INTRODUCTION

Historically, Richard Caton (1875) discovered evoked potentials and the electroencephalogram (EEG) at the same time. The visual evoked potentials (VEPs) result from change of brain activity following application of intermittent visual stimulus to the visual system. They provide a quantitative measure of the functional integrity of the visual pathways. Visual evoked potentials (VEPs) are massed electrical signals generated by occipital cortical areas 17, 18, and 19 in response to visual stimulation. VEPs can be used to assess the integrity or maturational state of the visual pathway in infants and preverbal children. The VEP to flash stimulation consists of a series of negative and positive waves. The most robust components of the flash VEP are the N2 (90 ms) latency and P2 (120 ms) latency. As demonstrated by Sokol and Jones, the latency of the VEP matures rapidly between three and five months. Latencies of all check sizes are at adult values by age 5, and show no significant change from then until about age 40.

On fundoscopic examination, foveal cone photoreceptors which subserve fine detail of vision, is not mature until at least 4 years of age. The robust nature of the chromatic responses have allowed researchers to apply the visual evoked response (VERP) technique as a sensitive and objective measure of neural integrity in the clinic. VEP responses have been worked out for adult population in various studies. However, in children the recording is...
difficult and their normatives are not consistent across the studies since the normative are very necessary for diseases like birth asphyxia, cortical blindness. Objective assessment of visual pathway is done in above diseases at our center. Therefore, this study is an attempt to document VEP responses among children population from 0.5 years to 4 years. The study is also likely to produce effect of age on VEP with the time of maturation of VEP.

MATERIALS AND METHODS

The study was conducted on healthy children (n=36, 72 eyes) of age 6 months to 4 years old in Electrodiagnosis labs (Neurophysiology lab) of Department of Basic and Clinical Physiology, BPKIHS. It was a descriptive cross sectional study. Subjects were taken from the well-baby clinic of the Department of Paediatric and Adolescent using convenient sampling method. All the subjects were age and sex matched. Further health status was assessed by detailed medical history and physical examination (pupillary light reflex-mesopic condition). This study was conducted from 15th August, 2010 to 14th August 2011. Recorded Variables were anthropometric variables: Age (year), height (m), weight (kg), body mass index (kg/m2) and VEP variables: N90 latency (ms) and P120 latency (ms) of right and left eye stimulation, P120 amplitude (peak to peak of N90-P120, μV) of right and left eye stimulation. Parents were instructed to bring the child with clean scalp and hair, with adequate last night sleep (minimum 6 hours) and with light food on the day of examination. Recordings were done in morning period.

Inclusion criteria

Healthy children of either sex between the ages of 6 months to 4 years were included.

Exclusion criteria

The children with history of head injury, stroke, medical illness, using mydriatic or miotic drugs, ophthalmic health= refractive errors, ocular diseases like congenital glaucoma, cataract, uveitis and so on were excluded.

Normal subjects criteria

Children with normal general health and with normal pupillary light reflex, normal pupillary diameter, no visual complaints, no refractive errors were considered normal.

Temperature of the laboratory was maintained at 26 ± 2°C by air condition. Children ≤ 4 years were selected for flash VEP and promethazine 0.5 mg/kg body weight was administered to induce sleep.

Pre-recording and recording procedure was started by consent of the parents, health status, recording setup. The sedation (promethazine 0.5 mg/kg body wt) of the child was done as recommended by the FDA (US Food and Drug Administration). Recording of VEP was done while the subject was sleeping. The scalp electrodes were placed relative to bony landmarks, in proportion to the size of the head, according to the international 10-20 system. All electrode sites were cleaned with Skinpure to clean and reduce the skin resistance. Midline-occipital (MO) electrode, the active electrode, is 5 centimetres (cms) above the inion for single channel recording. Recommended montages used were right-occipital (RO) = 5 cm right of MO (midline-occipital), left-occipital (LO) = 5 cm left of MO and midline-frontal (MF), the reference electrode, were placed 12 cm above the nasion. Earthing electrode was placed on vertex (CZ). The electrodes were filled with Nihon Kohden Elefix gel, which acts as electrical conductor and was stabilized using the electrodes in place. Then electrodes were pressed firmly on the scalp.

Recording was done by placing the electrodes on the subject's scalp according to 10/20 international system. LED goggles connected with a LED visual stimulator was put on the child's eye while the subject were sleeping and red flash of strobe light was given monocularly through goggles to both right and left eye one at a time. Signals were displayed on the monitor of Nihon Kohden machine (NM-420S; H636, Japan). The record of VEP for each eye was done from channel-1 (RO-MF) and channel-2 (LO-MF) designated as A1, A2 waves respectively. Recording was repeated for reliability with waves designated as B1, B2. The flash reversal rate was 1/s (1.0 Hz ± 20%) for 300 ms. The signals were averaged for 200 times. The filter was provided creating window of 1-100 Hz. Recordings of VEPs (Figures 1 and 2).

Data analysis and statistical analysis

Statistical analysis was done by plotting the data in Microsoft Excel. Mean, median (inter quartile
range), mode and SD were analyzed in Excel for both anthropometric and VEP variables and normative values for each subgroups were expressed. Then all the VEP variables were entered in SPSS version 18. Post hoc (Bonferroni) analysis was done for comparing VEP variables of subgroups separately for flash group and pattern group. Pearson correlation were done for each variables of VEP with respect to age at significance value of p <0.05. Normative values for each subgroups are expressed in terms of mean ± SD.

The research proposal was submitted to the Institutional Ethical Review Board (IERB) of BPKIHS and ethical clearance was taken. Verbal and written consent was taken from the parent after giving detail information regarding the procedure.

RESULTS

Anthropometric variables were recorded and mean and standard deviation were calculated for total subjects and for subgroups (Tables 1 and 2).

Comparing flash VEP variables among subgroups of children (Table 3)
Comparing VEP latencies of children, the subgroup 3 children showed significantly less N90 latency on right eye stimulation as compared to subgroup 1 (103.75 ± 22.54 vs. 79.09 ± 17.77 ms, p=0.012). Similarly, on combining VEP N90 latencies of right and left eyes, the subgroup 3 children also showed significantly less as compared to subgroup 2 (95.29 ± 15.9 vs. 81.39 ± 16.36 ms, p=0.03) and subgroup 1 (99.26 ± 20.12 vs. 81.39 ± 16.36 ms, p=0.001). P120 latency on right eye stimulation (137.70 ± 12.04 vs. 116.67 ± 13.65 ms, p=0.004) and on left eye stimulation (137.18 ± 11.7 vs. 115.75 ± 12.69 ms, p=0.004) was significantly less in subgroup 3 as compared to subgroup 2 children. P120 latency was significantly less on right eye stimulation (148.5 ± 15.38 vs. 116.67 ± 13.65 ms, p=0.001) and on left eye stimulation (145.91 ± 18.11 vs. 116.67±13.65 ms, p=0.001) in subgroup 3 children as compared to subgroup 1. Combining VEP P120 latencies of right and left eyes were significantly less in subgroup 3 as compared to subgroup 2 (137.43 ± 11.57 vs. 116 ± 12.86 ms, p=0.001) and subgroup 1 (147.26 ±16.4 vs. 116 ± 12.86 ms, p=0.001).

Pearson correlation (2-tailed) of flash VEP variables with age (Table 4)
Flash VEP N90 latency of right eye stimulation showed significant negative correlation (r= -0.503, p = 0.003) with respect to age. When both eyes were combined flash VEP N90 latencies showed more significant negative correlation (r= -0.424, p= 0.001) with respect to age. The VEP P120 latency of right eye (r= -0.733, p=0.001) and left eye (r= -0.722, p=0.001) also showed significant negative correlation with respect to age. Similarly, significant negative correlation for both eyes combined VEP P120 latency (r= -0.728, p= 0.001) was shown with respect to age.

DISCUSSION

The most common indication for VEP clinically is evaluation of the visual pathway function. It is part of the work-up for children with abnormal visual development or behaviour, for children or adults with visual symptoms not explained by physical examination, for cases where a psychogenic visual disorder or malingering is suspected, and in cases of optic
nerve disease, especially inflammation. Our centre, Basic and Clinical Physiology of BPKIHS is regularly reporting the cases of Birth Asphyxia, Cortical Blindness, etc. Apart from these established indications, some centres also use specialised VEP methods for the objective assessment of visual acuity according to Tyler et al. 1979.11

In our study, we have subjected flash stimulus to 0.5 to 4 years children. These children were divided into subgroups 1 (0.5 to ≤2 years, n=13), 2 (2 to ≤3 years, n=11), and 3 (3 to ≤4 years, n=12) according to their age. The study showed the continuation of maturation of VEP around 4 years. Comparing VEP latencies of children, age group 3-4 yr children showed significantly less N90 latency on right eye stimulation as compared to age-group 0.5-2 yr (103.75 ± 22.54 vs. 79.09 ± 17.77 ms, p= 0.012). Amplitude also showed decreasing trend in our study in the age group 0.5 year to ≤4 years. Since the study showed...
the decreasing trend of the latency up to 4 yrs supporting the incomplete VEP maturation. Lippe et al\textsuperscript{12} have found similar result on VEP analysis of sixty-three infants aged 27 days to 5.5 years comparing among six age subgroups. All components follow a significant decrease in latency with age, reaching their adult values around 7–12 months. Langrova et al\textsuperscript{13} have mentioned that maturation of the flash VEPs and pattern-related VEPs finish by 6 years of age.

In our study (check size 66'), latency has not reached to adult level up to 4 years among flash subgroups children. Mean latencies and amplitudes obtained for mean age (each subgroup) in given stimulus conditions showed age related decline from 6 months to 4 years. N90 latency showed significant negative correlation with age for right eye ($r= -0.503$, $p=0.003$) and both eyes combined ($r=-0.424$, $p=0.001$). P120 latency also showed significant decreasing trend ($r=-0.728$, $p=0.001$) for both eyes combined. The study of Moskowitz et al\textsuperscript{14} have found that P1 (first positive wave) latency decreases rapidly during the first year of life for both large and small checks and that the time course of the latency change differs as a function of check size. VEPs to large checks attain adult-like P1 latency values by 27 days to 5.5 years comparing among six age subgroups children.

Our results are consistent with the results of Jelka B\textsuperscript{15} using pattern reversal and pattern onset stimulus. Maturation changes were rapid in infants and gradual in school children. Age-related changes in pattern electroretinograms in infants was seen as a decrease in latency. Pattern VEP showed age-related decrease in latency, and increase in amplitude and the development of the waveform as seen in our results. In schoolchildren, pattern VEP changes showed more gradual decrease in latency suggest that the electrophysiological maturation proceeds until adulthood. Mean latencies and amplitudes obtained for mean age (each subgroup) in given stimulus conditions showed age related decline from 6 months to 4 years in our study. Normative values in infants and schoolchildren are an important factor in differentiating maturation of the visual system from pathological processes.

Concluding the various findings, it has been found small variations due to the difference of the stimulus. Most of the studies including the present one showed the results that the maturation of the visual pathway is around 4 years.

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### REFERENCES