

Some Notes on the Vértesszöllös Occipital¹

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ABSTRACT An examination of the Vértesszöllös occipital bone indicates the need for taking its condition and preservation as well as certain aspects of its morphology into account prior to a metric assessment of its features. This analysis confirms the presence of *extrasutural bones* on the lambdoidal suture as well as some *distortion* in the sagittal curvature of this region. The advisability of using the endocranial sutures as well as other data for *defining lambda* in cases such as this is discussed, and a procedure for determining the *position of opisthion* is indicated. A metric and morphological analysis is used to support the claim that the specimen can be considered a large late representative of *Homo erectus* in Europe.

The occiput from Vértesszöllös is the only hominid cranial fragment known from the European Mindel at the present time. Because there is only a single specimen, the desire to obtain as much information from it as possible must be balanced against the fact that it cannot represent the range of the population from which it was drawn and may or may not approximate the mean. The potential importance of the specimen lies in the knowledge it provides concerning evolutionary changes at the very end of the *Homo erectus* lineage or the very beginning of the *Homo sapiens* one, depending on how it is interpreted taxonomically; other important facets include the question of whether European hominids are geographically distinct from their contemporaries and whether there is evidence for multiple hominid lineages in Europe during the Mindel. It is possible that the potential importance of the specimen has led to its overinterpretation. However, another factor underlying the continuing discussions regarding the specimen results from the damaged and incomplete condition of the bone. I had the opportunity to examine this specimen during the summer of 1976. The purpose of this report is to ascertain the influence of damage and distortion on the interpretation of the specimen, to discuss the problems involved in delineating some of the important landmarks, and to compare it with occiputs from other

Riss and pre-Riss European hominids with the intent of determining its relationship to them.

SOME PROBLEMS IN PRESERVATION

While the occiput recovered from the Hungarian site of Vértesszöllös is in surprisingly good condition, given its age (Kretzoi and Vértes, '65) and the lack of substantial internal mineralization, some elements of distortion combined with the preserved morphology underlie the continuing discussions concerning its phylogenetic position (Thoma, '66, '69, '72; Howells, '66; Wolpoff, '71). Metric comparisons have been hampered by the absence of opisthion and questions concerning the position of lambda, while morphological evaluation must contend with the absence of virtually all of the outer bone table in the region of the nuchal torus and elsewhere, as well as with what appear to be numerous breaks and angled joins in the superior occipital plane.

The main distortions in the bone occur towards the top of the occipital plane (fig. 1). Here, reconstruction (or its absence) has resulted in an unusual angulation at the most superior aspect of the bone, sharply setting the curvature off from the relatively smoothly curving occipital plane. This acts to make the

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Fig. 1 Lateral view of the occiput, showing the deep sulcus below the nuchal torus, and the bulging effect above it.

occipital curvature appear too high. Inferior and posterior to this, the occipital plane swells convexly and gives the superficial appearance of a "Neandertal-like" bun. The top of this swelling is some 13 mm below lambda (see discussion below) and 36.8 mm above

inion. However, unlike the condition in these later European hominids there is no corresponding fossa internally. Instead, the internal surface is smoothly curved so that the apparent bulge is the result of thickened bone.

No sulcus separates this convex region from the nuchal torus below, although a deep sulcus lies inferior to the torus. However, virtually no outer bone table remains on the torus so that its true size is easily underestimated. Observed where the full bone surface remains (as in the position of inion), the torus is projecting and is rather thick (fig. 2). Its position is high above the internal inion (table 2).

Opisthion

While prior metric analyses of the specimen (Thoma, '66, '69, '72; Wolpoff, '71) utilized measures of both nuchal and occipital planes, procedures regarding the definitions of the positions for both lambda and opisthion must be explicitly stated since there are problems regarding both measuring points. In the case of opisthion, the problem is quite simply that the bone is broken away posterior to the foramen magnum border, and in fact not far anterior to the position of internal inion in some positions.

The position of opisthion can be estimated from other landmarks. The relatively high position of the external inion makes it difficult

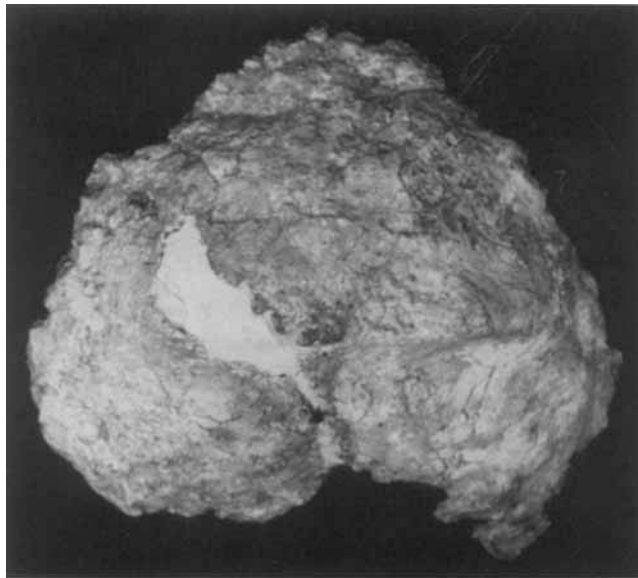


Fig. 2 Posterior view of the occiput.

to use in estimating opisthion. For instance, if a Swanscombe model is used, emphasizing the more derived features of the specimen, opisthion would be placed 51 mm from inion. However, this would position it only 25 mm from the internal occipital protuberance (endinion), in contrast to the corresponding Swanscombe value of 30.6 mm.

Other features can more accurately be used to position the posterior border of the foramen magnum. A notch visible on the broken edge of the inferior right surface seems to represent the posterior portion of the condylar fossa border; a smaller portion of the posterior border appears in the symmetric position on the left (fig. 3). Medial and anterior to the right notch border a small ridge appears to mark the lateral base of the condyle on the jugular process. Both of these features lie anterior to opisthion, and thus help define its position.

Comparison with the internal morphology of Swanscombe (the only cast available) reveals a virtually identical position for the features described above, as well as an equivalent position for the divergence of the lines laterally bounding the occipital sinus as measured from endinion. Use of the Swanscombe occiput as a model seems justified by these comparisons, and the position of opisthion was placed some 30.5 mm below endinion. Although this estimate provides a *minimum* inion-opisthion dimension (table 1), the nuchal plane of Vértesszöllös appears to have been very long. The 57 mm value thus determined is virtually the same as the 56 mm dimension reported by Thoma ('69: p. 234).

Lambda

There are also some difficulties concerning the true position of lambda. Contra Thoma ('72) lambda is not at the most superior and anterior point on the midline of the bone as preserved. As I previously stated on the basis of a cast examined at the Harvard Peabody Museum, extrasutural bones on the anterior border of the occipital confuse the definition of lambda. Thoma regards these as "small fractures of the tabula externa" (p. 464). My examination of the original specimen suggests that both statements are probably correct; there is considerable breakage in the area and extrasutural bones are present. Several different factors lead me to conclude that more than breakage characterizes this area (fig. 4). (1)



Fig. 3 Inferior view of the occiput. The posterior condylar fossa border is on the superior edge of the large broken notch on the lower right.

Many of the fractures, if they are such, extend through to the internal bone surface and could just as easily define extrasutural bones. (2) The termination of the cerebral fossa is clearly observable, marking the superior border of the occiput. However this termination lies below the superior border of the *preserved* internal surface. (3) The most superior extension of the sagittal sulcus falls some 21.5 mm short of the most superior anterior point on the midline (Thoma's lambda) although it normally extends to lambda. (4) In the region superior and anterior to the termination of the sulcus the unusual angulation of the occipital plane occurs, described above as a consequence of breaks.

Yet, if the region anterior and superior to the border of the sagittal sulcus is not part of the occiput, *the absence of sagittal suture indicates that it is also not part of the parietal*. Instead, the region consists of at least two definable extrasutural bones. The condition is similar to the large extrasutural bone found in the pre-Riss cranium from Petralona (fig. 5), although in this case the extrasutural bone involves only the occiput. These bones should not be used in metric or morphologic descriptions of either occipital or parietal; including them as part of the occipital leads to the unusual length of the occipital plane reported by Thoma ('69). Thoma's occipital length is not homologous with the dimension as normally defined since it includes bone surface

TABLE 1

Comparison of dimensions on the sagittal plane for Vertesszöllös and Swanscombe, Petralona, and a cast of La Chaise 2

	La Chaise	Petralona	Swanscombe	Vertesszöllös				
				a	b	c	d	e
Lambda-opisthion								
Chord	84.7	92.9	93.5	91.6	107.3	99.0	100.0	102.0
Arc	108.0	130.0	117.0	111.0	132.0			
Index	127.5	139.9	125.1	121.2	123.0			
Occipital plane (inion-lambda)								
Chord	49.5	63.6	58.0	49.8	71.3			73.0
Arc	55.0	68.0	63.0	53.0	74.0			79.0
Index	111.1	107.9	108.6	107.1	103.8			108.2
Nuchal plane (opisthion-inion)								
Chord	51.0	60.6	51.0	57.0	57.0		53.0	56.0
Arc	53.0	62.0	54.0	58.0	58.0			
Index	103.0	102.3	105.9	101.8	101.8			
Occipital-nuchal index								
Chord	97.1	105.0	113.7	87.4	125.1			130.4
Arc	103.8	109.7	116.7	91.4	127.6			
Inion angle	114.9°	96.8°	118.0°	118.0°	113.0°			103.7°
Lambda angle	33.1°	40.4°	28.8°	33.3°	29.3°			32.2°
Opisthion angle	32.0°	42.8°	33.2°	28.7°	37.7°			44.0°

All dimensions are in millimeters and with the exception of La Chaise were taken on the original specimens. The three angles were calculated from the dimensions of the sides of a triangle defined by the occipital plane, nuchal plane, and lambda-opisthion chord. Under the Vertesszöllös heading are the following:

- position of lambda and opisthion as discussed in text, determined on the original specimen by Wolpoff
- position of opisthion as above, lambda defined as the point used by Thoma (the most anterior preserved point on the midline)
- taken by Wolpoff on a cast ('71)
- measured by Tobias, reported in Thoma ('69)
- measured by Thoma ('69).

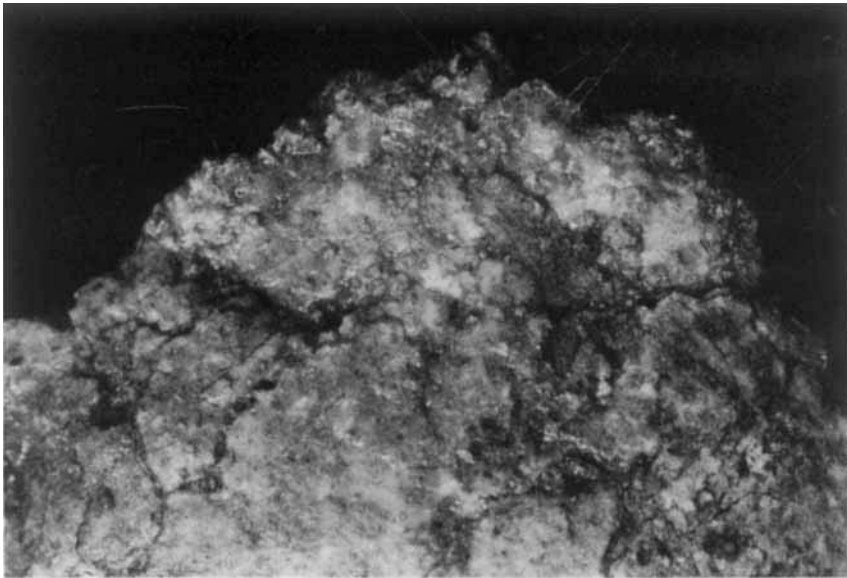


Fig. 4 View of the region surrounding the most superior aspect of the lambdoidal suture. This area seems to be characterized by both breaks and the presence of extra sutural bones.



Fig. 5 Posterior view of the Petralona cranium. Note the large extrasutural bone just below lambda.

covering more than the occipital lobes of the brain.

The homologous measurement can be defined by treating the extrasutural bones as neither part of the occiput nor the parietal, and defining lambda on the basis of the internal morphology as in MLD 2 (Wolpoff, '74). This would place lambda internally some 57.5 mm above the endinion position, and externally 49.8 mm above inion (table 2). The resulting height of the left cerebral fossa is 56 mm while on the right it is 47 mm from the transverse sinus to the internal position of lambda. The asymmetry is exactly the opposite of Swanscombe, where the right cerebral fossa is the larger.

AFFINITIES

Evaluation of the metrics (tables 1, 2) indicates that Vértesszöllös is a large specimen. Generally, its dimensions exceed those of *Homo erectus* occiputs in the total height of the bone, as indicated by the lambda-opisthion chord as reconstructed by the author. Since both nuchal and occipital plane lengths fall well within the erectus range, the greater height of the bone is the result of a higher angle measured at inion; the value of 118° reported here exceeds the 103° value given by Thoma ('69) and both values may be incorrect because of the presence of sagittal distortion in this region discussed above.

In spite of its size, the occiput shows a series of affinities to the *Homo erectus* sample,

matched by differences from the other fairly early occiputs from Europe; I interpret these to suggest that it probably should be included in the former taxon. Moreover, no features indicate that it represents a population ancestral to only some of the later European hominids while not to others (i.e., a pre-Riss lineage divergence of European hominids, or the "Presapiens Hypothesis"). This interpretation is now also agreeable to Thoma ('76).

My reasons for considering the Vértesszöllös occiput as a late representative of *Homo erectus* include the many detailed resemblances discussed by both Thoma ('66, '69) and myself ('71), and the position of the specimen with respect to trends in the evolution of the mid-Pleistocene occipital. Apart from some specific morphological characters, the main aspects that characterize the occiputs of the earliest members of *Homo sapiens* result from selection reducing the length of the nuchal plane (while generally increasing its breadth) and increasing the length of the occipital plane. The index formed by these two lengths changes markedly, passing from a value of about 70 in *Homo erectus* (i.e., the occipital plane is 70% the length of the nuchal plane) to 113 in six Solo crania and considerably higher values in the Würm Neandertals. Values for European hominids from the mid-Pleistocene are all above the erectus range (maximum index is 94) except for Vértesszöllös (table 1) which falls close to the Choukoutien mean of 83.

Other characteristics which progressively change with the appearance and subsequent evolution of *Homo sapiens* include the approach of internal and external inion as the nuchal plane decreases in length (the Vértesszöllös inion is far above endinion); the loss of the sulcus inferior to the nuchal torus (Vértesszöllös retains the sulcus); the "filling out" of the occipital plane resulting in a high arch-chord index (the Vértesszöllös index is like both *Homo erectus* and the earlier European hominids); thinning of the occipital in the regions of asterion and inion (the Vértesszöllös values for bone thickness are close to the erectus mean); and the more open occipital angle measured at inion (here alone is Vértesszöllös more like the later hominids and outside of the known *Homo erectus* range). In some respects, the Petralona occiput resembles Vértesszöllös. There is a similar ap-

TABLE 2

Comparison of *Vertesszöllös* dimensions (mm) with the specimens used in table 1

	La Chaise	Petralona	Swanscombe	Vertesszöllös	
				a	b
Biasterionic breadth	109.0	119.0	121.8	128.3	
Nuchal plane breadth	105.5	106.5	84.5	108.0	
Biasterionic arc		155.0	160.0	150.0	
Lambda-asterion					
Chord					
L		87.3	91.6	90.3	100.5
R		80.0		85.8	96.0
Arc					
L		97.0	104.0	101.0	115.0
R	101.0	90.0		97.0	111.0
Index					
L					
R					
Inion-internal protuberance	11.0		15.8	27.8	
Maximum thickness at					
Asterion	11.3			14.3	
Lambda	11.6		10.4	10.0	
Inion	14.0		9.0	16.0	
Internal protuberance	15.0		14.5	10.0	
Minimum thickness	3.5		6.0	3.0	

"a" and "b" are as defined in table 1.

parent bun-like configuration to the occipital plane and a sulcus below it, although not as deep. The occipital plane exceeds the nuchal plane in length, and the lambda-opisthion chord is greater (and the arc much greater) in spite of the lower occipital angle. A greater contrast is provided by the Bilzingsleben occiput (Mania, '75) which is smaller and less strongly angled (about 113°), with little curvature in the occipital sagittal plane. Steinheim has a longer and more rounded occipital plane in its present condition (Weinert, '36). How an eventual reconstruction will effect this is unknown, and in any event the specimen may be younger than normally assumed.

CRANIAL CAPACITY

Finally, the capacity of the specimen has been used in assessing its phylogenetic position, although a direct determination is impossible from the occiput alone. Estimation of the *Vertesszöllös* cranial capacity has been attempted on the basis of linear regressions from the lambda-opisthion chord (Thoma, '69; Wolpoff, '71). However, for a regression based on the lambda-opisthion chord to be used in calculating cranial capacity, two different problems must be settled. First which specimens should provide the basis for defining the regression (Olivier and Tissier, '75)? A regression based on Neandertals gives quite different results from one based on *Homo erectus* (Wolpoff, '71). Thus, using capacity to argue

for a particular taxonomic position is bound to be circular, since one must make a taxonomic decision in order to determine the capacity. Second, which of the different lambda-opisthion estimates should be used; values have been provided by Thoma, Tobias, and myself in various publications as well as this one.

In table 3, all of the lambda-opisthion estimates are used to calculate capacity in a series of regressions based on the seven *Homo erectus* specimens with lambda-opisthion dimensions that can be determined without reconstruction. Since Thoma ('72) has suggested that an exponential curve instead of a line be used to relate the chord to capacity, least square formulae are determined for a line, a second degree polynomial, a power curve, and an exponential curve. As it turns out, none of these have dramatically different percent errors when applied to the samples from which they were derived, and the estimates of the *Vertesszöllös* capacity likewise are not particularly sensitive to which model is used.

The *Vertesszöllös* estimate is much more sensitive to which lambda-opisthion chord estimate is used; a 10 mm difference in the chord can effect the capacity estimate between 224 cc in the linear regression and 310 cc in the exponential regression. While I would support the capacity estimate based on the lambda-opisthion chord determined between the points as defined here, the different

TABLE 3

Derivation of four curves for estimating cranial capacity from the lambda-opisthion chord in *Homo erectus* specimens with a known lambda-opisthion dimension

	Lambda opisthion chord	Actual or estimated capacity	Y = AX + B Linear regression capacity	B = AX ² + BX + C = AX ^B Polynomial regression capacity	= AX ^B Power curve capacity	= Ae ^{BX} Exponential curve capacity
Sangiran 2	75	775	735	739	740	745
Sangiran 4	78	750	802	802	802	802
Sangiran 12	83	900	914	911	909	905
Sangiran 17	93	1,125	1,138	1,143	1,145	1,154
Peking 3	84	915	936	934	932	928
Peking 11	86	1,015	981	979	977	974
Peking 12	87	1,030	1,004	1,002	1,000	998
Percent Error ¹			3.3%	3.2%	3.3%	3.2%
Values A			22.37	0.09	0.12	120.37
B			-942.82	8.05	2.03	0.02
C				-345.93		
Vértesszöllös estimate	92		1,115	1,119	1,120	1,127
	99		1,272	1,289	1,300	1,336
	100		1,294	1,314	1,327	1,369
	102		1,339	1,365	1,381	1,437

¹ Based on absolute difference.

All of the curves were determined through a least squares procedure. In turn, each curve is used to estimate the Vértesszöllös capacity given various determinations of the lambda-opisthion chord (see table 1), and the percent error of the curve as applied to calculating known erectus capacities is given.

capacity estimates should not mask the fact that if one treats the Vértesszöllös cranium as a late *H. erectus*, its capacity is in the uppermost portion of the *erectus* range whichever chord is correct.

SUMMARY

In sum, this occiput likely belonged to an individual with a cranial capacity near or perhaps even slightly above the known range for *Homo erectus*, although probably not much different from the capacity that can be estimated for the Ternifine juvenile parietal. Both the high occipital angle and the large capacity result from the value of the lambda-opisthion chord, and thus are not independent of each other. Moreover, the fact that both lambda and opisthion must be reconstructed or estimated in this specimen detracts from the value of estimates made on the basis of the chord. All other features of the specimen, both individually considered and in combination, are consistent with the interpretation that Vértesszöllös is a late and distinctively European *Homo erectus* which cannot be excluded from the ancestry of any later European hominid on the basis of morphology or metrics.

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