

# Study of the growth and shape of the brain and cranial base during the first two years of life

C. Delteil, E. Lesieur, L. Tuchtan, A. Carballeira Alvarez, K. Chaumoitre, B. Saliba, P. Adalian, M.-D. Piercecchi-Marti

# ▶ To cite this version:

C. Delteil, E. Lesieur, L. Tuchtan, A. Carballeira Alvarez, K. Chaumoitre, et al.. Study of the growth and shape of the brain and cranial base during the first two years of life. Morphologie, 2021, 105(348), pp.45-53. 10.1016/j.morpho.2020.09.007. hal-03570651

HAL Id: hal-03570651

https://hal.science/hal-03570651

Submitted on 14 Feb 2023

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Version of Record: https://www.sciencedirect.com/science/article/pii/S1286011520300898 Manuscript b6422dcf9cc2056fa7cbf69d5124482e

Study of the growth and shape of the brain and cranial base during the first two years of life

Etude de la croissance et de la conformation de l'encéphale et de la base du crâne durant les deux

premières années de vie.

C. Delteil a,b, E. Lesieur c, L. Tuchtan A, A. Carballeira Alvarez d, K. Chaumoitre e, B. Saliba b, P. Adalian b, M-D

Piercecchi-Marti a

a Forensic Department, Assistance Publique-Hôpitaux de Marseille, La Timone, 264 rue St Pierre, 13385

Marseille Cedex 05, France

b Aix Marseille Univ, CNRS, EFS, ADES, Marseille, France

c Department of Gynaecology and Obstetrics, Gynépole, Assistance Publique-Hôpitaux de Marseille, AMU,

Aix-Marseille Université, Marseille, France

d Department of Radiology, Conception Hospital, Assistance Publique-Hôpitaux de Marseille, 147 Boulevard

Baille, 13005 Marseille Cedex 05, France

e Department of Radiology, CHU Nord, Assistance Publique-Hôpitaux de Marseille, France, Chemin des

Bourrely, 13015 Marseille

Corresponding author:

Clémence Delteil

https://orcid.org/0000-0001-5171-8520

Forensic Department, APHM, La Timone, 264 rue St Pierre, 13385 Marseille Cedex 05, France

Tel. (33) 4 91 38 63 85

Fax (33) 4.91.92.33.31

E-mail address: clemence.delteil@ap-hm.fr

1

© 2020 published by Elsevier. This manuscript is made available under the CC BY NC user license https://creativecommons.org/licenses/by-nc/4.0/

KEYWORDS: GROWTH, MATURATION, BRAIN, SKULL BASE, INFANT

MOTS-CLES: CROISSANCE, MATURATION, CERVEAU, BASE DU CRANE, NOURRISSON

**Abstract** 

Ontogeny of the cranial base and the brain integrates data on growth, maturation and ontogenetic allometry of

these two systems in the course of development.

The aim of our work was to study the ontogeny of the cranial base and the brain in order to understand their

growth dynamic and shape changes using a traditional morphometric approach in individuals with normal (non-

pathological) development.

Materiel and Method: Forty-seven infants having been included in the unexpected infant death french protocol

were analyzed. Medical imaging (CT and MRI) exams, followed by an autopsy and pathology examination

allowed us to include only infants free from brain disease or pathology affecting growth.

Results: Testing of measurement reliability validated 12 distances and 3 angles as well as the positioning of the

landmarks that had been used to obtain the distances and the angles.

No correlation between sex and the various variables studied was found. However, a correlation was observed

between these variables and age, making it possible to propose a growth curve. A medium to strong correlation

was found between brain variables and the bone variables of the cranial base, underlining the parallel

development of the two systems.

Conclusion: Our study, carried out in a rigorously selected population of infants, presents a fundamental

approach to the study of ontogenesis.

Résumé:

L'ontogénie de la base du crâne et de l'encéphale intègre les données de croissance, de maturation et d'allométrie

ontogénétique de ces deux systèmes au cours du développement.

But de l'étude : Etudier l'ontogénie de la base du crâne et de l'encéphale afin de comprendre leur dynamique de

croissance ainsi que les modifications de conformation à partir d'une approche en morphométrie traditionnelle.

Matériel et méthodes : Quarante-sept nourrissons ayant été inclus dans le protocole français de mort subite du

nourrisson ont été analysés. Les examens d'imagerie médicale (tomodensitométrie et IRM), suivis d'une autopsie

et d'un examen anatomopathologique nous ont permis de n'inclure que les nourrissons exempts de maladie

cérébrale ou de pathologie affectant la croissance.

2

**Résultats:** Le test de fiabilité des mesures a permis de valider 12 distances et de 3 angles ainsi que la pose des points de repères ayant servi à obtenir les distances et les angles. Aucune corrélation entre le sexe et les différents paramètres étudiés n'a été mise en évidence. En contrepartie, une corrélation entre les différents paramètres étudiés et l'âge a été attestée, permettant de proposer une courbe de croissance. Une corrélation dont la valeur fluctue de moyenne à forte entre les paramètres encéphaliques et les paramètres osseux de la base du crâne a été mise en évidence, soulignant une évolution parallèle des deux systèmes.

Conclusion : Notre étude, conduite sur une population de nourrissons rigoureusement sélectionnés, présente une approche fondamentale dans l'étude de l'ontogénèse.

#### 1. Introduction

Brain volume doubles in the first 6 months of life, reaching 50% of adult brain volume. Brain development is heterogeneous. The rate of maturation is different depending on the brain region. Thus, the sensory and motor regions organize themselves first, while the regions involved in more associative functions are extremely slow and will not mature until after puberty and even continue to change throughout life. The integration of active growth systems of the brain and cranial base suggest that these systems mutually influence each other, although this influence has never been proven to be an exclusive one [1]. This reciprocal influence may occur during the the fetal period and the first three years of life, at the time when brain growth is greatest [2].

However, evaluation of growth of the brain and the cranial base relies on incomplete data [3–5]. Radiological charts exist for the fetus [6], but there are none for infants with no pathological condition. Only radiological comparisons between pathological [7] or premature infants [8,9] and infants considered as "normal" have been reported, with no mention of "normal" measurements.

In view of this lack of data in the infant, we aimed to study the development of the cranial base and the brain during the first two years of life through an ontogenic study of these two systems using conventional morphometrics.

Conventional morphometrics consists of the application of multivariate statistical analyses to a series of quantitative variables: linear distances (between two points: length, width, height), angles, ratios, surface and volume [10]. It is used in comparative biology to study growth, shape and allometry [10].

The aim of our work was to study the ontogeny of the cranial base and the brain in order to understand their normal (non-pathological) growth dynamic and age-related shape changes.

## 2. Material and Methods

## Sample

The samples studied consisted of a retrospective series of infants who had died between the ages of 0 and 24 months and who had been included in the sudden infant death (SID) protocol in the sudden infant death unit of the Hôpital de la Conception in the geographical reference region of west Provence-Côte d'Azur, France.

We studied 58 infants who had died between January 1, 2010, and 31 December, 2016, and had been managed in the public hospitals of Marseille (Assistance Publique des Hôpitaux de Marseille). The majority of

deaths were due to sleeping accidents (positional mechanical asphyxia), infectious disorders (viral and bacterial) and surgical abdominal disorders (acute intestinal intussusception, peritonitis, volvulus).

Only infants with no disorder affecting growth and no brain disorder identified on imaging followed by autopsy and histopathological examination were included in the study.

The non-inclusion criteria were incomplete or insufficient quality imaging exams (CT-scan or magnetic resonance imaging).

Autopsy completed by histopathological examination was considered the gold standard to exclude any disorder. Autopsy was performed after the written consent of the parents had been obtained. Any external or internal abnormalities were described. Information on pregnancy, term at birth, and medical and surgical history was also recorded.

For each infant, anonymized anthropological data were available: chronological age (or corrected age if the infant was born before 38 weeks amenorrhea), weight, crown-heel length, head circumference and sex.

# **Imaging protocols**

As part of the recommendations of the High Authority for Health (Haute Autorité de la Santé, HAS) [11] regarding sudden infant death, each infant underwent:

- CT scan: whole-body scans were obtained and recorded in Digital Imaging and Communications in Medicine (DICOM) format. The native scans (overlapping 0.50–5 mm sections) were the raw data used in our study.
- Magnetic resonance imaging (MRI): whole-body scans recorded in DICOM format. The native scans (overlapping 0.9-5 mm sections) were the raw data used in our study.

## Conventional morphometric measurement protocols

Seventeen linear measurements and 5 angles were obtained using OsiriX version 5.6 software developed by Pixmeo. Three-dimensional multiplanar reconstruction (MPR) enabled optimal selection of the landmarks and thus of the measurements obtained. The measurements were selected according to literature relating to normal cranio-cerebral development in the foetus: brainstem (1), cerebral (1, 6), cerebellum (1, 7), cranial base (1, 12). All measurements and the methods used to obtain them are described in Table 1 (figures in Annex).

Table 1

| Imaging | Measurements | Abbreviation | Definition |
|---------|--------------|--------------|------------|
|         |              |              |            |

| type |                        |         |   |  |
|------|------------------------|---------|---|--|
| CT   | AP length of the PP    | PP AP   | Axial section perpendicular to middle cranial fossa: most       |  |
|      |                        |         | anterior and most posterior point of the right PP (12)          |  |
| CT   | T PP height PP H       |         | Axial section perpendicular to right middle cranial fossa:      |  |
|      |                        |         | most medial point, passing through the external auditory        |  |
|      |                        |         | canal (12)  |  |
| CT   | Surface of the PP      | PP Suf  | Axial section perpendicular to right middle cranial fossa,      |  |
|      |                        |         | delimiting the region of interest (ROI)                         |  |
| CT   | Maximum length of      | PB ML   | Axial section through the axis: maximum length of the PB        |  |
|      | the PB                 |         | of the occipital bone (12)                                      |  |
| CT   | Sagittal length of the | PB SL   | Axial section through the axis: sagittal length of PB the       |  |
|      | PB                     |         | occipital bone (12)   |  |
| CT   | Maximum width of       | PB MW   | Axial section through the axis: maximum width of PB the         |  |
|      | the PB                 |         | occipital bone (12)   |  |
| MRI  | Length of anterior     | Ant L   | Median sagittal section: distance from the foramen caecum       |  |
| (T1) | level                  |         | to the area between the two anterior clinoid processes          |  |
| CT   | Length of the middle   | Mid L   | Horizontal section: distance between the most lateral part of   |  |
|      | level                  |         | the anterior border of the right PP to the most lateral part of |  |
|      |                        |         | the small wing of the right sphenoid                            |  |
| MRI  | Length of the          | Post L  | Median sagittal section: distance from the region between       |  |
| (T1) | posterior level        |         | the two posterior clinoid processes to the internal occipital   |  |
|      |                        |         | protuberance  |  |
| MRI  | Total length of the    | Total L | Median sagittal section: distance from the foramen caecum       |  |
| (T1) | cranial base           |         | to the internal occipital protuberance                          |  |
| MRI  | Sphenoid angle         | Sph A   | Median sagittal section: nasion-sphenoid-occipital basion       |  |
| (T1) |                        |         | angle (1)   |  |
| MRI  | Interpetrous angle     | IPA     | Axial section: angle of the intersection of lines drawn         |  |
| (T1) |                        |         | through the posterior border of the Crista partis petrosae (7)  |  |
| MRI  | Occipitofrontal        | OFD     | Median sagittal section: distance between the extremities of    |  |
| (T2) | diameter               |         | the occipital and frontal lobes (6)                             |  |

| MRI  | Bone biparietal           | Bone BPD   | Coronal section through the temporal horns of the lateral      |  |
|------|---------------------------|--|--|--|
| (T2) | diameter                  |  | ventricles: greatest diameter of the brain: maximum distance   |  |
|      |                           |  | between the two internal tables of the cranium (6)             |  |
| MRI  | Brain biparietal          | Brain BPD  | Coronal section through the temporal horns of the lateral      |  |
| (T2) | diameter                  |  | ventricles: greatest diameter of the brain (6)                 |  |
| MRI  | Length of corpus          | LCC  | Median sagittal section: distance between the genu and the     |  |
| (T2) | callosum                  |  | posterior extremity of the splenium (6)                        |  |
| MRI  | AP vermis diameter        | P vermis diameter APVD Median sagittal section: diameter taken p |  |  |
| (T2) |                           |  | median part of the roof of the 4 <sup>th</sup> ventricle (6)   |  |
| MRI  | Height of the vermis      | HV   | Median sagittal section: maximum height of the vermis,         |  |
| (T2) |                           |  | parallel to the brainstem (6)                                  |  |
| MRI  | Transverse cerebellar TCD |  | Coronal section through the atrium: maximum cerebellar         |  |
| (T2) | diameter                  |  | diameter (6)   |  |
| MRI  | Brainstem-tentorium       | BSTA   | Median sagittal section: intersection between the line         |  |
| (T2) | angle                     |  | tangent to the posterior aspect of the pons and the line       |  |
|      |                           |  | tangent to the cerebellar tentorium                            |  |
| MRI  | Basion-tentorium          | BTA  | Median sagittal section: intersection between the line joining |  |
| (T2) | angle                     |  | the most internal part of the roof of the cerebellum and the   |  |
|      |                           |  | line tangent to the cerebellar tentorium (7,13)                |  |
| MRI  | Pontomedullary angle PMA  |  | Median sagittal section: angle between the upper 2/3 of the    |  |
| (T2) |                           |  | anterior aspect of the pons, the posterior aspect of the       |  |
|      |                           |  | bulbomedullary junction and the posterior aspect of the        |  |
|      |                           |  | lower cervical spinal cord                                     |  |

PP: petrous pyramid; AP: anteroposterior; PB: pars basilaris; BS: brainstem; T1: T1-weighted sequence; T2: T2-weighted sequence.

# Statistical analysis

Statistical analysis was carried out using the RStudio statistical environment [14] and R software version 3.2. Repeatability (intra-observer) and reproducibility (inter-observer) were analyzed from the coefficient of precision (mean of the differences observed between the two observations), Lin's concordance coefficient (Lin's CC) and the Bland and Altman method using an Excel add-in [15] in 20 infants. Repeatability was checked blind after an interval of two months for age, sex and the results previously obtained. Reproducibility was checked blind by two observers for age, sex and the results previously obtained. The values obtained for Lin's CC were interpreted according to Partik et al. [16]. Only results with a Lin's CC greater than 71 for repeatability and reproducibility were retained as we considered that their measurement error was acceptable (Table 2).

Table 2

| Interpretation of the value of the correlation |                         |  |  |  |  |  |
|--|-------------------------|--|--|--|--|--|
| interpretation of the value of the correlation |                         |  |  |  |  |  |
| coefficient                                    |                         |  |  |  |  |  |
| < 0.50   | Unacceptable            |  |  |  |  |  |
| 0.51-0.60                                      | Poor                    |  |  |  |  |  |
| 0.61-0.70                                      | Mediocre                |  |  |  |  |  |
| 0.71-0.80                                      | Moderately satisfactory |  |  |  |  |  |
| 0.81-0.90                                      | Fairly good             |  |  |  |  |  |
| 0.91-0.95                                      | Very good               |  |  |  |  |  |
| > 0.95   | Excellent               |  |  |  |  |  |

The normality of distribution of each of the biometric variables and the equality of variance were studied using the Shapiro-Wilk test. The null hypothesis of this test is that the population is normally distributed. If the p-value is lower than the alpha level selected, the null hypothesis is rejected [17].

For normally distributed variables, the correlation between the different variables was studied using the Pearson correlation (multivariate statistical test applied to a series of quantitative variables). Growth curves were established for the variables correlated with age.

A correlation between sex and the different variables analyzed was sought using an ANOVA test.

For all tests, the alpha risk was set at 0.05.

## 3. Results

# **Study population**

The retrospective cohort included 47 infants (22 girls, 25 boys).

# Measurement reliability

Repeatability and reliability were studied in 20 infants selected at random (11 boys, 9 girls).

The precision coefficient was low.

Lin's correlation coefficient was used to select the variables presented in Table 3.

Table 3

|                                | Reproducibility: agreement   |
|--------------------------------|--|
| moderately satisfactory (0.74) | fairly good (0.90)   |
| moderately satisfactory (0.90  | fairly good (0.83 and 0.88)  |
| and0.81)                       |  |
| very good (0.92)               | fairly good (0.81)   |
| very good (0.95)               | fairly good (0.85)   |
| fairly good (0.84)             | moderately satisfactory (0.78)   |
| moderately satisfactory (0.76) | moderately satisfactory (0.76)   |
| very good (0.95)               | very good (0.95)   |
| very good (0.93 and 0.91)      | Very good and fairly good (0.91  |
|                                | et 0.85)   |
| moderately satisfactory (0.78) | moderately satisfactory (0.78)   |
| Fairly good (0.83)             | moderately satisfactory (0.78)   |
|                                |  |
| excellent (0.99)               | excellent (0.98)   |
| very good (0.91)               | very good (0.92)   |
|                                | moderately satisfactory (0.90 and 0.81)  very good (0.92)  very good (0.95)  fairly good (0.84)  moderately satisfactory (0.76)  very good (0.95)  very good (0.93 and 0.91)  moderately satisfactory (0.78)  Fairly good (0.83)  excellent (0.99) |

The difference between the two measurements (intra and interobserver) were normally distributed and 95% of the values of measurement differences were within the limits of agreement. Bias (mean of the differences) was low. Among the variables retained, regarding inter-observer variability, none was not within the confidence

interval of bias for measurement of petrous pyramid height and vermis height: there was a systematic difference and bias was therefore significant.

# **Conventional morphometrics**

All variables were normally distributed.

#### Correlation between sex and the variables studied

None of the variables studied showed a significant difference according to sex.

# Correlation between age and the variables studied

Pearson's test showed a correlation between age and all the variables studied. The principal findings were a strong correlation between age and crown-heel length (p<0.01, R=0.855), head circumference (p<0.01, R=0.799), maximum length (p<0.01, R=0.747) and width (p<0.001, R=0.738) of the pars basilaris, petrous pyramid height (p<0.01, R=0.714), bone biparietal diameter (p<0.01, R=0.788), brain biparietal diameter (p<0.01, R=0.807) and cerebellar variables: transverse cerebellar diameter (p<0.01, R=0.817), anteroposterior diameter and height of the vermis ((p<0.01, R=0.664 and R=0.677).

Notably, there was a negative correlation between age and the sphenoid angle (p<0.01, R=-0.353).

Growth curves were drawn for each of these variables (Fig. 1). There was positive linear growth during the first two years of life, with the exception of the interpetrous angle, which did not develop after 12 months. The value of the sphenoid angle decreased during the period studied.

#### Correlation between the cranial base and the brain

Overall, the correlations between the dimensions of the cranial base and the brain were very strong.

Head circumference was strongly correlated with crown-heel length (p<0.01, R=0.909), maximum length (p<0.01, R=0.841) and maximum width (p<0.01, R=0.843) of the pars basilaris, height of the petrous pyramid (p<0.01, R=0.718), occipitofrontal diameter (p<0.01, R=0.865), bone biparietal diameter (p<0.01, R=0.932) and brain biparietal diameter (p<0.01, R=0.926), transverse cerebellar diameter (p<0.01, R=0.870) and interpetrous angle (p<0.01, R=0.615).

Occipitofrontal diameter was very strongly correlated with the total length of the cranial base (p<0.01, R=0.911). Brain biparietal diameter was very strongly correlated with bone biparietal diameter (p<0.01,

R=0.988) and transverse cerebellar diameter (p<0.01, R=0.937). The relation between transverse cerebellar diameter and biparietal diameter showed a linear increase, reflecting relatively greater increase in cerebellar width than in biparietal diameter in the infant (Fig. 2).

The sphenoid angle was correlated with transverse cerebellar diameter (p=0.01, R=0.386), brain and bone biparietal diameter (p<0.05, R= -0.367 and -0.376, respectively) and also with maximum length (p<0.01, R=-0.381), and maximum width (p<0.01, R=-0.420) of the pars basilaris. The sphenoid angle was correlated with crown-heel length (p<0.02, R=-0.405).

The cephalic index (CI) corresponding to the ratio of brain biparietal diameter to occipitofrontal diameter showed relative widening of the brain (Fig. 3). According to the classification of Cohen and Maclean [18], for our cases, up to the age of 8 months the CI is between 81.0–85.4 and indicates a brachycephalic cranium. After 8 month the CI is superior at 85.5, indicated a hyperbrachycephalic cranium.

There was a strong correlation between transverse cerebellar diameter and maximum length (p<0.01, R=0.861) and maximum width (p<0.01, R=0.895) of the pars basilaris, as well as with the height (p<0.01, R=0.801) and volume (p<0.01, R=0.75) of the petrous pyramid.

There was also a strong correlation between the height of the vermis and maximum height (p<0.01, R=0.764) and maximum width (p<0.01, R=0.809) of the pars basilaris.

The interpetrous angle, corresponding to the opening of the posterior fossa, was correlated with occipitofrontal diameter (p<0.01, R=0.459), brain biparietal diameter (p<0.01, R=0.604), transverse cerebellar diameter, anteroposterior diameter and height of the vermis (p<0.01, R=0.355 and 0.494, respectively). It became stable after the first year of life.

Transverse cerebellar diameter was also strongly correlated with occipitofrontal diameter (p<0.01, R=0.851) and bone biparietal diameter (p<0.01, R=0.937). The length of the corpus callosum was correlated with occipitofrontal diameter (p<0.01, R=0.779).

## 4. Discussion

As far as we are aware, this study is the first to propose a joint approach to analysis of the cranial base and the brain during the first years of life while taking into account the possibility of sexual dimorphism.

It revealed that the growth dynamic of these two systems was strongly correlated and synchronous. Studies in variations of the angles enabled analysis of shape changes of the cranial base.

Lastly, our study is original in that it uses two types of data to allow an overall approach to ontogeny in a well-defined population of infants. We were thus able to study a population that was unique both by the size of the series and by our use of multiple data obtained by imaging (CT and MRI) and by autopsy.

# Study population and imaging techniques

Although our series was large compared with other studies, the population included in the SID protocol was heterogeneous in distribution but coherent with the epidemiological data for SID [11]. In France, cases of SID are managed in a reference center (specialized pediatric hospital department) following a protocol based on the recommendations of the High Authority for Health (Haute Autorité de la Santé, HAS). This protocol enables rapid medical investigations in order to identify a cause of death and includes, in particular, medical imaging and full autopsy. There was a marked predominance of infants aged under 10 months and an unbalanced sex ratio with a majority of boys. This led to non-representativity of certain age groups, limiting the scope of our conclusions for certain variables.

The infants we included were rigorously selected (only 80% of the initial population was retained) in order to exclude any disorder that might affect the biometric variables examined.

Our study interested a wide range of measurements obtained through the combined use of CT and MRI. This combination is, to the best of our knowledge, not used previous literature on its use for study of ontogeny, in the infant in particular. The combination of these two investigations enabled study of 15 validated quantitative variables (12 linear measurements and 3 angles) of both the brain and the cranial base.

This study is as yet the most voluminous collection of biometric data on the head segment.

## Measurement reliability

More than half (15 of 23) of the biometric variables included in our study were found to be reliable and valid after study of their reproducibility and repeatability. Error could be explained by the thickness of the MRI sections studied and by lack of precision in placing landmarks in this context where distances are 14often less than a millimeter.

McBride [19] proposed a descriptive scale for values of Lin's correlation coefficient in biology. It is considered as poor when below 0.90. However, in the medical domain this evaluation is arbitrary and unjustified. Partik et al. [16] proposed threshold values that are more pertinent for medical research, and notably for medical imaging.

## Metric approach

All biometric variables studied showed variance that increased during the first two years of life, with the exception of the interpretrous angle which remained stable after the 12<sup>th</sup> month of life. Brain biparietal diameter, transverse cerebellar diameter, crown-heel length and head circumference were the variables that were most strongly correlated with age. Our approach to study of brain growth and of elements of the cranial base yielded growth curves that may be used to assess normal development in the infant. However, the relatively small size of our sample did not enable us to establish true norms for growth or speed of growth.

We demonstrated that overall, the correlations between the brain and the cranial base were strong in the infant. We confirmed certain findings of the literature [20,21], notably with regard to the relation between age and closure of the sphenoid angle after birth. Like Lieberman, we showed a negative correlation between closure of the sphenoid angle and crown-heel length [21, 22]. We also showed that closure of sphenoid angle and opening of the interpetrous angle was significantly correlated with widening of the cerebellum and with increase of biparietal diameter. The angle variation could be explained because of the rapid growth of the posterior fossa growth from birth to 3.58 years [23]. The rapid growth phase is explained by the absence of ossification of occipitomastoidal synchondrosis, petro-occipital synchondrosis and spheno-occipital synchondrosis [23].

In the adult, it has been shown that variations in the width, length and flexion of the cranial base are independent and that only the maximum width of the cranial base has a significant effect on overall cranial proportions [22]. However, in our study we underlined, in addition to these data, a close relationship between occipitofrontal diameter on the one hand and total length of the cranial base, sphenoid angle, biparietal diameter and transverse cerebellar diameter on the other.

Our study confirmed that measurements of the vermis and cerebellum were strongly correlated with age in the infant. Some neonatology teams use these measurements to assess gestational age [24]. We also observed that transverse cerebellar diameter (representing cerebellar growth in width), which is part of fetal development after 28 weeks amenorrhea [1,25], increased faster than biparietal diameter (growth in width of the cerebral hemispheres).

Several authors have shown with traditional morphometry that basioccipital dimensions evolve during foetal development [26-28] and its characteristics intensify with age [27]. The basioccipital are used not only in anatomy but also in biological anthropology: it can give an idea about the foetal age [29] and as our study showed about the post-natal age.

As Paetz et al, we find that the temporal bone grows study is also relevant during the first 2 years of life [30] in particular the petrous pyramid height, strongly correlated, with age and head circumference.

The cephalic index, based on MRI imaging of brain structures, showed in the infant a brain of brachycephalic type (width greater than length). These data are coherent with those of the literature [31,32] obtained from CT imaging.

The influence of sex on growth of the immature child aged less than 2 years was studied for each biometric variable. We found no sex-related difference in growth in any of the variables.

Few studies have looked for dimorphism of brain structures in this age group. Choe et al. [33] showed sexual dimorphism only in the caudate nucleus and the vermis. Their findings are not in agreement with ours. Matthews et al also showed craniofacial dimophism in 1 year-olds [34]. Moreover Rees and al. [35] described in the sudden infant death syndrome morphological facial characteristics, it could be extrapolate to the skull base. The sample size is very minimal once distributed into the particular age categories. The lack of a correlation between sex and the variables examined could merely reflect the small sample size.

With regard to the cranial base, some studies have shown sexual dimorphism of the anterior part of the cranial base (width) [36,37]. However, sexual dimorphism was not observed at the posterior level or the sphenoid angle [36].

## Interactions between growth of the cranial base and the brain

Our methodological approach did not enable us to determine whether growth of the brain was dependent on that of the cranial base, or the contrary. We did however demonstrate that there was a parallel dynamic of the one in relation to the other. In order to study a true cause-effect relationship, it would be necessary to examine a brain or bone disorder and observe its impact on adjacent structures. However, there is no sufficiently prevalent disorder that would enable pertinent study.

The effect of abnormalities of cranial suture closure on the brain is known [18], but congenital abnormalities of the cranial base are rare (encephalocele, dermoid cyst) [38], which limits the possibilities of extrapolating a possible interrelation of the systems in development.

#### 5. Conclusion

Our study demonstrated that growth of the cranial base and the brain develop in parallel during the first two years of life. It was carried out in a large series of rigorously selected infants and proposes an innovatory approach to the study of ontogenesis through the combined use of medical imaging, autopsy and

histopathological data. Lastly, it contributes to better knowledge of overall growth and normal development,

improving our understanding of pathological mechanisms that affect the brain and cranial segment.

Conflict of interest: The authors declare that they have no conflict of interest.

For this type of study formal consent is not required.

# **Declaration:**

Funding: none

Conflicts of interest/Competing interests: none

Ethics approval (include appropriate approvals or waivers): by the ethics committee of our establishment

CIL/AP-HM 2017-41

**Consent to participate**: Not applicable (death infant)

**Consent for publication** Not applicable (death infant)

# **Authors' contributions**

C. Delteil: •Protocol/project development; Data curation; Writing

E. Lesieur: Data curation

L. Tuchtan: Reviewing

A. Carballeira Alvarez: Resources; Software

C. Chaumoitre: Resources; Software

B. Saliba: Statistical analysis

P. Adalian: Project administration; Reviewing

M.D. Piercecchi-Marti: Supervision; Validation; Reviewing

### References

[1] G. Captier, L.J. Boë, P. Badin, A.M. Guihard-Costa, F. Canovas, J.C. Larroche, Geometrical growth

models of the fetal forebrain, cerebellum, brainstem and change of the cranial base angles during fetal

period, Morphologie 97 (2013) 38-47. doi: 10.1016/j.morpho.2012.12.002

[2] A.M. Guihard-Costa, F. Ramirez-Rozzi, Growth of the human brain and skull slows down at about 2.5

years old, Comptes Rendus Palevol. 3 (2004) 397-402. doi.org/10.1016/j.crpv.2004.03.004

15

- [3] J.N. Giedd, F.X. Castellanos, J.C. Rajapakse, D. Kaysen, A.C. Vaituzis, Y.C. Vauss, et al., Cerebral MRI of human brain development: Ages 4–18, Biol. Psychiatry 37 (1995) 657. doi.org/10.1016/0006-3223(95)94641-9.
- [4] R.K. Lenroot, N. Gogtay, D.K. Greenstein, E.M. Wells, G.L. Wallace, L.S. Clasen, et al., Sexual dimorphism of brain developmental trajectories during childhood and adolescence, Neuroimage 36 (2007) 1065–1073. doi: 10.1016/j.neuroimage.2007.03.053
- [5] R.K. Lenroot, J.N. Giedd, Brain development in children and adolescents: insights from anatomical magnetic resonance imaging, Neurosci. Biobehav. Rev. 30 (2006) 718–729. doi: 10.1016/j.neubiorev.2006.06.001.
- [6] C. Garel, MRI of the Fetal Brain: Normal Development and Cerebral Pathologies, Springer-Verlag, Berlin, Heidelberg, 2004.
- [7] R. Calandrelli, G. D'Apolito, M. Panfili, G. Zampino, T. Tartaglione, C. Colosimo, Costello syndrome: Analysis of the posterior cranial fossa in children with posterior fossa crowding, Neuroradiol. J. 28 (2015) 254–258. doi:10.1177/1971400915592549.
- [8] J.M. Walsh, L.W. Doyle, P.J. Anderson, K.J. Lee, Cheong JL, Moderate and late preterm birth: effect on brain size and maturation at term-equivalent age, Radiology 273 (2014) 232–240. doi: 10.1148/radiol.14132410.
- [9] S.N. Tich, P.J. Anderson, R.W. Hunt, K.J. Lee, L.W. Doyle, T.E. Inder, Neurodevelopmental and perinatal correlates of simple brain metrics in very preterm infants, Arch. Pediatr. Adolesc. Med. 5 (2011) 216–222. doi: 10.1001/archpediatrics.2011.9.
- [10] F. James Rohlf, L.F. Marcus, A revolution morphometrics, Trends Ecol. Evol. 8 (1993) 129–132. doi: 10.1016/0169-5347(93)90024-J.
- [11] Haute Autorité de Santé (HAS), Prise en charge en cas de mort inattendue du nourrisson (moins de 2 ans), Haute Autorité de Santé; 2007 p. 46.
- [12] I. G. Fazekas, F. Kósa, Forensic Fetal Osteology, Akadémiai Kiadó, Budapest, 1978. fourth ed., Longman, New York, 2000.
- [13] D. Vatansever, V. Kyriakopoulou, J.M. Allsop, M. Fox, A. Chew, J.V. Hajnal, et al., Multidimensional analysis of fetal posterior fossa in health and disease, Cerebellum 12 (2013) 632–644. doi: 10.1007/s12311-013-0470-2.

- [14] R Development Core Team, R: A language and environment for statistical computing. [Internet], R Foundation for Statistical Computing, Vienna, Austria, 2005. Available from: http://www.R-project.org.
- [15] L. Desquilbet, Répétabilité, reproductibilité, et concordance de méthodes de mesure, version 9. 2019. Available at https://loicdesquilbet.wixsite.com/biostat epidemio/documents]
- [16] B.L. Partik, A. Stadler, S. Schamp, A. Koller, M. Voracek, G. Heinz, et al., 3D versus 2D ultrasound: accuracy of volume measurement in human cadaver kidneys, Invest. Radiol. 37 (2002) 489–495. doi: 10.1097/01.RLI.0000023573.59066.43.
- [17] S.S. Shapiro, M.B. Wilk, An analysis of variance test for normality (complete samples), Biometrika 52 (1965) 591–611. doi:org/10.1093/biomet/52.3-4.591.
- [18] M.M. Cohen, R.E. MacLean, Craniosynostosis: Diagnosis, Evaluation, and Management, second ed.,
  Oxford University Press, New York, Oxford, 2000. Available from:
  http://trove.nla.gov.au/version/39761935
- [19] McBride GB. A proposal for strength-of-agreement criteria for Lin's Concordance Correlation Coefficient. 2005 NIWA Client Report: HAM2005-062. <a href="http://www.niwa.co.nz">http://www.niwa.co.nz</a>
- [20] I.S. Watanabe, M.C. Madeira, I.S. Watanabe, M.C. Madeira, [The clivus-sphenoidale angle in children (postnatal flexion of the cranial base)], Rev. Bras. Pesqui. Med. Biol. 10 (1977) 325–329.
- [21] D.E. Lieberman, R.C. McCarthy, The ontogeny of cranial base angulation in humans and chimpanzees and its implications for reconstructing pharyngeal dimensions, J. Hum. Evol. 36 (1999) 487–517.
- [22] D.E. Lieberman, O.M. Pearson, K.M. Mowbray, Basicranial influence on overall cranial shape, J. Hum. Evol. 38 (2000) 291–315.
- [23] G. Coll, J.-J. Lemaire, F. Di Rocco, I. Barthélémy, J.-M. Garcier, E. De Schlichting, et al. Human foramen magnum area and posterior cranial fossa volume growth in relation to cranial base synchondrosis closure in the course of child development Neurosurgery, 79 (2016), pp. 722-735, 10.1227/NEU.000000000001309.
- [24] E. Co, T.N. Raju, O. Aldana, Cerebellar dimensions in assessment of gestational age in neonates, Radiology 181 (1991) 581–585.
- [25] N. Jeffery, F. Spoor, Brain size and the human cranial base: a prenatal perspective, Am. J. Phys. Anthropol. 118 (2002) 324–340.

- [26] Scheuer L, MacLaughlin-Black S. Age estimation from the pars basilaris of the fetal and juvenile occipital bone. International Journal of Osteoarchaeology. 1994;4(4):377-380. https://doi.org/10.1002/oa.1390040412
- [27] Scheuer L, Black S, Cunningham C (2000) Developmental Juvenile Osteology. San Diego: Elsevier Academic Press.
- [28] Nagaoka T, Kawakubo Y, Hirata K. Estimation of fetal age at death from the basilar part of the occipital bone. Int J Legal Med. 2012;126(5):703-711. doi:10.1007/s00414-012-0718-2
- [29] Niel M, Chaumoître K, Corny J, Lalys L, Adalian P. Maturation of the human foetal basioccipital: quantifying shape changes in second and third trimesters using elliptic Fourier analysis. J Anat. 2019;235(1):34-44. doi:10.1111/joa.12997
- [30] Paetz P, Goetz GF, Lanfermann H, Giesemann AM. The developing temporal bone: computed tomography measurements and assessment of suture closure from birth to 18 years of age. Surg Radiol Anat. 2017;39(6):663-671. doi:10.1007/s00276-016-1786-7
- [31] P. Hummel, D. Fortado, Impacting infant head shapes, Adv. Neonatal Care 5(2005) 329–340. doi: 10.1016/j.adnc.2005.08.009
- [32] W. Likus, G. Bajor, K. Gruszczyńska, J. Baron, J. Markowski, M. Machnikowska-Sokołowska, et al., Cephalic index in the first three years of life: study of children with normal brain development based on computed tomography, ScientificWorldJournal 2014;2014:502836. doi: 10.1155/2014/502836.
- [33] M. S. Choe, S. Ortiz-Mantilla, N. Makris, M. Gregas, J. Bacic, D. Haehn, et al., Regional Infant brain development: an MRI-based morphometric analysis in 3 to 13 month olds, Cereb. Cortex 23 (2013) 2100–2117. doi: 10.1093/cercor/bhs197.
- [34] Matthews H, Penington T, Saey I, Halliday J, Muggli E, Claes P. Spatially dense morphometrics of craniofacial sexual dimorphism in 1-year-olds. J Anat. 2016;229(4):549-559. doi:10.1111/joa.12507
- [35] Rees, K., Wright, A., Keeling, J. W., & Douglas, N. J. (1998). Facial structure in the sudden infant death syndrome: case-control study. BMJ (Clinical research ed.), 317(7152), 179–180. https://doi.org/10.1136/bmj.317.7152.179
- [36] W.J. Ursi, C.A. Trotman, J.A. McNamara, R.G. Behrents, Sexual dimorphism in normal craniofacial growth, Angle Orthod. 63 (1993) 47–56.

[37] E. Bulygina, P. Mitteroecker, L. Aiello, Ontogeny of facial dimorphism and patterns of individual

development within one human population, Am. J. Phys. Anthropol. 131 (2006) 432-443. doi:

10.1002/ajpa.20317

[38] A. Di Ieva, E. Bruner, T. Haider, L.F. Rodella, J.M. Lee, M.D. Cusimano, et al., Skull base

embryology: a multidisciplinary review, Childs Nerv. Syst. 30 (2014) 991-1000. doi: 10.1007/s00381-014-

2411-x.

Légends

Table 1 : Conventional morphometric measurement protocols

Table 2: Interpretation of the value of the correlation coefficient

Table 3: Lin's correlation coefficient

**Figures** 

Fig. 1. Growth curves according to age (in months) for the (a) sphenoid angle, (b) interpetrous angle, (c)

anteroposterior diameters: OFD (black) and total cranial base (gray), (d) biparietal diameters (black and

mid gray) and transverse cerebellar diameter (clear gray), (e) anteroposterior height (black) and height

(gray) of the vermis, (f) height of the petrous pyramid, (g) maximum length (black) and width (clear gray)

of the pars basilaris, (h) body height (i) weight, (j) head circumference. The circle corresponds to the girls

and the triangles to the boys.

Fig. 2. Relation between transverse cerebellar diameter and biparietal diameter according to age group:

TDC/brain BPD x 100 versus age in months. The circle corresponds to the girls and the triangles to the

boys.

Fig. 3. Cephalic index according to age group (in months). The circle corresponds to the girls and the

triangles to the boys

19























