The association between hearing impairment and polymorphisms of genes encoding inflammatory mediators in Japanese aged population

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Abstract

Background: Aging process is accompanied by a chronic sub-clinical systemic inflammation. This study aimed to assess the association between hearing impairment and polymorphisms of genes encoding cytokines deeply-committed to the inflammatory response and immune homeostasis in an elderly Japanese population. Data were collected in the Longitudinal Study of Aging surveyed biennially between 1997 and 2010. The participants without any missing information at baseline were 1,957 individuals, and the gross accumulated number of 8,675 subjects (40–89 years of age) was analyzed. Two hearing impairment criteria were taken as the better ear pure-tone average (PTABE) greater than 25 dB and greater than 40 dB. We analyzed cumulative data using generalized estimating equations to investigate the effect of 9 polymorphisms, namely, tumor necrosis factor (TNF) α, rs1800630; TNF receptor super family (TNFRSF) 1B, rs1061624; interleukin (IL)-1A, rs1800587; IL-1B, rs16944; IL-4R, rs1801275; IL-6, rs1800796; IL-10, rs1800872; IL-1 receptor-associated kinase 1 (IRAK1), rs1059702; C reactive protein (CRP), rs1130864.

Results: The odds ratios for the hearing impairment (PTABE >25 dB) risk under additive genetic model were significant in TNF-α rs1800630 and TNFRSF1B rs1061624, which were respectively 1.172 (confidence interval [CI]: 1.005-1.367), 1.211 (CI: 1.053-1.392) in model after adjustment for possible confounders. Using the criterion of PTABE >40 dB as disabling hearing impairment, the association remains significant in TNFRSF1B rs1061624, but not in TNF-α rs1800630. No other polymorphisms showed a significant association.

Conclusions: The present population-based cohort study demonstrated that TNF-α rs1800630 and TNFRSF1B rs1061624 contributed to the incremental risk of hearing impairment in the elderly. TNF-α and TNF receptor interactions play a pivotal role in the pathogenesis of the inflammatory response, and also cause programmed cell death and cell proliferation. The present observation implied the signalling cascades of TNF were involved in ear aging.

Keyword: Age-related hearing loss, Polymorphism, Longitudinal study, TNFα

Background

Chronic low-grade inflammation is the hallmark of the aging process and is now accepted as a fundamental pathogenic mechanism in the development of several age-related degeneration including cardiovascular disease [1], atherosclerosis [2], cognitive disorders [3], and type 2 diabetes [4].

The low-grade proinflammatory status appearing during the aging process has become termed ‘inflamming’, coined by Franceschi et al. [5]. Physiological aging is characterized by elevated levels of inflammatory mediators. Serum pro-inflammatory cytokines such as interleukin (IL)-6 and tumor necrosis factor (TNF) α and acute phase proteins such as C-reactive protein (CRP) show an increase with age [5]. Further, the levels of cytokines that counteract the inflammatory state, such as IL-10, are reduced with age [6] compounding the inability to maintain immune homeostasis.

The subjects in the present analyses were derived from the National Institute for Longevity Sciences - Longitudinal Study of Aging (NILS-LSA), a population-based biennial...
survey of a cohort of approximately 2,200 community dwellers. In the present study, we intended to clarify the relationship between hearing impairment and nine polymorphisms of genes encoding inflammatory mediators, namely, \textit{TNF-\alpha} C-863A, rs1800630; \textit{TNF} receptor superfamily member (\textit{TNFRSF}) 1B G593A, rs1061624; \textit{IL-1\alpha} -889C/T, rs1800587; \textit{IL-1\beta} -511C/T, rs16944; \textit{IL-4R} G1902A, rs1801275; \textit{IL-6} C-572G, also known as C-634G, rs1800796; \textit{IL-10} A-592C, rs1800872; \textit{IL-1} receptor-associated kinase 1 (\textit{IRAK1}) T587C, rs1059702; \textit{CRP} +1444 C>T, rs1130864. Our ultimate aim is to get closer to the pathophysiological elucidation of presbycusis as an age-related disease.

Results and discussion

Table 1 shows the demographic profiles of the study subjects at baseline and cumulative data. Of the 1957 participants at baseline, the mean age was 59.2 years, and rates of male and hearing impairment with the World Health Organization (WHO) criteria were 50.4% and 20.1%, respectively. Allelic and genotypic distribution of 9 single-nucleotide polymorphisms (SNPs) in subjects for the present analyses at baseline are indicated in Table 2.

The results of multivariable analysis using logistic regression with generalized estimating equations (GEEs) were demonstrated in Table 3. Of the 9 SNPs analyzed, there were significant associations of polymorphic alleles in \textit{TNF-\alpha} and \textit{TNFRSF1B} with risk increments for hearing impairment, based on the criterion of PTABE >25 dB. The odds ratios (ORs) for hearing impairment (PTABE >25 dB) risk in \textit{TNF-\alpha} rs1800630 were 1.120 (95% confidence interval [CI]: 0.979-1.281) in model 1, 1.192 (CI: 1.020-1.392) in model 2, and 1.275 (CI: 1.026-1.584) in model 3.

Using the criterion of PTABE >40 dB as disabling hearing impairment, the association between polymorphic allele and hearing impairment remained significant in \textit{TNFRSF1B} rs1061624, but not in \textit{TNF-\alpha} rs1800630. The ORs for hearing impairment (PTABE >40 dB) risk in \textit{TNFRSF1B} rs1061624 were 1.272 (CI: 1.052-1.538) in model 1, 1.293 (CI: 1.045-1.599) in model 2, and 1.275 (CI: 1.026-1.584) in model 3.

In the other 7 SNPs, no significant risks of hearing impairment in either criterion were observed in models with different combination of moderator variables.

In the present study, we assessed the contribution of polymorphic genes encoding inflammatory mediators to hearing loss in an elderly Japanese community-dwelling population. The results suggested that \textit{TNF-\alpha} rs1800630 and \textit{TNFRSF1B} rs1061624 may be involved in risk increment of age-related hearing impairment. The significant ORs observed both in two hearing impairment criteria implied a substantial impact of \textit{TNFRSF1B} rs1061624 on the susceptibility of age-related hearing impairment, since hearing impairment of the better ear means bilateral impairment and thus WHO defines disabling hearing impairment.

\textit{TNF-\alpha} is a proinflammatory cytokine that is a master regulator of vascular proatherogenic changes and the pathogenesis of inflammatory diseases [7]. It serves a function in the regulation of cell differentiation, proliferation and apoptosis, also in inflammation, innate and adaptive immune responses. \textit{TNF-\alpha} does not work independently at the cellular level. It activates intracellular signaling cascades via binding to two types of receptors, \textit{TNF} receptor superfamily member 1A (\textit{TNFRSF1A}) also known as p55 and \textit{TNFR1}) and member 1B (\textit{TNFRSF1B/ also known as p75 and TNFR2}), then exerts its pleiotropic role. \textit{TNFRSF1A} is expressed on most human cell types, while \textit{TNFRSF1B} expression is restricted to certain subpopulations of specific neuronal subtypes, endothelial cells, and cells of the immune system [8,9].

Many researchers have substantiated emergence of \textit{TNF-\alpha} and other mediators in the inflammatory response of inner ear. \textit{TNF-\alpha} is secreted mainly by activated macrophages, monocytes, T cells, B cells, and fibroblasts [10]. In the cochlea, it is reported that spiral ligament fibrocytes have been shown to release chemokines including \textit{TNF-\alpha} when challenged with otitis media pathogens [10]. Other authors have reported the role of inflammatory cytokines, particularly \textit{TNF-\alpha}, in inner ear inflammation [11-13]. Studies using immunohistological and genetic amplification analyses from University of Debrecen in Hungary reported that elevated \textit{TNF-\alpha} receptor expression correlates with

| Table 1 Demographics of study subjects |
|---------------------------------------|
| Number                              | 1957 | 8675 |
| Sex, % male                         | 50.4 | 51.5 |
| Mean age (SD), year                  | 59.2 (10.9) | 62.0 (10.4) |
| Average hearing threshold for frequencies (dB) | 17.3 (12.0) | 19.7 (12.1) |
| Hearing impairment with the WHO criteria, % | 20.1 | 25.0 |
| Ear disease history, %               | 27.4 | 30.2 |
| Occupational noise exposure, %      | 22.8 | 21.4 |
| Hypertension, %                     | 24.9 | 29.6 |
| Diabetes, %                         | 7.5  | 8.2  |
| Dyslipidemia, %                     | 16.7 | 19.7 |
| Stroke, %                           | 2.6  | 3.8  |
| Cardiac disease, %                  | 11.8 | 13.4 |
| Smoking, %                          | 44.2 | 43.2 |
histologic activity of otosclerosis [14]. Also they reported that histologic otosclerosis exhibits a strong correlation with measles virus presence in the bone and related inflammation with TNF-α and speculated that the TNF-α chronically released from the foci enters the inner ear fluid spaces in histologically active stages of otosclerosis and may cause outer hair cell functional disorder and subsequent sensorineural hearing loss [15].

Recently inflammatory responses in inner ear under various damaging conditions including noise-overstimulation have been reported [16,17]. Fujioka, et al. evaluated the time-dependent expression of proinflammatory cytokines in noise-exposed rat cochlea, and suggested that proinflammatory cytokines, including TNF-α, IL-1 β, and IL-6 might initiate an inflammatory response and could have some role in the mechanism of noise-induced cochlear damage. Hwang, et al. evaluated tinnitus and mRNA expression levels of TNF-α, IL-1 β, and N-methyl D-aspartate receptor subunit 2B (NR2B) genes in cochlea and inferior colliculus of mice after intraperitoneal injections of salicylate, then concluded that salicylate treatment resulting in tinnitus augmented expression of the TNF-α, IL-1 β genes in cochlea and inferior colliculus [18].

One study in experimental animals demonstrated that the use of a TNF-α-neutralizing agent could prevent the sensorineural hearing loss secondary to induced pneumococcal meningitis [19]. Meanwhile, the potential cytotoxicity of TNF-α, itself on cochlear sensory cells is open to debate. Keithley, et al. tested for circulating leukocytes recruitment induced by TNF-α in the guinea pig inner ear and examined hair cells in culture directly exposed to TNF-α [20]. TNF-α infused into the guinea pig cochlear scala tympani in the absence of antigen or pathogens resulted in infiltration of leukocytes around the venules and within scala tympani. In their interpretation, TNF-α is sufficient to recruit inflammatory cells to the cochlea. However, there was no associated hearing loss as measured with a click stimulus. And TNF-α at concentrations used in the in-vivo experiments applied directly to organ of Corti explants caused minimal hair cell death. They concluded that the profound hearing loss and severe loss of inner hair cells observed in antigen-induced labyrinthitis cannot be explained by direct TNF-α mediated cytotoxicity alone.

Through extensive examinations of expression and function, some genetic variations have been shown to explain inter-individual variation. SNPs in the TNF-α, TNFRSF1A and TNFRSF1B genes have been identified, however functional data pertaining to these polymorphisms in scarce. Rs1800630 is a common functional polymorphism, involving a C to A substitution at position −863. It is located in the proximal promoter of the TNF-α gene. The presence of DNA sequence variations in regulatory region might interfere with transcription of TNF-α gene, and with increase in the susceptibility to human wide range of diseases (infectious, cancer, autoimmune, neurodegenerative and other

| Gene                                      | rs No  | Allele1  | Allele N= 3914 | Genotype | N= 1957 | Number (%) |
|-------------------------------------------|--------|----------|----------------|----------|---------|------------|
| Tumor necrosis factor-α (TNF-α)           | rs1800630 | A (C)  | C              | A        | CC      | CA         | AA         |
|                                           |        |          | 3229 (82.5)    | 685 (17.5)| 1420 (72.5)| 389 (19.9) | 148 (7.6)  |
| Tumor necrosis factor receptor superfamily member 1B (TNFRSF1B) | rs106124 | A (G)  | G              | A        | GG      | GA         | AA         |
|                                           |        |          | 1916 (49.0)    | 1998 (51.0)| 473 (24.2)| 970 (49.6) | 514 (26.2) |
| Interleukin-1A (IL1A)                     | rs1800587 | T (C)  | C              | T        | CC      | CT         | TT         |
|                                           |        |          | 3525 (90.1)    | 389 (9.9)| 1587 (81.1)| 351 (17.9) | 19 (1.0)   |
| Interleukin-1B (IL1B)                     | rs16944 | T (C)  | C              | T        | CC      | CT         | TT         |
|                                           |        |          | 2094 (53.5)    | 1820 (46.5)| 546 (27.9)| 1002 (51.2)| 409 (20.9) |
| Interleukin 4 receptor (IL-4R)            | rs1801275 | G (A)  | A              | G        | AA      | AG         | GG         |
|                                           |        |          | 3375 (86.2)    | 539 (13.8)| 1454 (74.3)| 467 (23.9) | 36 (1.8)   |
| Interleukin-6 (IL-6)                      | rs1800796 | G (C)  | C              | G        | CC      | CG         | GG         |
|                                           |        |          | 2969 (75.9)    | 945 (24.1)| 1134 (58.0)| 701 (35.8) | 122 (6.2)  |
| Interleukin-10 (IL-10)                    | rs1800872 | C (A)  | A              | C        | AA      | AC         | CC         |
|                                           |        |          | 2648 (67.7)    | 1266 (32.3)| 896 (45.8)| 856 (43.7) | 205 (10.5) |
| Interleukin 1 receptor-associated kinase-1 (IRAK-1) | rs1059702 | C (T)  | T              | C        | TT      | TC         | CC         |
|                                           |        |          | 3000 (76.6)    | 914 (23.4)| 1312 (67.0)| 376 (19.2) | 269 (13.8) |
| C-reactive protein 2 (CRP2)               | rs1130864 | T (C)  | T              | C        | CC      | CT         | TT         |
|                                           |        |          | 3670 (93.8)    | 244 (6.2)| 1721 (87.9)| 228 (11.7) | 8 (0.4)    |

1Reference alleles are indicated in parentheses.
diseases). Qidwai & Khan reviewed TNF-α gene polymorphism and disease prevalence comprehensively [7]. In the review, several mechanisms playing important role in gene regulation were discussed. Two most important factors that interfere with the gene regulation are promoter hypermethylation and the presence of polymorphism in regulatory region, and their hypothesis was that during disease conditions, there was upregulation or downregulation of gene (transcriptional dysregulation).

TNF-α polymorphism rs1800630 A-allele was associated with lower specific anti-pneumococcal IgG levels compared with children carrying C/C genotype of rs1800630, in one instance [7].

In the present analyses, significant odds ratios for hearing impairment risk were obtained even after adjustment for possible influential variables. The minor A-allele of rs1800630 polymorphism increased the risk of hearing impairment.

There are limited data regarding the functionality of the various SNPs in the TNFRSF1B gene. One clinical cancer investigation demonstrated that the TNFRSF1B rs1061624 genotype was a predictive factor of clinical response i.e., a complete response or not, to treatment with a 5-fluorouracil (5-FU)/cisplatin (CDDP) - based chemoradiotherapy in Japanese esophageal squamous cell carcinoma.

### Table 3 Hearing impairment risk of inflammatory mediators related SNPs analyzed using logistic regression with generalized estimating equations

| Gene    | rs No       | PTABE >25 dB | PTABE >40 dB |
|---------|-------------|--------------|--------------|
| TNF-α   | rs1800630   | 1.120 0.979-1.281 0.100 | 1.192 1.020-1.392 0.027 |
|         |             | 1.090 0.885-1.344 0.417 | 1.109 0.884-1.390 0.372 |
| TNFRSF1B| rs1061624   | 1.151 1.021-1.297 0.022 | 1.210 1.052-1.393 0.008 |
|         |             | 1.272 1.052-1.538 0.013 | 1.293 1.045-1.599 0.018 |
| IL1A    | rs1800587   | 1.093 0.894-1.335 0.386 | 1.061 0.845-1.331 0.611 |
|         |             | 0.975 0.692-1.373 0.885 | 0.946 0.660-1.354 0.760 |
| IL1B    | rs16944     | 0.941 0.834-1.060 0.316 | 0.915 0.796-1.052 0.212 |
|         |             | 0.924 0.762-1.121 0.422 | 0.960 0.774-1.190 0.706 |
| IL-4    | rs1801275   | 0.986 0.829-1.72 0.872 | 1.038 0.851-1.265 0.715 |
|         |             | 0.968 0.729-1.85 0.822 | 1.010 0.746-1.367 0.949 |
| IL-6    | rs1800796   | 1.011 0.881-1.161 0.872 | 1.083 0.924-1.269 0.327 |
|         |             | 1.034 0.829-1.289 0.770 | 1.042 0.822-1.320 0.734 |
| IL-10   | rs1800872   | 0.981 0.864-1.114 0.764 | 0.983 0.844-1.145 0.826 |
|         |             | 0.887 0.714-1.102 0.280 | 0.860 0.572-1.101 0.231 |
| IRAK-1  | rs1059702   | 1.026 0.912-1.55 0.666 | 1.012 0.889-1.151 0.862 |
|         |             | 0.887 0.718-1.097 0.268 | 0.816 0.707-1.051 0.141 |
| CRP2    | rs1130864   | 0.976 0.758-1.257 0.81 | 1.099 0.816-1.479 0.535 |
|         |             | 0.924 0.624-1.363 0.692 | 1.081 0.711-1.643 0.715 |

PTABE: pure-tone average threshold at 0.5, 1, 2 and 4 kHz for the better ear.
Moderating variables:
model 1: No variables.
model 2: Age and sex.
model 3: Any history of noise exposure, ear disease, hypertension, diabetes, dyslipidemia, stroke and cardiac disease, and lifetime smoking habit in addition to those of model 2.
patients [21]. Carrying the rs1061624 variant was reported to affect the phenotype of inflammatory bowel diseases in a New Zealand Caucasian population (decreased the risk of ulcerative colitis in the left colon) [22]. In our results, TNFRSF1B rs1061624 was associated with risk increment of age-related hearing impairment defined as a hearing level of PTABE > 25 dB and PTABE > 40 dB.

Recently, we found a significant association of risks for both sudden sensorineural hearing loss (SSNHL) and Ménière's disease with IL-1A rs1800587 polymorphism [23], and also reported that carriers of the IL-6 rs1800796 minor-allele were potentially more susceptible to SSNHL than noncarriers [24]. From one Korean case–control investigation, the presence of the IL-4R polymorphism Q576R, also known as G1902A, rs1801275 [25], was reported to be a risk factor for SSNHL [26]. In the present results, however, no associations of the presence of the 7 SNPs other than TNF-α rs1800630 and TNFRSF1B rs1061624, were found with the risk of age-related hearing impairment.

Several potential limitations of the present study should be cited. The genes in the selection of the present analyses were highly polymorphic respectively. To cite an example, 32 SNPs lie in transcription factor–binding sites of 20 transcription factors have been detected in the TNF-α upstream region [7]. But restrictions in budget and efficiency allowed us only to investigate widely and shallowly. Selection criterion of SNPs in the NILS-LSA study was based on candidate polymorphisms analysis for aging or geriatric diseases, such as senescence, geriatric syndrome, dementia, and osteoporosis. We focused the association between candidate SNPs for aging and hearing, then previously published the effects on hearing in the elderly, of EDN1, MTHFR, MTR (folate metabolism), UCP1, UCP2, and FABP2, oxidative stress related SNPs [27-30].

In addition, we were not able to obtain each inflammatory cytokine level in individuals. However, recent experimental observation, which explored the cochlear cytokine expression pattern under normoxia and hypoxia, suggested presence of tissue-specific regulatory pathways even among three different cochlear tissues; the organ of Corti, stria vascularis together with spiral ligament, and modiolus [31]. Given the tissue-specificity of cochlear tissue, even if blood cytokine level during common condition is obtained, it cannot be directly linked to pathogenetic elucidation. Further basic and applied researches will be required to interpret the role of each SNP in hearing deterioration.

From the aspect of methodology, there may be some other way to define a criterion for hearing impairment by different age groups, such as comparison with normal hearing levels according to the International Organization for Standardization (ISO) standard 7029:2000 [32]. ISO-7029 is based on a linear model of the hearing threshold for populations of otologically normal persons within the age limits of 18 years to 70 years, not including the older ages. Because the age range of the present subjects was distributed up to 89 years of age, we did not use a hearing assessment according to the ISO-7029.

The strength of this study is found in a feature of the NILS-LSA subject group. The NILS–LSA cohort consists of inhabitants who were selected randomly from resident registrations and stratified according to both age and sex. The risk of genotypic bias in subjects was therefore likely to be minimal. There were no significant differences in genotype frequencies of TNF-α rs1800630 and TNFRSF1B rs1061624 between the HapMap-JPT values and those obtained for the current participants. The ancestral allele of the rs1061624 single nucleotide variation is A-allele. The reference allele in the present analyses was G-allele, and G-allele frequencies of the rs1061624 polymorphism were released in the frequency reports as 0.535 in HapMap-JPT (Japanese in Tokyo), 0.539 in 1000 Genomes Project JPT (Japanese in Tokyo), 0.390 in AoD_Japanese, 0.507 in JBIC-allele (752 anonymous unrelated Japanese volunteers), and 0.580 in JPT_GENO_PANEL.

Conclusion
The present population-based cohort study demonstrated that TNF-α rs1800630 and TNFRSF1B rs1061624 contributed to the incremental risk of hearing impairment in the elderly Japanese population. TNF-α and TNF receptor interactions play a pivotal role in the pathogenesis of the inflammatory response, and also cause programmed cell death and cell proliferation. The present observation implied that the signaling cascades of TNF-α were involved in age-related hearing loss.

Materials and methods
Subjects
The NILS-LSA is a comprehensive and interdisciplinary study to observe age-related changes and consists of various gerontological and geriatric measurements, such as medical examinations, blood chemical analysis, body composition, anthropometry, physical function, nutritional analysis, psychological tests, and visual and auditory function. The participants were community-dwelling adults of Aichi Prefecture in central Japan, who were randomly selected from resident registrations, stratified by both age and sex in cooperation with the local government. The numbers of men and women recruited were similar, and age at baseline was 40 years to 79 years, with similar numbers of participants in each decade of age (40s, 50s, 60s, and 70s). Most of medical check-up are examined repeatedly every two years, and venous blood samples for genetic test were collected only at the first-wave examination. Sera and DNA samples were stored in deep freezers for later examination. The study protocol was approved by the Committee of Ethics of Human Research of the
| Gene                                      | rs No       | Labeled primers               | Sequence(5'→3')    | Amplicon (F1/R)/(F2/R) (bp) | Annealing temp.(°C) | Mg(mM) | Probe(5'→3')   |
|-------------------------------------------|-------------|-------------------------------|--------------------|-----------------------------|---------------------|--------|----------------|
| Tumor necrosis factor-α (TNF-α)           | rs1800630   | F1(FITC)                      | ATG GCC CTG TCG TTA Axt G | 75/77                      | 62.5                | 3      |                |
|                                           |             | F2(Texas Red)                 | GGC CTC GTG TCG GTA Axt GG |                           |                     |        |                |
|                                           |             | R(Biotin)                     | ACG GCA ATG GGT AAG AGA ATG TC |                         |                     |        |                |
| Interleukin-1A (IL1A)                     | rs1800587   | F1(FITC)                      | AATAATAAAGTACCAAGGCAACxtC | 91/92                      | 60                  | 2.5    |                |
|                                           |             | F2(Texas Red)                 | TAAATAATAAAGTACCAAGGCAACxCC |                       |                     |        |                |
|                                           |             | R(Biotin)                     | AGTGGCTAAGTTGGAATGGA |                         |                     |        |                |
| Interleukin-1B (IL1B)                     | rs16944     | F1(FITC)                      | GGTGCTGTTTCTGTGCTxGG | 50                         | 60                  | 1.5    |                |
|                                           |             | F2(Texas Red)                 | GGTGCTGTTTCTGTGCTxAG |                           |                     |        |                |
|                                           |             | R(Biotin)                     | TCAGAGGCTCCTGTCAATTGAC |                         |                     |        |                |
| Interleukin 4 receptor (IL4R)             | rs1801275   | F1(FITC)                      | GCC CCC ACC AGT GCC TAT xGG | 49/50                      | 67.5                | 2.8    |                |
|                                           |             | F2(Texas Red)                 | GCC CCC CAC CAG TGG CTA TXA G |                       |                     |        |                |
|                                           |             | R(Biotin)                     | CAC CCT GTG CCA CGG CAT GTA |                 |                     |        |                |
| Interleukin-6 (IL-6)                      | rs1800796   | F1(FITC)                      | GCC AGT TCT ACA ACA Gx CC | 43/42                      | 60                  | 1.5    |                |
|                                           |             | F2(Texas Red)                 | GCA GTT CTA CAA CAG CxG C |                           |                     |        |                |
|                                           |             | R(Biotin)                     | CTG TGT TCT GCC TCT CC CG TG |                         |                     |        |                |
| Interleukin-10 (IL10)                     | rs1800872   | F1(FITC)                      | CAG AGA CTG CTC TCC TACAxG A | 51/53                      | 62.5                | 2.5    |                |
|                                           |             | F2(Texas Red)                 | CCA GAG ACT GCC TGC TCA CA TAx T |                  |                     |        |                |
|                                           |             | R(Biotin)                     | GCC TGG AAC ACA TCC TGT GA |                         |                     |        |                |
| Interleukin 1 receptor-associated kinase-1 (IRAK-1) | rs1059702 | F1(FITC)                      | AGGGGCGAGAAGGxAAA | 99                         | 67.5                | 2.8    |                |
|                                           |             | F2(Texas Red)                 | AGGGGCGAGAAGGxAGA |                           |                     |        |                |
|                                           |             | R(Biotin)                     | GGGAGCCCCACCTGATCCAGGC |                         |                     |        |                |
| Melting curve method                      |             |                               |                    |                            |                     |        |                |
| Tumor necrosis factor receptor superfamily member 1B (TNFRSF1B) | rs1061624 | F                             | CCC TCT GAC CTG CAG GCC AAG | 61                         | 65                  | 2      | AGC AGA GCC AGC GAG TGT GCC |
National Center for Geriatrics and Gerontology (#14, #52, #74). Written informed consent was obtained from all participants. The details of the NILS-LSA have been described elsewhere [30,33].

Data were collected from the NILS-LSA participants who partook at least in the first-wave examination. Those who did not consent to have blood samples taken, and those who lacked results for any of the nine SNP genotype analyses were excluded. The participants at baseline were 2,149 adults, and the cumulative data were 9,358 samples in accumulated total, which were compiled from the first through the sixth-wave examination surveyed between 1997 and 2010. Participants filled out a series of questionnaires in advance of the examination visit. Ear disease history and experience of occupational noise exposure were taken account as basic informations in an evaluation of age-related hearing impairment. Histories of hypertension, diabetes, dyslipidemia, stroke, and cardiac disease were also taken from the self-reported account. Lifetime smoking history was categorized as never or ever. Those who had a deficiency of any required information for the present analyses and those who did not complete the hearing test were excluded. The subjects for the present analyses without any missing information were 1,957 individuals at baseline, and the gross accumulation number of subjects, regardless of repetitive visits, was 8,675 adults 40 years to 89 years of age. The mean number of repetitive visits was 4.4.

Genetic analyses and other measures
Genomic DNA was extracted from peripheral blood lymphocytes using the standard procedure and polymerase chain reaction (PCR) amplification was performed. Nine polymorphisms of genes encoding inflammatory mediators,

| Gene     | rs No  | Allele | Genotype | N= 2149, Number (%) | p-value |
|----------|--------|--------|----------|----------------------|---------|
| TNF-α    | rs1800630 | A (C) | CC       | 1551 (72.2)          |         |
|          |        |       | CA       | 431 (20.0)           |         |
|          |        |       | AA       | 167 (7.8)            |         |
|          |        |       |          | NS                   |         |
| HapMap-JPT |       |       |          |          |         |
| TNFRSF1B | rs1061624 | A (G) | GG       | 519 (24.1)           |         |
|          |        |       | GA       | 1070 (49.8)          |         |
|          |        |       | AA       | 560 (26.1)           |         |
|          |        |       |          | NS                   |         |
| HapMap-JPT |       |       |          |          |         |
| IL-1A    | rs1800587 | T (C) | CC       | 1745 (81.2)          |         |
|          |        |       | CT       | 385 (17.9)           |         |
|          |        |       | TT       | 19 (0.9)             | NS      |
| HapMap-JPT |       |       |          |          |         |
| IL-1B    | rs16944  | T (C) | CC       | 600 (27.9)           |         |
|          |        |       | CT       | 1096 (51.0)          |         |
|          |        |       | TT       | 453 (21.1)           | NS      |
| PGA-UW-FHCRC: JPT_GENO_PANEL2) | |       |          |                      |         |
| IL-4R    | rs1801275 | G (A) | AA       | 1585 (73.8)          |         |
|          |        |       | AG       | 522 (24.3)           |         |
|          |        |       | GG       | 42 (1.9)             | NS      |
| HapMap-JPT |       |       |          |          | 0.034 |
| IL-6     | rs1800796 | G (C) | CC       | 1257 (58.5)          |         |
|          |        |       | CG       | 762 (35.5)           |         |
|          |        |       | GG       | 130 (6.0)            | NS      |
| HapMap-JPT |       |       |          |          | 0.007 |
| IL-10    | rs1800872 | C (A) | AA       | 985 (45.8)           |         |
|          |        |       | AC       | 938 (43.7)           |         |
|          |        |       | CC       | 226 (10.5)           | NS      |
| HapMap-JPT |       |       |          |          |       |
| IRAK-1   | rs1059702 | C (T) | TT       | 1444 (67.2)          |         |
|          |        |       | TC       | 406 (18.9)           |         |
|          |        |       | CC       | 299 (13.9)           | NS      |
| HapMap-JPT |       |       |          |          |       |
| CRP2     | rs1130864 | T (C) | CC       | 1889 (87.9)          |         |
|          |        |       | CT       | 251 (11.7)           |         |
|          |        |       | TT       | 9 (0.4)              | NS      |
| HapMap-JPT |       |       |          |          |       |

1Reference alleles are indicated in parentheses.
2Variation Discovery Resource. A collaboration between the University of Washington and the Fred Hutchinson Cancer Research Center.
namely, TNF-α C-863A, rs1800630; TNF receptor superfamily member 1B (TNFRSF1B) G593A, rs1061624; IL-1A -889C>T, rs1800587; IL-1B -511C>T, rs16944; IL-4R G1902A, rs1801275; IL-6 C-572G, also known as C-634G, rs1800796; IL-10 A-592C, rs1800872; IL-1 receptor-associated kinase 1 (IRAK1) T587C, rs1059702; CRP +1444 C>T, rs1130864 were investigated.

Genotyping was carried out using a fluorescence method and melting curve method. Details including of primer sequences and PCR conditions are listed in Table 4.

Genotype distributions of 9 SNPs in participants at first-wave examinations of NILS-LSA before application of exclusion criteria are shown in Table 5. There were no significant differences between genotypic frequencies of our participants and public database samples but in IL-4R and IL-6 (chi-square test, level of significance: 0.05).

Air-conduction pure-tone thresholds at octave intervals from 0.5 kHz to 8 kHz were measured in a sound-proof booth by trained examiners using a standardized protocol and a diagnostic audiometer (AA-73A and AA-78; Rion, Tokyo, Japan). The pure-tone average threshold level of the better ear at frequencies of 0.5, 1, 2, and 4 kHz (PTABE) was used as an index of hearing status. Hearing impairment was defined as a PTABE greater than 25 dB according to WHO grades [34]. The WHO defines disabling hearing impairment in adults as a permanent unaided PTABE of 41 dB or greater. In view of this, we adopted two hearing impairment criteria for the statistical analyses; PTABE >25 dB and PTABE >40 dB.

Statistical analyses
Statistical analyses were conducted using the Statistical Analysis System (SAS) version 9.1.3 (SAS Institute, Cary, NC, USA). For univariate analysis, a t-test was used to assess differences of continuous variables between two groups, and comparisons of categorical variables were performed using the chi-square test. Fisher’s exact test was also used, as and when the number of cell samples was less than five. Unless otherwise noted, values are expressed as mean ± standard deviation.

Cumulative data were analyzed using GEEs, which take into account the dependency of repeated observations within subjects. GEE is a common statistical model to analyze epidemiological longitudinal data. GEE models were fitted using the GENMOD procedure of Statistical Analysis System version 9.1.3. The GENMOD procedure fits generalized linear models. The correlation structure was specified as autoregressive. We took mode of inheritance in analyses as additive genetic model, which is the prevailing analytical model in genetic epidemiology, assumes that there is a linear gradient in risk with increasing numbers of variant alleles (reference allele homozygotes base). ORs for the additive genetic models of 9 SNPs with the risk of hearing impairment were analyzed. As a combination of moderator variables, three models were analyzed. In model 1, no moderator variables were adjusted; in model 2, age and gender were taken as moderator variables; and in model 3, any history of noise exposure, ear disease, hypertension, diabetes, dyslipidemia, stroke and cardiac disease, and lifetime smoking habit was adjusted in addition to age and gender.

Abbreviations
PTABE: Better ear pure-tone average; TNF: Tumor necrosis factor; TNFRSF: TNF receptor superfamily; IL: Interleukin; IRAK1: IL-1 receptor-associated kinase 1; CRP: C reactive protein; CI: Confidence interval; NILS-LSA: The National Institute for Longevity Sciences – Longitudinal Study of Aging; WHO: World Health Organization; SNP: Single-nucleotide polymorphism; GE: Generalized estimating equation; OR: Odds ratio; CI: Confidence interval; NLR2B: N-methyl D-aspartate receptor subunit 2B; 5-FU: 5-fluorouracil; CDDP: Cisplatin; SSNHL: Sudden sensorineural hearing loss; PCR: Polymerase chain reaction; SAS: Statistical analysis system.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
YU performed the data analyses, interpretation and manuscript writing; SS contributed to analytical design and assisted in the statistical analyses; FA and HS were responsible for conducting the cohort study; HU and TN assisted in interpretation of the data; All authors read and approved the final manuscript.

Acknowledgments
We would like to give our sincere thanks to all concerned: the participants of NILS-LSA and the health professionals and researchers from the department of the Development of Preventive Medicine, National Center for Geriatrics and Gerontology who were involved in data collection and analyses. This study was supported in part by the Research Grant for Longevity Sciences, [20shi-2], [21A-17] and [2-25] from the Ministry of Health, Labour and Welfare of Japan.

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Received: 19 October 2014 Accepted: 7 November 2014
Published online: 26 November 2014

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26. Cite this article as: Uchida et al: The association between hearing impairment and polymorphisms of genes encoding inflammatory mediators in Japanese aged population. Immunity & Ageing 2014 11:18.