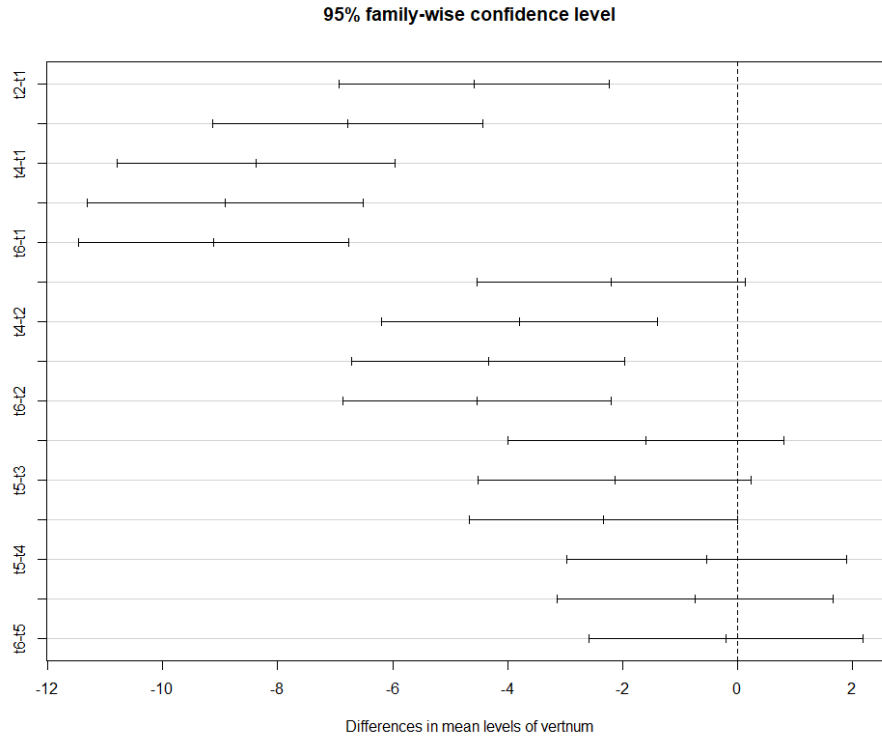
**Figure 34:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Transverse Process Length measurement.



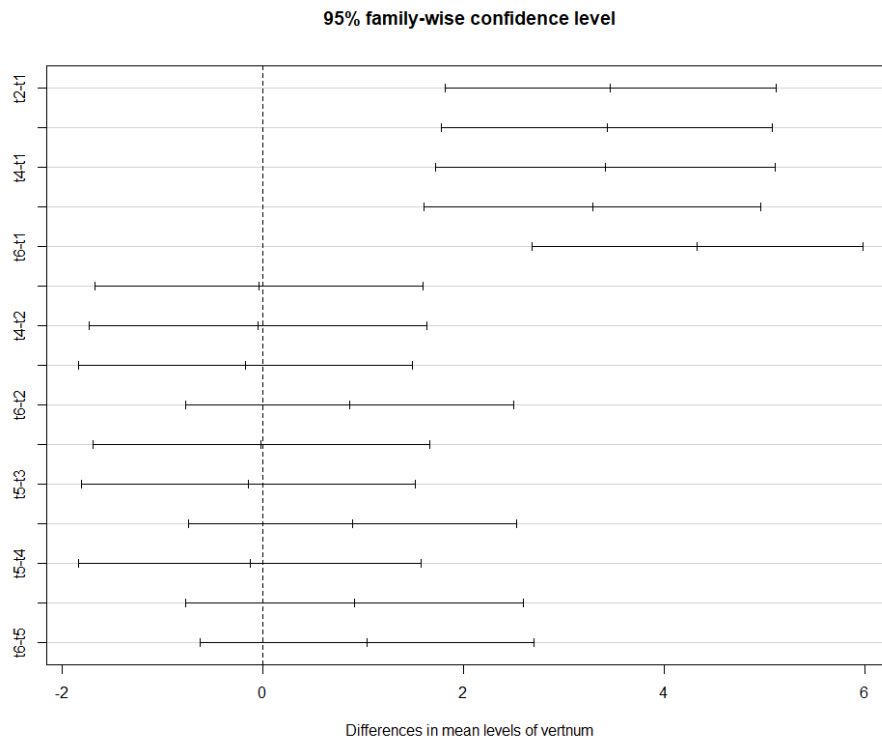
**Figure 35:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Transverse Process Width measurement.



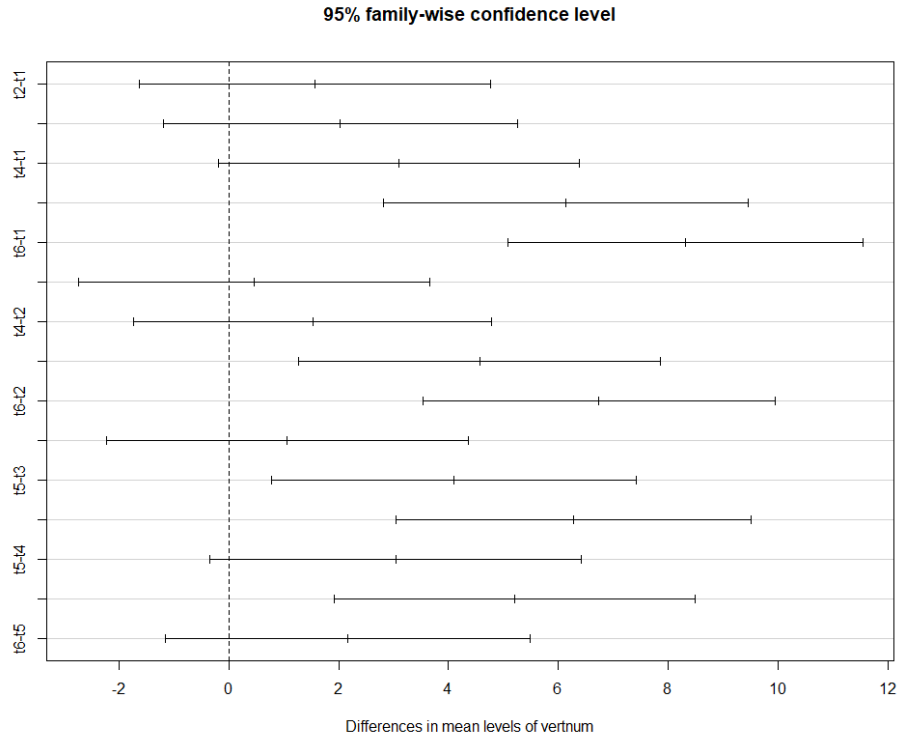
**Figure 36:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Superior Articular Process Width measurement.



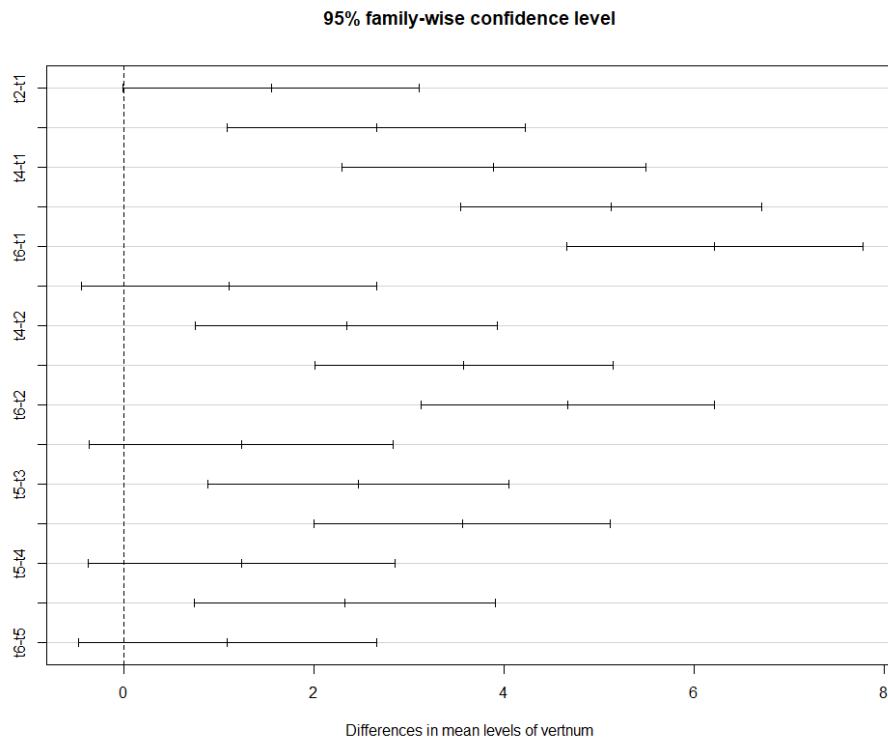
**Figure 37:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Inferior Articular Process Width measurement.



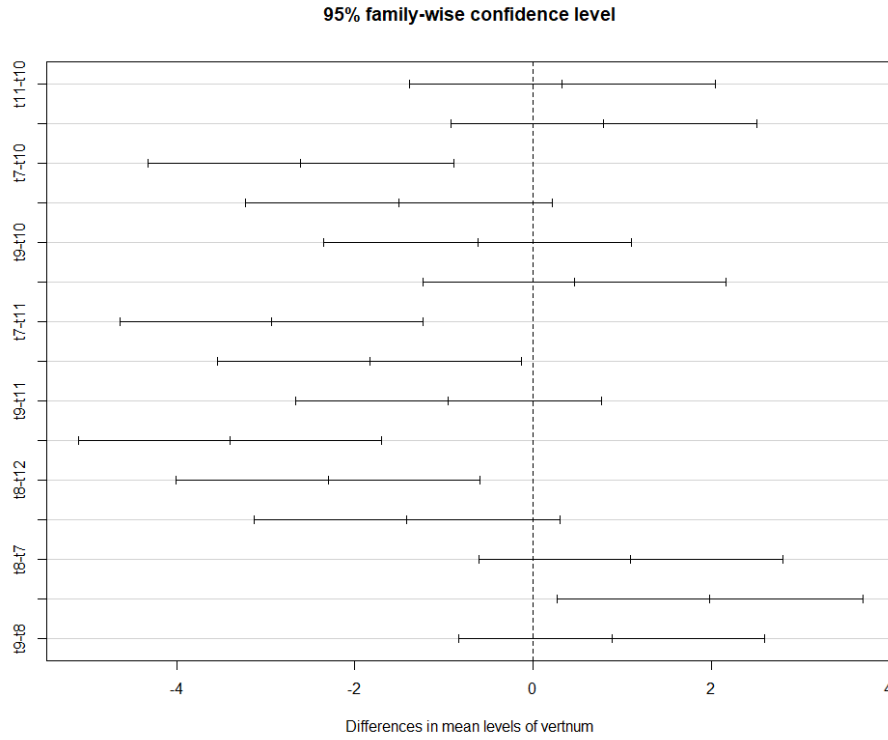
**Figure 38:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Articular Process Height measurement.



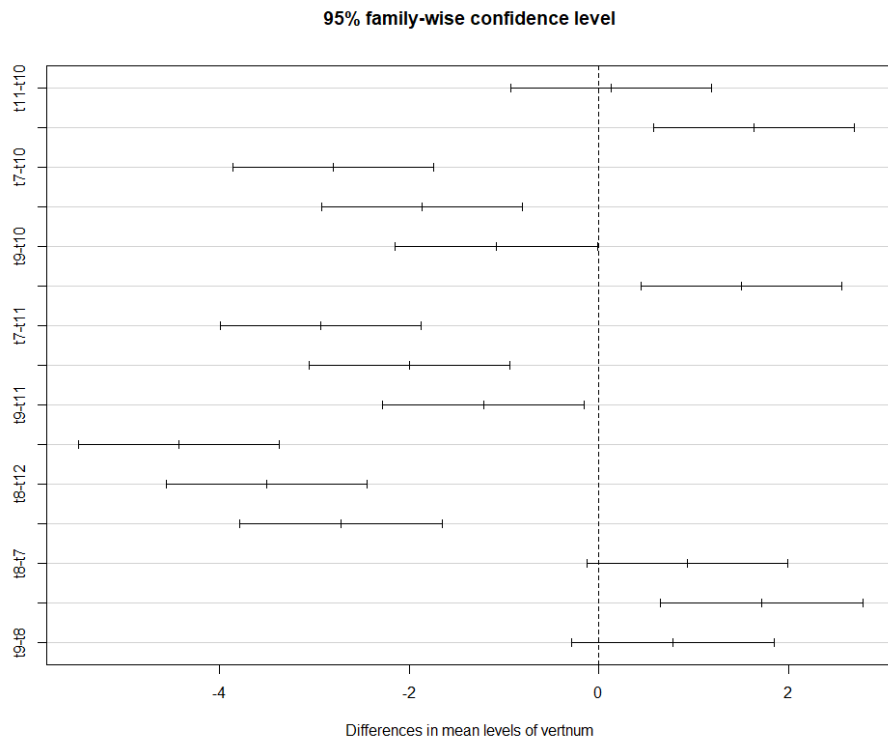
**Figure 39:** Results for the Tukey's HSD illustrating differences among the upper thoracic vertebrae (T1-T6) Maximum Spinous Process Length measurement.



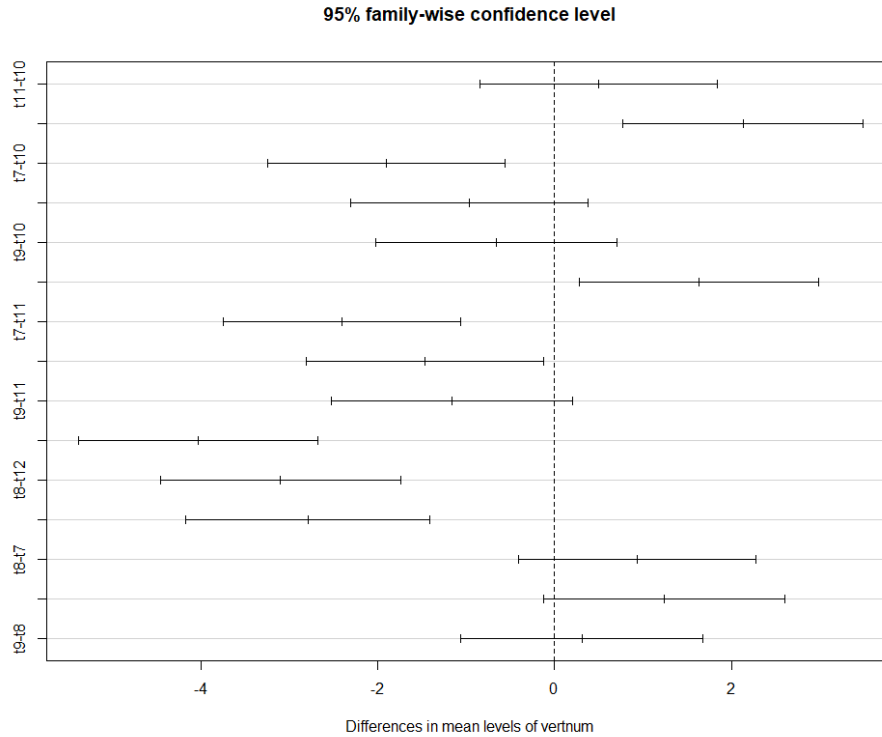
**Figure 40:** Results for the Tukey's HSD illustrating sub-type differences and variable overlap for the upper thoracic vertebrae (T1-T6) Maximum Inferior Sagittal Length measurement.



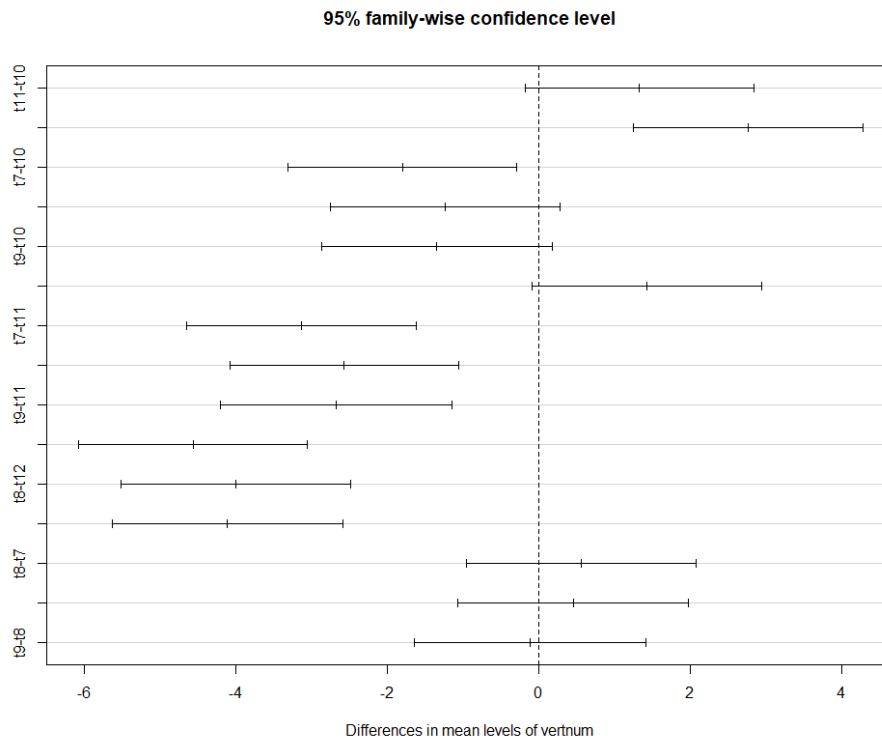
**Figure 41:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Superior Sagittal Length measurement.



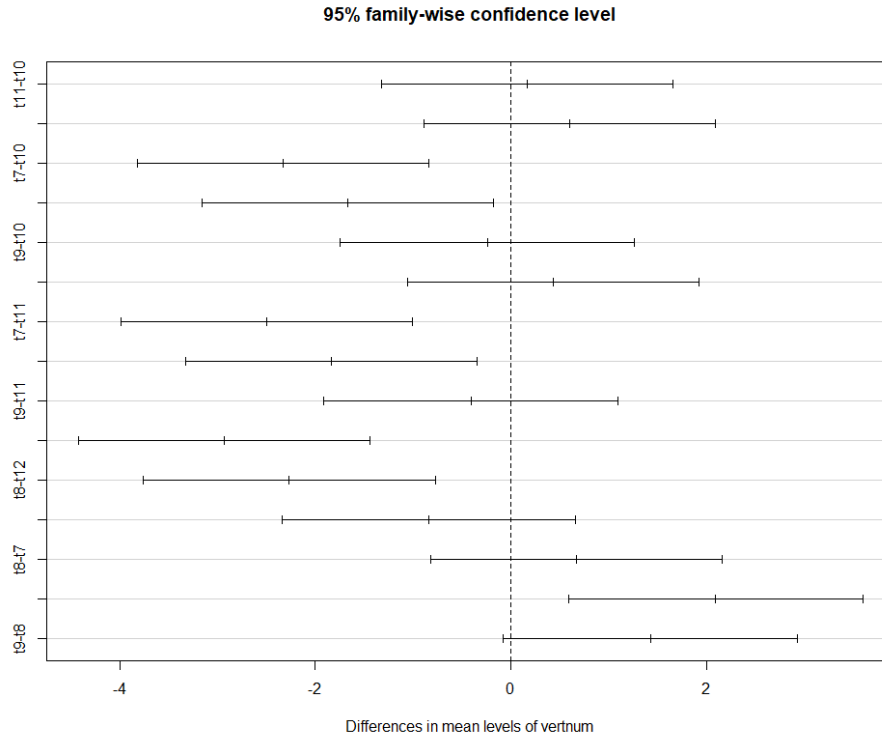
**Figure 42:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Centroid Height measurement.



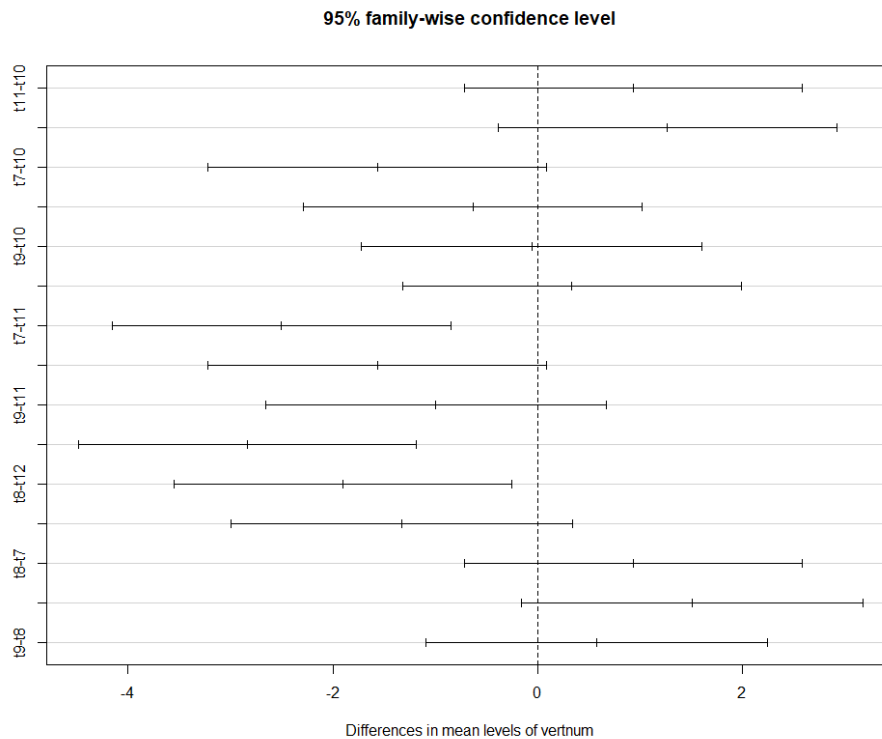
**Figure 43:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Anterior Height measurement.



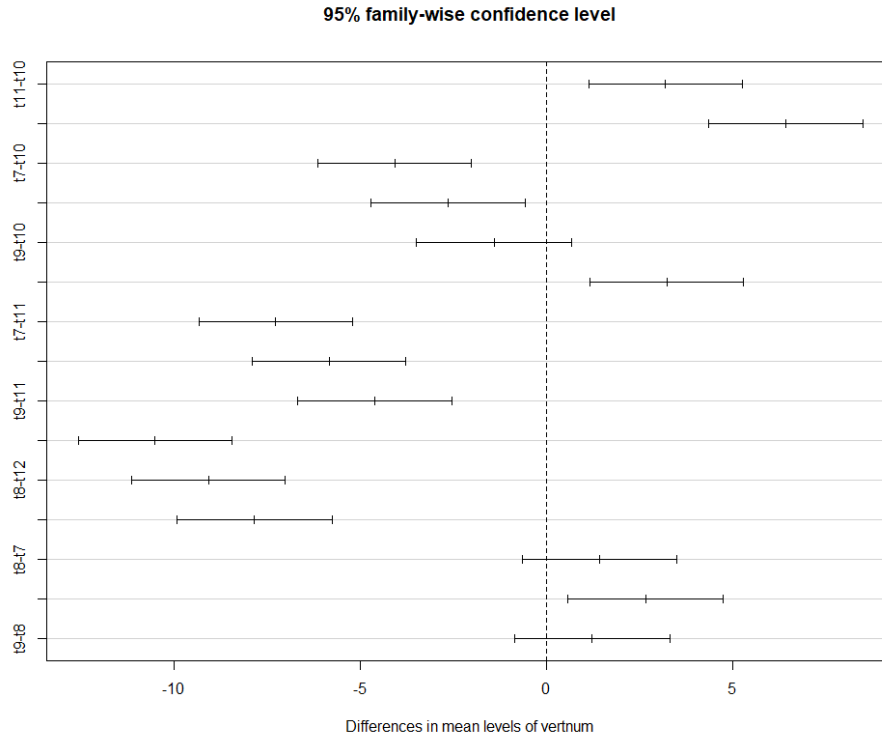
**Figure 44:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Posterior Height measurement.



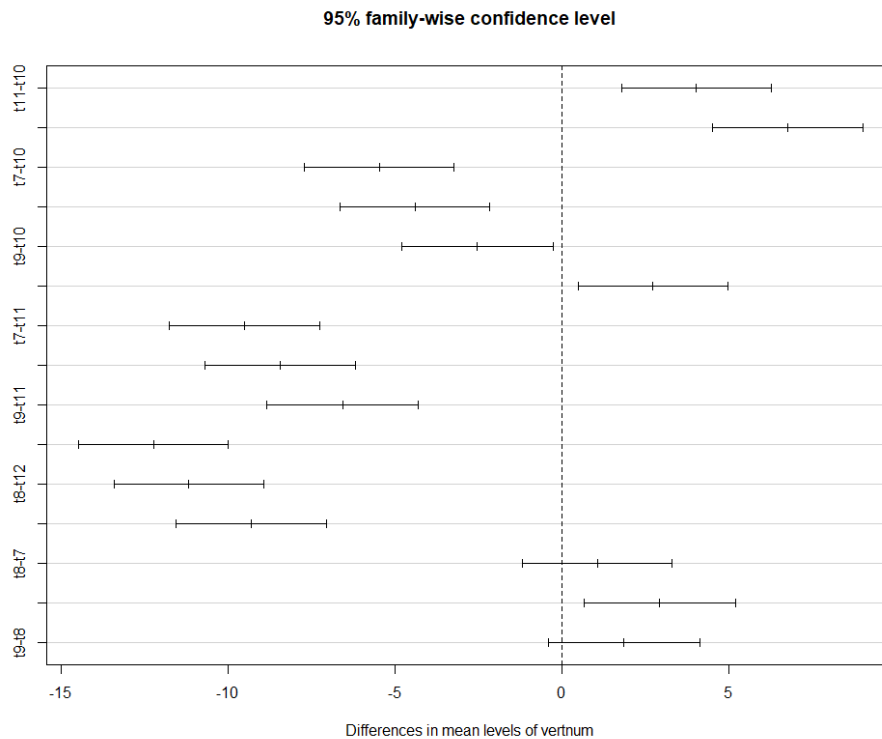
**Figure 45:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Superior Articular Length measurements.



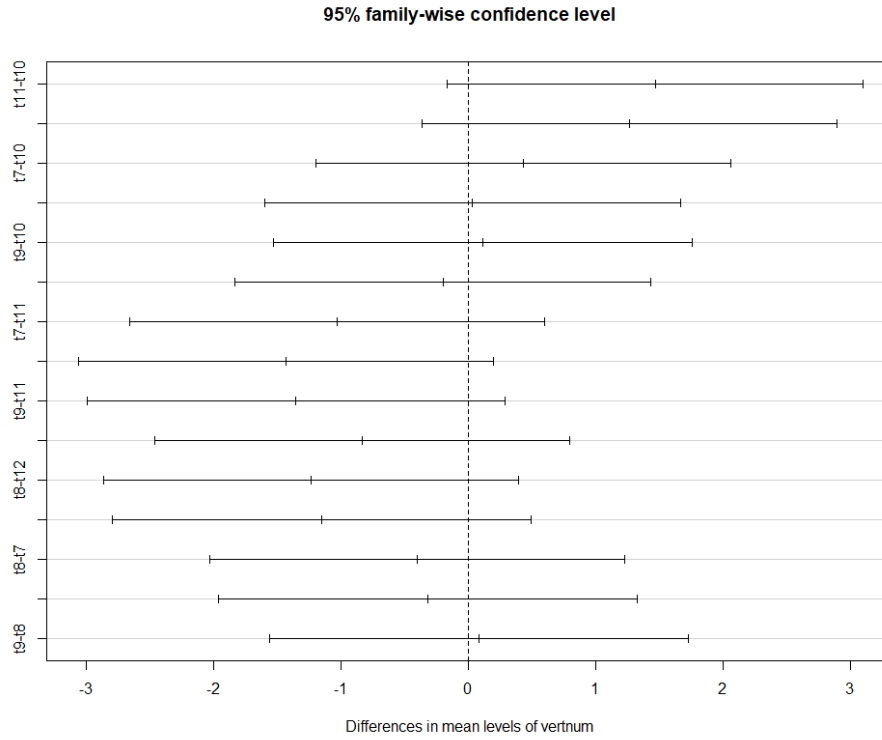
**Figure 46:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Superior Articular Length measurement.



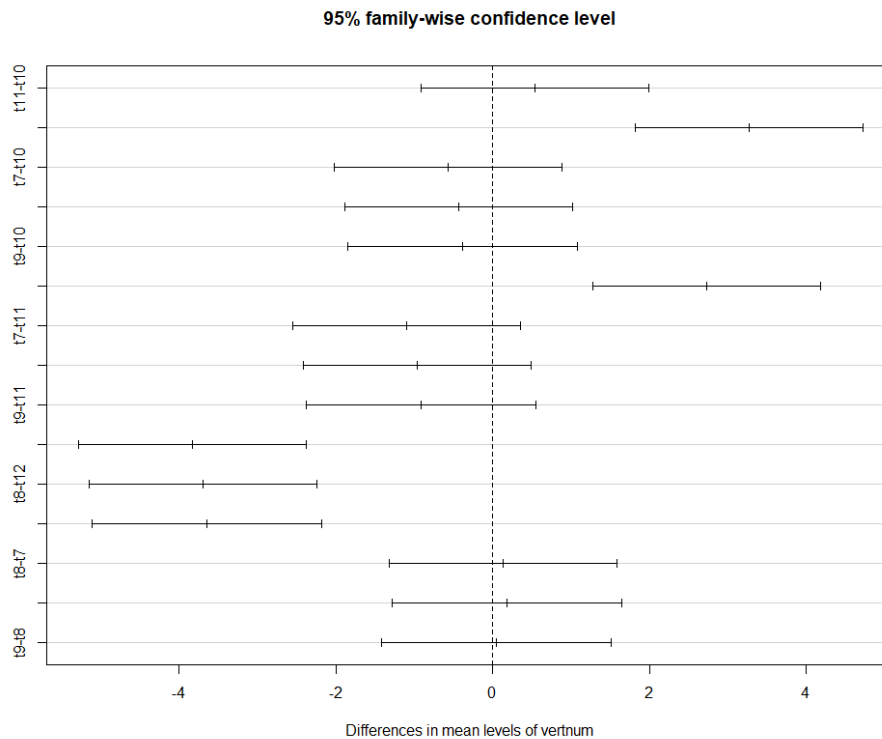
**Figure 47:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Superior Articular Width measurement.



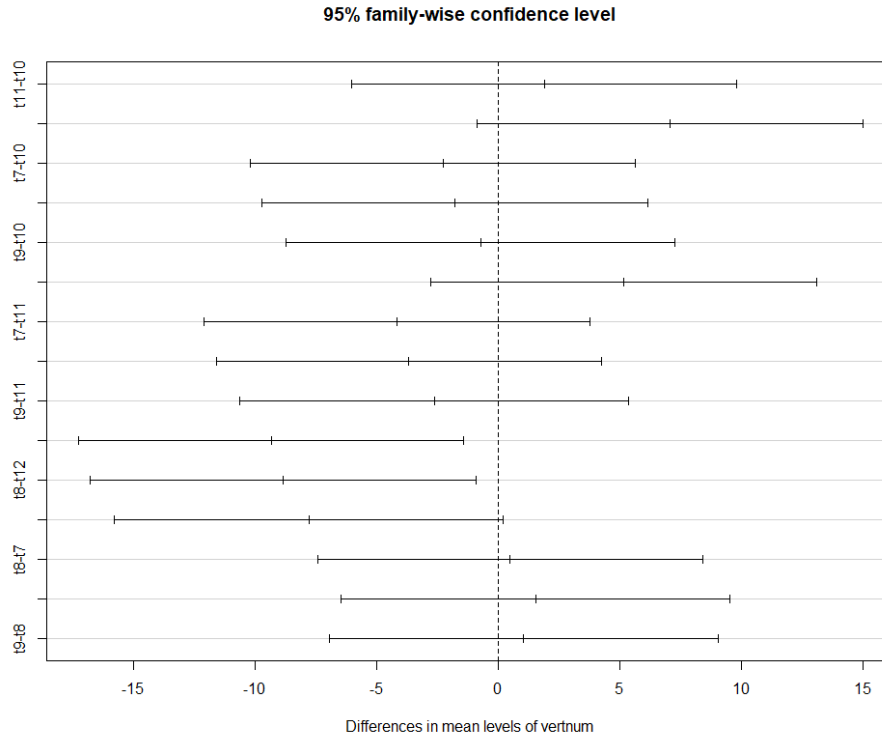
**Figure 48:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Inferior Articular Width measurement.



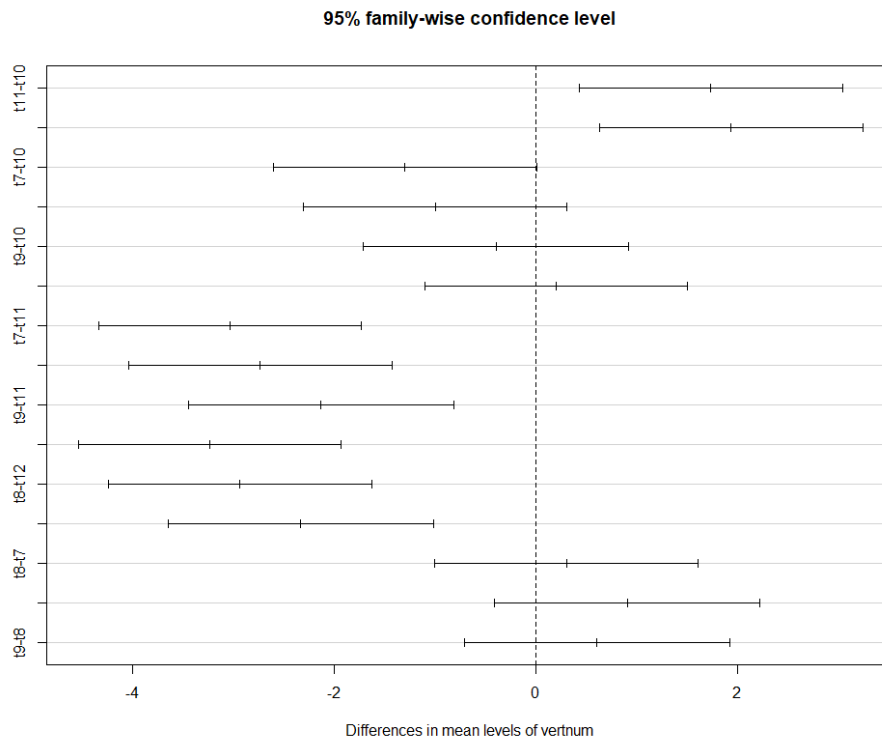
**Figure 49:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Foramen Sagittal Length measurement.



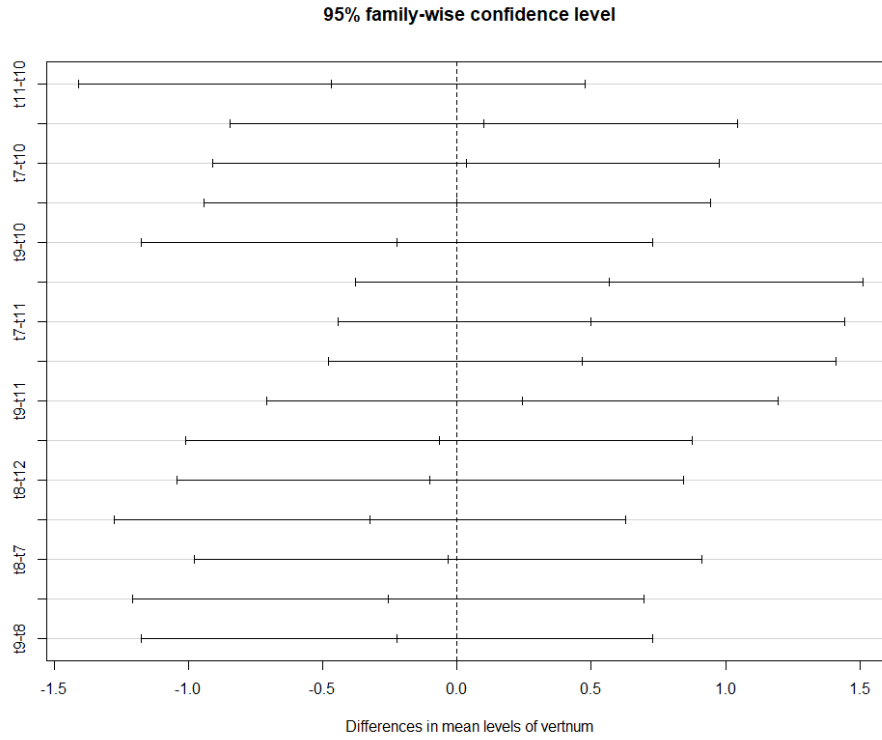
**Figure 50:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Foramen Coronal Width measurement.



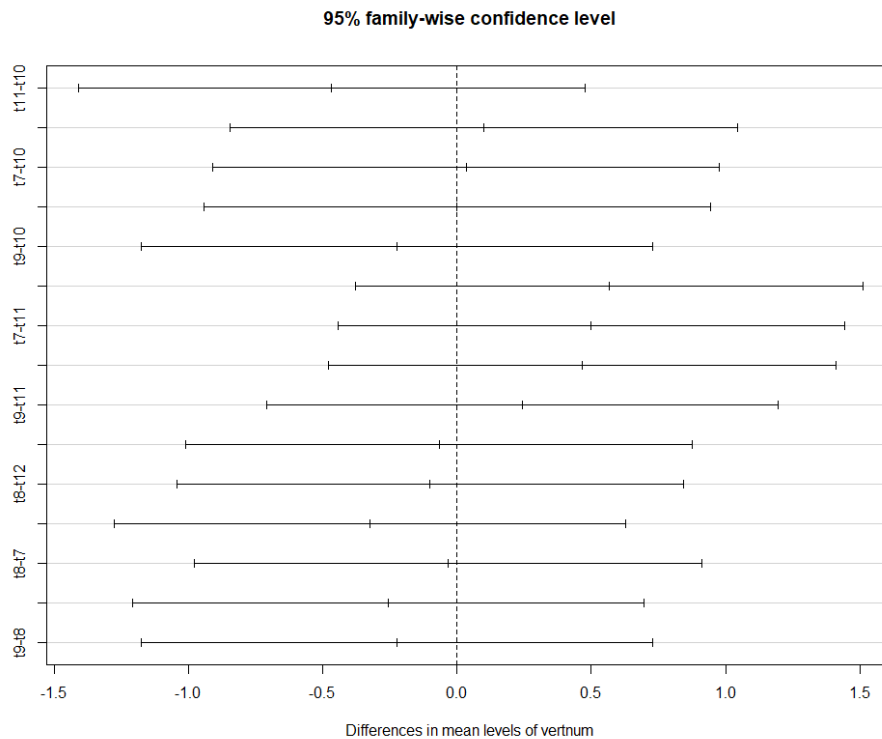
**Figure 51:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Pedicle Height measurement.



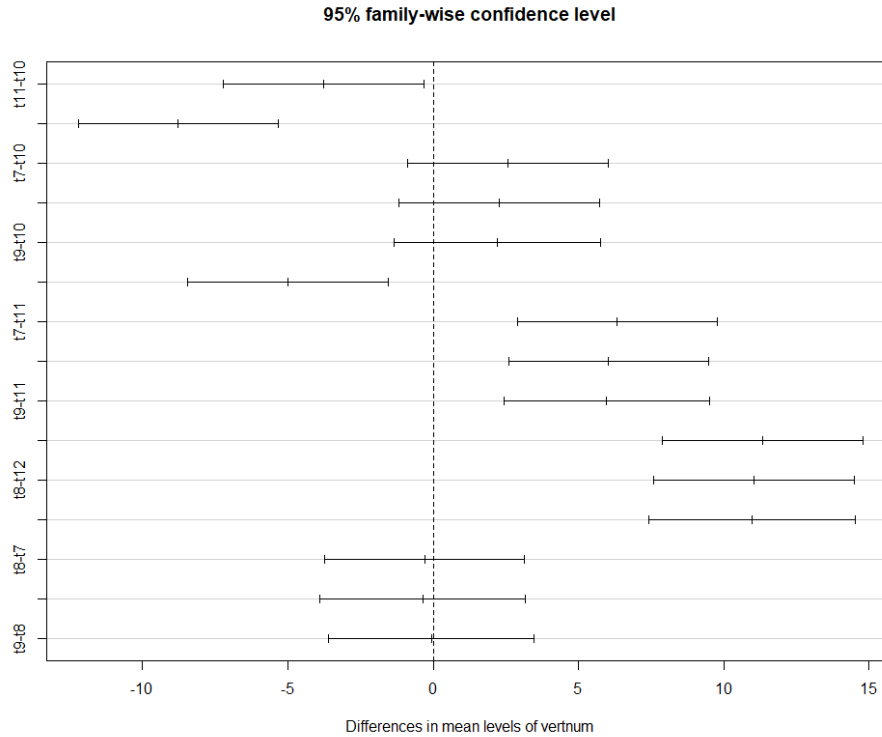
**Figure 52:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Pedicle Width measurement.



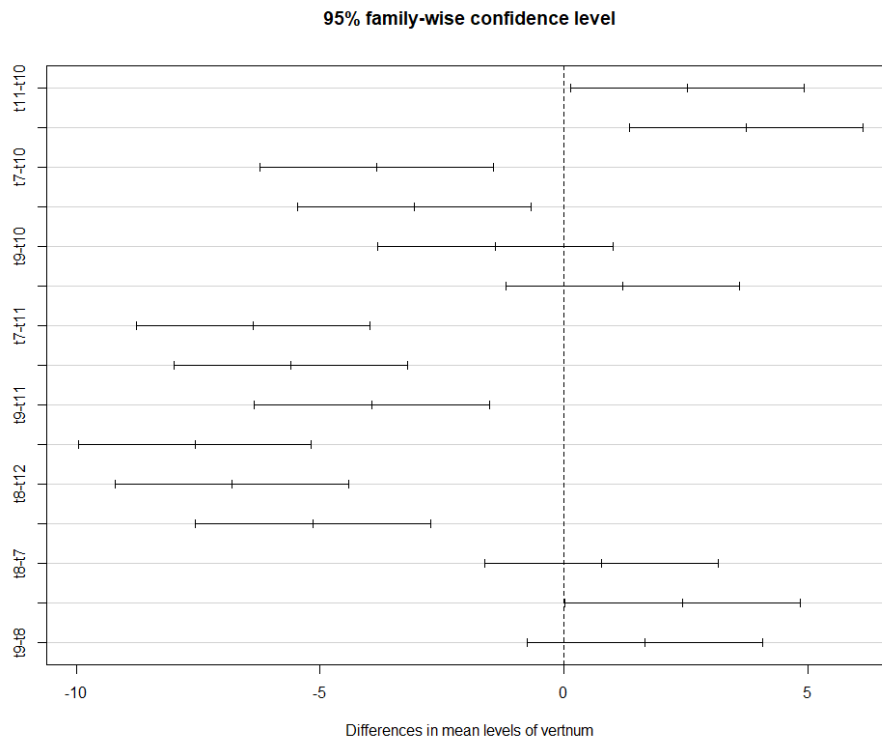
**Figure 53:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Pedicle Length measurement.



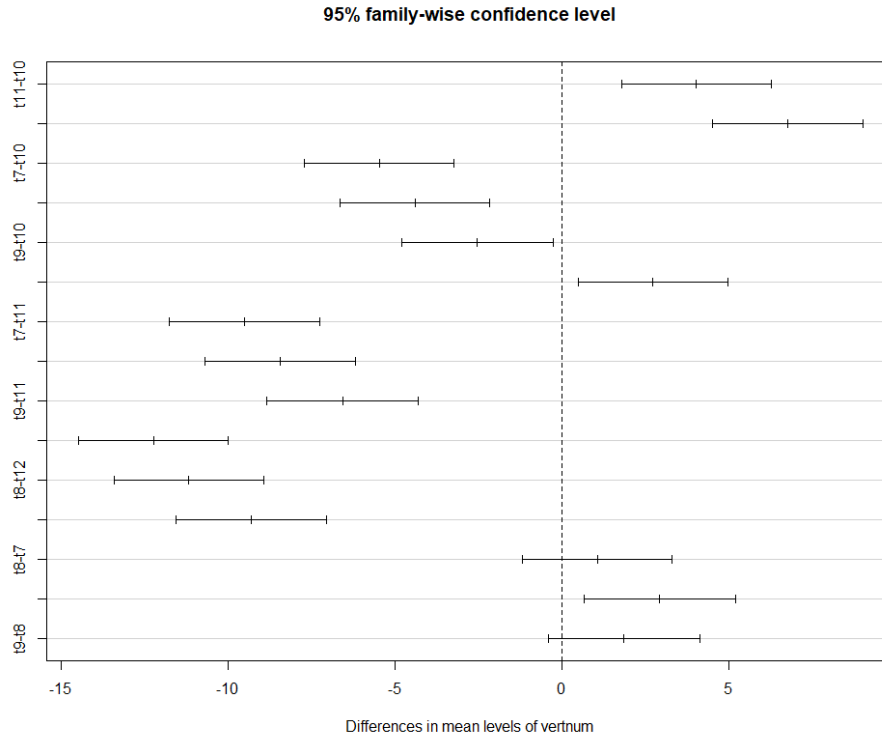
**Figure 54:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Transverse Process Length measurement.



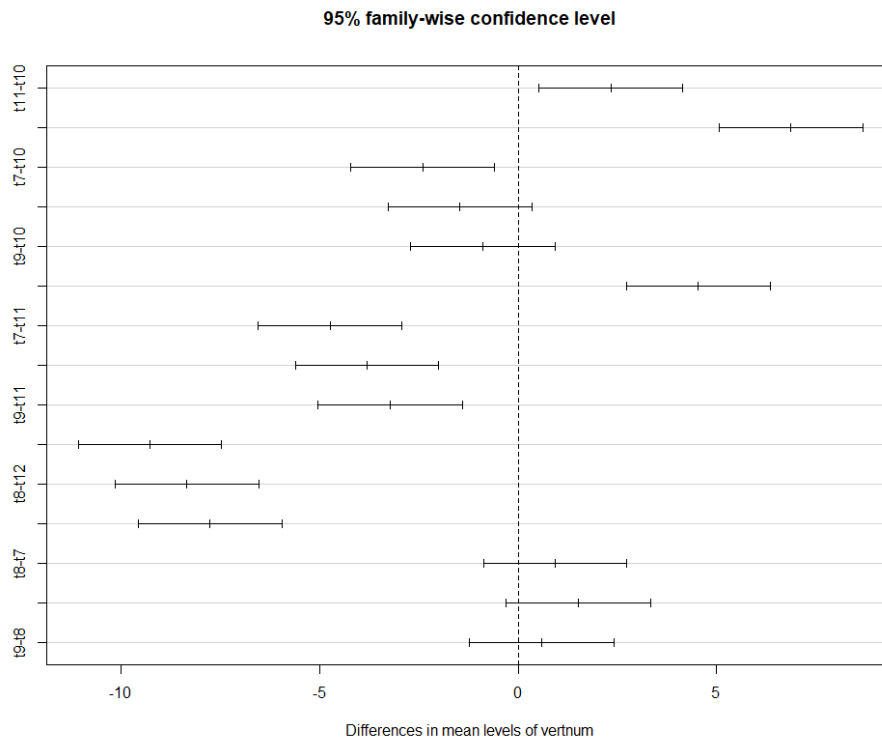
**Figure 55:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Transverse Process Width measurement.



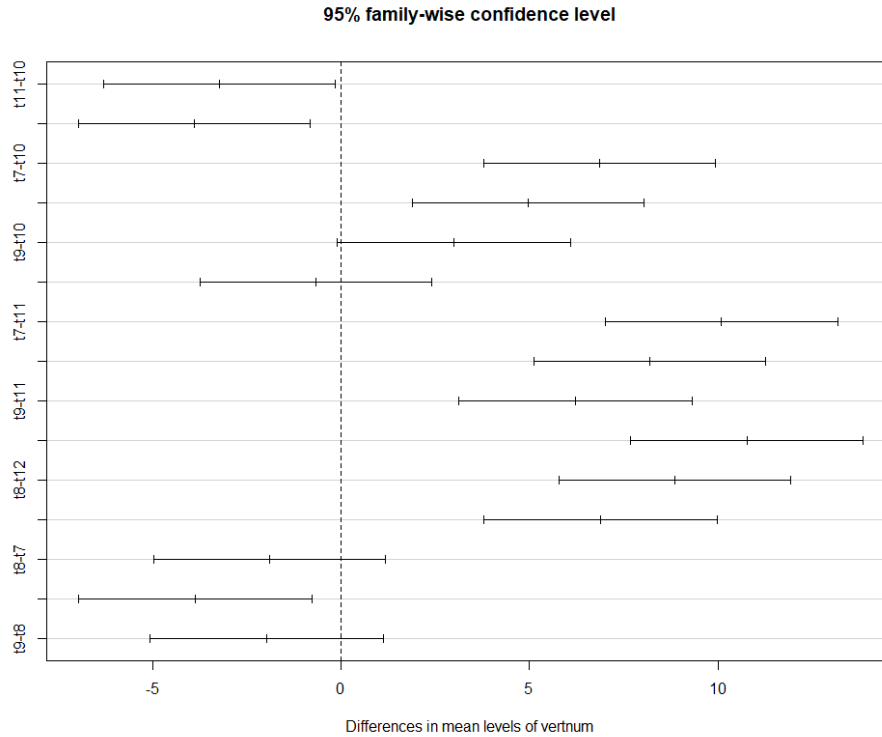
**Figure 56:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Superior Articular Process Width measurement.



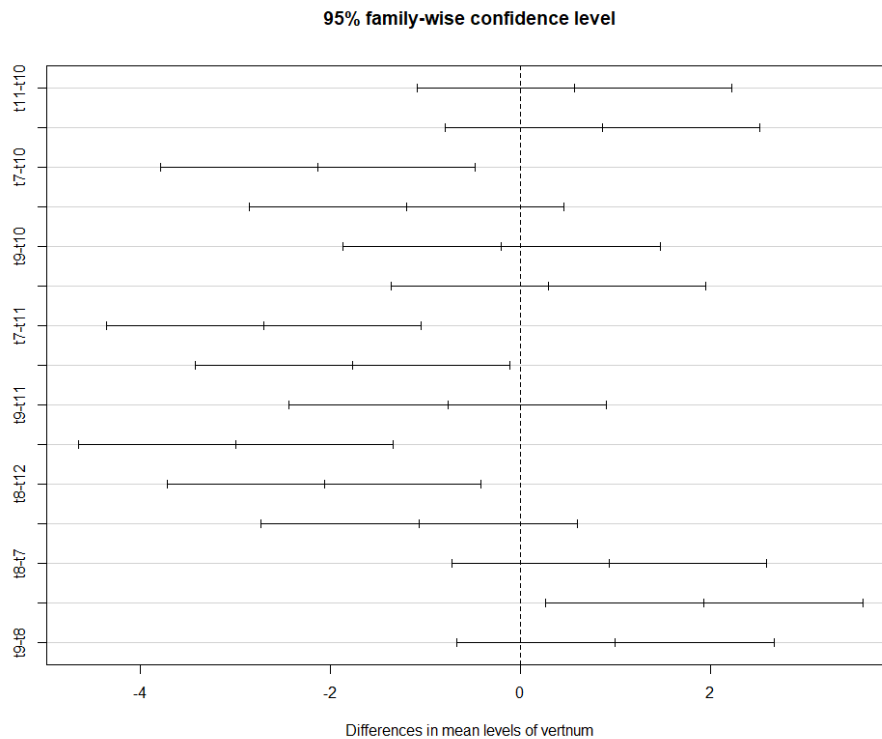
**Figure 57:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Inferior Articular Process Width measurement.



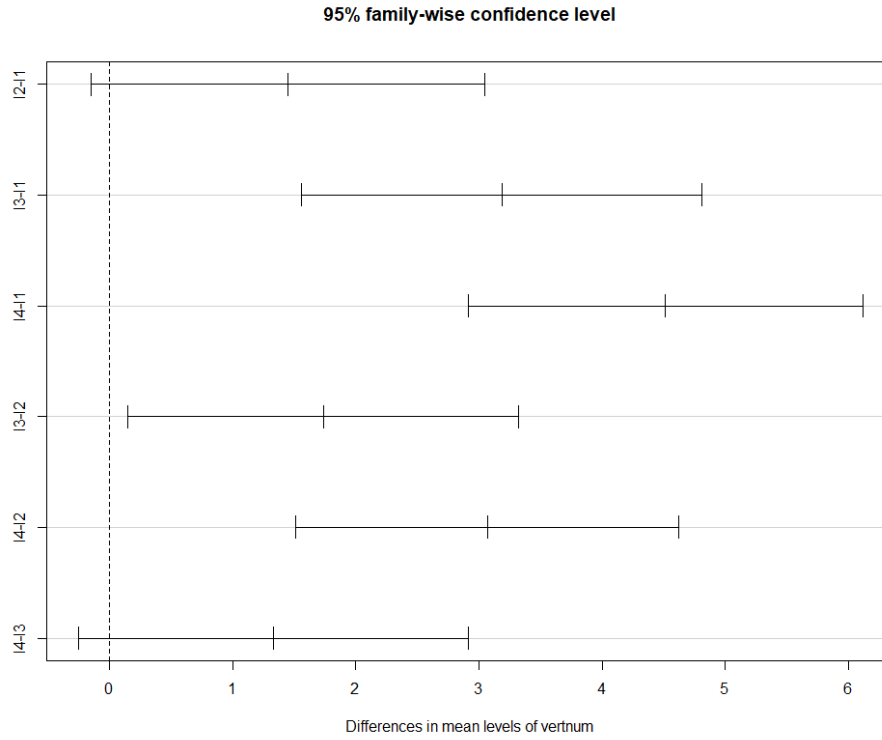
**Figure 58:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Articular Process Height measurement.



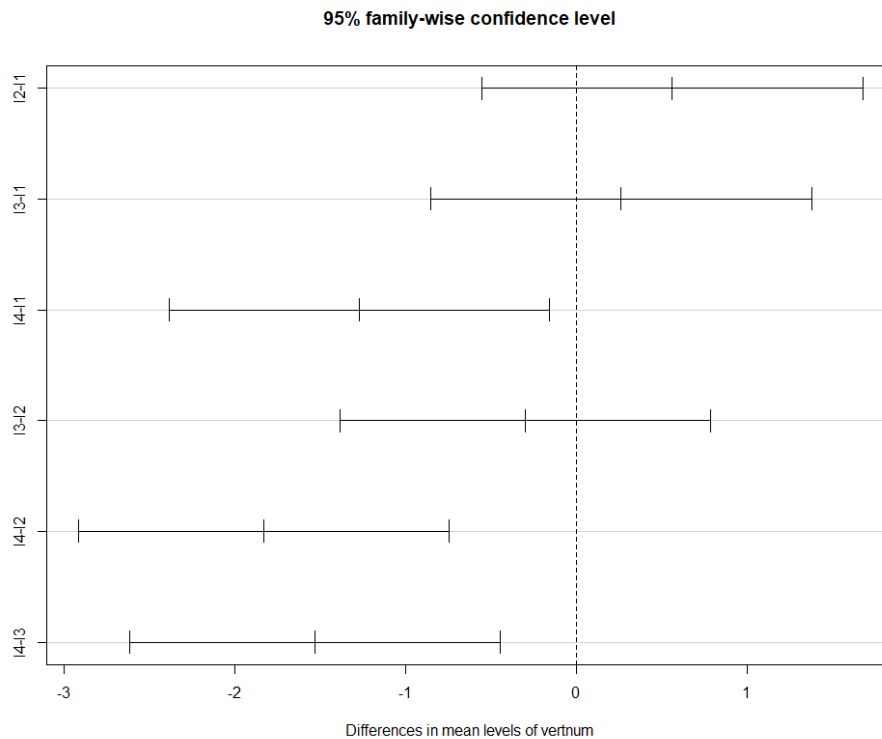
**Figure 59:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Spinous Process Length measurement.



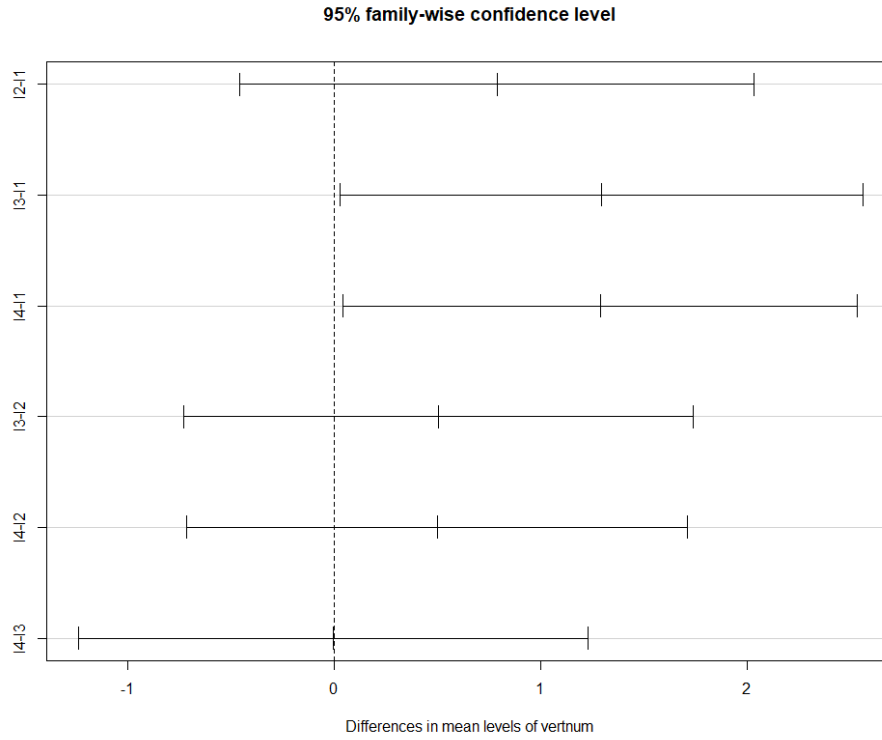
**Figure 60:** Results for the Tukey's HSD illustrating differences among the lower thoracic vertebrae (T7-T12) Maximum Inferior Sagittal Length measurement.



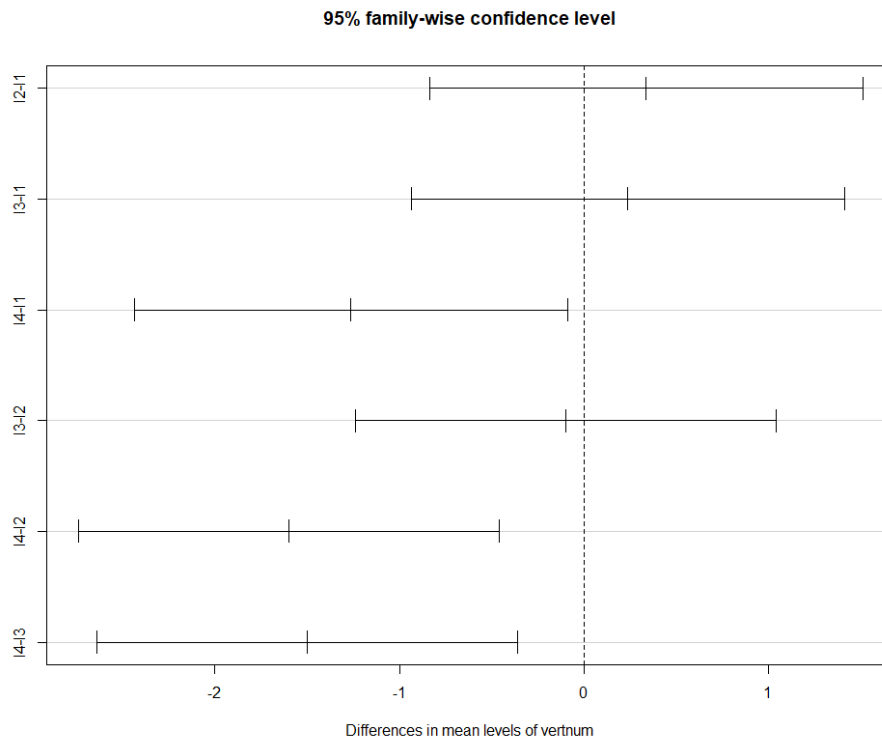
**Figure 61:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Superior Sagittal Length measurement.



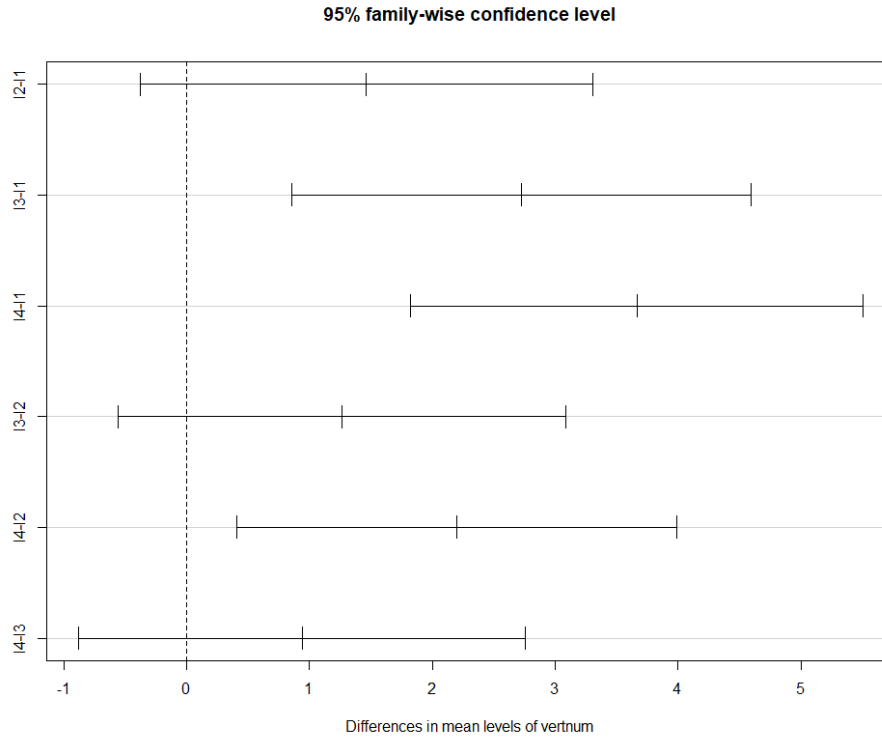
**Figure 62:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Centroid Height measurement.



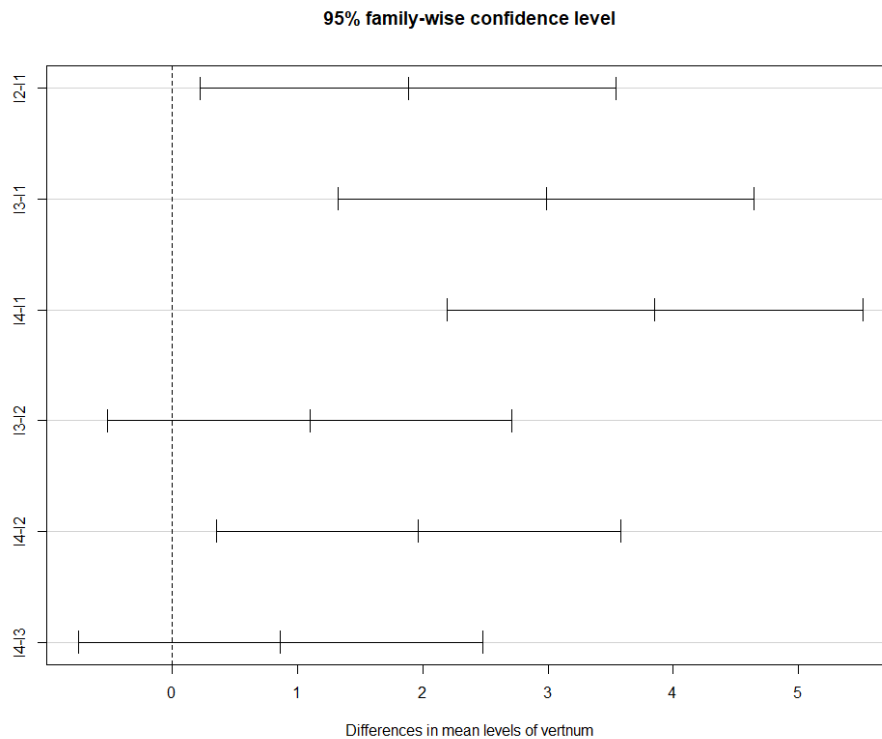
**Figure 63:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Anterior Height measurement.



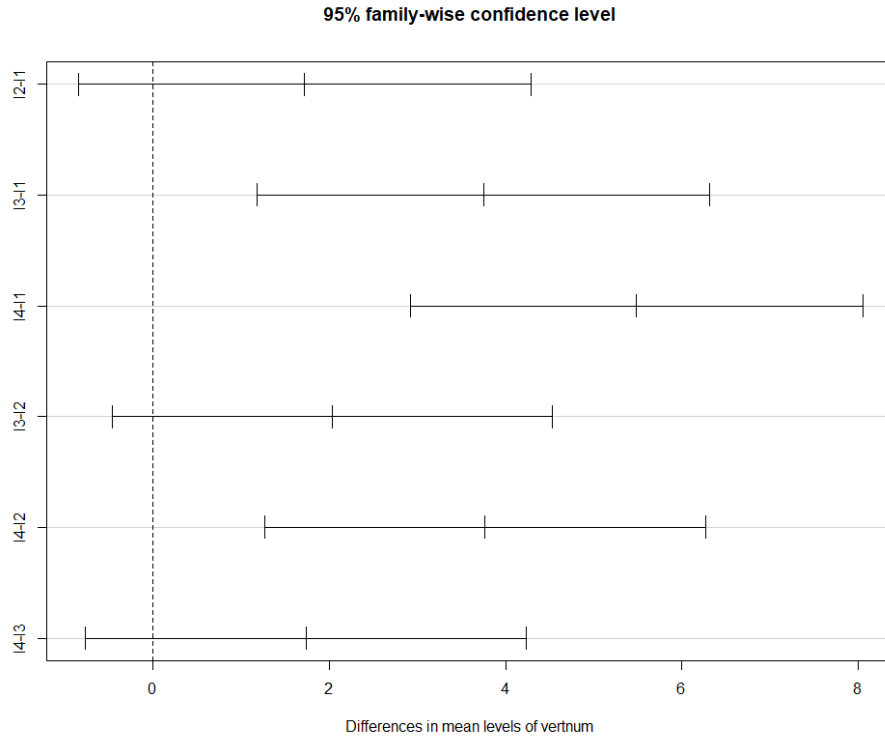
**Figure 64:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Posterior Height measurement.



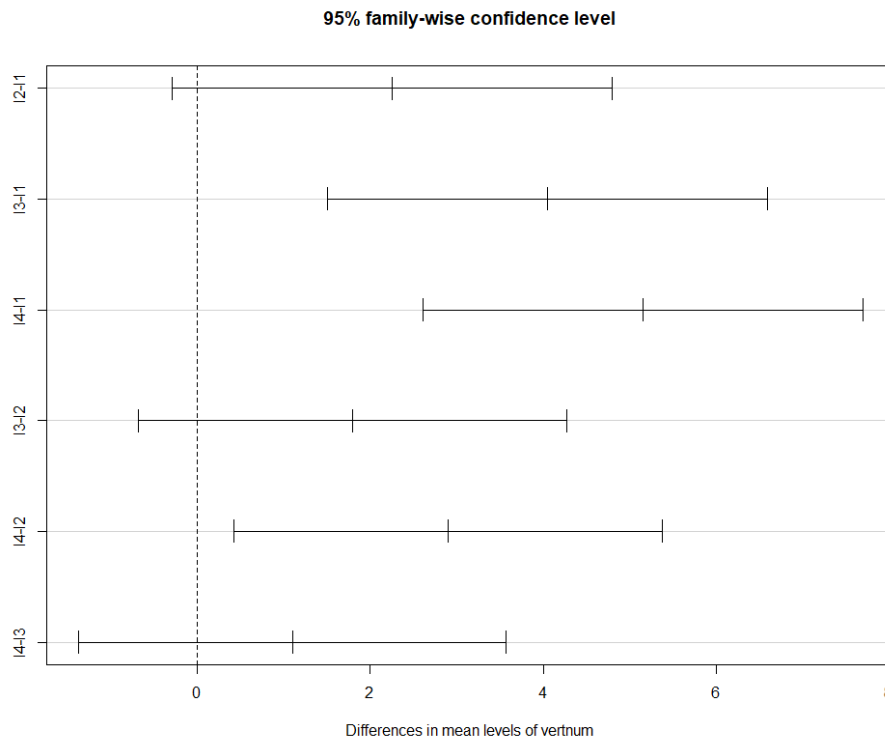
**Figure 65:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Superior Articular Length measurement.



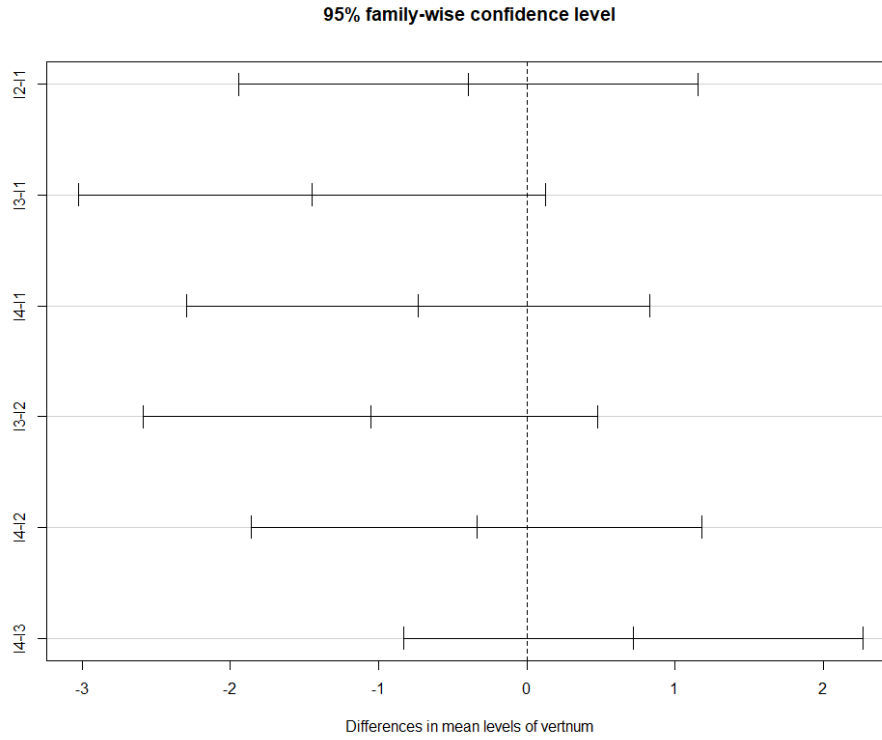
**Figure 66:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Superior Articular Length measurement.



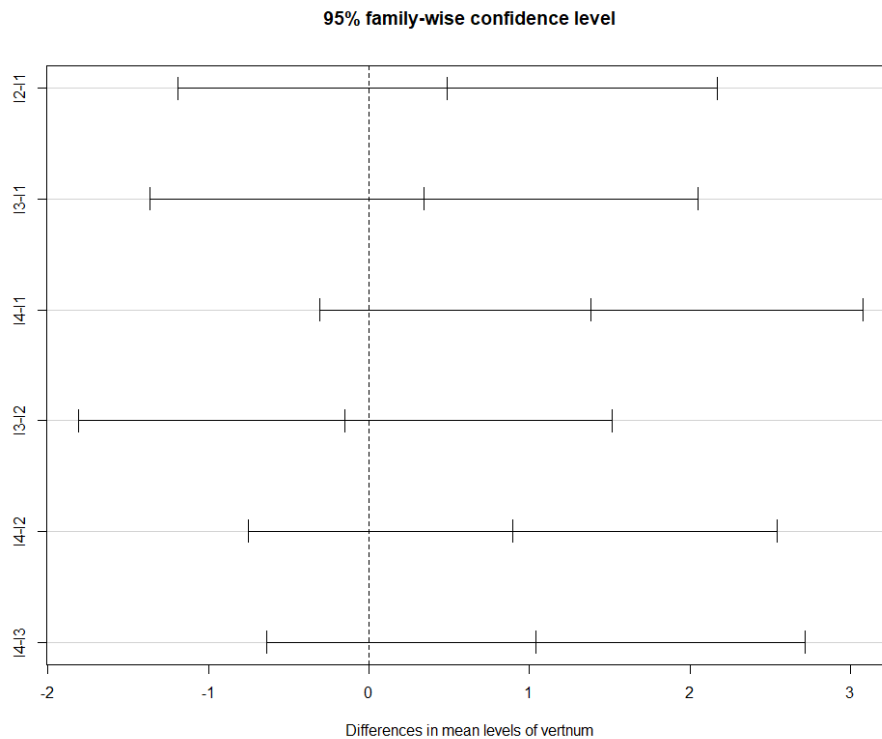
**Figure 67:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Superior Articular Width measurement.



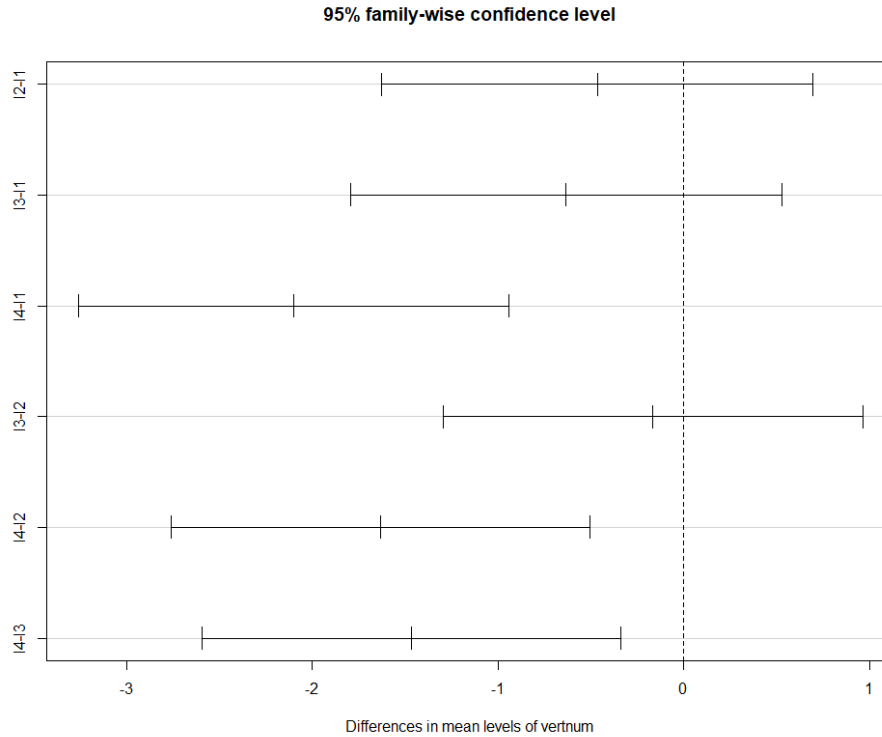
**Figure 68:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Inferior Articular Width measurement.



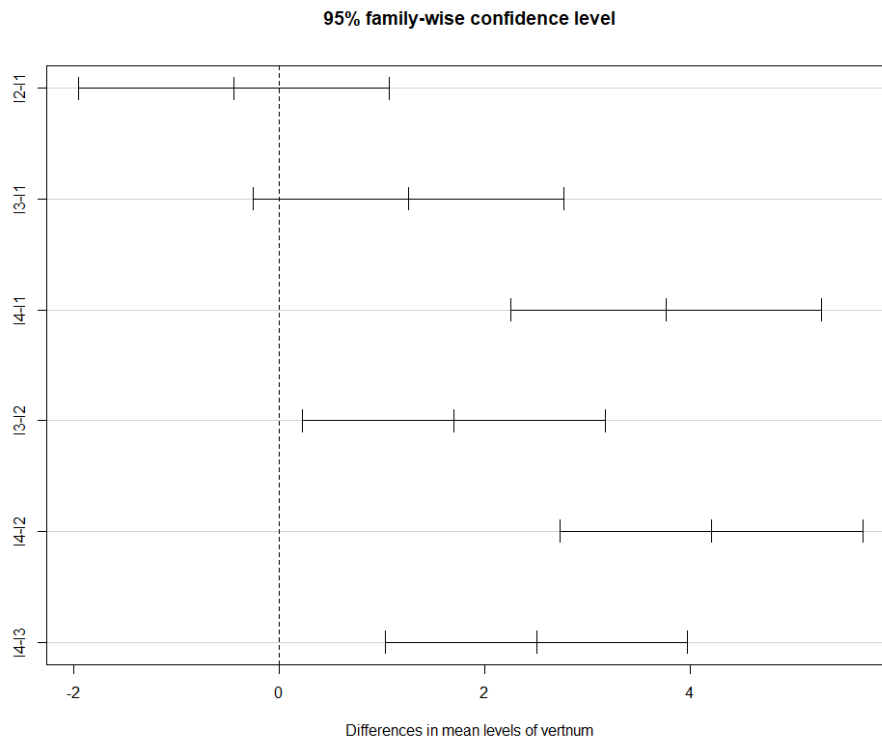
**Figure 69:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Foramen Sagittal Length measurement.



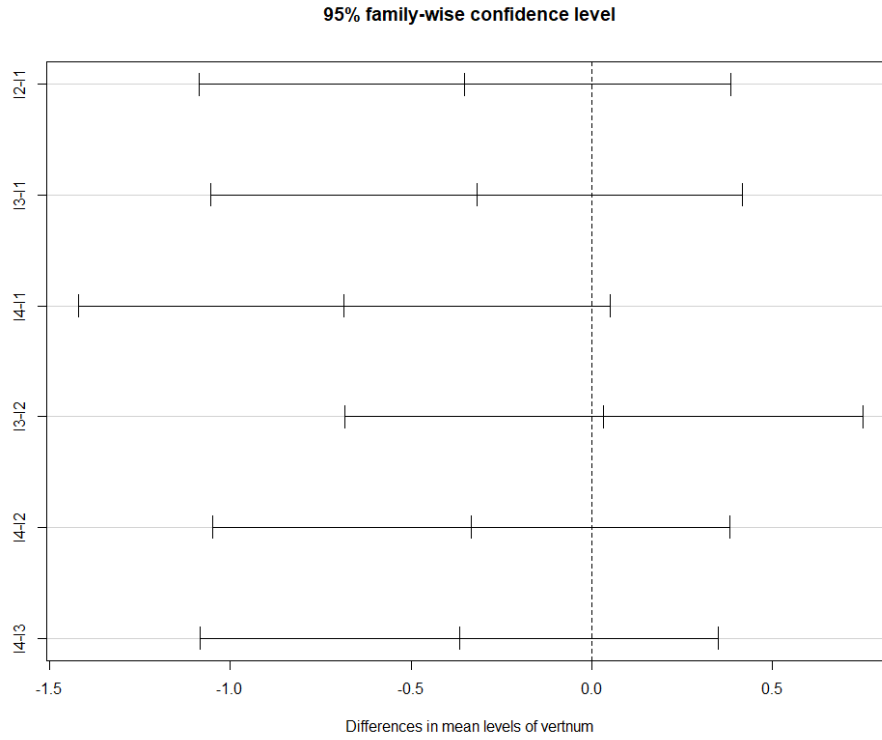
**Figure 70:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Foramen Coronal Width measurement.



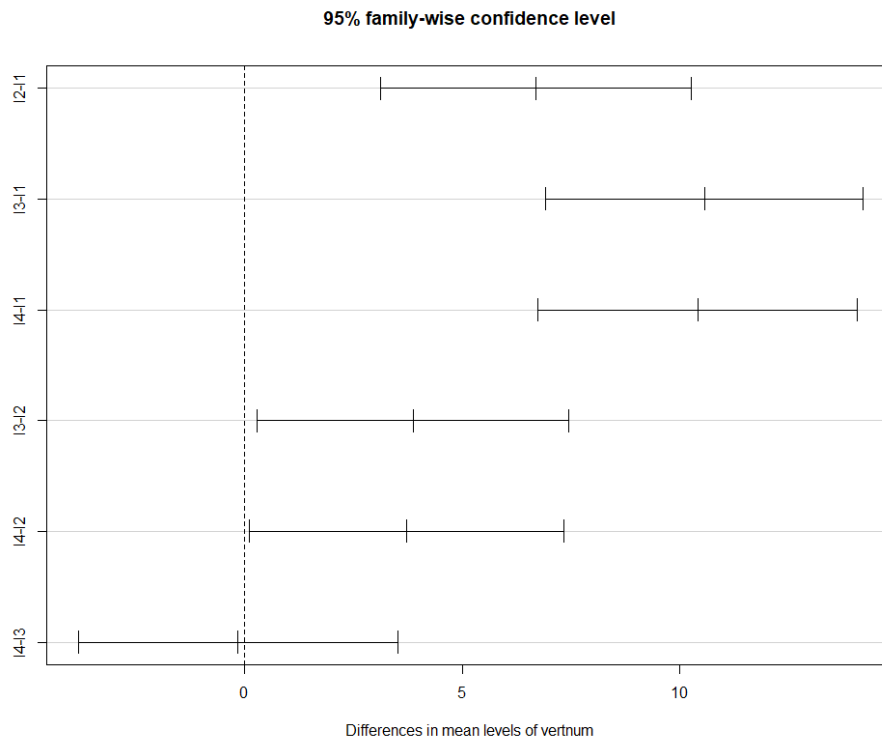
**Figure 71:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Pedicle Height measurement.



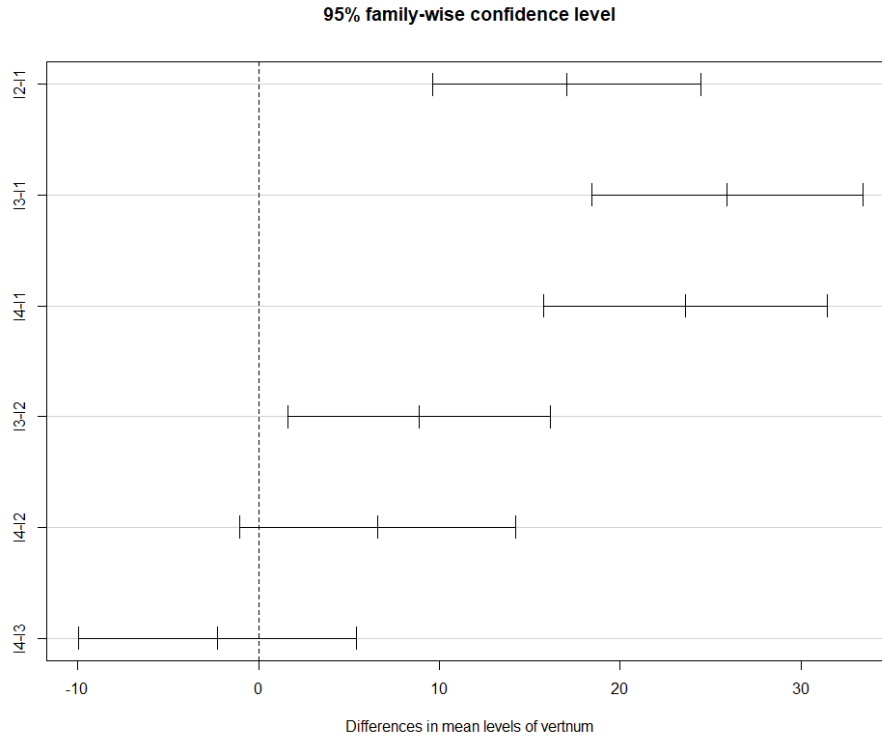
**Figure 72:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Pedicle Width measurement.



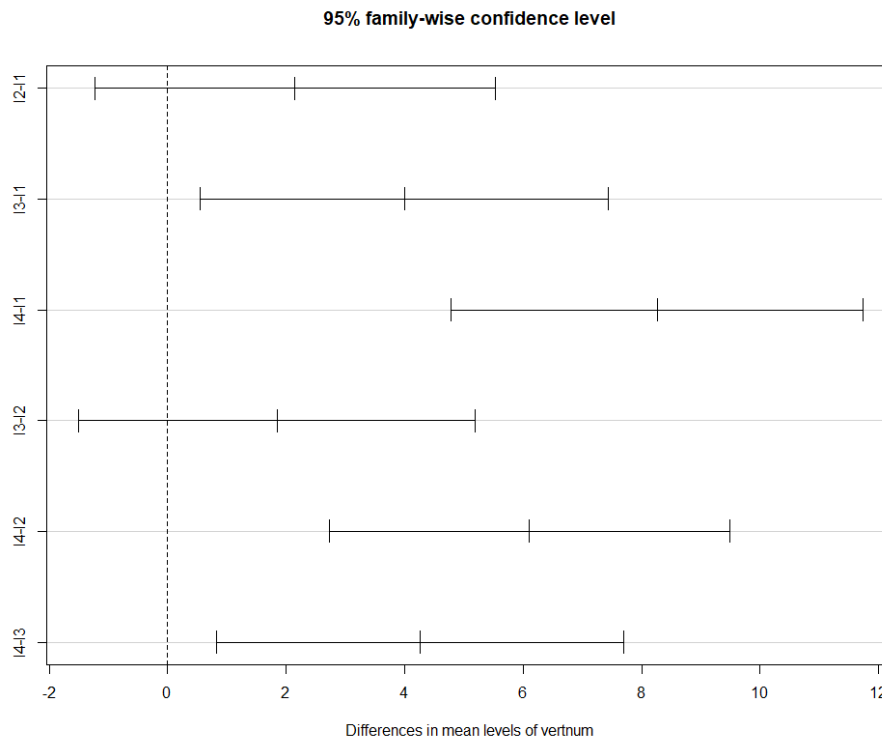
**Figure 73:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Pedicle Length measurement.



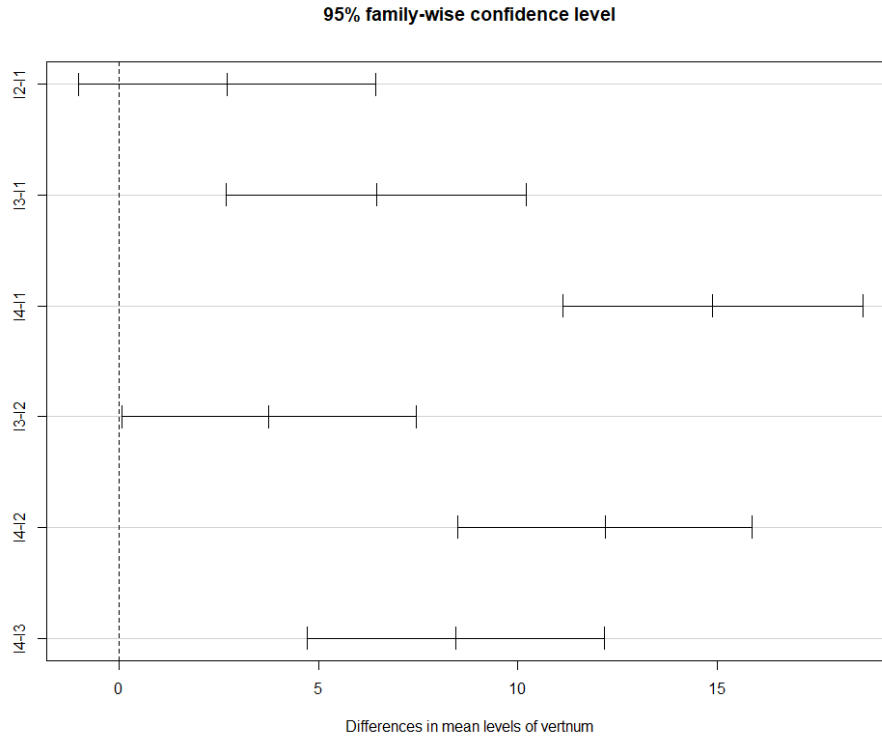
**Figure 74:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Transverse Process Length measurement.



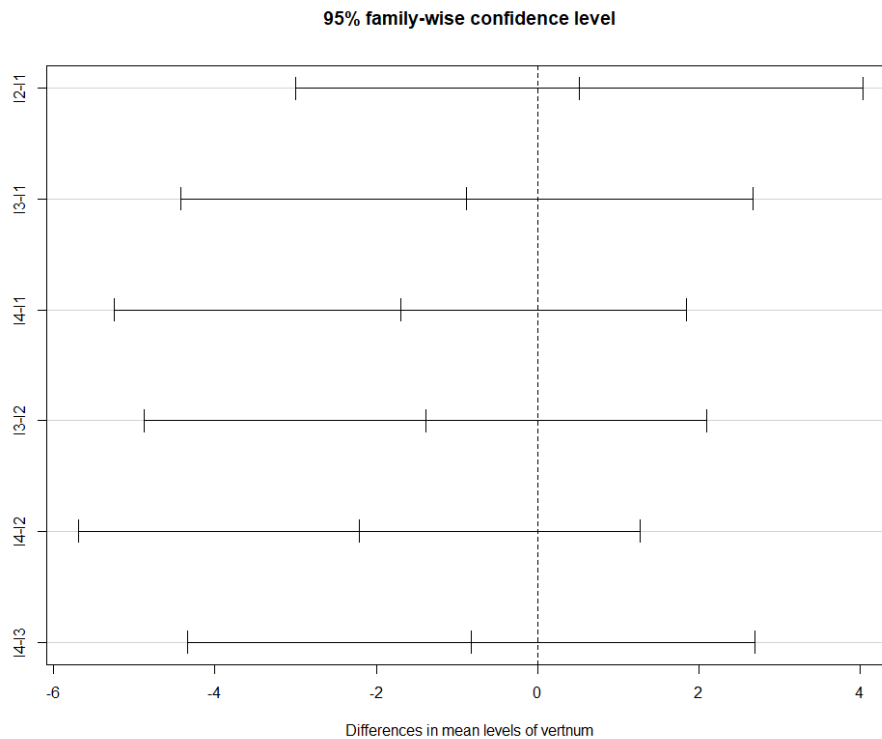
**Figure 75:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Transverse Process Width measurement.



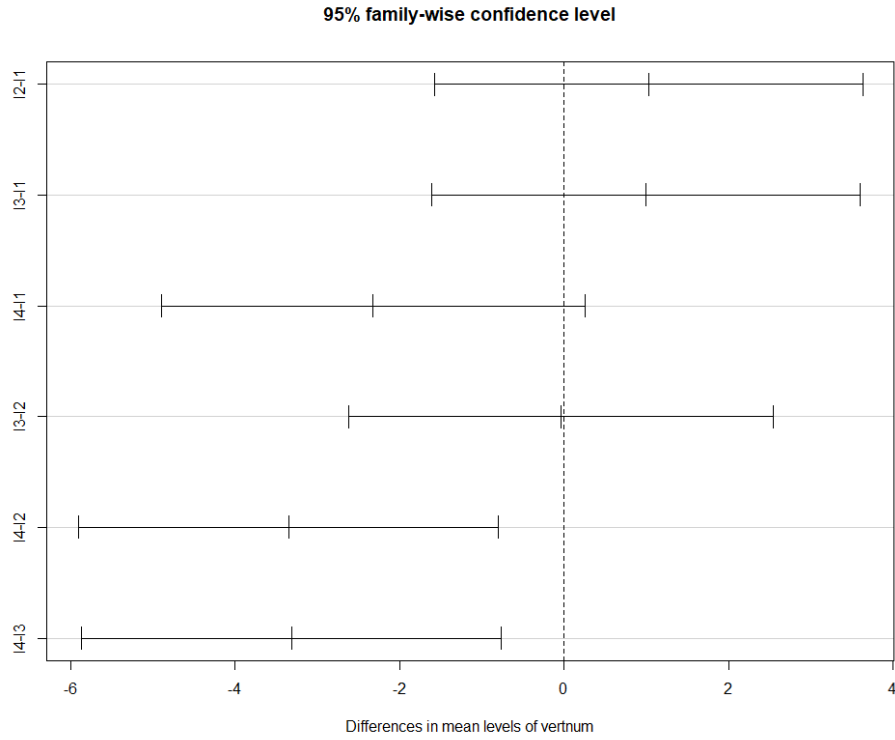
**Figure 76:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Superior Articular Process Width measurement.



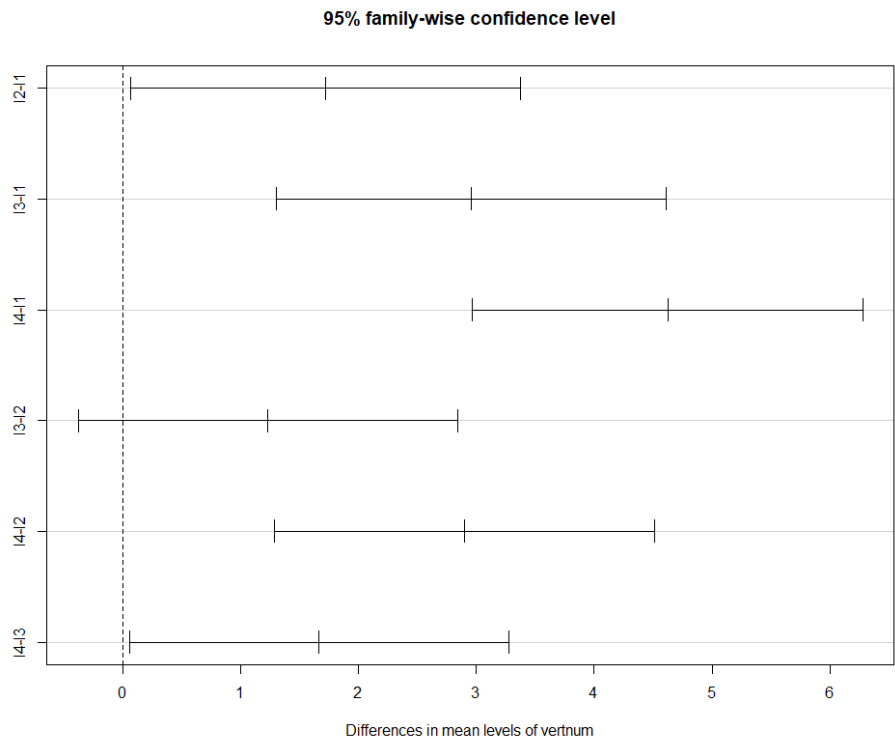
**Figure 77:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Inferior Articular Process Width measurement.



**Figure 78:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Articular Process Height measurement.



**Figure 79:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Spinous Process Length measurement.



**Figure 80:** Results for the Tukey's HSD illustrating differences among the lumbar vertebrae (L1-L4) Maximum Inferior Sagittal Length measurement.

## APPENDIX IV – DESCRIPTIVE ANALYSIS

**Tables A1 – A4:** Summary statistics of the cervical vertebrae (C3 – C6) showing the means and standard deviations for population affinity and sex.

**Table A1:** Summary statistics of the cervical vertebrae (C3-C6) showing the means and standard deviation for black South Africans. Refer to Appendix I for measurement names and definitions.

	C3			C4			C5			C6		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
XSLs	55	15.24	1.51	58	15.28	1.65	58	15.22	1.93	55	15.69	1.78
XAH	55	12.60	1.47	57	12.74	1.17	57	11.88	1.27	55	12.04	1.05
XPH	55	12.85	1.19	58	12.62	1.04	58	12.38	1.11	55	12.55	1.12
XALs	54	12.43	1.31	58	12.78	1.53	58	12.72	1.37	56	12.98	1.46
XALi	55	13.20	1.38	57	13.51	1.76	57	13.93	1.50	55	13.76	1.53
XAWs	54	17.19	1.68	59	18.24	1.45	58	19.26	2.00	56	19.89	1.87
XAWi	54	16.52	1.37	58	17.14	1.49	58	18.17	2.14	56	19.68	2.20
XPL	54	4.44	0.77	58	4.50	0.73	57	4.95	0.85	56	5.30	0.85
TPL	52	31.63	2.37	57	31.79	2.40	57	32.56	2.58	52	34.31	2.95
XTPW	49	49.98	3.89	53	51.00	3.87	50	53.72	4.21	39	58.38	5.14
XAPi	54	47.52	3.18	58	49.26	3.24	58	50.88	3.38	54	50.17	3.95
APH	54	19.91	1.93	59	20.53	1.99	58	20.57	2.07	55	21.95	1.85
XSPL	54	14.87	3.07	56	14.50	2.65	58	17.29	3.24	55	24.13	4.68
XSLi	55	16.22	1.82	58	16.38	2.05	58	16.57	1.82	56	16.61	1.80

**Table A2:** Summary statistics of the cervical vertebrae (C3-C6) showing the means and standard deviation for white South Africans. Refer to Appendix I for measurement names and definitions.

	C3			C4			C5			C6		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
XSLs	51	15.06	1.71	57	15.89	1.85	56	15.91	2.01	51	17.35	2.30
XAH	52	14.02	1.32	53	13.77	1.58	54	13.43	1.44	50	12.88	1.21
XPH	50	14.18	1.29	56	13.88	1.28	56	13.84	1.36	50	13.80	1.18
XALs	52	12.96	1.41	57	13.46	1.36	54	13.54	1.67	50	14.40	1.70
XALi	50	13.96	1.51	54	14.28	1.65	55	15.11	2.42	50	15.38	1.46
XAWs	52	18.58	1.58	58	19.78	1.81	56	20.77	1.94	51	22.82	2.00
XAWi	52	17.71	1.93	53	18.75	1.75	53	19.32	2.12	50	21.68	2.00
XPL	51	4.65	0.80	57	5.11	0.96	54	5.44	1.18	52	6.04	0.95
TPL	48	34.10	1.95	54	34.52	1.94	52	34.92	2.39	37	35.03	2.32
XTPW	34	53.79	2.71	40	54.67	3.33	34	55.91	3.64	29	58.21	3.12
XAPi	52	52.04	3.02	57	53.19	2.82	53	54.08	3.26	50	53.52	3.50
APH	50	21.96	2.49	54	23.00	2.50	53	23.11	2.75	51	24.22	2.71
XSPL	47	17.94	3.20	50	18.82	3.61	46	20.15	3.37	48	24.71	3.82
XSLi	49	16.27	1.75	53	16.85	1.83	53	17.51	2.09	49	17.98	1.98

**Table A3:** Summary statistics of the cervical vertebrae (C3-C6) showing the means and standard deviation for males. Refer to Appendix I for measurement names and definitions.

	C3			C4			C5			C6		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	51	16.22	1.38	57	16.60	1.69	57	16.33	2.00	54	17.22	2.23
<b>XAH</b>	52	13.77	1.58	54	13.80	1.46	57	12.91	1.56	55	12.73	1.28
<b>XPH</b>	51	14.10	1.40	56	13.77	1.24	57	13.56	1.50	53	13.64	1.33
<b>XALs</b>	52	13.52	1.31	58	13.74	1.58	56	13.79	1.69	54	14.22	1.90
<b>XALi</b>	51	14.41	1.42	55	14.78	1.62	58	15.53	2.19	54	15.02	1.72
<b>XAWs</b>	51	18.57	1.81	58	19.34	1.98	57	20.60	2.15	55	21.76	2.65
<b>XAWi</b>	50	17.88	1.78	55	18.53	1.84	56	19.25	2.30	55	21.40	2.48
<b>XPL</b>	51	4.67	0.82	56	4.88	0.95	54	5.22	1.21	55	5.73	1.06
<b>TPL</b>	49	33.59	2.32	55	34.16	2.29	54	34.81	2.71	44	36.09	2.59
<b>XTPW</b>	41	53.49	3.28	45	54.51	3.85	45	56.67	3.86	35	60.54	4.03
<b>XAPi</b>	52	51.04	3.64	57	52.37	3.39	56	54.09	3.40	52	54.08	3.29
<b>APH</b>	52	21.38	2.46	55	22.20	2.50	54	22.65	2.62	55	23.69	2.62
<b>XSPL</b>	48	17.52	3.56	51	17.67	3.94	51	19.80	3.47	50	25.82	4.53
<b>XSLi</b>	49	17.20	1.65	55	17.49	1.93	56	17.93	1.99	53	17.77	1.85

**Table A4:** Summary statistics of the cervical vertebrae (C3-C6) showing the means and standard deviation for females. Refer to Appendix I for measurement names and definitions.

	C3			C4			C5			C6		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	55	14.16	1.10	58	14.59	1.20	57	14.79	1.67	52	15.73	1.91
<b>XAH</b>	55	12.84	1.42	56	12.70	1.28	54	12.33	1.50	50	12.12	1.02
<b>XPH</b>	54	12.91	1.14	58	12.72	1.20	57	12.63	1.20	52	12.63	1.07
<b>XALs</b>	54	11.89	0.90	57	12.47	1.05	56	12.45	1.09	52	13.06	1.29
<b>XALi</b>	54	12.76	1.04	56	13.00	1.39	54	13.41	1.22	51	14.02	1.52
<b>XAWs</b>	55	17.22	1.46	59	18.66	1.56	57	19.40	1.90	52	20.79	2.05
<b>XAWi</b>	56	16.41	1.45	56	17.30	1.56	55	18.18	1.96	51	19.78	1.84
<b>XPL</b>	54	4.43	0.74	59	4.73	0.85	57	5.16	0.88	53	5.58	0.86
<b>TPL</b>	51	32.08	2.46	56	32.09	2.44	55	32.58	2.32	45	33.16	1.95
<b>XTPW</b>	42	49.64	3.57	48	50.77	3.40	39	52.23	2.99	33	55.94	3.38
<b>XAPi</b>	54	48.48	3.63	58	50.07	3.49	55	50.69	3.14	52	49.48	3.48
<b>APH</b>	52	20.40	2.33	58	21.24	2.54	57	20.96	2.58	51	22.33	2.32
<b>XSPL</b>	53	15.19	3.01	55	15.49	3.37	53	17.36	3.28	53	23.06	3.60
<b>XSLi</b>	55	15.38	1.43	56	15.73	1.55	55	16.09	1.54	52	16.71	2.02

**Table A5:** The mean of all cervical vertebrae (C3-C6) combined for population and sex. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Black	White	Male	Female
<b>XSLs</b>	15.35	16.05	16.59	14.8
<b>XAH</b>	12.31	13.53	13.29	12.51
<b>XPH</b>	12.60	13.92	13.76	12.72
<b>XALs</b>	12.73	13.58	13.82	12.46
<b>XALi</b>	13.60	14.68	14.95	13.28
<b>XAWs</b>	18.66	20.46	20.09	18.99
<b>XAWi</b>	17.88	19.34	19.3	17.88
<b>XPL</b>	4.8	5.31	5.13	4.97
<b>TPL</b>	32.56	34.62	34.62	32.45
<b>XTPW</b>	52.96	55.51	56.11	51.89
<b>XAPi</b>	49.48	53.21	52.90	49.69
<b>APH</b>	20.73	23.08	22.5	21.22
<b>XSPL</b>	17.69	20.4	20.22	17.75
<b>XSLi</b>	16.44	17.15	17.61	15.97

**Tables A6-A9:** Summary statistics of the upper thoracic vertebrae (T2-T6) showing the means and standard deviations for population and sex. Refer to Appendix I for measurement names and definitions.

**Table A6:** Summary statistics of the upper thoracic vertebrae (T2-T6) showing the means and standard deviation for black South Africans. Refer to Appendix I for measurement names and definitions.

	T2			T3			T4			T5			T6		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	59	16.97	1.63	60	18.53	1.83	57	19.65	2.16	58	20.95	1.96	59	22.32	2.05
<b>XAH</b>	58	17.17	1.35	58	17.26	1.19	53	17.21	1.26	55	17.15	1.16	56	17.66	1.27
<b>XPH</b>	58	17.24	1.26	60	17.75	1.22	57	17.96	1.22	57	18.37	1.23	59	18.81	1.27
<b>XALs</b>	59	13.69	1.45	60	15.20	1.88	57	16.04	1.84	58	17.47	1.84	59	18.58	1.98
<b>XALi</b>	58	15.31	1.64	58	16.29	1.71	54	17.20	1.82	56	18.62	1.93	58	19.64	1.88
<b>XAWs</b>	58	22.45	2.33	60	22.15	2.39	57	21.61	1.73	58	22.38	1.86	59	23.07	1.75
<b>XAWi</b>	59	24.15	2.50	60	23.27	2.26	56	23.23	2.40	58	23.76	2.07	58	24.72	1.95
<b>XPL</b>	59	4.76	1.15	60	4.18	0.98	57	4.39	0.88	58	4.41	0.88	59	4.64	1.01
<b>TPL</b>	59	31.22	3.59	60	29.17	2.84	57	28.33	2.92	58	28.34	2.89	59	28.83	2.65
<b>XTPW</b>	59	63.24	7.08	60	58.17	5.60	55	55.85	4.65	54	56.30	5.04	56	56.80	4.89
<b>XAPi</b>	58	33.66	4.03	60	31.48	3.71	56	30.27	2.96	57	29.46	3.42	59	29.61	3.29
<b>APH</b>	59	30.32	2.49	59	30.29	2.35	56	30.16	2.43	56	30.52	2.07	58	31.67	2.24
<b>XSPL</b>	56	36.32	4.23	58	37.12	4.40	56	37.59	5.06	53	41.02	5.20	55	43.67	5.07
<b>XSLi</b>	58	18.14	1.74	59	19.25	1.86	55	20.35	2.07	57	21.58	2.06	58	22.67	2.16

**Table A7:** Summary statistics of the upper thoracic vertebrae (T2-T6) showing the means and standard deviation for whites South Africans. Refer to Appendix I for measurement names and definitions.

	T2			T3			T4			T5			T6		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	54	18.00	1.97	58	20.45	2.49	54	23.13	2.13	56	25.00	2.44	52	26.40	2.25
<b>XAH</b>	52	18.17	1.22	57	18.37	1.41	53	15.62	1.13	57	18.88	1.55	51	18.94	1.46
<b>XPH</b>	57	18.72	1.32	59	18.92	1.35	57	19.47	1.34	57	20.25	1.46	52	20.88	1.38
<b>XALs</b>	53	15.45	1.87	59	17.02	2.43	55	19.33	2.25	57	21.02	2.45	52	22.35	2.33
<b>XALi</b>	53	17.06	2.39	56	19.07	2.16	54	20.85	2.07	57	22.51	2.85	50	23.82	2.78
<b>XAWs</b>	56	25.02	1.77	59	23.75	1.84	57	23.72	1.68	57	24.42	2.09	52	25.77	2.29
<b>XAWi</b>	56	26.16	1.85	58	25.10	2.12	57	25.09	2.25	57	26.32	2.38	51	27.73	2.60
<b>XPL</b>	58	6.60	1.43	59	5.10	1.17	57	4.56	0.91	56	4.61	1.02	51	4.84	1.10
<b>TPL</b>	58	35.19	3.11	59	32.31	2.90	56	31.32	2.80	56	31.66	2.35	51	32.10	2.14
<b>XTPW</b>	55	70.15	5.74	55	64.47	5.22	55	62.60	4.56	51	63.22	3.96	45	64.64	4.14
<b>XAPi</b>	58	37.19	3.56	59	35.02	2.73	56	34.27	2.40	57	34.19	2.51	50	34.36	2.80
<b>APH</b>	58	31.64	2.33	59	32.19	2.14	57	32.25	2.32	56	33.02	2.48	51	34.12	2.23
<b>XSPL</b>	57	39.32	4.74	56	40.14	4.96	53	43.11	5.36	51	47.49	6.62	50	49.58	5.79
<b>XSLi</b>	52	19.71	2.37	55	22.24	2.25	54	24.31	2.12	54	26.00	2.56	51	27.35	2.47

**Table A8:** Summary statistics of the upper thoracic vertebrae (T2-T6) showing the means and standard deviation for males. Refer to Appendix I for measurement names and definitions.

	T2			T3			T4			T5			T6		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	58	18.31	1.91	60	20.40	2.53	54	22.54	2.64	57	24.26	3.10	55	25.20	3.01
<b>XAH</b>	56	18.16	1.35	58	18.34	1.34	53	18.42	1.23	57	18.56	1.72	55	18.84	1.46
<b>XPH</b>	59	18.64	1.35	60	18.88	1.39	55	19.33	1.23	57	19.88	1.62	55	20.40	1.51
<b>XALs</b>	58	15.47	1.83	60	17.30	2.39	55	18.91	2.63	57	20.58	2.93	55	21.49	3.01
<b>XALi</b>	57	17.18	2.33	56	18.71	2.53	54	20.17	2.69	57	21.91	3.30	54	22.70	3.45
<b>XAWs</b>	59	24.69	2.21	60	24.15	2.28	55	23.62	2.01	57	24.56	2.15	55	25.35	2.35
<b>XAWi</b>	60	26.32	2.27	60	25.50	2.34	54	25.50	2.35	57	26.25	2.50	53	27.26	2.90
<b>XPL</b>	60	5.60	1.89	60	4.78	1.32	55	4.76	0.84	57	4.88	0.96	55	5.11	1.07
<b>TPL</b>	60	35.05	3.51	60	32.50	2.95	54	31.46	3.00	57	31.37	2.64	55	31.78	2.86
<b>XTPW</b>	57	70.02	6.73	58	64.62	5.52	55	62.02	5.04	56	62.55	4.56	53	62.85	5.19
<b>XAPi</b>	60	37.48	3.75	60	34.83	2.98	55	33.62	2.72	57	33.49	3.25	55	33.36	3.52
<b>APH</b>	60	31.93	2.36	60	32.23	2.07	55	32.20	2.21	56	32.77	2.49	55	33.45	2.49
<b>XSPL</b>	59	39.97	4.24	57	40.81	4.93	53	42.57	5.83	51	46.90	6.64	53	48.49	5.96
<b>XSLi</b>	57	19.86	2.28	56	21.66	2.65	54	23.39	2.79	56	25.04	3.24	55	25.78	3.18

**Table A9:** Summary statistics of the upper thoracic vertebrae (T2-T6) showing the means and standard deviation for females. Refer to Appendix I for measurement names and definitions.

	T7			T8			T9			T10			T11		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	55	16.56	1.33	58	18.52	1.76	57	20.21	2.38	57	21.61	2.22	56	23.29	2.60
<b>XAH</b>	54	17.11	1.21	57	17.26	1.28	53	17.42	1.36	55	17.47	1.30	52	17.67	1.31
<b>XPH</b>	56	17.27	1.29	59	17.26	1.18	59	18.15	1.48	57	18.74	1.47	56	19.18	1.62
<b>XALs</b>	54	13.52	1.33	59	14.88	1.54	57	16.44	2.00	58	17.90	1.89	56	19.21	2.20
<b>XALi</b>	54	15.06	1.41	58	16.64	1.70	54	17.89	2.12	56	19.23	2.22	54	20.44	2.30
<b>XAWs</b>	55	22.65	2.23	59	21.71	1.49	59	21.78	1.53	58	22.24	1.60	56	23.34	2.07
<b>XAWi</b>	55	23.84	1.84	58	22.79	1.44	59	22.95	1.95	58	23.83	2.02	56	25.05	2.05
<b>XPL</b>	57	5.75	1.20	59	4.49	0.99	59	4.20	0.87	57	4.14	0.79	55	4.36	0.91
<b>TPL</b>	57	31.23	3.28	59	28.92	2.49	59	28.31	2.63	57	28.58	2.93	55	28.91	2.43
<b>XTPW</b>	57	63.12	6.20	57	58.23	4.95	55	56.44	4.94	49	56.35	5.08	48	57.48	5.60
<b>XAPi</b>	56	33.96	4.23	59	31.61	3.67	57	30.96	3.41	57	30.16	3.64	54	30.19	3.57
<b>APH</b>	57	29.96	2.23	58	30.21	2.36	58	30.28	2.58	56	30.77	2.32	54	32.17	2.45
<b>XSPL</b>	54	35.50	4.10	57	36.40	3.79	56	38.11	5.09	53	41.58	5.79	52	44.44	5.72
<b>XSLi</b>	53	17.83	1.54	58	19.76	2.05	55	21.25	2.59	55	22.40	2.56	54	23.93	3.16

**Table A10:** The mean of all upper thoracic vertebrae (T2-T6) combined for population and sex. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Black	White	Male	Female
<b>XSLs</b>	19.63	22.54	22.08	20.00
<b>XAH</b>	17.31	18.60	18.46	17.40
<b>XPH</b>	18.01	19.62	19.41	18.20
<b>XALs</b>	16.21	19.01	18.70	16.43
<b>XALi</b>	17.43	20.64	20.11	17.87
<b>XAWs</b>	22.36	24.51	24.47	22.36
<b>XAWi</b>	23.84	25.96	26.15	23.61
<b>XPL</b>	4.48	5.26	5.03	4.69
<b>TPL</b>	29.19	32.66	32.48	29.30
<b>XTPW</b>	57.86	64.93	64.46	57.88
<b>XAPi</b>	30.94	35.18	34.61	31.42
<b>APH</b>	30.68	32.65	32.50	30.80
<b>XSPL</b>	38.83	43.70	43.60	38.81
<b>XSLi</b>	20.41	23.91	23.12	21.05

**Tables A11-A14:** Summary statistics of the lower thoracic vertebrae (T7-T11) showing the means and standard deviations for population and sex. Refer to Appendix I for measurement names and definitions.

**Table A11:** Summary statistics of the lower thoracic vertebrae (T7-T11) showing the means and standard deviation for black South Africans. Refer to Appendix I for measurement names and definitions.

	T7			T8			T9			T10			T11		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	57	23.30	2.28	57	24.61	2.51	59	25.39	2.37	59	26.05	2.31	56	26.12	2.78
<b>XAH</b>	57	18.14	1.38	55	18.85	1.56	59	19.66	1.56	59	20.92	1.69	56	21.32	1.45
<b>XPH</b>	57	19.58	1.36	57	20.12	1.30	58	20.71	1.56	60	21.68	1.51	56	23.02	1.81
<b>XALs</b>	57	19.68	2.02	57	20.58	1.82	59	21.73	2.18	60	22.15	2.11	56	22.11	2.20
<b>XALi</b>	57	20.77	2.18	56	21.62	2.23	59	22.22	2.08	59	22.34	1.93	57	22.93	2.49
<b>XAWs</b>	57	24.18	2.11	57	25.72	2.05	59	26.98	2.28	59	22.68	1.93	56	31.39	2.61
<b>XAWi</b>	56	25.73	2.07	56	27.00	2.30	58	28.98	2.56	60	31.87	2.89	57	34.72	3.63
<b>XPL</b>	57	4.93	1.22	57	5.05	1.19	59	5.19	0.96	60	5.83	1.12	57	6.11	1.62
<b>TPL</b>	57	29.00	2.67	57	28.65	2.75	58	28.31	2.84	59	27.10	2.99	56	24.91	3.23
<b>XTPW</b>	55	56.93	4.57	56	56.23	5.11	56	55.84	5.31	60	53.78	5.38	56	49.93	6.28
<b>XAPi</b>	57	30.49	3.27	57	31.30	3.44	59	33.75	3.57	60	34.52	4.00	56	34.12	4.53
<b>APH</b>	55	32.76	2.21	56	33.88	2.04	58	34.59	2.38	58	36.02	2.46	57	38.23	2.61
<b>XSPL</b>	53	43.13	4.65	54	41.39	3.62	57	38.79	3.82	60	35.78	4.09	56	32.93	3.98
<b>XSLi</b>	57	23.98	2.31	56	25.00	2.33	58	25.86	2.31	59	25.98	2.26	57	26.42	2.54

**Table A12:** Summary statistics of the lower thoracic vertebrae (T7-T11) showing the means and standard deviation for whites South Africans. Refer to Appendix I for measurement names and definitions.

	T7			T8			T9			T10			T11		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	52	27.63	2.94	54	29.11	3.13	56	29.88	3.20	55	30.58	3.07	55	31.27	2.98
<b>XAH</b>	52	19.23	1.72	53	19.40	1.59	56	20.32	1.85	56	21.73	1.84	54	22.44	1.93
<b>XPH</b>	52	21.17	1.70	55	21.58	1.57	56	22.29	1.49	55	23.33	1.87	56	25.12	1.81
<b>XALs</b>	51	23.82	2.81	54	24.80	2.83	57	26.14	3.24	55	26.95	2.95	55	27.05	2.98
<b>XALi</b>	51	24.88	2.99	53	26.26	3.28	54	27.15	3.24	55	27.78	3.25	56	28.27	3.31
<b>XAWs</b>	52	27.44	2.59	54	29.06	2.86	56	30.34	2.74	55	32.25	2.72	56	35.00	3.16
<b>XAWi</b>	52	29.23	3.10	54	30.74	2.97	58	32.86	3.12	54	35.43	2.98	57	38.11	3.55
<b>XPL</b>	52	5.06	1.35	55	5.22	1.30	58	5.74	1.16	57	6.49	1.57	56	7.34	1.91
<b>TPL</b>	51	31.90	2.59	53	31.09	2.61	55	30.80	2.56	53	29.04	3.08	52	27.08	3.25
<b>XTPW</b>	50	63.38	4.90	50	62.42	5.02	52	61.17	4.47	49	58.49	5.10	50	54.14	6.14
<b>XAPi</b>	52	34.98	2.69	55	36.29	2.55	56	36.75	2.42	57	37.40	3.15	57	35.72	4.91
<b>APH</b>	52	34.67	2.37	54	35.54	2.21	57	36.19	2.49	54	38.04	2.96	55	40.98	3.41
<b>XSPL</b>	43	49.47	5.82	49	46.78	5.47	50	42.94	4.78	54	38.85	5.34	55	35.22	5.07
<b>XSLi</b>	51	28.55	2.90	53	29.85	3.50	54	30.65	3.19	57	31.37	3.40	56	31.96	3.44

**Table A13:** Summary statistics of the lower thoracic vertebrae (T7-T11) showing the means and standard deviation for males. Refer to Appendix I for measurement names and definitions.

	T7			T8			T9			T10			T11		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	55	26.53	3.30	57	27.91	3.55	58	29.05	3.64	56	29.61	3.65	58	29.98	3.62
<b>XAH</b>	55	19.24	1.68	56	19.62	1.43	57	20.42	1.75	57	21.89	1.69	57	22.11	1.83
<b>XPH</b>	55	21.05	1.67	57	21.42	1.51	57	22.09	1.69	57	22.88	1.96	58	24.52	2.05
<b>XALs</b>	55	22.98	3.31	57	23.75	3.42	58	25.60	3.69	57	25.77	3.66	58	26.00	3.63
<b>XALi</b>	54	24.00	3.61	56	25.25	3.94	56	26.18	3.95	58	26.47	3.98	57	27.11	4.09
<b>XAWs</b>	55	26.98	2.91	57	28.60	2.92	58	30.00	3.12	55	31.38	3.43	57	34.65	3.45
<b>XAWi</b>	54	28.78	3.23	56	30.23	3.40	58	32.31	3.67	58	34.83	3.70	58	38.33	3.86
<b>XPL</b>	55	5.62	1.16	57	5.61	1.26	58	5.72	1.21	58	6.21	1.55	57	6.28	2.09
<b>TPL</b>	55	31.93	2.60	56	31.52	2.55	57	31.21	2.40	55	29.93	2.66	55	27.67	2.80
<b>XTPW</b>	54	62.83	5.16	56	62.32	5.11	53	61.57	4.51	54	59.07	4.76	54	55.06	5.44
<b>XAPi</b>	55	34.05	3.56	57	35.44	3.39	57	39.91	2.86	58	38.12	2.78	58	36.74	4.28
<b>APH</b>	55	34.55	2.22	57	35.58	2.37	58	36.07	2.59	56	37.62	3.10	57	40.16	3.56
<b>XSPL</b>	48	48.69	6.20	56	45.55	5.25	53	42.68	4.98	56	39.52	4.89	56	35.84	4.79
<b>XSLi</b>	55	27.31	3.51	56	28.59	3.94	56	29.50	3.56	58	30.21	3.93	57	30.63	4.04

**Table A14:** Summary statistics of the lower thoracic vertebrae (T7-T11) showing the sample size, means and standard deviation for females. Refer to Appendix I for measurement names and definitions.

	T7			T8			T9			T10			T11		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	54	24.19	3.09	54	25.63	3.04	57	26.07	2.85	58	26.91	2.84	53	27.25	3.62
<b>XAH</b>	54	18.07	1.39	52	18.58	1.58	58	19.55	1.61	58	20.74	1.75	53	21.62	1.72
<b>XPH</b>	54	19.61	1.46	55	20.24	1.49	57	20.88	1.51	58	22.07	1.71	54	23.59	2.05
<b>XALs</b>	53	20.25	2.35	54	21.44	2.38	58	22.19	2.34	58	23.14	2.79	53	22.98	2.86
<b>XALi</b>	54	21.43	2.38	53	22.43	2.60	57	23.00	2.49	56	23.41	2.89	56	24.02	3.16
<b>XAWs</b>	54	24.46	2.17	54	26.02	2.44	57	27.21	2.14	59	29.37	2.44	55	31.69	2.63
<b>XAWi</b>	54	26.06	2.39	54	27.39	2.32	58	29.53	2.57	56	32.23	2.48	56	34.43	2.99
<b>XPL</b>	54	4.35	1.07	55	4.64	1.01	59	5.20	0.91	59	6.10	1.23	56	7.16	1.51
<b>TPL</b>	53	28.75	2.50	54	28.07	2.21	56	27.80	2.47	57	26.18	2.46	53	24.17	3.05
<b>XTPW</b>	51	57.00	4.66	50	55.60	4.64	55	55.36	4.79	55	52.78	4.85	52	48.65	5.99
<b>XAPi</b>	54	31.19	3.38	55	32.00	3.70	58	33.53	3.06	59	33.76	3.59	55	33.02	4.55
<b>APH</b>	52	32.79	2.42	53	33.74	1.73	57	34.68	2.33	56	36.36	2.53	55	38.98	2.95
<b>XSPL</b>	48	43.25	4.57	47	42.04	4.77	54	38.81	3.65	58	35.03	3.92	55	32.25	4.62
<b>XSLi</b>	53	24.92	2.98	53	26.06	3.24	56	26.84	3.26	58	27.05	3.29	56	27.68	3.62

**Table A15:** The mean of all lower thoracic vertebrae (T7-T11) combined for population and sex. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Black	White	Male	Female
<b>XSLs</b>	19.62	22.55	22.09	19.99
<b>XAH</b>	17.31	18.62	18.48	17.41
<b>XPH</b>	18.01	19.62	19.41	18.20
<b>XALs</b>	16.23	19.01	18.71	16.45
<b>XALi</b>	17.44	20.66	20.12	17.90
<b>XAWs</b>	22.31	24.49	24.46	22.30
<b>XAWi</b>	23.59	25.89	26.08	23.36
<b>XPL</b>	4.88	5.41	5.17	5.10
<b>TPL</b>	29.30	32.60	32.50	29.31
<b>XTPW</b>	57.66	64.69	64.23	57.70
<b>XAPi</b>	31.12	35.27	34.72	31.58
<b>APH</b>	30.66	32.71	32.55	30.79
<b>XSPL</b>	38.78	43.58	43.49	37.74
<b>XSLi</b>	20.39	23.98	23.18	21.04

**Tables A16-A19:** Summary statistics of the lumbar vertebrae (L1-L4) showing the means and standard deviations for population and sex. Refer to Appendix I for measurement names and definitions.

**Table A16:** Summary statistics of the lumbar vertebrae (L1-L4) showing the means and standard deviation for black South Africans. Refer to Appendix I for measurement names and definitions.

	L1			L2			L3			L4		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	57	27.26	2.30	59	29.05	2.35	56	30.88	2.55	53	32.00	2.85
<b>XAH</b>	56	24.48	1.80	58	25.62	1.68	56	26.05	1.79	53	26.00	1.88
<b>XPH</b>	57	26.40	1.92	59	26.86	1.69	58	26.67	1.89	54	25.41	1.92
<b>XALs</b>	57	23.05	2.47	59	24.93	2.31	55	26.09	2.15	53	27.42	2.55
<b>XALi</b>	56	24.66	2.38	58	26.84	2.63	58	27.71	2.78	53	28.87	2.56
<b>XAWs</b>	56	37.23	3.23	59	39.64	3.40	58	41.95	3.92	54	43.56	2.91
<b>XAWi</b>	57	39.65	3.89	59	42.10	3.78	58	44.29	3.46	54	45.57	3.31
<b>XPL</b>	57	6.96	1.24	59	6.76	1.34	58	7.64	2.13	54	7.89	2.68
<b>TPL</b>	56	31.86	5.55	54	38.33	3.82	49	42.94	4.69	45	42.96	4.88
<b>XTPW</b>	47	62.45	10.30	41	77.10	6.03	41	86.83	8.94	30	87.03	10.46
<b>XAPWi</b>	57	26.82	3.90	58	29.52	4.16	54	33.17	4.82	50	40.34	6.40
<b>APH</b>	57	44.54	2.92	53	46.13	2.50	50	45.02	3.49	51	43.45	2.54
<b>XSPL</b>	56	33.30	3.61	56	34.73	3.63	51	34.67	4.52	52	32.08	3.25
<b>XSLi</b>	55	28.33	2.57	58	30.24	2.64	58	31.64	2.61	53	33.17	2.99

**Table A17:** Summary statistics of the lumbar vertebrae (L1-L4) showing the means and standard deviation for whites South Africans. Refer to Appendix I for measurement names and definitions.

	L1			L2			L3			L4		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	56	31.93	2.92	59	33.36	3.35	55	34.45	2.87	53	35.23	2.91
<b>XAH</b>	56	26.09	2.07	56	27.96	2.04	54	28.78	2.06	55	29.13	2.13
<b>XPH</b>	57	28.35	1.73	57	28.88	1.81	53	28.72	1.95	55	28.04	2.20
<b>XALs</b>	57	28.60	3.19	59	29.88	3.26	55	30.58	3.22	53	31.19	2.67
<b>XALi</b>	56	29.23	2.97	54	30.15	3.08	53	30.72	2.79	52	31.62	2.87
<b>XAWs</b>	58	40.71	3.48	58	42.66	3.66	54	44.44	3.66	56	46.61	3.83
<b>XAWi</b>	56	43.64	3.69	56	42.20	3.66	56	47.38	3.74	55	49.45	3.81
<b>XPL</b>	57	7.65	1.47	59	7.85	1.40	57	9.19	1.56	56	11.38	1.86
<b>TPL</b>	47	35.36	6.74	37	39.65	4.57	42	43.19	5.02	35	43.31	4.80
<b>XTPW</b>	32	67.12	13.30	25	78.48	6.65	21	84.67	5.01	16	86.06	11.13
<b>XAPi</b>	57	28.63	3.15	55	31.49	3.85	52	34.23	4.77	53	43.57	6.68
<b>APH</b>	53	47.26	3.60	53	48.49	3.30	47	48.55	3.81	49	45.73	3.87
<b>XSPL</b>	55	35.89	2.79	50	38.36	3.19	45	40.49	3.31	47	38.32	4.27
<b>XSLi</b>	56	32.88	3.05	56	33.73	3.17	53	34.23	2.71	53	34.75	2.74

**Table A18:** Summary statistics of the lumbar vertebrae (L1-L4) showing the means and standard deviation for males. Refer to Appendix I for measurement names and definitions.

	L1			L2			L3			L4		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	54	31.24	3.31	59	32.73	3.54	54	33.78	3.15	58	34.90	3.14
<b>XAH</b>	55	25.44	2.16	57	26.72	2.08	55	27.53	2.51	59	27.76	2.87
<b>XPH</b>	55	28.05	2.02	57	28.32	1.91	55	28.20	2.04	59	27.36	2.50
<b>XALs</b>	55	27.78	3.84	59	29.14	<b>3.97</b>	54	29.72	3.73	59	30.44	3.30
<b>XALi</b>	55	28.69	3.37	56	29.95	3.32	57	30.79	2.94	57	31.65	3.02
<b>XAWs</b>	55	41.33	3.43	59	43.14	3.69	56	44.96	4.25	59	46.92	3.50
<b>XAWi</b>	54	44.06	4.15	57	45.77	3.79	57	47.82	3.96	58	49.40	3.98
<b>XPL</b>	55	7.44	1.50	59	7.29	1.64	58	7.93	2.15	59	8.93	3.46
<b>TPL</b>	51	35.57	6.81	45	41.00	3.68	48	45.15	4.47	47	44.87	4.55
<b>XTPW</b>	38	67.32	12.73	33	80.06	4.68	33	89.15	7.84	29	89.07	10.19
<b>XAPi</b>	55	28.71	3.88	55	31.38	3.95	53	34.26	4.76	56	44.05	6.55
<b>APH</b>	52	47.13	3.45	54	48.13	2.71	51	47.67	3.84	54	45.17	3.42
<b>XSPL</b>	54	36.07	3.23	52	37.85	3.93	52	38.52	4.54	59	36.04	5.25
<b>XSLi</b>	55	32.33	3.25	56	33.29	3.04	56	34.23	2.80	57	35.42	2.69

**Table A19:** Summary statistics of the lumbar vertebrae (L1-L4) showing the means and standard deviation for females. Refer to Appendix I for measurement names and definitions.

	L1			L2			L3			L4		
	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>	<i>n</i>	<i>mean</i>	<i>sd</i>
<b>XSLs</b>	59	28.05	2.98	59	29.68	2.98	57	31.58	2.99	48	32.06	2.79
<b>XAH</b>	57	25.14	2.04	57	26.82	2.34	55	27.25	2.21	49	27.39	2.10
<b>XPH</b>	59	26.75	1.91	59	27.41	2.03	56	27.11	2.16	50	26.00	2.18
<b>XALs</b>	59	24.00	3.17	59	25.68	2.56	56	27.00	2.76	47	27.87	2.46
<b>XALi</b>	57	25.26	2.80	56	26.93	2.49	54	27.41	2.37	48	28.54	2.04
<b>XAWs</b>	59	36.83	2.62	58	39.10	2.77	56	41.34	2.70	51	43.02	2.78
<b>XAWi</b>	59	39.41	3.02	58	41.48	3.02	57	43.79	2.61	51	45.41	2.97
<b>XPL</b>	59	7.19	1.29	59	7.32	1.29	57	8.89	1.76	51	10.51	1.69
<b>TPL</b>	52	31.38	5.10	46	36.78	3.53	43	40.72	4.11	33	40.61	4.08
<b>XTPW</b>	41	61.59	10.16	33	75.18	6.74	29	82.62	6.38	17	82.65	10.29
<b>XAPi</b>	59	26.81	3.18	58	29.62	4.12	53	33.11	4.82	47	39.55	6.12
<b>APH</b>	58	44.71	3.20	52	46.46	3.36	46	45.70	4.04	46	43.87	3.36
<b>XSPL</b>	57	33.18	3.09	54	35.09	3.32	44	36.07	5.10	46	33.89	4.20
<b>XSLi</b>	56	28.95	3.17	58	30.67	3.22	55	31.49	2.42	49	32.27	2.30

**Table A20:** The means of all lumbar vertebrae (L1-L4) combined for population and sex. Refer to Appendix I for measurement names and definitions.

	Population		Sex	
	Black	White	Male	Female
<b>XSLs</b>	29.78	33.73	33.20	30.28
<b>XAH</b>	25.55	27.97	26.87	26.63
<b>XPH</b>	26.36	28.51	27.99	26.84
<b>XALs</b>	25.36	30.02	29.27	26.07
<b>XALi</b>	26.98	30.49	30.37	26.95
<b>XAWs</b>	40.36	43.58	44.14	39.74
<b>XAWi</b>	42.86	46.22	46.63	42.41
<b>XPL</b>	7.72	8.97	7.87	8.83
<b>TPL</b>	38.23	39.93	41.38	36.35
<b>XTPW</b>	76.68	77.01	80.42	72.84
<b>XAPi</b>	32.17	34.29	34.59	31.84
<b>APH</b>	44.75	47.49	46.99	45.15
<b>XSPL</b>	33.76	38.22	37.18	34.55
<b>XSLi</b>	30.85	33.89	33.81	30.82