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Nerve Conduction Study in Healthy Elderly Subjects in Central India: A Cross-Sectional Study

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Abstract

Background

There are many physiological changes that accompany aging. Slowing of muscle contraction, alteration of muscle metabolism and neuromuscular junction, and reduction of nerve conduction velocity (NCV) are among these physiological changes. The present study was conducted to elucidate the effect of physiological factors like gender, height, and Body Mass Index (BMI) on motor and sensory nerve conduction study (NCS) of the upper limb and find out the normal data for healthy elderly subjects in central India.

Methods

A cross-sectional study among 382 healthy adult participants of central India aged 60 years and above. The study was carried out in the department of Physiology, Acharya Vinoba Bhave Rural Hospital, Wardha, India, from July 2017 to June 2022. An NCS was performed using the Neuron Spectrum 5 machine (Neurosoft, Ivanovo, Russia). A Microsoft Excel spreadsheet (Microsoft Corporation, Redmond, Washington, United States) was used to tabulate the information gathered. For statistical analysis, IBM SPSS Statistics for Windows, Version 22.0 (Released 2013; IBM Corp., Armonk, New York, United States) was used.

Results

All NCS parameters were greater in males as compared to females. As age advances, longer distal motor latency (DML) and F-wave minimum latency, decreasing amplitude, and slowing of conduction velocity (CV) were observed. As height increases, increasing DML and F-wave minimum latency, decreasing amplitude, and slowing of CV were observed. Higher BMI was found to be associated with lower amplitudes and slowing of CVs.

Conclusion

Age and height have a negative influence on amplitude and CV is a positive influence on DML and F-min latency. Higher BMI has a negative influence on amplitude and CV.

Categories: Internal Medicine, Neurology, Anatomy

Keywords: f-wave minimum latency, distal motor latency, effect of bmi on conduction velocity, conduction velocity, age, latency, central india, healthy elderly, nerve conduction

Introduction

Electrical pulses applied to the skin overlying peripheral nerves are usually sufficient to bring them to an action potential [1]. An NCS and needle electromyography (EMG) can be used to supplement the clinical examination of the peripheral nervous system [2]. It can help distinguish between neuropraxia and axonal transection in patients with extremity trauma and track their clinical progress [3]. There are many physiological changes that accompany aging. The slowing of muscle contraction, alteration of muscle metabolism and neuromuscular junction, and reduction of NCV are some of these physiological changes [4-6]. Neurological illnesses in older adults are a major cause of disability and dependence and account for a disproportionate number of patients treated in neurological outpatient and hospital settings [7]. Therefore, this study aims to elucidate the effect of these physiological factors (gender, height, and BMI) on motor and sensory nerve conduction study of the upper limb and find out the normal data for healthy elderly subjects in central India.

Objectives

To establish NCV values of the commonly tested upper limb nerves and to study the effects of gender, height, and BMI on NCV in healthy elderly subjects.

Materials And Methods

How to cite this article

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This is a cross-sectional study conducted at the Department of Physiology, Acharya Vinoba Bhave Rural Hospital, Wardha, India, between July 2017 and June 2022. Adults over the age of 60 who were otherwise healthy were recruited for the study.

Individuals over the age of 60 years make up 7.7% of India's population [8], and a margin of error of 5% was used to arrive at the sample size we used in the study. No previous history of trauma, neurological deficiency, or systemic illness that could cause neuropathy was found in the study's participants. Medical history was taken from each participant. Cases that met our inclusion criteria were included in the study after signing a written informed consent form. Parameters such as distal motor latency (DML), compound muscle action potential (CMAP) amplitude, and CV of the motor nerve and sensory nerve action. potential (SNAP) amplitude and CV for the sensory nerves as well as the motor nerve's shortest possible F-wave have been used to establish norms. An investigation into the transmission of nerve impulses was carried out with the Neuron Spectrum 5 apparatus (Neurosoft, Ivanovo, Russia). A Microsoft Excel spreadsheet (Microsoft Corporation, Redmond, Washington, United States) was used to tabulate the information gathered. For statistical analysis, IBM SPSS Statistics for Windows, Version 22.0 (Released 2013; IBM Corp., Armonk, New York, United States) was used. For comparison, Chi-squared and Fisher's exact tests were used.

Results

Demographic characteristics of the study population

Table 1 shows that among participants, the majority were in the age group of 60-65 years (35.34%) with a mean age of 68.43 ± 5.67 years. The majority of the participants were male (82.46%) as compared to females (17.54%).

Variables		No. of participants (n=382)	Percentage
Age group (years)	60-65	135	35.34
	66-70	108	28.27
	71-75	88	23.04
	76-80	48	12.56
	>80	03	0.78
Gender	Male	315	82.46
	Female	67	17.54

TABLE 1: Demographic variables among participants

Electrophysiological distribution

Table 2 shows normative data of NCS for median and ulnar nerves (motor and sensory) in the form of distal motor latency (DML), amplitude, CV, and F minimum latency.

Nerves	Parameters	Mean ±SD
Median nerve	Distal latency (msec)	3.91 ±0.76
	Amplitude (mV)	12.68 ±3.81
	Nerve conduction velocity (m/s)	54.66 ±5.87
	Sensory amplitude (µ)	32.17 ±14.43
	Sensory nerve conduction velocity (m/s)	57.88 ±6.83
	F minimum latency (msec)	27.61 ±3.73
	Distal latency (msec)	2.75 ±0.67
	Amplitude (mV)	12.88 ±2.86
Liner per/o	Nerve conduction velocity (m/s)	58.16 ±4.91
Ulnar nerve	Sensory amplitude (µ)	25.88 ±13.71
	Sensory nerve conduction velocity (m/s)	57.72 ±7.76
	F minimum latency (msec)	27.70 ±3.16

TABLE 2: Electrophysiological distribution among the participants

Electrophysiological distribution according to gender

Table 3 shows the electrophysiological distribution according to gender for the median and ulnar nerve.

Nerves	Parameters	Male	Female	P-value
Median nerve	DML (msec)	3.91 ±0.74	3.90 ±0.84	0.92
	AMP (mV)	12.74 ±3.84	12.41 ±3.70	0.52
	CV (m/s)	54.80 ±5.90	53.99 ±5.80	0.31
	S-AMP (µ)	30.48 ±12.47	40.63 ±19.50	<0.0001*
	S-CV (m/s)	58.20 ±6.84	56.59 ±6.39	0.08
	F min (msec)	27.76 ±3.33	26.74 ±5.11	0.04*
	DML (msec)	2.69 ±0.63	3.02 ±0.82	<0.001*
	AMP (mV)	12.91 ±2.98	12.87 ±2.09	0.92
	CV (m/s)	58.01 ±4.81	58.96 ±5.31	0.15
Unar nerve	S-AMP (µ)	24.35 ±11.83	33.19 ±18.91	<0.0001*
	S-CV (m/s)	57.36 ±7.67	59.69 ±7.72	0.02*
	F min (msec)	27.97 ±3.02	26.38 ±3.45	<0.001*

TABLE 3: Electrophysiological distribution according to gender

DML: distal motor latency; AMP: amplitude; CV: conduction velocity; S-AMP: sensory amplitude; S-CV: sensory conduction velocity; F min: F minimum latency

Electrophysiological distribution according to age

Table 4 shows the electrophysiological distribution according to age for the median and ulnar nerve.

Nerves	Parameters	60-65	66-70	71-75	76-80	>80	P-value
	DML (msec)	3.79 ±0.71	3.85 ±0.71	3.90 ±0.75	4.24 ±0.83	5.51 ±0.86	<0.001
	AMP (mV)	14.05 ±3.96	13.09 ±3.37	10.73 ±3.11	11.34 ±3.56	12.73 ±8.33	<0.001
Madian nanya	CV (m/s)	55.31 ±5.78	54.39 ±6.21	53.56 ±5.79	54.88 ±5.12	56.18 ±3.63	0.26
Median nerve	S-AMP (µ)	37.43 ±14.70	32.39 ±11.86	27.46 ±14.64	26.90 ±13.57	9.99 ±3.60	<0.001
	S-CV (m/s)	59.51 ±7.44	57.33 ±6.12	57.19 ±6.31	55.98 ±6.79	54.88 ±7.33	0.008
	F min (msec)	26.75 ±2.90	27.05 ±2.95	27.05 ±2.95	29.19 ±5.25	25.73 ±1.15	<0.001
	DML (msec)	2.86 ±0.74	2.68 ±0.66	2.65 ±0.54	2.73 ±0.70	2.98 ±0.96	0.13
	AMP (mV)	12.88 ±3.01	12.98 ±3.10	12.83 ±2.70	12.74 ±2.25	13.87 ±1.32	0.96
Lilper popyo	CV (m/s)	58.48 ±5.12	58.34 ±4.44	57.81 ±4.96	57.58 ±5.29	56.40 ±5.99	0.69
	S-AMP (µ)	29.04 ±15.29	26.03 ±12.31	22.45 ±12.89	23.88 ±11.44	10.77 ±4.76	0.001
	S-CV (m/s)	59.17 ±6.50	56.67 ±6.70	56.69 ±7.84	57.89 ±11.85	57.97 ±3.50	0.08
	F min (msec)	29.97 ±2.63	27.27 ±2.77	28.52 ±3.35	29.33 ±4.06	25.83 ±1.17	<0.001

TABLE 4: Electrophysiological distribution according to age

DML: distal motor latency; AMP: amplitude; CV: conduction velocity; S-AMP: sensory amplitude; S-CV: sensory conduction velocity; F min: F minimum latency

Electrophysiological distribution according to height

Table 5 shows the electrophysiological distribution according to height for the median and ulnar nerve.

Nerves	Parameters	≤150	151-160	161-170	171-180	>180	P-value
	DML (msec)	4.10 ±0.94	3.83 ±0.80	3.88 ±0.71	4.01 ±0.77	3.74 ±0.60	0.36
	AMP (mV)	12.58 ±4.67	12.35 ±3.67	12.61 ±4.01	12.99 ±3.43	13.74 ±4.23	0.72
Median nerve	CV (m/s)	54.39 ±5.91	54.02 ±5.82	54.77 ±5.74	55.43 ±6.30	52.05 ±4.12	0.32
Median nerve	S-AMP (µ)	39.23 ±24.18	37.39 ±17.59	31.27 ±11.86	27.77 ±11.92	27.97 ±11.92	<0.001*
	S-CV (m/s)	58.03 ±6.22	57.24 ±6.23	58.33 ±6.49	57.45 ±8.37	59.21 ±3.30	0.68
	F min (msec)	27.38 ±3.47	27.06 ±4.99	27.42 ±3.27	28.45 ±2.91	28.74 ±4.01	0.09
	DML (msec)	3.07 ±0.54	2.88 ±0.85	2.70 ±0.65	2.64 ±0.49	2.88 ±0.65	0.03*
	AMP (mV)	12.85 ±2.74	13.23 ±2.29	12.88 ±2.98	12.57 ±3.22	12.48 ±2.19	0.63
l linar nerve	CV (m/s)	55.10 ±3.35	58.63 ±5.28	58.57 ±4.65	57.96 ±5.10	53.15 ±0.68	0.001*
Unial herve	S-AMP (µ)	30.78 ±17.66	31.07 ±18.54	25.16 ±10.95	21.45 ±10.37	22.48 ±8.49	<0.001*
	S-CV (m/s)	59.28 ±5.54	58.26 ±7.35	57.11 ±7.22	58.20 ±9.71	56.81 ±3.10	0.62
	F min (msec)	27.05 ±3.07	26.96 ±3.88	27.66 ±3.15	28.45 ±2.13	29.60 ±1.23	0.006*

TABLE 5: Electrophysiological distribution according to height

DML: distal motor latency; AMP: amplitude; CV: conduction velocity; S-AMP: sensory amplitude; S-CV: sensory conduction velocity; F min: F minimum latency

Electrophysiological distribution according to BMI

Table 6 shows the electrophysiological distribution according to BMI for the median and ulnar nerve.

Nerves	Parameters	<18.5	18.5-24.9	≥25	P-value
	DML (msec)	3.75 ±0.75	3.95 ±0.76	3.87 ±0.70	0.10
	AMP (mV)	13.35 ±4.12	12.62 ±3.64	11.09 ±4.15	0.02*
Median nenve	CV (m/s)	54.44 ±5.30	54.74 ±6.07	54.55 ±5.84	0.83
Wedian neive	S-AMP (µ)	35.72 ±14.48	30.74 ±13.88	34.35 ±17.71	0.01*
	S-CV (m/s)	57.49 ±6.35	58.08 ±7.03	57.28 ±6.55	0.69
	F min (msec)	27.82 ±3.66	27.40 ±3.69	28.76 ±4.09	0.58
	DML (msec)	2.81 ±0.62	2.72 ±0.69	2.81 ±0.69	0.73
	AMP (mV)	13.34 ±3.03	12.85 ±2.87	11.94 ±1.98	0.06
l llnar nenve	CV (m/s)	57.63 ±5.11	58.43 ±4.94	57.33 ±4.01	0.58
	S-AMP (µ)	28.71 ±16.14	24.87 ±12.69	26.55 ±13.80	0.07
	S-CV (m/s)	56.46 ±7.13	58.09 ±7.95	58.09 ±7.72	0.51
	F min (msec)	27.87 ±3.25	27.60 ±3.10	28.08 ±3.75	0.64

TABLE 6: Electrophysiological distribution according to BMI

DML: distal motor latency; AMP: amplitude; CV: conduction velocity; S-AMP: sensory amplitude; S-CV: sensory conduction velocity; F min: F minimum latency

Discussion

Clinical evaluations of individual patients and epidemiological studies rely on normal values. The present cross-sectional study was carried out to generate comparative normative data on NCS of healthy elderly subjects in central India in the tertiary care center. The findings of the NCV in the upper limb of the present study was shown in Table 7.

Nerves	Parameters	Shehab et al.	Thakur et al.	Palve et al.	Present study
	DML (msec)	3.1+0.3	2.68±0.29	3.95 ±0.6	3.91 ±0.76
	AMP (mV)	11.1+2.8	9.68±3.11	-	12.68 ±3.81
	CV (m/s)	56.5+3.5	-	56.7 ±1.1	54.66 ±5.87
Median nerve	S-AMP (µ)	-	-	-	32.17 ±14.43
	S-CV (m/s)	-	-	54.5 ±7.5	57.88 ±6.83
	F min (msec)	-	23.82±1.48	25.3 ±1.6	27.61 ±3.73
	DML (msec)	2.4+0.3	2.25±0.54	-	2.75 ±0.67
	AMP (mV)	9.2+2.2	9.09±2.65	-	12.88 ±2.86
l llnar nanva	CV (m/s)	60.4+5.2			58.16 ±4.91
Unital herve	S-AMP (µ)	-	-	-	25.88 ±13.71
	S-CV (m/s)	-	-	-	57.72 ±7.76
	F min (msec)	-	24.88±1.79	-	27.70 ±3.16

TABLE 7: Values of NCS in upper limb

DML: distal motor latency; AMP: amplitude; CV: conduction velocity; S-AMP: sensory amplitude; S-CV: sensory conduction velocity; F min: F minimum latency; NCS: nerve conduction study

Shehab [9]; Thakur et al. [10]; Palve and Palve [11]

The mean age among male and female subjects was 68.33 ±5.69 and 68.85 ±5.08 years, respectively. The findings showed that significantly longer DML was observed as age advances except for ulnar DML. There was a clear decline in CMAP amplitude as well as SNAP amplitude as a person gets older. This decline is significant in the motor median nerve. Slowing of motor and sensory CV was observed in the older age group. The slowing is significant in sensory ulnar and sensory median nerves. F-wave demonstrated a significant (p<0.05) rise in F-wave minimum latency as age progressed. Research has shown that CV begins to decline around the age of 30-40 years, but by the time people reach their 60s or even their 80s, the values dropped by about 10 m/s. The loss of myelinated and unmyelinated nerve fibers in peripheral nerves with aging may be responsible for the decline in nerve conduction and the increase in sensory latency that occurs with age [9-11]. The effect of sex on NCS can be explained on the basis of gender-wise differences in anatomical and physiological factors [12]. We observed greater values of SNAP amplitude and sensory CV in females as compared to males, especially in ulnar nerves. This might be because of sampling error (less number of females). Robinson et al. [13] explained that gender differences in nerve conduction parameters could be due to differences in height.

In the present study, the majority of subjects were in the height group 161-170 cms (46.33%) with the mean height of the subjects being 166.08 ±8.48 cms. There was an increase in DML as the height increased. This rise was significant (p<0.05) in the ulnar nerve. CMAP amplitude showed a non-significant decrease as the heights increased whereas SNAP amplitude depicted a significant decrease in the median and ulnar nerves with an increase in height. Motor CV showed a significant (p<0.05) decrease as the height advances in ulnar nerves whereas in other nerves this decrease is non-significant. Sensory CV showed no significant decrease with an increase in height in the median and ulnar. F-wave minimum latencies showed a significant positive association with height, i.e., as the height increased F-wave minimum latency also increased. Thus we observed a negative association of CV and amplitude and a positive association of DML and F-wave minimum latency with height as shown in Table 5. Our observations were in accordance with findings by Soudmand et al. [4], who found an inverse correlation of CV with height (r=-0.46 (p<0.01); r=-0.36 (p<0.05)); a positive correlation between height and median, peroneal F-wave minimum latency (r=0.74 and 0.69, respectively; p<0.001). However, they observed no relationship between height and median (motor and sensory) CV (r=-0.04 (p>0.25); r=-0.14(p>0.25)), which is in contrast to our observation.

In the present study, the majority of subjects were with normal BMI 18.5-24.9 kg/m2 (69.11%) with the mean BMI of the subjects being 20.76 \pm 2.92 Kg/m2. No observable fixed trend could be seen in regard to DML and F-wave minimum latencies varying BMI. (Table 6) Only the median nerve was found to be associated with a

lower amplitude (sensory and motor). Table 6 shows a non-significant slowing of both motor and sensory CV with increasing BMI. Our findings are in line with those of Awang et al. [14], who found a decrease in a CV in the median nerve (motor and sensory) and the ulnar with increasing BMI (motor). Despite this, they found no discernible change in the ulnar nerve's sensory properties. In motor ulnar and peroneal nerves, Buschbacher et al. [15] found a longer latency association with lower BMI. They also reported a significant association between latency and BMI in sensory radial and ulnar nerves. In all other sensory nerves and in the median nerves, no statistically significant difference was observed in latency, amplitude, and CV with varying BMI. In all sensory nerves, the amplitude was found to be varying with BMI (increased BMI association with lower amplitude).

Conclusions

The present study concludes that age and height negatively influence amplitude and CV, whereas they positively influence DML and F-min latency. Significantly longer DML was observed as the age advances, except for ulnar nerve DML. There was an apparent decline in CMAP and SNAP amplitude. The reduction is significant in the motor median nerve, and slowing of motor and sensory CV was observed in the older age group that is significant in sensory ulnar and sensory median nerves. This rise was significant in the ulnar nerve. SNAP amplitude depicted a substantial decrease in the median and ulnar nerves with an increase in height. Motor CV showed a significant reduction as the height advances in the ulnar nerves. We observed a negative association between CV and amplitude and a positive association between DML and F-wave minimum latency with height. Higher BMI has a negative influence on amplitude and CV.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. Datta Meghe Institute of Medical Sciences (Deemed to be University) Institutional Ethics Committee, Wardha, India issued approval DMIMS (DU)/IEC/2016-17/6143. Animal subjects: All authors have confirmed that this study did not involve animal subjects or tissue. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- Giacomini PS: Electromyography and neuromuscular disorders: clinical electrophysiologic correlations. McGill University, Montreal, Canada; 2006.
- Lee DH, Claussen GC, Oh S: Clinical nerve conduction and needle electromyography studies. J Am Acad Orthop Surg. 2004, 12:276-87. 10.5435/00124635-200407000-00008
- Robinson LR: Role of neurophysiologic evaluation in diagnosis. J Am Acad Orthop Surg. 2000, 8:190-9. 10.5435/00124635-200005000-00006
- Soudmand R, Ward LC, Swift TR: Effect of height on nerve conduction velocity. Neurology. 1982, 32:407-10. 10.1212/wnl.32.4.407
- Campbell MJ, McComas AJ, Petito F: Physiological changes in ageing muscles. J Neurol Neurosurg Psychiatry. 1973, 36:174-82. 10.1136/jnnp.36.2.174
- Roos MR, Rice CL, Vandervoort AA: Age-related changes in motor unit function. Muscle Nerve. 1997, 20:679-90. 10.1002/(sici)1097-4598(199706)20:6<679::aid-mus4>3.0.co;2-5
- Sirven JI, Malamut BL: Clinical Neurology of the Older Adult. Wolters Kluwer Health/Lippincott Williams & Wilkins, Philadelphia; 2008.
- 8. Elderly in India 2021 . Ministry of Statistics and Programme Implementation, Government of India, New Delhi, India; 2021.
- https://www.mospi.gov.in/documents/213904/301563/Elderly%20in%20India%2020211627985144626.pdf/a4647f03bca1-1ae2-6c0f....
- Shehab DK: Normative data of nerve conduction studies in the upper limb in Kuwait: are they different from the western data?. Med Principles Pract. 1998, 7:203-8. 10.1159/000026043
- 10. Thakur D, Paudel BH, Jha CB: Nerve conduction study in healthy individuals: a preliminary age based study . Kathmandu Univ Med J (KUMJ). 2010, 8:311-6. 10.3126/kumj.v8i3.6218
- Palve SS, Palve SB: Impact of aging on nerve conduction velocities and late responses in healthy individuals . J Neurosci Rural Pract. 2018, 9:112-6. 10.4103/jnrp.jnrp_323_17
- 12. Stetson DS, Albers JW, Silverstein BA, Wolfe RA: Effects of age, sex, and anthropometric factors on nerve conduction measures. Muscle Nerve. 1992, 15:1095-104. 10.1002/mus.880151007
- Robinson LR, Rubner DE, Wahl PW, Fujimoto WY, Stolov WC: Influences of height and gender on normal nerve conduction studies. Arch Phys Med Rehabil. 1993, 74:1134-8.
- Awang MS, Abdullah JM, Abdullah MR, Tahir A, Tharakan J, Prasad A, Razak SA: Nerve conduction study of healthy Asian Malays: the influence of age on median, ulnar, and sural nerves. Med Sci Monit Int Med J Exp Clin Res. 2007, 13:330-2.
- 15. Buschbacher RM: Body mass index effect on common nerve conduction study measurements . Muscle Nerve. 1998, 21:1398-404. 10.1002/(sici)1097-4598(199811)21:11<1398::aid-mus6>3.0.co;2-4