

Disorders of Peripheral Implications for the study of auditory adaptation to spectral shift: vowel spaces

Richard W and Kujawa DV

Department of Pediatrics, University of Wisconsin Medical School, Madison, Wisconsin

***Corresponding Author:** Kujawa DV, Department of Pediatrics, University of Wisconsin Medical School, Madison, Wisconsin

Citation: Disorders of Peripheral Implications for the study of auditory adaptation to spectral shift: vowel spaces. Am J Rhin and Otol. 2019; 1(1): 001-009.

Submitted: 08 February 2019; **Approved:** 16 February 2019; **Published:** 18 February 2019

Abstract

Cochlear implant (CI) users differ in their ability to perceive and recognize speech sounds. Two possible reasons for such individual differences may lie in their ability to discriminate formant frequencies or to adapt to the spectrally shifted information presented by cochlear implants, a basalward shift related to the implant's depth of insertion in the cochlea. In the present study, we examined these two alternatives using a method-of-adjustment (MOA) procedure with 330 synthetic vowel stimuli varying in F1 and F2 that were arranged in a two-dimensional grid. Subjects were asked to label the synthetic stimuli that matched ten monophthongal vowels in visually presented words. Subjects then provided goodness ratings for the stimuli they had chosen. The subjects' responses to all ten vowels were used to construct individual perceptual "vowel spaces." If CI users fail to adapt completely to the basalward spectral shift, then the formant frequencies of their vowel categories should be shifted lower in both F1 and F2. However, with one exception, no systematic shifts were observed in the vowel spaces of CI users. Instead, the vowel spaces differed from one another in the relative size of their vowel categories. The results suggest that differences in formant frequency discrimination may account for the individual differences in vowel perception observed in cochlear implant users.

INTRODUCTION

Although cochlear implants allow profoundly deaf people to hear, cochlear implant users show a very wide range of speech perception skills. The most successful cochlear implant users can easily hold a face-to-face conversation, and they can even communicate on the telephone, a difficult task because there are no visual cues available and because the acoustic signal itself is highly degraded (Gstoettner et al., 1997). On the other hand, the least successful cochlear implant users have a difficult time communicating even in a face-to-face situation, and can barely perform above chance on auditory-alone speech perception tasks (Dorman, 1993).

One long term goal of our research is to understand the mechanisms that underlie speech perception by cochlear implant (CI) users and, in so doing, gain an understanding of the individual differences in psychophysical characteristics which may explain individual differences in speech perception with a CI. It is important to remember that electrical hearing as provided by a cochlear implant is quite

different from normal acoustic hearing. One important difference lies in listeners' ability to discriminate formant frequencies. Kewley-Port and Watson (1994) report difference limens between 12 and 17 Hz in the F1 frequency region for highly practiced normal hearing listeners. In the F2 frequency region, they found a frequency resolution of approximately 1.5%. For cochlear implant users, formant frequency discrimination depends on two factors: the frequency-to-electrode map that is programmed into their speech processor, and the individual's ability to discriminate stimulation pulses delivered to different electrodes. It is not uncommon for some cochlear implant users to have formant frequency difference limens that are one order of magnitude larger than those of listeners with normal hearing, or even more (Nelson et al., 1995; Kewley-Port and Zheng, 1998). It is reasonable to hypothesize that cochlear implant users with such limited frequency discrimination skills will find it quite difficult to identify vowels accurately because formant frequencies are important cues for vowel recognition.

Cite this article: Disorders of Peripheral Implications for the study of auditory adaptation to spectral shift: vowel spaces. Am J Rhin and Otol. 2019; 1(1): 001-009.

Another important difference between acoustic and electric hearing is related to the finding that cochlear implants do not stimulate the entire neural population of the cochlea but only the most basal 25 mm at best, because the electrode array cannot be inserted completely into the cochlea. Therefore, cochlear implants stimulate cochlear locations that are more basal and thus elicit higher pitched percepts than normal acoustic stimuli. For example, when the input speech signal has a low frequency peak (e.g., 300 Hz), the most apical electrode is stimulated, regardless of the particular frequency-to-electrode table employed. The neurons stimulated in response to this signal may have characteristic frequencies of 1000 Hz or even higher. This represents a rather extreme modification of the peripheral frequency map. To the extent that the auditory nervous system of CI users is adaptable enough to successfully “re-map” the place frequency code in the cochlea, the basalward shift provided by a CI should not hinder speech perception. On the other hand, an inability to adapt and re-map the place frequency code may severely limit speech perception in CI users and may be an important source of individual differences in speech perception (Fu and Shannon, 1999a, b). Although there is consensus in the literature that cochlear implants stimulate neurons with higher characteristic frequencies than those stimulated by the same sound in normal ears, there is controversy about the amount of this possible basalward shift. For example, Blamey et al. (1996), in a study where CI users with some residual hearing were asked to match the electrical percepts in one ear to the acoustic percepts in the other ear, concluded that the electrode positions that matched acoustic pure tones were more basal than predicted from the characteristic frequency coordinates of the basilar membrane in a normal human cochlea. However, Blamey et al. (1996) also acknowledge that “the listeners may have adapted to the sounds that they hear through the implant and hearing aid in everyday life so that simultaneously occurring sounds in the two ears are perceived as having the same pitch.” On the other hand, Eddington et al. (1978) conducted the only experiment we are aware of, where a unilaterally deaf volunteer received a cochlear implant and was asked to match the pitch of acoustic and electric stimuli while still in the operating room, before he had much of a chance to adapt to the basalward shift in the way described by Blamey et al. (1996). Eddington et al. (1978) concluded that their pitch-matching data were “consistent with frequency versus distance relationships derived from motion of the basilar membrane.”

Several previous studies have addressed the issue of adaptation to changes in frequency-to-electrode

assignments for cochlear implant users. Skinner et al. (1995) showed that users of the SPEAK stimulation strategy identified vowels better with a frequency-to-electrode table that mapped a more restricted acoustic range into the subject's electrodes than the default frequency-to-electrode table. The experimental table that resulted in better vowel perception represented a more extreme basalward shift in spectral information than the default table, suggesting that listeners with cochlear implants can indeed adapt to such shifts, at least within certain limits. Another study that demonstrates the adaptation ability of human listeners in response to spectral shifts was conducted by Rosen et al. (1999), who used acoustic simulations of the information received by a hypothetical cochlear implant user who had a basalward spectral shift of 6.5 mm on the basilar membrane (equivalent to 1.3–2.9 octaves, depending on frequency). Initially, the spectral shift reduced word identification in normal-hearing subjects (1% correct, as compared to 64% for the unshifted condition), but after only three hours of training, subjects' performance improved to 30% correct. Whether or not such performance represents the maximum possible by CI users was not addressed in this study, given the relatively short time spent in training.

Recently, Fu et al. (submitted) performed an experiment in which the frequency-to-electrode tables of three cochlear implant users were shifted one octave with respect to the table they had been using daily for at least three years. It is important to note that this one-octave shift was in addition to the original shift imposed by the cochlear implant. After three months of experience with the new table, it was apparent that adaptation was not complete because, on average, subjects did not reach the same levels of speech perception that they had achieved before the table change. Taken together, these previous studies show that auditory adaptation to a modified frequency map is possible but it may be limited, depending on the size of the spectral shift that listeners are asked to adapt to.

In the present study, we investigated the adaptation of human listeners to a basalward shift using a new paradigm, a method-of-adjustment (MOA) procedure. This methodology was used to obtain maps of the perceptual vowel spaces of adult, post-lingually deafened cochlear implant users. Similar tasks have been used with normal hearing listeners as well as CI users (Johnson et al., 1993; Hawks and Fourakis, 1998). In this task, subjects select the region of the F1 – F2 plane that sounds to them like a given vowel, and the procedure is repeated for ten English vowels. This task simultaneously assesses

a cochlear implant user’s auditory adaptation ability, by comparing the locations of his/her selected regions to those selected by normal hearing listeners, and his/her frequency discrimination skills, by examining the spread of the selected regions. More specifically, a listener who was unable to adapt to the basalward spectral shift introduced by their cochlear implant would be expected to select regions whose centers are systematically shifted to lower frequencies with respect to the regions of the vowel space selected by normal hearing listeners in mapping their vowel categories. The extent to which cochlear implant listeners show relatively normal vowel category centers could be used as a measure of their adaptation to basalward spectral shift.

An alternative to this hypothesis predicts that the spread of selected regions (category sizes), as well as category centers, would increase (or “smear”) as a result of perceptual adaptation. The resulting vowel categories may be larger (show greater spread) reflecting the cochlear implant listener’s need to map a greater range of frequencies to a given vowel. To differentiate between the spectral smearing hypothesis and the frequency discrimination explanation would require a longitudinal study of the changes in vowel spaces and frequency discrimination. For the purposes of this study, a simple shift hypothesis limited to vowel category centers was tested and compared with the frequency discrimination hypothesis.

In addition to the MOA task, two other perceptual tests were administered to CI users, an F1 jnd test with synthetic vowel stimuli and a closed set identification test with naturally produced vowels. Taken together, these measures were intended to investigate the role of formant frequency discrimination and auditory adaptation in vowel perception by CI users.

EXPERIMENT

Methods

Participants

Forty-three Indiana University undergraduates with no reported history of speech or hearing problems and eight adult cochlear implant (CI) users, all monolingual speakers of English, participated in this experiment. The normal-hearing participants consisted of 20 males and 23 females ranging in age between 18 and 28. The normal-hearing participants were recruited to represent the dialect of American English spoken in central Indiana with a common inventory of vowels. Only normal-hearing listeners who reported living their entire lives in central Indiana were included in this experiment. Central Indiana was defined in terms of a 60-mile radius around Indianapolis, roughly covering the

Midland dialect as described by Wolfram and Schilling-Estes (1998). This criterion was used to exclude two other regional dialects found at the northern and southern extremes of the state. These other regional dialects are reported to differ from the Midland dialect in terms of vowel quality and degree and type of diphthongization (Labov, 1991). For participating in two 1-h sessions, the participants received either \$7.50 per hour or two credits towards their research requirement if they were enrolled in an undergraduate psychology class.

The CI users were recruited from the population of adult patients served by the Department of Otolaryngology-Head and Neck Surgery at the Indiana University School of Medicine in Indianapolis. The demographics of the CI users are given in Table I, while information concerning their cochlear implants is provided in Table II. All of the CI users were native speakers of American English, with the exception of CI1, who was a native speaker of British English. British and Midland American English are not reported to differ substantively from one another in vowel quality for the ten vowels used in this study (Gimson, 1962; Pilch, 1994). Thus the American English vowel spaces were deemed an acceptable benchmark for CI1 as well as the other CI users.

All of the CI users had received their cochlear implants at least one year prior to participating in this study. The CI users differed from one another in terms of the type of co-chlear implant they received: Five were users of the Nucleus-22 or Nucleus-24 device, programmed with either the SPEAK strategy or the MPEAK strategy, while three were users of the Clarion device, programmed with the CIS strategy. The SPEAK strategy (Skinner et al., 1994) filters the incoming speech signal into a maximum of 20 frequency bands, which are associated with different intracochlear stimulation channels. Typically, six channels are sequentially stimulated in a cycle, and this cycle is repeated 250 times per second. The channels to be stimulated during each cycle are chosen based on the frequency bands with the highest output amplitude. In contrast, the CIS strategy (Wilson et al., 1991) as implemented in the Clarion device filters the signal into eight bands, one for each stimulation channel. All channels are sequentially stimulated with pulses whose amplitudes are determined by the filters’ outputs. The stimulation cycle is repeated at a rate of at least 833 times per second. The CIS strategy differs from the SPEAK strategy in its use of a different stimulation rate, fewer stimulation channels, and in its stimulation of all channels in a cycle rather than only a subset of the available channels.

The CI users also differed from one another in terms of the depth of insertion of the electrode ar-

ray in the cochlea. The array's depth of insertion, in turn, determines the magnitude of the basalward spectral shift induced by the implant. It is possible to roughly estimate the size of this basalward shift for an individual CI user with three pieces of information: the location of the electrodes, the frequency to electrode mapping used by the cochlear implant's speech processor, and estimates of the characteristic frequency of the neurons stimulated by a given electrode pair. The speech processors of the participants in this study divide the acoustic frequency spectrum into channels. Each channel is specified by an acoustic frequency range that is assigned to a pair of intra-cochlear electrodes. Low frequencies are mapped to the apical electrodes, while high frequencies are mapped to the basal electrodes.

In the present study, the basalward spectral shift was calculated for two channels for every subject, the channels corresponding to 475 Hz and 1500 Hz. These frequencies correspond to the F1 and F2 of a neutral vowel for an average male speaker. First, the place of stimulation for a specific channel was defined as occurring half way between the electrodes for that channel. When stimulation was bipolar (i.e., both electrodes were intracochlear), the place of stimulation was defined as occurring half way between the electrodes for that channel. For patients who received monopolar stimulation (i.e., the return electrode was extracochlear) the location of electrical stimulation was considered to be at the intracochlear electrode. Second, the intraoperative report of insertion depth was used to adjust this place estimate based on the depth of electrode insertion. Cochlear lengths of the subjects were not measured individually. Instead, the basalward shift was calculated assuming that all subjects had average sized cochleas with a length of 35 mm (Hinojosa and Marion, 1983). Third, the electrically assigned frequency for this location was calculated as the geometric mean of the frequency boundaries defining the channel being studied in the speech processor. Finally, the characteristic frequency of the neurons stimulated by a given electrode was calculated from Greenwood (1961) and compared to the frequency calculated in step three. This discrepancy for each CI user, measured in Hz and octave, is shown in Table III. Alternatively, the shift is also reported in terms of the location difference between electrical and acoustic stimulation, in mm. The use of Greenwood's equation is based on the assumption that the average characteristic frequency of neurons stimulated by an electrode pair placed x mm from the round window is the frequency that would cause maximum displacement of the basilar membrane at the same x mm from the round window. Clearly, these are only rough estimates of the amount of basalward

shift. In particular, the estimate of a 35-mm cochlea may lead to substantial overestimates or underestimates of the actual basalward shift. Future studies may improve the precision of these estimates by measuring the length of each individual cochlea using 3D reconstructions of CAT scans (Skinner et al., 1994; Ketten et al., 1998); by using the same 3D reconstructions to obtain more precise estimates of electrode location in the cochlea; and by obtaining physiological and behavioral data that may help determine the characteristic frequency of the neurons stimulated by different electrode pairs.

Stimulus materials

Method-of-adjustment task

The stimulus set consisted of 330 synthetic, steady-state isolated vowels that varied from one another in their first and second formants in 0.377 Bark increments. The vowels were generated using the Klatt 88 synthesizer (Klatt and Klatt, 1990). The Bark increment size was chosen as a close approximation of the just noticeable difference for vowel formants of Flanagan (1957). The F1 and F2 values for this stimulus set ranged between 2.63 Z (250 Hz)–7.91 Z (900 Hz) and 7.25 Z (800 Hz)–15.17 Z (2800 Hz). These ranges were chosen to represent the full range of possible values for speakers of American and British English, and were successfully used in piloting the present experiment and in an earlier method-of-adjustment study of vowel perception in normal-hearing listeners (Johnson et al., 1993). All of the other synthesis parameters for this stimulus set also followed Johnson et al. (1993). The formulas for calculating the values of the most relevant synthetic parameters are summarized in Table IV. The F0 parameter was varied to generate two sets of the 330 stimuli, one representing a male voice and one representing a female voice. All of the synthetic speech sounds were presented at a 70 dB C-weighted SPL listening level. The stimuli were presented over Beyer Dynamic DT-100 headphones for normal-hearing listeners, and over an Acoustics Research loudspeaker for CI users.

Vowel identification task

The vowel identification task employed a closed-set procedure that used nine vowels in an “h-vowel-d” format. The stimuli were digitized from the female vowel tokens of the Iowa laserdisc (Tyler et al., 1987). Only steady-state vowels (no diphthongs) were used from this stimulus set. There were three separate productions of each vowel. Listeners were administered three lists that consisted of five repetitions of each vowel and three practice tokens. The stimuli were presented at a level of 70 dB C-weighted SPL over an Acoustics Research loudspeaker.

Vowel identification task

The vowel identification task, administered only to the CI users, was a closed-set speech perception task in which three separate tokens of each of nine /hVd/ tokens were presented in random order, one at a time. The CI users had to say which one of the nine stimuli they thought they heard by responding verbally. They were instructed to guess if they did not know which vowel was presented. All subjects heard a total of at least 15 presentations of each vowel (except CI5 who heard 10). The subjects' responses were tabulated and scored for total percentage of correct responses.

F1 jnd task

The F1 jnd task, administered only to the CI users, required listeners to make an absolute judgment. The stimuli for this task were labeled "1" through "7" in order of increasing F1. In this task, all stimuli were played in sequence several times, so the subjects could become familiar with the stimuli. The stimuli were then presented ten times each in random order and subjects were asked to identify the stimulus that was presented using one of the seven responses. The subject's response and the correct response were displayed on the computer monitor before the presentation of the next stimulus. After each block of 70 trials (10 presentations of each of 7 stimuli), the mean and standard deviations of the responses to each of the 7 stimuli were calculated. The d' for each pair of successive stimuli was calculated as the difference of the two means divided by the average of the two standard deviations. These d' measurements were then cumulated to calculate a cumulative d' curve, which provided an overall measure of the subject's ability to discriminate and pitch rank the seven stimuli (Durlach and Braida, 1969; Levitt, 1972). To calculate the cumulative d' curves, we followed the common assumption that the maximum possible value of d' was three. The average jnd (defined as the mean stimulus difference resulting in d'=1) was calculated based on the cumulative d' curve. Given that the F1 range spanned by the seven stimuli was 600 Hz, the jnd was defined as (600/cumulative d'). At least eight blocks of 70 trials were administered in order for all subjects to reach a plateau in performance as measured by the cumulative d'. The cumulative d' reported here is the average of the best two blocks for each subject. The normative mean cumulative d' from this procedure was calculated to be 53 Hz, based on a pilot study with six normal-hearing listeners.

Results

Normal-hearing participants

The normal-hearing listeners were expected to

select vowel category centers with first and second formant values that corresponded to those typical in vowel production in American English. Figure 1 shows the mean vowel categories obtained from the group of normal-hearing subjects, for the male-voice stimulus set.² Each category center is shown along with error bars denoting the "size" of each category, that is, the relative spread of the category in both formant dimensions. The error bars represent the standard deviation from the category centers in both formant dimensions. In panel (A) of this figure, on the left, all of the ratings have been used to calculate the center and size of all ten vowel categories. Panel (B) of this figure, on the right, shows the vowel space of normal-hearing subjects calculated using only ratings of four and above. The rating of four was chosen because it was the highest rating that still allowed for category sizes to be calculated for all ten vowels of all of the normal-hearing and CI participants. The center of each category was determined by weighting the Bark values (in each dimension) of all synthetic stimuli that were chosen by their goodness ratings, and then averaging the weighted values.

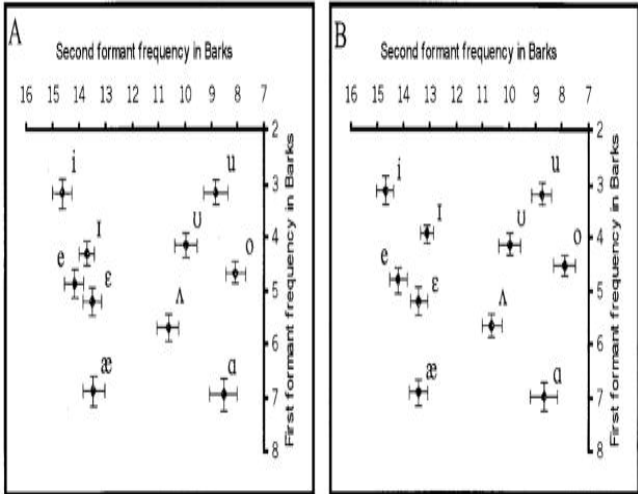


FIG. 1: The mean vowel space of normal-hearing listeners, calculated (a) using all of the ratings provided and (b) using only ratings of four or above. The perceptual spaces shown in Fig. 1 for the normal-hearing listeners demonstrate that the method-of-adjustment technique for measuring vowel categories can be successfully used to generate vowel spaces that display the typical intervowel relationships that have been observed in F1 by F2 spaces generated from vowel production data (Peterson and Barney, 1952; Hillenbrand et al., 1995).

presentation condition, suggests to us that these children relied primarily on visual-spatial encoding of the target sequence to perform the task. These results were obtained despite the fact that many of these cochlear implant children did well on the auditory WISC digit span task and on the auditory-only presentation condition of the memory game.

In summary, the present results suggest that even those cochlear implant children who are able to accurately identify speech signals in isolation, may not have phonological working memory mechanisms or processing strategies that are developed to a point equivalent to chronologically age-matched normal-hearing children. This outcome would not exactly be surprising, as many important milestones in the development of speech perception and memory are reached during the first 2 yr of life (Aslin, Jusczyk, & Pisoni, 1998; Jusczyk, 1997). Despite their prelingually deafened status, most of the cochlear implant users reported on in this paper received their implant at a point in time when the FDA did not permit implantation of children under 2 yr of age. Additionally, because the implantation procedure requires that candidates show a demonstrated failure to benefit from conventional hearing aids, we can be fairly certain that most of these 8- and 9-yr-old children had received only minimal auditory input for at least one quarter to one third of their lives. It should not be surprising, then, that the encoding strategies and working memory mechanisms of pediatric cochlear implant users seem to differ measurably from those of normal-hearing children.

Ongoing research in our lab is attempting to describe in more detail how these encoding/rehearsal mechanisms differ, and what kind of developmental changes can be observed or effected in these children. Increasingly, clinicians are beginning to see pediatric cochlear implant users that have reached ceiling levels of performance on the traditional standardized measures of speech perception and spoken word recognition that are typically used with this population—and yet these children are still clearly having problems with reading and other more advanced language skills that are based on listening, phonological encoding, and other metalinguistic abilities. Further investigation of how pediatric cochlear implant users engage in cognitive processing of information originating from this reintroduced sensory input modality may help us develop new assessment and treatment techniques (Pisoni, 2000). Eventually we would like to answer the question of whether individual differences in the function of particular components of working memory within the pediatric cochlear implant pop

ulation might have a meaningful causal relation to the level of verbal language skill attained by individual children. The present research begins to address this important issue because it provides some of the first behavioral data on working memory in pediatric cochlear implant users involving tasks in which the potential contribution of each available sensory modality was varied.

References

1. Early Alzheimer’s Disease: Patterns of Functional MRI Activation-The Neural Substrates of Semantic Memory Deficits. Am J Bra Dis and Tum. 2018; 1(1): 001-010.
2. H Chahal, S W D Souza, A J Barson and P Slater. How to develop human brain using magnesium of N-methyl-D-aspartate receptors, Am J Bra Dis and Tum. 2018; 1(1): 001-005.
3. F S LaBella, et al. Concepts and correlations related to general anaesthesia and cytochrome P450 oxygenases. Am J Anest and Pai med. 2018; 1(1): 01-05.
4. Hazim J Safi, et al. The long term method with the elephant trunk for the repair of aortic aneurysms. Am J Anest and Pai med. 2018; 1(1): 001-008.
5. Yoshitaka Fujii, et al. Diaphragmatic Fatigue is treated with Inhaled Aminophylline Therapy in an Experimental Canine procedure. Am J Anest and Pai med. 2018; 1(1): 001-003.
6. O Demirkiran, et al. Complications in patients with Crush syndrome after the Marmara earthquake. Am J Anest and Pai med. 2018; 1(1): 001-005.
7. Qi Wei, et al. Laparoscopic choledochotomy after Biliary drainage: Study. Am J Anest and Pai med. 2018; 1(1): 001-007.
8. Mark Palazzo, et al. Unilateral Babinski/Plantar Reflex - Acute Inflammatory Demyelinating Polyneuropathy. Am J Anest and Pai med. 2018; 2(1): 01-02.
9. Hakan Alfredson, et al. Achilles and patellar tendon operations performed in local anesthesia, Am J Anest and Pai med. 2018; 1(1): 001-002.
10. Naemeh Nikvarz, et al. Evaluation The Analgesic Effect of Duloxetine Drug in Burn Patients. Am J Anest and Pai med. 2019; 2(1): 01-07.
11. Chuandong Zheng, et al. Intravascular Plaque: Cause for Radial Arterial Catheterization Failure. Am J Anest and Pai med. 2019; 2(1): 01-05.
12. Laura Tyler Perryman, et al. Wireless Dorsal Root Ganglion Stimulation: An Introduction and Early Experience with the New Approach for Chronic Pain Management. Am J Anest and Pai med. 2019; 2(1): 01-04.
13. Lazraq Mohamed, et al. Pediatric Pre-Anesthesia Consultation: What are Parents Expectations?. Am J Anest and Pai med. 2019; 2(1): 01-02.
14. Alaa Ali M. Elzohry, et al. Safety and Efficacy of Intraperitoneal Irrigation of Levo-Bupivacaine plus Morphine in Patients Undergoing Major Abdominal Cancer Surgeries. Am J Anest and Pai med. 2019; 2(1): 01-07.
15. Yildiz K, et al. Comparison between Anesthesia Methods In Orthopaedics Initiatives of Upper Extremity. Am J Anest and Pai med. 2019; 2(2): 01-03.
16. Jianming Liu, et al. The Analgesic Effects Nalbuphine Hydrochloride Combined With Sufentanil for Patients after Thoracoscopic Lobectomy. Am J Anest and Pai med. 2019; 2(2): 01-03.
17. Fudong Shi, et al. The Patient Controlled Intravenous Analgesia of Dezocine on the Elderly Patients After Orthopedic Surgery. Am J Anest and Pai med. 2019; 2(1): 01-04.
18. GE Meglia, et al. Investigation in blood Leukocytes and Neutrophils in Periparturient Dairy Cow. Sci J of Ani and Vet Sci. 2018; 1(1): 001-009.

19. G E Duhamel, et al. DNA Sequence Analysis of an Immunogenic Glucose-Galactose Mglb. *Sci J of Ani and Vet Sci.* 2018; 1(1): 001-009.
20. David G. White, et al. Chloramphenicol and Florfenicol Resistance in *Escherichia Coli* of Characterization . *Sci J of Ani and Vet Sci.* 2018; 1(1): 001-006.
21. N B Alhaji, et al. Anophthalmia and Choanal Atresia In Two Months Old Kid. *Sci J of Ani and Vet Sci.* 2018; 1(1): 001-004.
22. Christopher W Olsen, et al. Isolation and Characterization of H4N6 Avian and Influenza Viruses. *Sci J of Ani and Vet Sci.* 2018; 1(1): 001-0025.
23. Teresa Lopez-Arteaga, et al. Apathy as a Psychiatric Manifestation of Meningioma. *Am J Bra Dis and Tum.* 2018; 1(1): 001-004.
24. David R Murdoch, et al. The Use of Brain Natriuretic Peptide- Whole Blood can be Measured, *Am J Bra Dis and Tum.* 2018; 1(1): 001-003.
25. Stefan Brocke, et al. Antibodies to Integrin $\alpha 4$ and CD44, but not CD62L, Prevent CNS Inflammation and Experimental Encephalomyelitis by Blocking Secondary Leukocyte Recruitment. *Am J Bra Dis and Tum.* 2018; 1(1): 001-006.
26. Andrew J Saykin, et al. Early Alzheimer's Disease: Patterns of Functional MRI Activation-The Neural Substrates of Semantic Memory Deficits. *Am J Bra Dis and Tum.* 2018; 1(1): 001-010.
27. P Slater, et al. How to develop human brain using magnesium of N-methyl-D-aspartate receptors, *Am J Bra Dis and Tum.* 2018; 1(1): 001-005.
28. Clyde W Hodge, et al. The Paraventricular Nucleus Interactively Modulate Ethanol Consumption -Norepinephrine and Serotonin Receptors, *Am J Bra Dis and Tum.* 2018; 1(1): 001-005.
29. Paulo C Carvalho, et al. Bioinformatics grid application in simple - Squid. *Sci J Biome and Biost.* 2018; 1(1): 001-004.
30. Mahmoud A E Abdelrahman, et al. On The New Exact Solutions for the Nonlinear Models Arising In Plasma Physics. *Sci J Biome and Biost.* 2018; 1(1): 001-004.
31. Weicheng Shen, et al. Based on Personal Identification- Automated Biometrics. *Sci J Biome and Biost.* 2018; 1(1): 001-002.
32. V Prasathkumar, et al. Fingerprint Biometric System - Using of Personal Authentication. *Sci J Biome and Biost.* 2018; 1(1): 001-003.
33. Savita Choudhary, et al. Software Development Environment : Design of Biometric Based Transaction System. *Sci J Biome and Biost.* 2018; 1(1): 001-003.
34. D J Lawrence, et al. Measuring the effectiveness in reliability and validity of a visual function outcomes instrument in cataract surgery. *Sci J Biome and Biost.* 2018; 1(1): 001-004.
35. Z Suvakovic, et al. Evaluation of early detection of gastric cancer requires more than gastroscopy. *Anna of Can Ther and Phar.* 2018; 1(1): 05.
36. Ho GY, et al. Informing and involving personalised computer based data for cancer patients. *Anna of Can Ther and Phar.* 2018; 1(1): 001-005.
37. Ray Jones, et al. Prostate Cancer Risk is associated with Polymorphism of Insulin gene. *Anna of Can Ther and Phar.* 2018; 1(1): 001-005.
38. : Jean-Pierre J. Issa, et al. Role of DNA Methylation in Tumor Suppressor Gene Silencing in Colorectal Cancer. *Anna of Can Ther and Phar.* 2018; 1(1): 001-008.
39. Jules J Berman, et al. Histological classification of tumour and molecular analysis meets Aristotle. *Anna of Can Ther and Phar.* 2018; 1(1): 001-005.
40. Kafil Akhtar, et al. Tuberculosis of the Tongue with Coexistent Squamous Cell Carcinoma: An Interesting Case Presentation, *Anna of Can Ther and Phar.* 2018; 1(1): 001-002.
41. Serafin Morales Murillo, et al. Vitamin D as A Prognostic Factor in Triple Negative Breast Cancer. *Anna of Can Ther and Phar.* 2019; 2(1): 01-08.
42. Ahmet Fuat, et al. A Qualitative Study of Accurate Diagnosis and Effective Management of Heart Failure in Primary Care. *Am J of Card and Cardiovas Disc.* 2018; 1(1): 01-05.
43. Jesús Millán Núñez-Cortés, et al. Prescription Habits for Statins in Patients with Impaired Glucose Metabolism. Results of a program with Focus Groups to Assess the Selection Criteria. *Am J of Card and Cardiovas Disc.* 2019; 1(1): 01-04.
44. G D Kolovou, et al. Evaluation of Postprandial hypertriglyceridaemia in patients with Tangier disease. *Am J of Card and Cardiovas Disc.* 2018; 1(1): 01-04.
45. Brian O rourke, et al. Determination of The Mitochondrial Redox Waves and Subcellular Metabolic Transients in Heart Cells. *Am J of Card and Cardiovas Disc.* 2018; 1(1): 01-04.
46. Shuixiang Yang, et al. Radiofrequency Ablation Treating Atrial Fibrillation Can Reverse the Changes of Mirnas Regulating Ion Channel Proteins. *Am J of Card and Cardiovas Disc.* 2018; 1(1): 01-08.
47. Hadi abdulsalam Abo Aljadayel, et al. Penetrating War Cardiac and Great Vessels Injury, Surgical Outcome Analysis in 24 Patients. *Am J of Card and Cardiovas Disc.* 2018; 1(2): 01-05.
48. Hatice Yorulmaz, et al. Assessment of the Death Anxiety and Death Depression Levels of Cardiac Patients. *Am J of Card and Cardiovas Disc.* 2019; 2(1): 01-06.
49. Camara Abdoulaye, et al. Cardiomyopathie Du Peripartum Complicee D'accident Vasculaire Cerebral Cas D'une Guinéenne De 19ans : Cas Clinique. *Am J of Card and Cardiovas Disc.* 2019; (1): 01-03.
50. Sergio F. Estrada-Orihuela, et al. Lasalocid, Interrupts and Reverses, Within One Minute, The Myocardial Damage Caused By Coronary Anoxia Reperfusion in Rat Heart. *Am J of Card and Cardiovas Disc.* 2019; (1): 01-05.
51. Jesus Millan Nunez-Cortes, et al. Prescription Habits for Statins in Patients with Impaired Glucose Metabolism. Results of a program with Focus Groups to Assess the Selection Criteria. *Am J of Card and Cardiovas Disc.* 2019; 1(1): 01-06.
52. Federico Cacciapuoti, et al. The Dilemma of Diastolic Heart Failure. *Am J of Card and Cardiovas Disc.* 2019; 1(1): 01-03.
53. Elad Boaz, Bowel Ischemia and Vascular Air-Fluid Levels. *Anna Cas Rep and Ima Surg.* 2018; 1(1): 001-00.
54. Sinisa Franjic, et al. A Patient With A Maxillofacial Problem. *Anna Cas Rep and Ima Surg.* 2018; 1(1): 001-004.
55. Davidson W, et al. Case Presentation: Hantavirus pulmonary syndrome [HPS]. *Anna Cas Rep and Ima Surg.* 2018; 1(1): 001-005.
56. Farid ZM, et al. Uropathy Secondary Chronic obstructive to Ureter Inguinal Herniation. *Anna Cas Rep and Ima Surg.* 2018; 1(1): 001-002.
57. De Letter DJ, et al. Cornual Molar Ectopic Pregnancy Diagnosis and Treatment. *Anna Cas Rep and Ima Surg.* 2018; 1(1): 001-003.
58. Ameni Touati, et al. Silver Russell Syndrome: Case Reports from North Africa and Review on The Literature. *Anna Cas Rep and Ima Surg.* 2019; 1(1): 001- 004.
59. Kunst WM, et al. Case Reports and Review of Spontaneous Rupture of Hyperreactive Malarial Splenomegaly [HMS]. *Anna Cas Rep and Ima Surg.* 2018; 1(1): 001-005.
60. F Hanefeld, et al. A Review of The Literature an Emerging Community Pathogen methicillin-Resistant *Staphylococcus*. *Anna Cas Rep and Ima Surg.* 2018; 1(1): 001-0011.
61. Page W Caufield, et al. Evidence for a Discrete Window of Infectivity. *Am J Den and Ora Car.* 2018; 1(1): 001-006.
62. Robert T Dirksen, et al. Dihydropyridine Receptors and Ryanodine Receptors: Bi-Directional Coupling . *Am J Den and Ora Car.* 2018; 1(1): 001-009.

Cite this article: Disorders of Peripheral Implications for the study of auditory adaptation to spectral shift: vowel spaces. *Am J Rhin and Otol.* 2019; 1(1): 001-009.

63. IJ Jacobs, et al. Cancer and Intraepithelial Neoplasia-Tissue-specific apoptotic effects of the p53 codon 72 polymorphism. *Am J Den and Ora Car.* 2018; 1(1): 001-003.
64. Iain L C Chapple, et al. Human Immunodeficiency Virus disease in oral health significances. *Am J Den and Ora Car.* 2018; 1(1): 001-007.
65. H Larjava, et al. Activity of $\alpha\beta 6$ Integrin in Oral Leukoplakia. *Am J Den and Ora Car.* 2018; 1(1): 001-005.
66. Siddharth Kothari, et al. Effectiveness of Counselling and Home Care Self-Management Strategies in Reducing Masticatory Muscle Pain: A Review. *Am J Den and Ora Car.* 2019; 2(1): 001-007.
67. Betania Maria Soares, et al. Use of Blue LED and Curcumin for Photosensitization of *Candida Albicans*. *Am J Den and Ora Car.* 2019; 2(1): 001-005.
68. Jing Guo, et al. Advances in Methods of Maxillary Transverse Expansion. *Am J Den and Ora Car.* 2019; 2(1): 01-05.
69. Dario C. Altieri, et al. Cell division by p34cdc2 phosphorylation of survivin- Regulation. *Sci J of Der and Ven.* 2018; 1(1): 001-005.
70. Axel Trautmann, et al. Eczematous dermatitis: T cell and keratinocyte apoptosis plays a key pathogenetic. *Sci J of Der and Ven.* 2018; 1(1): 001-007.
71. JD Fine, et al. Epidermolysis bullosa Cardiomyopathy in inherited. *Sci J of Der and Ven.* 2018; 1(1): 001-004.
72. NE Fusenig, et al. Human Skin Angiogenic Switch Occurs Squamous Cell Carcinomas. *Sci J of Der and Ven.* 2018; 1(1): 001-007.
73. Tapani Tuomi, et al. Water- Damaged Building and Mycotoxins in Crude Building Materials. *Sci J of Der and Ven.* 2018; 1(1): 001-005.
74. John S Davies, et al. The Use of Social Media among Doctors Under taking a Post-Graduate Endocrinology Diploma. *Sci J Endo and Meta.* 2018; 1(1): 001-004.
75. Juan J Gagliardino, et al. By Short-Term Dietary Manipulation: The Endocrine Pancreas Activity of Tyrosine Hydroxylase. *Sci J Endo and Meta.* 2018; 1(1): 001-005.
76. Colin A. Leech, et al. The Glucose Dependent in Pancreatic β -Cells : Voltage-Independent Calcium Channels Mediate Slow Oscillations of Cytosolic CalciumPancreatic β -Cells. *Sci J Endo and Meta.* 2018; 1(1): 001-009.
77. Colin A. Leech, et al. The Voltage-Independent Activation of Inward Membrane Currents and Elevation of Intracellular Calcium in HIT-T15 Insulinoma CellsPituitary Adenylate Cyclase-Activating Polypeptide Induces. *Sci J Endo and Meta.* 2018; 1(1): 001-008.
78. Suhail AR Doi, et al. Making Use Of Combined Criteria - Diagnostic Criteria For Diabetes. *Sci J Endo and Meta.* 2018; 1(1): 001-006.
79. Maria I Borelli, et al. Effect Of Endogenous Islet Catecholamines Possible Modulatory On Insulin Secretion. *Sci J Endo and Meta.* 2018; 1(1): 001-005.
80. Louis Irwin, et al. Effect of exercise in combination with dietary nopal and zucchini on chronic and acute glucohomeostasis in genetically obese mice. *Inte J Expe Bio.* 2018; 1(1): 001-005.
81. Vijaya Saradhi Settaluri, et al. Validation of Non Essential Amino Acids and Total Protein Content in Different Categories of Tea. *Inte J Expe Bio.* 2018; 1(1): 01-04.
82. Patrick D Craig, et al. T Antigen: Polyomavirus Middle of Natural Biology. *Inte J Expe Bio.* 2018; 1(1): 001-007.
83. Yoshinori Ohsumi, et al. The HIV Coreceptor CCR5 - Recycling and Endocytosis. *Inte J Expe Bio.* 2018; 1(1): 001-008.
84. Marino Zerial, et al. Elicitation of the Angiogenic Phenotype1: Transforming Myc Protein for In Vivo. *Inte J Expe Bio.* 2018; 1(1): 001-008.
85. Zhang Y, et al. Odorant Receptor In Mammali : The *Caenorhabditis Elegans* Seven-Transmembrane Protein ODR-10 Functions on Cells. *Inte J Expe Bio.* 2019; 1(1): 001-008.
86. Kazuo Maeda, et al. Improved Outcome with Novel Studies in Fetal Monitoring. *Sci J of Gyne and Obste.* 2019; 2(1): 001-004.
87. Sunil J. Wimalawansa, et al. Vitamin D Deficiency-Related Reproductive Consequences. *Sci J of Gyne and Obste.* 2019; 2(1): 001-006.
88. Munch A, et al. Investigation in blood Leukocytes and Neutrophils in Periparturient Dairy Cow. *Sci J of Gas and Hepa.* 2018; 1(1): 001-006.
89. Jie Song Hua, et al. Primary *Helicobacter Pylori* Resistance to Clarithromycin and Metronidazole in Singapore. *Sci J of Gas and Hepa.* 2018; 1(1): 001-003.
90. Paul Moayyedi, et al. A Systematic Review and Economic Analysis: Proton Pump Inhibitors in Nonulcer Dyspepsia Efficacy. *Sci J of Gas and Hepa.* 2018; 1(1): 001-003.
91. Zhen-Ning Wang, et al. Gastric Cancer: Collagen IV Expression and Biological Behavior. *Sci J of Gas and Hepa.* 2018; 1(1): 001-002.
92. Zhen-Ning Wang, et al. A Possible Pathophysiologic Contribution to Necrotizing Enterocolitis: Human Intestine Inflammation. *Sci J of Gas and Hepa.* 2018; 1(1): 001-006.
93. Paul M Wassarman, et al. Egg Interaction during Mammalian Fertilization in the Molecular Basis of Sperm. *Sci J of Gyne and Obste* 2018; 1(1): 001-006.
94. Mary Lou Moore, et al. Breastfeeding Benefits Support -Research. *Sci J of Gyne and Obste* 2018; 1(1): 001-002.
95. Pepita Gimenez-Bonafe, et al. Preservation of Fertility in Patients with Cancer. *Sci J of Gyne and Obste* 2018; 1(2): 001-006.
96. Yueyang F Fei, et al. Non-Hemorrhagic Unilateral Adrenal Infarct In Pregnancy: A Case Report. *Sci J of Gyne and Obste.* 2019; 1(1): 001-002.
97. Karen Pierre, et al. Protein-Energy Adequacy of Dialysis Patients in Trinidad and Tobago. *Am J of Nep and Ther.* 2018; 1(1): 01-05.
98. Balakrishna N, Tenckhoff Catheter Surgical under Local Anesthesia. *Am J of Nep and Ther.* 2018; 1(1): 001-003.
99. J T Ohlsson, et al. Man in angiotensin and noradrenaline inhibits the Endothelin. *Am J of Nep and Ther.* 2018; 1(1): 001-005.
100. David J, et al. Apoptosis and Ischemic Renal Injury Reduce the Guanosine Supplementation. *Am J of Nep and Ther.* 2018; 1(1): 001-005.
101. R W Baldeweg, et al. Tumor-induced osteomalacia : Cloning and characterization of Fibroblast Growth Factor 23. *Am J of Nep and Ther.* 2018; 1(1): 001-006.
102. Amitabh Arya, et al. Post Pyeloplasty Follow Up In Children And Adolescents: Diuretic Renography Or Renal Ultrasonography? *Am J of Nep and Ther.* 2019; 2(1): 001-005.
103. Amitabh Arya, et al. Post Pyeloplasty Follow Up In Children And Adolescents: Diuretic Renography Or Renal Ultrasonography? *Am J of Nep and Ther.* 2019; 2(1): 001-005.
104. Richard Lechtenberg, et al. Tau Interferon in Multiple Sclerosis. *Amer J Neur & Neurophysi.* 2018; 1(1): 001-002.
105. Eva Guy Rodriguez, et al. Discussion of the differential diagnosis of bilateral thalamic lesions-Bilateral thalamic infarcts due to occlusion of the Artery of Percheron. *Amer J Neur & Neurophysi.* 2018; 1(1): 001-004.
106. Yhashi Chang, et al. IVIg for Miller Fisher syndrome: Cerebral infarction. *Amer J Neur & Neurophysi.* 2018; 1(1): 001-002.
107. Fredrick J. Seil, et al. T cell responses to Myelin Antigens and Antimyelin Antibodies. *Amer J Neur & Neurophysi.* 2018; 1(1): 001-005.
108. Y Niimi, et al. Embolization of Spinal Cord AVMs: Neurophysiologic Provocative Testing. *Amer J Neur & Neurophysi.* 2018; 1(1): 001-002.

109. Ameni Touati, et al. Some Reducibility Results for Differentiable Sets. *Amer J Neur & Neurophysi.* 2019; 1(1): 001-005.
110. Chrisostomos Sofoudis, et al. Sofoudis C. Septic Abortion Accompanied with Dessiminated Intravascular Coagulation and Acute Cardiomyopathy Presentation of a Rare Case and Mini Review. *Am J Nur & Pract.* 2018;1(1): 001-00.
111. Nick Jones, et al. Nurse Practitioners and Family Physicians Ethics Health Care Services. *Am J Nur & Pract.* 2018; 1(1): 001-005.
112. Thomas R A, et al. Human Infants Learning by Prenatal and Postnatal Flavor . *Am J Nur & Pract.* 2018; 1(1): 001- 006.
113. D. J Wise, et al. A Randomized, Double-Blind, Placebo-Controlled - Milk Production in Mothers of PremaTure Newborns Domperidone Drug effect. *Am J Nur & Pract.* 2018; 1(1): 001-005.
114. Bronagh Bufton, et al. Effects of Nursing Homes Ownership Compromise the Quality of Care. *Am J Nur & Pract.* 2018; 1(1): 001-005.
115. Kerstin Ekberg, et al. How Physicians Deal With the Task of Sickness Certification in Cause-Based and Comprehensive Disability Systems – A Scoping Review. *Am J Nur & Pract.* 2019; 2(1): 01-10.
116. Michael J. Vives, et al. Factors in Choosing the Surgical Approach: Cervical Spondylotic Myelopathy. *Am J Orth and Rhe.* 2018; 1(1): 001-004.
117. M. Runge, et al. Geriatric Patients in Balance Training and Exercise. *Am J Orth and Rhe.* 2018; 1(1): 001-003.
118. Ukoha Ukoha Ukoha, et al. Nutrient Foramina in Long Bones : Study. *Am J Orth and Rhe.* 2018; 1(1): 001-003.
119. Zhiquan An, et al. Human Humeral Diaphysis of the Nutrient Foramina : Anatomical Study : Study. *Am J Orth and Rhe.* 2018; 1(1): 001-007.
120. K-P Günther, et al. Hip Replacement in Rates: International Variation: Study. *Am J Orth and Rhe.* 2018; 1(1): 001-005.
121. Saeed Taj din, et al. Level of Physical Activity among Diabetic Patients of Rural and Urban Areas. *Am J Orth and Rhe.* 2019; 2(1): 001-004.
122. Carolina Caleza Jiménez, et al. Breastfeeding, Bed-Sharing and Early Childhood Caries. Is There An Association? A Review of the Literature. *Am J Pedi and Heal care.* 2018; 1(1): 001-004.
123. Katarzyna Niewiadomska-Jarosik, et al. Lipid Profile in Children Born As Small for Gestational Age . *Am J Pedi and Heal care* 2018; 2(1): 01-03.
124. Mustafa Aydin, et al. Antibiotic Susceptibility Pattern and Clinical Features of Klebsiella Sepsis in Newborn Infants. *Am J Pedi and Heal care* 2019; 1(1): 01-04.
125. H Dele Davies, et al. Necrotizing Fasciitis- Flesh-Eating Bacteria Disease. *Am J Pedi and Heal care* 2019; 1(1): 01-06.
126. Marie Westwood, et al. The diagnosis of urinary tract infection (UTI) in children under five years: Rapid tests and urine sampling techniques. *Am J Pedi and Heal care* 2019; 1(1): 01-09.
127. Folkert Fehr, et al. What Entrustable Professional Activities Add To a Primary Care Residency Curriculum. *Am J Pedi and Heal care* 2019; 2(1): 01-06.
128. Sonya Martin, et al. Spatially Modulated Illumination Microscopy□D measures the size of Biological Nanostructures . *Ann of Phar Nano Tech and Nanomedi.* 2018; 1(1): 01-05.
129. Sonya Martin, et al. Genetic analysis of Fis interactions with their binding sites. *Ann of Phar Nano Tech and Nanomedi.* 2018; 1(1): 01-07.
130. John H Reif, et al. Nucleation assembly of DNA tile complexes is directed by barcode-patterned lattices. *Ann of Phar Nano Tech and Nanomedi.* 2018; 1(1): 01-07.
131. Thomas H LaBean, et al. Self assembly of DNA nanotubes from triple-crossover tiles as templates for conductive nanowires. *Ann of Phar Nano Tech and Nanomedi.* 2018; 1(1): 01-05.
132. Ulrich Kettling, et al. Dual-Photon Fluorescence Coincidence Analysis: Rapid quantification of Enzyme activity. *Ann of Phar Nano Tech and Nanomedi.* 2018; 1(1): 01-05.
133. Ahmed R. Gardouh, et al. Design, Optimization and In-Vitro Evaluation of Antifungal Activity of Nanostructured Lipid Carriers of Tolnaftate *Ann of Phar Nano Tech and Nanomedi.* 2019; 2(1): 01-05.
134. Mohammed Khalid, et al. Khalid M. Predictors of Prognosis in Pulmonary Hypertension. *Anna Pul and Crit Car Med.* 2018; 1(1): 001-004.
135. Abdullah Alsaeeedi, et al. The Prevalence of Smoking among sample of Kuwait Asthmatics and its impact on the response of the treatment, *Anna Pul and Crit Car Med.* 2018; 1(2): 001-002.
136. Nicolau Beckmann, et al. Resolving the Oedematous Signals Induced by OVA Challenge in the Lungs of Actively Sensitised Rats. *Anna Pul and Crit Car Med..* 2018; 1(1): 01-06.
137. Thomas J walsh, et al. Investigate the performance of non-invasive diagnostic tests such as galactomannan enzyme immunoassay and quantitative Caspofungin in the early diagnosis of invasive aspergillosis (IA). *Anna Pul and Crit Car Med..* 2018; 1(1): 01-06.
138. Charles B. Huddleston, et al. Lung Transplantation in pediatrics. *Anna Pul and Crit Car Med..* 2018; 1(1): 01-05.
139. Jeffrey P. Lamont, et al. Comparison of valved vs non-valved implantable ports for vascular access:A randomized trial. *Anna Pul and Crit Car Med..* 2018; 1(1): 01-03.
140. D Inwald, et al. Risk and relevance of open lung biopsy in Nonneonatal extracorporeal membrane oxygenation (ECMO) patients. *Anna Pul and Crit Car Med..* 2018; 1(1): 01-04.
141. Guillaume Mortamet, et al. Does Esophageal Pressure Monitoring Reliably Permit To Estimate Trans pulmonary Pressure In Children?. *Anna Pul and Crit Car Med..* 2018; 2(2): 01-05.
142. Yang Jin, et al. Extracellular Vesicle-Shuttling MicroRNAs Regulate the Development of Inflammatory Lung Responses. *Anna Pul and Crit Car Med..* 2018; 1(2): 01-04.
143. Nicola Clemente, et al. Pneumonectomy As A Salvage Therapy: A Rare Indication For A Gastric Malt Lymphoma Disseminated To The Lung. *Anna Pul and Crit Car Med..* 2018; 1(2): 01-04.
144. Nicola Clemente, et al. Pneumonectomy As A Salvage Therapy: A Rare Indication For A Gastric Malt Lymphoma Disseminated To The Lung. *Anna Pul and Crit Car Med..* 2018; 1(2): 01-04.
145. Victor Chew, et al. Pulmonary Cement Embolism. *Anna Pul and Crit Car Med.* 2019; 2(1): 01-02.
146. Victor Chew, et al. An Unusual Cause of a Tension Pneumothorax. *Anna Pul and Crit Car Med.* 2019; 2(1): 01-03.
147. Mark C. Lavigne, et al. A Performance Summary of Agents Used in Oral Care for Non-Ventilated and Mechanically-Ventilated Patients. *Anna Pul and Crit Car Med.* 2019; 2(2): 01-34.
148. Elisangela Hermes, et al. Psychomotricity in Vestibular Dysfunction Therapy (VDT): A Collective Health Question. *Am J Rhin and Otol.* 2018; 1(1): 001- 005.
149. Ramtej J Verma, et al. Diethanolamine-Induced Hepatic Injury and Its Amelioration by Curcumin. *Am J Toxi and Res.* 2018; 1(1): 001-004.
150. Chee Kong Yap, et al. A Preliminary Screening of Cd and Pb Concentrations in the Some Traditional Chinese Herbal Medicines Bought From Selected Shops in Peninsular Malaysia. *Am J Toxi & Res.* 2018; 1(1): 001-004.
151. Geza Bozoky, et al. Acute Silent Non-Massive (submassive) Pulmonary Embolism. *Am J Ang and Surg .* 2018; 1(1): 001-003.

Cite this article: Disorders of Peripheral Implications for the study of auditory adaptation to spectral shift: vowel spaces. *Am J Rhin and Otol.* 2019; 1(1): 001-009.