The morphometric measurement of the brainstem in Turkish healthy subjects according to age and sex

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**Background:** This paper determined the morphometric measurements of the brainstem including mesencephalon, pons and medulla using magnetic resonance imaging (MRI) in Turkish healthy population.

**Materials and methods:** Two hundred sixty-three (263; 158 females and 105 males) subjects aged from 18 to 65 years were included in this study. The measurements were taken from subjects having brain MRI in the Radiology Department. Statistical analysis was done by SPSS 21.00 package programme. ANOVA test and $\chi^2$ test were used to determine the relation between measurements and age and sex groups. The $p < 0.05$ value was considered as significant.

**Results:** The overall means and standard deviations of the measurements were: pons anteroposterior diameter, 15.41 ± 1.27 mm; pons vertical diameter, 22.02 ± 2.07 mm; mesencephalon anteroposterior diameter 9.39 ± 1.00 mm; mesencephalon vertical diameter, 15.20 ± 1.53 mm; distance between the interpeduncular fissure and aqueduct, 11.72 ± 1.58 mm; distance from cerebral peduncles to aqueduct, 13.64 ± 1.66 mm; anterior surface of the pons midway between the mesencephalon and medulla to the fourth ventricular floor, 21.62 ± 1.64 mm; the shortest anteroposterior diameter of the medulla at the pontomedullary junction, 13.46 ± 1.28 mm, and the shortest anteroposterior diameter of the medulla at the medullospinal junction, 10.24 ± 1.43 mm in females, respectively, whereas the corresponding values were 15.58 ± 1.53 mm; 22.64 ± 2.35 mm; 9.37 ± 1.66 mm; 15.64 ± 1.52 mm; 11.14 ± 1.31 mm; 13.01 ± 1.30 mm; 21.97 ± 1.65 mm; 13.47 ± 1.19 mm; 9.91 ± 1.35 mm in males, respectively. There were significant differences in some parameters such as pons vertical diameter, mesencephalon vertical diameter, distance between the interpeduncular fissure and aqueduct, and distance between cerebral peduncles to aqueduct between sexes.

**Conclusions:** The brainstem dimensions of healthy population provide important and useful knowledge in terms of comparison of abnormalities clinically. These data may be valuable for the representatives of clinical disciplines. (Folia Morphol 2020; 79, 1: 36–45)

**Key words:** brainstem morphometry, age and sex changes, Turkish population
INTRODUCTION

The brainstem which is divided in three parts; mesencephalon, pons and medulla, is one of the critical small anatomic structures and it includes many important cranial nerve nuclei, tracts and fibres [4, 5, 29]. The brainstem directs vital functions including breathing, blood pressure, heart rate, swallowing, movement ability, sensation of face, mouth, tongue, extremities [10]. Moreover, it is surrounded by the clivus anteriorly; by the temporal bone petrosus part laterally; by the cerebellum posteriorly; by the diencephalon superiorly [4, 5]. Furthermore, the brainstem structures play an important role, because brain functions including motor, sensory, sympathetic, and parasympathetic are integrated and pass through the medulla, pons and mesencephalon [5]. There can be difficulty in surgery because of brainstem complex structure [5, 10]. Additionally, surgical procedures of the brainstem lesions make difficulties due to morbidity and mortality risk. Moreover, brainstem gliomas usually are settled into pons, sometimes they originate from mesencephalon or medulla [13]. Therefore, it requires the knowledge of detailed functional and topographical anatomy and microsurgical information [6, 10, 29]. Imaging techniques such as computed tomography (CT) and magnetic resonance imaging (MRI) provide an assessment opportunity of intracranial structures [1, 3]. It has been reported CT were insufficient in revealing atrophic changes in the brainstem region [23]. Therefore, the MRI using was more suitable for examination and it has become possible to achieve reliable and accurate neuroanatomical views [2, 23]. In this respect, the MRI of brain stem regions gives critical detailed knowledge about brain related with normal aging or in neuromuscular diseases [7, 8, 11, 15, 28], and normal brain growth and atrophy [20]. Also, MRI determines shrinkage in the brainstem and proves to be a strong predictor of disease progression [1]. On the other hand, the studies of MRI have associated with mesencephalon morphology in many neurologic disorders like Parkinson disease (PD) and Wilson disease (WD). Therefore, the morphometric data performed using MRI, may explain the underlying neurochemical or pathologic situations [20]. Previous studies investigated age- and sex-related changes in the brainstem [7, 8, 15, 17, 21, 24]. In that studies results of age- and sex-related differences in brainstem may give different results in general [7, 8, 15, 17, 21]. However, the results of age-dependent brainstem measurement in studies are still controversial. Besides, the brainstem linear measurements with MRI were rare in Turkish healthy population in the literature [1, 2, 20]. The linear measurements can be performed more quickly, are practical and suitable for work and do not require additional hardware and software [23]. On the other hand, the interpeduncular fissure and cerebral peduncles’ localisation surface could be difficult. Because of the volume between brain and surrounding cerebrospinal fluid, these measurements were difficult for identification of the interpeduncular fissure [14]. This is the first study considering the morphometric measurements and ratios of the brainstem using MRI in detailed to investigate age and sex differences in the brainstem region in Turkish healthy population. In order to understand the pathologies of brainstem structures and the effects of neurodegenerative diseases on brain stem, measurements of age and sex differences in normal healthy individuals should be well known. Therefore, in this study, it was aimed to determine the age and sex differences of brainstem structures in healthy Turkish individuals using MRI.

MATERIALS AND METHODS

This study was carried out on 263 healthy adult subjects (158 females; 105 males) aged 18–65 years over a period of 2 years between January 2017 and 2019. All the test procedures were approved by Ethics Committee. Cranial MRI findings were evaluated by radiologists and anatomists. Moreover, inclusion criteria for adult subjects selected by criteria of optimal health were: no history of receiving a diagnosis of cancer, hemiplegia, intracranial tumoural mass, any neurologic disease (such as multiple sclerosis, amyotrophic lateral sclerosis or dementia, PD), having surgical operation related to brain, and psychiatric illness.

Magnetic resonance imaging was performed using a 1.5 T MRI system (Siemens; Essen, Erlangen, Germany). Brain MRI protocol including axial T2-weighted turbo spin echo (TR: 3600, TE: 87 ms; slice thickness: 5 mm; gap: 1.5 mm) and sagittal T2-weighted spin echo (TR: 3600, TE: 87 ms; slice thickness: 5 mm; gap: 1.5 mm) was used. The measurements were performed from digital MRI images at a hospital using caliper function with \( \times 2 \) magnification. Using the midsagittal T2-weighted spin echo image, the following parameters of brainstem dimensions were evaluated [8, 12, 14, 29]:
— DIPFA — the distance between the interpeduncular fissure and the aqueduct;
— DCPA — the distance from the anterior surface of the cerebral peduncles to the aqueduct;
— DPMM — the distance between the anterior surface of the pons midway between the mesencephalon and medulla to the fourth ventricular floor;
— MOPMJ — the shortest anteroposterior diameter of the medulla at the pontomedullary junction;
— MOMSJ — the shortest anteroposterior diameter of the medulla at the medullospinal junction;
— PAP — pons anteroposterior diameter as the distance between two points perpendicular to the midline along the tangent lateral to the pons;
— PVD — pons vertical diameter was measured midpoint between its upper and lower borders and perpendicular to its long axis to the fourth ventricle;
— MBAPD — mesencephalon anteroposterior diameter at a midline level on a sagittal section;
— MBVD — mesencephalon vertical diameter was measured across the mesencephalon to the tectum.

The ratio values of the DPMM/MOPMJ, DPMM/MOMSJ, DPMM/DIPFA, DPMM/DCPA and midbrain/pons calculated.

The data were divided into two groups: healthy adult female and male subjects (Tables 1, 2). Furthermore the data were divided also into five groups according to age:
— group 1 — subjects aged between 18 and 30 years were assigned to group 1;
— 31–40 years — group 2;
— 41–50 years — group 3;
— 51–60 years — group 4;
— and 61–65 years — group 5 (Tables 3–6).

The measurements were made on the computer screen with an electronic calliper and estimations were expressed as millimetres.

### Table 1. The distribution of Brainstem diameters according to sex

| Parameters                                      | Females (n = 158)          | Males (n = 105)         | Total          | P    |
|------------------------------------------------|---------------------------|------------------------|----------------|------|
| Pons anteroposterior diameter (PAP)            | 15.41 ± 1.27 (12.30–22.60) | 15.58 ± 1.53 (12.20–19.30) | 15.47 ± 1.38 (12.20–22.60) | 0.326 |
| Pons vertical diameter (PVD)                   | 22.02 ± 2.07 (13.80–29.00) | 22.64 ± 2.35 (13.70–27.60) | 22.27 ± 2.20 (13.70–29.00) | 0.025 |
| Mesencephalon anteroposterior diameter (MBAPD) | 9.39 ± 1.00 (6.50–12.30)   | 9.37 ± 1.66 (7.30–13.10)  | 9.38 ± 1.07 (6.50–13.10)   | 0.854 |
| Mesencephalon vertical diameter (MBVD)         | 15.20 ± 1.53 (11.70–19.40) | 15.64 ± 1.52 (11.70–19.10) | 15.38 ± 1.54 (11.70–19.40) | 0.022 |
| Distance between the interpeduncular fissure and aqueduct (DIPFA) | 11.72 ± 1.58 (8.00–16.90) | 11.14 ± 1.31 (8.00–14.70)  | 11.49 ± 1.51 (8.00–16.90)   | 0.002 |
| Distance from cerebral peduncles to aqueduct (DCPA) | 13.64 ± 1.66 (9.30–18.50) | 13.01 ± 1.30 (9.30–17.10)  | 13.39 ± 1.55 (9.30–18.50)   | 0.001 |
| Anterior surface of the pons midway between the mesencephalon and medulla to the fourth ventricular floor (DPMM) | 21.62 ± 1.64 (17.50–28.60) | 21.97 ± 1.65 (18.10–29.90)  | 21.76 ± 1.65 (17.50–29.90)   | 0.100 |
| The shortest anteroposterior diameter of the medulla at the pontomedullary junction (MOPMJ) | 13.46 ± 1.28 (10.00–18.00) | 13.47 ± 1.19 (10.90–18.10)  | 13.46 ± 1.24 (10.00–18.10)   | 0.917 |
| The shortest anteroposterior diameter of the medulla at the medullospinal junction (MOMSJ) | 10.24 ± 1.43 (7.20–15.10)  | 9.91 ± 1.35 (7.30–13.40)   | 10.11 ± 1.40 (7.20–15.10)    | 0.064 |

### Table 2. The sex related changes of brainstem ratio measurements

| Groups                          | Females (n = 158)          | Males (n = 105)         | Total          | P    |
|---------------------------------|---------------------------|------------------------|----------------|------|
| DPMM/MOPMJ                     | 1.617 ± 0.156 (1.15–2.04)  | 1.639 ± 0.148 (1.23–2.14)  | 1.626 (1.15–2.14) | 0.261 |
| DPMM/MOMSJ                     | 2.146 ± 0.289 (1.46–3.01)  | 2.253 ± 0.330 (1.76–3.78)  | 2.162 (1.15–2.14) | 0.006 |
| DPMM/DIPFA                     | 1.870 ± 0.228 (1.34–2.89)  | 1.993 ± 0.225 (1.45–2.56)  | 1.626 (1.15–2.14) | < 0.001 |
| DPMM/DCPA                      | 1.602 ± 0.184 (1.26–2.14)  | 1.700 ± 0.170 (1.34–2.23)  | 1.626 (1.15–2.14) | < 0.001 |
| Midbrain/Pons                   | 0.613 ± 0.074 (0.44–0.94)  | 0.606 ± 0.087 (0.40–0.84)  | 0.610 (0.40–0.94) | 0.505 |

Abbreviations — see text
Table 3. The distribution of the brainstem diameters according to age groups of healthy females and males

| Groups                                              | Group 1 (n = 91) | Group 2 (n = 99) | Group 3 (n = 42) | Group 4 (n = 23) | Group 5 (n = 8) | P      |
|-----------------------------------------------------|------------------|------------------|------------------|------------------|----------------|--------|
| Pons anteroposterior diameter (PAP)                 | 15.38 ± 1.31     | 15.67 ± 1.46     | 15.24 ± 1.41     | 15.47 ± 1.18     | 15.35 ± 1.48    | 0.440  |
| (12.60–18.70)                                       | (12.30–22.60)    | (13.10–18.10)    | (12.20–17.10)    | (13.20–17.30)    |                |        |
| Pons vertical diameter (PVD)                        | 22.51 ± 1.98     | 22.03 ± 2.24     | 22.00 ± 2.33     | 22.60 ± 2.56     | 22.60 ± 2.34    | 0.410  |
| (14.30–27.60)                                       | (13.80–27.20)    | (13.70–24.60)    | (15.10–29.00)    | (19.80–27.40)    |                |        |
| Mesencephalon anteroposterior diameter (MBAPD)       | 9.63 ± 1.08      | 9.34 ± 1.05      | 9.11 ± 1.08      | 9.19 ± 0.97      | 9.09 ± 0.92     | 0.052  |
| (7.30–13.10)                                        | (6.50–12.90)     | (7.00–12.00)     | (7.40–11.40)     | (7.40–10.00)     |                |        |
| Mesencephalon vertical diameter (MBVD)              | 15.43 ± 1.52     | 15.52 ± 1.48     | 15.05 ± 1.52     | 15.43 ± 1.78     | 14.61 ± 1.73    | 0.297  |
| (11.70–18.80)                                       | (11.70–19.40)    | (11.80–18.90)    | (13.00–18.40)    | (11.90–17.20)    |                |        |
| Distance between the interpeduncular fissure and     | 11.89 ± 1.48     | 11.41 ± 1.56     | 11.18 ± 1.50     | 11.09 ± 1.15     | 10.58 ± 1.05    | 0.010  |
| aqueduct (DIPFA)                                    | (8.80–15.70)     | (8.70–16.90)     | (8.00–14.60)     | (9.20–13.00)     | (9.10–12.00)    |        |
| Distance from cerebral peduncles to aqueduct (DCPA) | 13.85 ± 1.60     | 13.25 ± 1.61     | 13.11 ± 1.44     | 12.96 ± 0.98     | 12.56 ± 1.01    | 0.006  |
| (9.30–18.50)                                        | (9.30–18.00)     | (10.50–17.00)    | (10.70–14.30)    | (11.30–14.30)    |                |        |
| Anterior surface of the pons midway between the     | 21.64 ± 1.46     | 21.90 ± 1.66     | 21.95 ± 2.11     | 21.71 ± 1.21     | 20.60 ± 1.73    | 0.220  |
| mesencephalon and medulla to the fourth ventricular | (18.00–25.90)    | (17.50–29.90)    | (17.50–26.60)    | (19.60–23.60)    | (18.20–22.80)  |        |
| floor (DPMM/MOPMJ)                                  |                 |                  |                  |                  |                |        |
| The shortest anteroposterior diameter of the        | 13.38 ± 1.13     | 13.56 ± 1.13     | 13.60 ± 1.59     | 13.20 ± 1.33     | 13.20 ± 1.64    | 0.570  |
| medulla at the pontomedullary junction (MOPMJ)       | (10.90–16.90)    | (10.20–18.20)    | (10.00–18.00)    | (10.80–15.30)    | (10.70–15.20)  |        |
| The shortest anteroposterior diameter of the        | 10.40 ± 1.54     | 9.98 ± 1.30      | 9.90 ± 1.41      | 9.69 ± 1.07      | 10.66 ± 1.20    | 0.059  |
| medulla at the medullospinal junction (MOMSJ)        | (7.20–15.10)     | (7.30–13.40)     | (7.60–12.60)     | (7.70–12.10)     | (9.40–12.70)   |        |

Table 4. The brainstem ratio measurements according to age groups of healthy females and males

| Groups                                              | Group 1 (n = 91) | Group 2 (n = 99) | Group 3 (n = 42) | Group 4 (n = 23) | Group 5 (n = 8) | P      |
|-----------------------------------------------------|------------------|------------------|------------------|------------------|----------------|--------|
| DPMM/MOPMJ                                          | 1.625 ± 0.138    | 1.623 ± 0.151    | 1.628 ± 0.180    | 1.658 ± 0.167    | 1.574 ± 0.161  | 0.736  |
| (1.24–1.96)                                         | (1.23–2.14)      | (1.15–2.01)      | (1.40–2.00)      | (1.34–1.74)      |                |        |
| DPMM/MOMSJ                                          | 2.119 ± 0.297    | 2.227 ± 0.311    | 2.255 ± 0.343    | 2.264 ± 0.248    | 1.942 ± 0.163  | 0.005  |
| (1.46–2.82)                                         | (1.63–3.78)      | (1.62–3.03)      | (1.79–2.70)      | (1.73–2.22)      |                |        |
| DPMM/DIPFA                                         | 1.840 ± 0.203    | 1.946 ± 0.249    | 1.986 ± 0.247    | 1.976 ± 0.201    | 1.960 ± 0.205  | 0.002  |
| (1.34–2.30)                                         | (1.36–2.56)      | (1.58–2.89)      | (1.56–2.39)      | (1.68–2.20)      |                |        |
| DPMM/DCPA                                          | 1.577 ± 0.163    | 1.671 ± 0.203    | 1.685 ± 0.170    | 1.686 ± 0.166    | 1.645 ± 0.140  | 0.002  |
| (1.26–1.98)                                         | (1.28–2.23)      | (1.36–2.10)      | (1.42–2.14)      | (1.46–1.83)      |                |        |
| Midbrain/Pons                                       | 0.630 ± 0.078    | 0.599 ± 0.077    | 0.601 ± 0.076    | 0.599 ± 0.096    | 0.594 ± 0.074  | 0.070  |
| (0.44–0.83)                                         | (0.46–0.94)      | (0.40–0.78)      | (0.46–0.84)      | (0.40–0.94)      |                |        |

Abbreviations — see text

Statistical analysis

The SPSS 21.0 programme was used for statistical analysis of the measurement results. From these measurements, means, standard deviations (SD), minimum (min) and maximum (max) values were calculated. In all statistical analyses; p value under 0.05 was considered statistically significant.

RESULTS

The values of minimum, maximum, mean and SD of diameters of the mesencephalon (distance between the interpeduncular fissure and aqueduct [DIPFA] and distance from cerebral peduncles to aqueduct [DCPA]), pons (anterior surface of the pons midway between the mesencephalon and medulla to the fourth ventricular floor [DPMM], and medulla (the shortest anteroposterior diameter of the medulla at the pontomedullary junction [MOPMJ], and the shortest anteroposterior diameter of the medulla at the medullospinal junction [MOMSJ]), and brain stem ratio measured in 263 healthy subjects (158 females and 105 males) were shown in Tables 1–6. There are significant differences in some measurements such as pons vertical diameter (PVD); mesencephalon ver-
DIPFA measurements such as DIPFA \((p = 0.009)\), DCPA \((p = 0.034)\), \(\text{DPMM/DIPFA} \) and \(\text{DPMM/DCPA} \) decreased \((p = 0.034)\), \(\text{DPMM/DIPFA} \) and \(\text{DPMM/DCPA} \) measurements increased \((p = 0.012)\), and \(\text{DPMM/DIPFA} \) \((p = 0.009)\). Also, DIPFA values were decreased with increased age. The lowest value was seen in the seventh decade (group 5). In females, the mesencephalon (midbrain) anteroposterior (AP) diameter value showed the reduction until the age of 61 years and after the age of 60 years this value increased. \(\text{DPMM/DIPFA} \) and \(\text{DPMM/DCPA} \) measurements increased until the age of 60 years, while after the age of the 61 this value showed decrease (Table 5). In males, there was no statistically significant difference in all measurements. The highest value of PAP was seen in the age of 31–40 years, while the lowest value was in the age of 61 years and after (Table 6).

**DISCUSSION**

Brainstem, which is a small area, has vital importance and contains many functionally primary nuclei, fibres, neurons and nerve tracts [29]. The brainstem consists of mesencephalon, pons and medulla oblonga [5], and it controls many functions such as breathing, blood pressure, heart rate control, swal-

**Table 5. The distribution of the brainstem diameters and brainstem ratios according to age groups in females \((n = 158)\)**

| Parameters | Group 1          | Group 2          | Group 3          | Group 4          | Group 5          | P    |
|-----------|------------------|------------------|------------------|------------------|------------------|------|
| PAP       | 15.37 ± 1.16     | 15.43 ± 1.54     | 15.14 ± 1.21     | 15.66 ± 0.92     | 15.82 ± 0.56     | 0.705|
|           | (12.90–17.50)    | (12.30–22.60)    | (13.20–17.20)    | (13.60–17.10)    | (15.00–16.30)    |      |
| PVD       | 22.31 ± 1.96     | 21.35 ± 2.15     | 22.57 ± 1.29     | 22.20 ± 2.63     | 22.26 ± 1.85     | 0.079|
|           | (14.30–24.90)    | (13.80–26.90)    | (19.90–24.60)    | (15.10–29.00)    | (19.80–24.20)    |      |
| MBAPD     | 9.65 ± 0.93      | 9.29 ± 1.04      | 9.19 ± 1.15      | 9.08 ± 0.87      | 9.30 ± 0.76      | 0.117|
|           | (7.30–11.60)     | (6.50–12.30)     | (7.00–12.00)     | (7.40–10.50)     | (8.40–10.00)     |      |
| MBVD      | 15.59 ± 1.51     | 15.28 ± 1.52     | 14.40 ± 1.47     | 15.27 ± 1.57     | 15.22 ± 1.42     | 0.136|
|           | (11.70–18.80)    | (11.80–19.40)    | (11.80–17.60)    | (13.20–17.50)    | (13.30–17.20)    |      |
| DIPFA     | 12.23 ± 1.45     | 11.54 ± 1.70     | 11.50 ± 1.69     | 11.03 ± 1.22     | 10.66 ± 0.96     | 0.009|
|           | (9.50–15.70)     | (8.70–16.90)     | (8.00–14.60)     | (9.20–13.00)     | (9.50–12.00)     |      |
| DCPA      | 14.24 ± 1.51     | 13.35 ± 1.81     | 13.47 ± 1.70     | 12.92 ± 1.05     | 12.58 ± 1.08     | 0.003|
|           | (11.30–18.50)    | (9.30–18.00)     | (10.50–17.00)    | (10.70–14.30)    | (11.40–14.30)    |      |
| DPMM      | 21.73 ± 1.21     | 21.50 ± 1.64     | 22.00 ± 2.69     | 21.64 ± 1.18     | 19.98 ± 1.70     | 0.157|
|           | (18.00–25.50)    | (17.50–25.60)    | (17.50–26.60)    | (19.60–23.10)    | (18.20–22.30)    |      |
| MOPMJ     | 13.44 ± 1.16     | 13.43 ± 1.07     | 13.90 ± 1.67     | 13.24 ± 1.38     | 12.80 ± 1.49     | 0.363|
|           | (10.90–16.90)    | (10.20–16.30)    | (10.00–18.00)    | (10.80–15.30)    | (10.00–18.00)    |      |
| MOMSJ     | 10.62 ± 1.55     | 9.97 ± 1.31      | 10.30 ± 1.40     | 9.57 ± 1.10      | 10.38 ± 1.13     | 0.005|
|           | (7.20–15.10)     | (7.50–12.60)     | (8.10–12.60)     | (7.70–12.10)     | (9.40–12.20)     |      |
| DPMM/MOMSJ | 1.625 ± 0.133   | 1.608 ± 0.148    | 1.599 ± 0.211    | 1.649 ± 0.178    | 1.574 ± 0.189    | 0.783|
|           | (1.24–1.96)      | (1.39–2.04)      | (1.15–2.01)      | (1.40–2.00)      | (1.34–1.74)      |      |
| DPMM/MOPSJ | 2.087 ± 0.301   | 2.185 ± 0.069    | 2.160 ± 0.318    | 2.283 ± 0.233    | 1.935 ± 0.189    | 0.034|
|           | (1.46–2.82)      | (1.63–3.01)      | (1.62–3.01)      | (1.69–2.70)      | (1.73–2.22)      |      |
| DPMM/DIPFA | 1.798 ± 0.210   | 1.890 ± 0.218    | 1.937 ± 0.283    | 1.978 ± 0.193    | 1.884 ± 0.208    | 0.012|
|           | (1.34–2.30)      | (1.36–2.32)      | (1.63–2.89)      | (1.67–2.39)      | (1.68–2.11)      |      |
| DPMM/DCPA  | 1.540 ± 0.162   | 1.632 ± 0.201    | 1.634 ± 0.179    | 1.686 ± 0.170    | 1.594 ± 0.184    | 0.009|
|           | (1.26–1.98)      | (1.28–2.13)      | (1.36–2.10)      | (1.42–2.14)      | (1.26–2.14)      |      |
| Midbrain/Pons | 0.631 ± 0.076 | 0.606 ± 0.079    | 0.607 ± 0.063    | 0.582 ± 0.066    | 0.588 ± 0.046    | 0.090|
|           | (0.44–0.83)      | (0.46–0.94)      | (0.51–0.74)      | (0.46–0.76)      | (0.53–0.65)      |      |

Abbreviations — see text
lowing and make sounds [10]. Moreover, this part constitutes the crossing for descending and ascending tracts which links medulla spinalis to the top of the nervous system. These tracts may be obstructed due to demyelinating diseases, neoplasm or vascular disorder. Furthermore, nuclei of the glossopharyngeal, vagal, accessory and hypoglossal nerves emerge between inferior cerebellar peduncle and olivary nucleus in medulla oblongata [27]. The pons lies in the posterior cranial fossa and forms superior part of the 4th ventricle. Additionally, descending and ascending tracts, which are the most critical structures, pass from the pons. Also, the childhood astrocytomas are the most common tumour seen in brainstem [19, 27]. The mesencephalon is the shortest, rostral and superior part of the brainstem and lies at the junction of the middle and posterior cranial fossae. The cerebral aqueduct is the cavity of the mesencephalon. It is one of the narrower structures of the ventricular system and surrounding cerebrospinal fluid passes by the cerebral aqueduct between 3rd ventricle to the 4th ventricle and reaches subarachnoid space by foramina. Also, the oculomotor and trochlear nerves arise from mesencephalon and there are two important nuclei: substantia nigra and red nucleus [9, 19, 27]. Additionally, the brainstem external anatomy knowledge helps in surgical planning due to relation between the external and internal structures, and the anatomical reference points on the brainstem [2]. Also, the DPMM to MOPMJ ratio, and DPMM to MOMSJ ratio were found to be abnormally small in subjects having tumours localised in pons and mesencephalon. Additionally, brainstem measurements help in diagnosing of tumours or atrophy of this part of brain. The brain stem normal values abbreviations — see text

| Parameters     | Group 1 | Group 2 | Group 3 | Group 4 | Group 5 | P    |
|---------------|--------|--------|--------|--------|--------|------|
| PAP           | 15.39 ± 1.59 | 15.95 ± 1.34 | 15.34 ± 1.60 | 14.82 ± 1.85 | 14.57 ± 2.37 | 0.182 |
| (12.60–18.70) | (12.60–19.30) | (13.10–18.10) | (12.20–16.60) | (13.20–17.30) |
| PVD           | 22.94 ± 1.98 | 22.77 ± 2.13 | 21.43 ± 2.97 | 24.06 ± 1.85 | 23.70 ± 3.22 | 0.068 |
| (15.80–27.60) | (15.30–27.20) | (13.70–24.30) | (21.60–26.10) | (21.50–27.40) |
| MBAPD         | 9.60 ± 1.37 | 9.40 ± 1.07 | 9.03 ± 1.02 | 9.58 ± 1.29 | 8.73 ± 1.22 | 0.413 |
| (7.40–14.10)  | (7.40–12.90) | (7.30–10.90) | (8.00–11.40) | (7.40–9.80) |
| MBVD          | 15.53 ± 1.57 | 15.78 ± 1.40 | 15.70 ± 1.29 | 16.02 ± 2.52 | 13.60 ± 2.00 | 0.175 |
| (11.90–18.80) | (11.70–19.10) | (13.00–18.90) | (13.00–18.40) | (11.90–18.50) |
| DIPFA         | 12.17 ± 1.27 | 11.27 ± 1.40 | 10.87 ± 1.25 | 11.28 ± 0.99 | 10.43 ± 1.41 | 0.682 |
| (8.80–13.50)  | (8.90–14.70) | (9.30–13.50) | (9.90–12.60) | (9.10–11.90) |
| DCPA          | 13.01 ± 1.50 | 13.15 ± 1.35 | 12.76 ± 1.06 | 13.10 ± 0.77 | 12.53 ± 1.10 | 0.797 |
| (9.30–17.10)  | (10.90–16.20) | (10.90–14.90) | (11.90–14.00) | (11.30–13.40) |
| DPMM          | 21.44 ± 1.91 | 22.33 ± 1.59 | 21.91 ± 1.38 | 21.98 ± 1.43 | 21.63 ± 1.46 | 0.249 |
| (18.10–25.90) | (19.90–29.90) | (19.20–24.70) | (19.90–23.60) | (20.00–22.80) |
| MOPMJ         | 13.24 ± 1.05 | 13.71 ± 1.18 | 13.31 ± 1.24 | 13.06 ± 1.27 | 13.87 ± 1.97 | 0.366 |
| (10.90–15.60) | (11.30–18.10) | (11.40–16.70) | (11.40–14.90) | (11.60–15.20) |
| MOMSJ         | 9.94 ± 1.44 | 9.96 ± 1.31 | 9.49 ± 1.33 | 10.12 ± 0.95 | 11.13 ± 1.28 | 0.309 |
| (7.80–12.90)  | (7.30–13.40) | (7.60–12.50) | (9.00–11.10) | (10.10–12.70) |
| DPMM/MOPMJ    | 1.625 ± 0.150 | 1.638 ± 0.155 | 1.657 ± 0.141 | 1.690 ± 0.130 | 1.573 ± 0.138 | 0.788 |
| (1.41–1.96)   | (1.23–2.14) | (1.28–1.88) | (1.56–1.89) | (1.45–1.72) |
| DPMM/MOMSJ    | 2.189 ± 0.262 | 2.274 ± 0.349 | 2.351 ± 0.348 | 2.195 ± 0.319 | 1.954 ± 0.147 | 0.219 |
| (1.78–2.68)   | (1.76–3.78) | (1.62–3.03) | (1.79–2.62) | (1.80–2.08) |
| DPMM/DIPFA    | 1.931 ± 0.152 | 2.001 ± 0.268 | 2.04 ± 0.201 | 1.965 ± 0.254 | 2.087 ± 0.150 | 0.442 |
| (1.65–2.28)   | (1.45–2.56) | (1.58–2.33) | (1.58–2.20) | (1.92–2.20) |
| DPMM/DCPA     | 1.658 ± 0.137 | 1.715 ± 0.198 | 1.726 ± 0.154 | 1.685 ± 0.169 | 1.728 ± 0.037 | 0.603 |
| (1.46–1.95)   | (1.34–2.23) | (1.52–1.94) | (1.42–1.83) | (1.70–1.77) |
| Midbrain/Pons | 0.627 ± 0.084 | 0.593 ± 0.077 | 0.595 ± 0.089 | 0.662 ± 0.163 | 0.603 ± 0.068 | 0.268 |
| (0.49–0.80)   | (0.47–0.94) | (0.40–0.78) | (0.48–0.84) | (0.56–0.68) |

Abbreviations — see text
morphometric data provides the early detection of the disorders [7]. Koehler et al. [14] measured the distance between the interpeduncular fissure and the aqueduct (DIPFA), the distance from the anterior surface of the cerebral peduncles to the aqueduct (DCPA), the distance between the anterior surface of the pons midway between the mesencephalon and medulla to the fourth ventricular floor (DPMM), the shortest anteroposterior diameter of the medulla at the pontomedullary junction (MOPMJ), and the shortest anteroposterior diameter of the medulla at the medullospinal junction (MOMSJ), DPMM to MOPMJ ratio, and DPMM to MOMSJ ratio, DPMM to DIPFA ratio and DPMM to DCPA ratio in healthy subjects, subjects having glioma and having brain atrophy (Table 7). All measurements except DPMM/MOPMJ, DPMM/DIPFA and DPMM/DCPA values were higher in subjects having glioma than in healthy subjects [14]. Subjects with brain atrophy have smaller diameters than healthy subjects. Also, in the same study, normal value of the DPMM was stated; it ranged from 26.0 mm to 38.0 mm [14]. The DPMM/DIPFA and DPMM/DCPA values were lower in Koehler et al.’s [14] findings of subjects having glioma than our findings of healthy population. The measurements of MOMSJ and the ratio of DPMM to DCPA were higher in our study than Koehler et al.’s [14] findings of the healthy population, whereas DPMM/MOPMJ, and DPMM to MOMSJ ratios were higher in subjects having glioma than our findings. The mesencephalon, pons and medulla measurements of our study were lower than those in Chinese population performed by Liang Xiaochun et al. [16].

Some studies on medulla, pons and mesencephalon are shown in Table 8. In Sudanese healthy subjects aged between 21 and 30 years, the mesencephalon and medulla diameters were decreased with increased age from the age of 31–40 years to over 60 years and beyond. However, the pons diameter was decreased over the age of 50 years. Additionally, there was no significant difference in brainstem measurements between sexes [8]. In Finland, Raininko et al. [23] declared mesencephalon diameter decreased over the age of 60 years, whereas the pons diameter did not decrease with age. Also, sagittal diameter of medulla decreased over the age of 50 years, while in coronal diameter such decrease was not seen. No significant differences in changes related to sex were found [23]. In all values of our study except medulla, there was a decrease over the age of 50 years similar to Sudanese and Finns, and there were no significant differences between sexes, similar to above studies [8, 23]. The reason of the age-related changes of brainstem was shown as mesencephalon and espe-

### Table 7. The comparison of the different studies’ data

| Comparison of studies [mm] | Parameters — mean (min–max) |
|---------------------------|-----------------------------|
|                           | DIPFA | DCPA | DPMM | MOPMJ | MOMSJ | DPMM/MOPMJ | DPMM/MOMSJ | DPMM/DIPFA | DPMM/DCPA |
| Koehler et al. [14]       |       |      |      |       |       |            |            |            |           |
| Healthy subjects          | 13.2  | 18.1 | 26.6 | 15.3  | 9.4   | 1.7        | 2.8        | 2.0        | 1.5       |
| (11.1–)                   | (15.0–)| (24.5–)| (14.0–)| (8.1–) | (1.7–) | (2.6–)     | (1.9–)     | (1.4–)     |           |
| (15.3)                    | (21.2)| (28.7)| (16.6)| (10.7)| (1.8)  | (3.0)      | (2.2)      | (1.6)      |           |
| Subjects having glioma    | 22.1  | 22.95| 29.5 | 17.58 | 11.55 | 1.7        | 2.9        | 1.25       | 1.3       |
| (20.2–)                   | (20.9–)| (26.5–)| (15.3–)| (6.9–) | (1.2–) | (1.4–)     | (1.1–)     | (1.2–)     |           |
| (24.0)                    | (25.1)| (32.2)| (22.0)| (19.4)| (1.9)  | (4.3)      | (1.4)      | (1.6)      |           |
| Subjects having brain atrophy | 12.83 | 17.77| 19.67| 11.67 | 8.6    | 1.73       | 2.33       | 1.53       | 1.1       |
| (11.5–)                   | (17.0–)| (16.0–)| (10.6–)| (7.4–) | (1.3–) | (2.2–)     | (1.4–)     | (0.9–)     |           |
| (13.8)                    | (18.2)| (21.5)| (12.8)| (10.0)| (1.9)  | (2.6)      | (1.6)      | (1.2)      |           |
| Liang Xiaochun et al. [16]|       |      |      |       |       |            |            |            |           |
| Healthy subjects          | 13.10 | –    | 23.20| 14.70 | 12.50 | –          | –          | –          | –         |
| Females                   | 11.72 | 13.64| 21.62| 13.46 | 10.24 | 1.617      | 2.146      | 1.870      | 1.602     |
| (8.00–)                   | (9.30–)| (17.5–)| (10.0–)| (7.20–)| (1.15–)| (1.46–)    | (1.34–)    | (1.26–)    |           |
| (16.90)                   | (18.50)| (28.6)| (18.0)| (15.10)| (2.04)| (3.01)     | (2.89)     | (2.14)     |           |
| Males                     | 11.14 | 13.01| 21.97| 13.47 | 9.91  | 1.639      | 2.253      | 1.993      | 1.700     |
| (8.90–)                   | (9.30–)| (18.10)| (10.9–)| (7.30–)| (1.23–)| (1.76–)    | (1.45–)    | (1.34–)    |           |
| (14.70)                   | (17.10)| (29.9)| (18.10)| (13.40)| (2.14)| (3.78)     | (2.56)     | (2.23)     |           |
| Total                     | 11.49 | 13.39| 21.76| 13.46 | 10.11 | 1.626      | 2.189      | 1.919      | 1.641     |
| (8.00–)                   | (9.30–)| (17.50)| (10.00)| (7.20–)| (1.15–)| (1.46–)    | (1.34–)    | (1.26–)    |           |
| (16.90)                   | (18.50)| (29.90)| (18.10)| (15.10)| (2.14)| (3.78)     | (2.89)     | (2.23)     |           |

**Abbreviations** — see text
cially, this decline might be due to loss of superior 
cerebellar fibre bundles such as brachium conjunc-
tivum, fasciculus cerebello thalamicus. Also, these 
measurements in brainstem of normal subjects might 
be important for indicating the differences between 
patients with neurological disorders such as PD, AD 
or schizophrenia [23]. These parameters were lower 
in our study than in the studies performed in Suda-
nese and Japanese, and Massey et al. [12, 18, 23]. 

In a study by Massey et al. [18], pons AP diameters 
of subjects with Multiple System Atrophy were lower 
than in healthy subjects, patients with PD and pro-
gressive supranuclear palsy (PSP). Also, the lowest 
value of mesencephalon AP diameter was obtained 
in subjects having PSP. Additionally, mesencephalon 
to pons ratio was used as a clinical biomarker of PSP 
in some studies [18, 26]. So, the mesencephalon 
to pons ratio might be lower in PSP. In this study, 
this ratio was 0.613 ± 0.074 and 0.606 ± 0.087 in 
females and males, respectively. On the other hand, 
in studies of age-related changes, the lowest value 
of mesencephalon to pons ratio was obtained in 
the sixth decades in females (group 4), whereas, the 
Messey et al. [18].

Table 8. The comparison of the different populations’ data

| Studies                  | Groups                        | Parameters [mm] |
|--------------------------|-------------------------------|-----------------|
|                          |                               | PAP | PVD | MBAPD | MBVD | Midbrain to pons ratio |
| Elhussein et al. [8]     | Healthy subjects              | –   | 22.40 | –     | 15.40 | –                       |
| Hara et al. [12]         | Healthy subjects              | 22.77 | 28.40 | 15.10 | –     | –                       |
| Massey et al. [18]       | Healthy subjects              | 17.8  | –   | 11.1  | –     | 0.62                    |
|                          | Progressive supranuclear palsy | 17.1  | –   | 7.55  | –     | 0.44                    |
|                          | Parkinson’s disease           | 18.3  | –   | 11.4  | –     | 0.63                    |
|                          | Multiple system atrophy       | 14.8  | –   | 10.8  | –     | 0.77                    |
| Present study            | Healthy females               | 15.41 | 22.02 | 9.39  | 15.20 | 0.613 (0.44–0.94)        |
|                          | Healthy males                 | 15.58 | 22.64 | 9.37  | 15.64 | 0.606 (0.40–0.84)        |
|                          | Total                         | 15.47 | 22.27 | 9.38  | 15.38 | 0.610 (0.40–0.94)        |

Abbreviations — see text
in the same age group [21]. In a study reported by Shah et al. [25], there was a significant age related decrease in the mesencephalon cross-sectional area and neuronal death and nuclei degeneration was responsible for age-related decrease in midbrain. In the literature, there are few studies on brain stem in healthy Turkish population. In a study of Murshed et al. [20], there were no significant differences between males and females aged 13–50 years in whole brainstem volume measurements. However, the significant differences between sexes were found in the age group 51–77 years. Whole brainstem area was larger in males than in females, whereas there were no significant differences in whole brainstem dimension in all age groups [20]. The reasons of the reduction in brainstem age- and sex-related changes were shown as intrinsic and extrinsic factors including sex hormones, or hypertension and habits such as smoking and alcohol [15, 20, 21]. Moreover, environmental factors including acquired diseases, toxin exposure, trauma can lead to developmental aberrations and loss of brain tissue [20, 22]. This process might be affected by genetic factors [25]. Also, the brain volume may be affected by racial differences [15]. In the Antar et al.’s study [2], the effects of the age on the brainstem volume based upon the same factors, such as regional factors, genetics or several demographical features, were documented. In addition, the authors noted that the slice thickness, size, subject’s form, and the mean age of the study samples may affect MRI measurement results [2]. In 42 fresh cadavers having mean age of 45 years, the mean lengths of brainstem, medulla oblongata, and pons were found as 54.37 mm, 16.43 mm and 29.6 mm, respectively. Also, there were no significant differences between sex in the brain structures [2]. In the present study, the dimensions of pons vertical diameter, mesencephalon vertical diameter, the DIPFA, and the DCPA were significantly different in both sexes (p < 0.05). However, pons dimensions, mesencephalon vertical diameter, the distance between the anterior surface of the pons midway between the mesencephalon and medulla to the fourth ventricular floor (DPMM), the shortest anteroposterior diameter of the medulla at the pontomedullary junction (MOPMJ), and the shortest anteroposterior diameter of the medulla at the medullospinal junction (MOMSJ) measurements, and the ratio of mesencephalon to pons were larger in males than females. In addition, the data regarding DIPFA, and the DCPA were decreased with increasing age (from 18 to 70). Pons anteroposterior diameter (PAP) and pons vertical diameter (PVD) were lowest in the age of 41–50 years, whereas PAP value was highest in 31–40-year-olds. PVA value was the highest in the age of 60–70 years. The MOPMJ were increased in the age of 51–60 years. This increase remains unchanged in the age of 51–60 years and 61–70 years. The MOMSJ decreased with age, while the highest value was seen in the age of 61–70 years. The dimensions of mesencephalon were decreased both between in the 4th and 5th decades, and in the 6th and 7th decades. In evaluation of measurements, some of brainstem measurements remain speculative. It was resulted from the features of the study groups such as genetic factors, mean age, race, or method features including slice thickness and size. Especially, the significant difference seen in values of DIPFA, DCPA, DPMM/MOMSJ, DPMM/DIPFA, DPMM/DCPA was caused by the results obtained in females. It is speculated that the reason of this situation might be various sex hormones and differences of aging process between sexes.

**CONCLUSIONS**

In conclusion, this is the first study considering the morphometric measurements and ratios of the brainstem in a Turkish healthy population using MRI in detailed. Our study give important knowledge and normative data related to brainstem measurements and it sorts out the studies like age-related changes or sex differences in Turkish population. These presented findings will provide an evaluation opportunity to data regarding-age and sex-related brainstem studies and will shed light on patients with neurologic disorders. The age- and sex-related normal dimension changes of the brain stem are necessary in evaluation of mesencephalon, pons and medulla in neurological disorders and ageing process.

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