Association between human T cell leukemia virus 1 (HTLV-1) infection and advanced periodontitis in relation to hematopoietic activity among elderly participants: a cross-sectional study

Yuji Shimizu1,2, Hirotomo Yamanashi3, Masayasu Kitamura4, Reiko Furugen4, Takahiro Iwasaki5, Hideki Fukuda4, Hideaki Hayashida4, Koji Kawasaki6, Kairi Kiyoura1, Shin-Ya Kawashiri1, Toshiyuki Saito4, Atsushi Kawakami7 and Takahiro Maeda1,3,8

Abstract

Background: We reported that human T cell leukemia virus 1 (HTLV-1) infection is positively associated with atherosclerosis. Recent evidence has revealed a close association of periodontitis with atherosclerosis, endothelial dysfunction, and disruption of the microcirculation. However, the association between HTLV-1 and advanced periodontitis has not been investigated to date. Since hematopoietic activity is closely linked to endothelial maintenance activity and is known to decline with age, we hypothesized that the state of hematopoietic activity influenced the association between HTLV-1 and advanced periodontitis in elderly participants.

Methods: A cross-sectional study was performed including 822 elderly participants aged 60–99 years who participated in a dental health check-up. Advanced periodontitis was defined as a periodontal pocket ≥ 6.0 mm. Participants were classified as having low or high hematopoietic activity according to the median values of reticulocytes.

Results: HTLV-1 infection was positively related to advanced periodontitis among participants with lower hematopoietic activity (lower reticulocyte count), but not among participants with higher hematopoietic activity (higher reticulocyte count). The adjusted odds ratio (95% confidence interval) considering potential confounding factors was 1.92 (1.05–3.49) for participants with a lower reticulocyte count and 0.69 (0.35–1.36) for participants with a higher reticulocyte count.

Conclusions: Among elderly participants, the association between HTLV-1 infection and advanced periodontitis is influenced by hematopoietic activity. Since hematopoietic activity is associated with endothelial maintenance, these findings provide an efficient tool for clarifying the underlying mechanism of the progression of periodontitis among elderly participants.

Keywords: Elderly participants, Hematopoietic activity, HTLV-1, Periodontitis, Reticulocyte
Background

Human T cell leukemia virus 1 (HTLV-1) is an oncogenic retrovirus affecting individuals worldwide, although the southern part of Japan is known as a highly endemic region for HTLV-1 infection [1]. In Japan, the number of HTLV-1 carriers was estimated to be approximately 1.2 million during the late 1980s [2]. The majority of HTLV-1 carriers remain asymptomatic throughout their lives [3–6]. However, we previously identified a positive association of asymptomatic HTLV-1 infection with atherosclerosis [7], which is the result of endothelial dysfunction [8].

In addition, recent studies have revealed that periodontitis is closely associated with atherosclerosis [9], endothelial dysfunction [10, 11], and disruption of microcirculation [12, 13]. Periodontitis is one of the major causes of tooth loss, resulting in a significant decrease in the quality of life of elderly participants [14]; however, the association between HTLV-1 infection and advanced periodontitis among elderly participants remains unknown.

In addition, bone marrow activity (hematopoietic activity) has been recently identified to be associated with endothelial maintenance [15–17]; reticulocyte levels, a marker of hematopoietic activity, were found to be associated with the activity of endothelial maintenance in elderly Japanese individuals [18]. Since aging is also well known to be associated with a reduction in hematopoietic activity as the bone marrow function declines [19–21], hematopoietic activity can be expected to influence the observed association between HTLV-1 infection and advanced periodontitis among elderly participants. Therefore, clarifying the role of reticulocyte levels in mediating the association between HTLV-1 infection and advanced periodontitis could provide an efficient tool for elucidating the mechanism underlying the progression of periodontitis in the elderly.

To evaluate this hypothesis, we conducted a cross-sectional study to examine the influence of hematopoietic activity (assessed by reticulocyte counts) on the association between HTLV-1 and advanced periodontitis among 822 elderly participants that received an annual dental health check-up from 2016 to 2018.

Methods

Study population

The study population comprised a total of 1925 participants (702 men and 1223 women) aged 60–99 years from Goto city in the western part of Japan who participated in an annual health check-up with oral assessment in 2016–2018. This annual check-up program is conducted by the local government and directed by the Ministry of Health, Labor and Welfare in Japan. Owing to the shortage of staff to conduct the health check-up in the present survey, the entire city could not be surveyed in 1 year. Therefore, we conducted the survey in different parts of the city over a period of 3 years to ensure that all areas of Goto city were covered. Nevertheless, no individual in the population overlapped among the 3 years of the study. Participants were excluded if there were no carotid intima-media thickness (CIMT) data (n = 4) or laboratory data (n = 3) available, or if they had less than 10 remaining teeth (n = 1078). Since anemia (low levels of hemoglobin, Hb) has been reported to be associated with periodontitis [22, 23] and HTLV-1 infection [24], anemia was considered to be a strong confounding factor for the present analysis; therefore, we excluded participants diagnosed with anemia (Hb < 11 g/dL; n = 15). We also excluded participants without available HTLV-1 data (n = 3). The remaining participants, comprising 326 men and 496 women, were ultimately included for analysis.

Data collection and laboratory measurements

A trained interviewer obtained the information on smoking status. Body weight and height were measured while the participant was in bare feet and wearing light clothing using an automatic body composition analyzer (BF-220; Tanita, Tokyo, Japan). Body mass index was calculated as weight (kg)/height (m)².

Blood samples were collected into EDTA-2K tubes and siliconized tubes, and all measurements were conducted according to standard automated laboratory procedures at SRL, Inc. (Tokyo, Japan). The parameters tested included red blood cell (RBC) count, reticulocyte count, Hb level, and serum creatinine level. Reticulocyte count was calculated with the following formula: reticulocytes (× 10⁴ cells/µL) = (reticulocytes, %) × RBC (× 10⁴ cells/µL)/1000. The glomerular filtration rate (GFR) was estimated according to an established method with three modifications recently proposed by a working group of the Japanese Chronic Kidney Disease Initiative [25]. According to this adaptation, GFR (mL min/1.73 m²)=1.094 × [serum creatinine (enzyme method)]⁻¹.094 × (age)⁻⁰.287 × (0.739 for women).

Measurement of CIMT

CIMT was measured by an experienced vascular technician based on ultrasonography of the left and right common carotid arteries using a LOGIQ Book XP with a 10-MHz transducer (GE Healthcare, Milwaukee, WI, USA). Mean values for the left and right CIMT were calculated with automated digital edge-detection software (Intimascope; MediaCross, Tokyo, Japan) according to a previously described protocol [26]. The values of right and left CIMT, not including plaque measurements, were then calculated, and the mean CIMT value was used for analysis.
Detection of HTLV-1
A chemiluminescent enzyme immunoassay (CLEIA) kit was used to detect the presence of HTLV-1 (Fujirebio Inc., Tokyo, Japan) according to the manufacturer’s instructions.

Oral examination
Trained dentists performed a periodontal examination according to a modified method of the Third National Health and Nutrition Examination Survey [27]. Probing pocket depth was measured using a periodontal probe at the mesio-buccal and mid-buccal sites for all present teeth, excluding the third molars. Advanced periodontitis was diagnosed as a probing pocket depth ≥ 6 mm. Prior to the start of the study, all examiners were trained in the same manner and their assessments were calibrated using a chart, periodontal models, and volunteers at Nagasaki University Hospital.

Statistical analysis
The median value for the reticulocyte count ($5.175 \times 10^4$ cells/$\mu$L for men and $4.926 \times 10^4$ cells/$\mu$L for women) was set as the cutoff point for classifying the participants into those with low and high hematopoietic activity. The differences in mean values or proportions of reticulocyte levels were analyzed in relation to HTLV-1 status. Significant differences were evaluated using analysis of variance (ANOVA) for continuous variables and chi-squared test for proportion data.

Logistic regression models were used to calculate odds ratios (ORs) and 95% confidence intervals (CIs) to determine the association between HTLV-1 infection and advanced periodontitis. Since aging influences the hematopoietic activity, which in turn influences erythropoietin production, the participants were stratified according to their hematopoietic activity (maintained or reduced hematopoiesis) based on the level of reticulocytes (high or low).

Periodontitis risk factors with a direct influence on the intra-oral environment, such as smoking [28] and number of remaining teeth, were regarded as potential confounding factors in the present analysis. Although no significant correlation between caries (decayed teeth) and periodontitis has been observed in previous studies, both conditions have a common etiology [29, 30]. Therefore, we also included the presence of decayed teeth as a confounding factor in the present analysis. In addition, a low level of Hb can stimulate reticulocyte production and was reported to be associated with more aggressive forms of periodontitis [22], while high levels of Hb are associated with increased arterial stiffness (atherosclerosis) [31]. Since CIMT was reported to be positively associated with periodontitis [9], we included both Hb and CIMT as confounding factors in the model.

In addition, hemoglobin level is known to be associated with renal function [32] as well as increased arterial stiffness, as evaluated by the cardio-ankle vascular index (CAVI) [31]. CIMT is known to be associated with hematopoietic activity [33]. Moreover, hematopoietic activity serves as a determining factor for the association between chronic kidney disease and CIMT [34] and for the association between CAVI and CIMT [35]. Therefore, we also included renal function as a confounding factor in the model, which was evaluated by the GFR. Although endothelial dysfunction is a common initial mechanism leading to atherosclerosis and renal dysfunction [8], the model further adjusted for GFR could present a risk of multicollinearity. Therefore, we established three distinct models to adjust for these confounding factors. The first model was adjusted for only sex and age (model 1); the second model (model 2) included the other potential confounding factors, namely smoking status (never-smoker, former smoker, or current smoker), number of remaining teeth, decayed teeth (yes/no), CIMT (mm), and Hb (g/dL). The last model (model 3) was the one further adjusted for GFR.

All statistical analyses were performed with the SAS system for Windows (version 9.4: SAS Inc., Cary, NC, USA). Values of $p < 0.05$ were regarded as statistically significant.

Results
General characteristics of the study population
Among the total 822 participants with a mean ± standard deviation (SD) age of 71.2 ± 6.7 years, 140 (17.0%) showed an HTLV-1-positive status and 181 (22.0%) were diagnosed as having advanced periodontitis; 411 participants had low hematopoietic activity (i.e., a low reticulocyte level).

Table 1 shows the reticulocyte level-specific characteristics of the study population according to the HTLV-1 infection status. No significant associations were observed for participants with lower hematopoietic activity, whereas participants with higher hematopoietic activity and HTLV-1 infection were significantly older than those without HTLV-1 infection.

Association between advanced periodontitis and HTLV-1
As shown in Table 2, there was no significant association between HTLV-1 infection and advanced periodontitis among the total participants. However, HTLV-1 infection was significantly positively associated with advanced periodontitis for participants with lower hematopoietic activity, but not for those with higher hematopoietic activity (Table 3). In addition, a significant interaction was observed in model 1 and model 2 between HTLV-1 status and hematopoietic activity level (low vs. high) with respect to advanced periodontitis. After further
adjustment for GFR (model 3), the interaction showed marginal significance.

Sensitivity analysis according to the quartile of reticulocyte levels demonstrated associations between HTLV-1 infection and advanced periodontitis, similar to the main results.

**Discussion**

The major finding of the present study is that HTLV-1 infection is positively associated with advanced periodontitis only among elderly participants with lower hematopoietic activity (a lower reticulocyte count).

A previous case-controlled cross-sectional study including 42 periodontally healthy individuals, 64 HTLV-1-seronegative individuals presenting with chronic periodontitis, and 36 individuals with chronic periodontitis with HTLV-1-seropositive HTLV-1-associated myelopathy (HAM)/symptomatic tropical spastic paraparesis (TSP) revealed that HTLV-1 may play a critical role in the pathogenesis of periodontal disease through deregulation of the local cytokine network [36].

In the present study, we found further evidence that HTLV-1 infection is positively associated with advanced periodontitis among participants with low reticulocyte levels, even though most of the participants were asymptomatic.

Periodontitis has been shown to be closely associated with atherosclerosis [9], endothelial dysfunction [10, 11], and disruption of the microcirculation [12, 13], and low levels of erythrocytes and Hb are positively associated with periodontitis [22, 23]. In addition, reticulocyte levels are reported to be inversely associated with CIMT in elderly Japanese individuals [18]. Therefore, maintaining higher levels of hematopoietic activity (i.e., a high reticulocyte level) should protect against the progression of periodontitis in elderly participants.

However, we found a similar prevalence of advanced periodontitis in participants with low (22.6%) and high (21.4%) hematopoietic activity, as assessed by the reticulocyte count.

Aging is known to be associated with a decline of bone marrow activity (reduction in hematopoietic activity) [19–21] and also contributes to endothelial dysfunction via increasing oxidative stress [37, 38]. Since the hematopoietic activity in elderly participants is determined by both of these age-related factors, higher

### Table 1 Characteristics of study population

| Reticulocyte count levels | Low | High |
|--------------------------|-----|------|
| No of subjects           | 345 | 66   |
| No of advanced periodontitis (%) | (−) | (+) |
| HTLV-1 infection         | 72 (20.9) | 21 (31.8) |
| p                        | 0.052 | 0.009 |
| Men, %                   | 40.9 | 33.3 |
| Age, year                | 71.7 ± 7.0 | 72.8 ± 6.5 |
| Current smoker, %        | 3.8  | 7.6  |
| Former smoker, %         | 25.8 | 19.7 |
| No of remaining teeth    | 22.3 ± 5.3 | 23.4 ± 4.9 |
| Individuals with decayed teeth, % | 28.1 | 28.8 |
| CIMT, mm                 | 0.71 ± 0.13 | 0.73 ± 0.12 |
| Hemoglobin, g/dL         | 13.6 ± 1.2 | 13.4 ± 1.2 |
| Reticulocyte, × 10⁴ cells/μL | 3.871 ± 0.744 | 3.874 ± 0.788 |
| GFR, mL/min/1.73m²       | 70.5 ± 13.1 | 67.9 ± 14.1 |

Values: mean ± 1 standard deviation (SD). Advanced periodontitis is defined as periodontal pocket ≥ 6.0 mm. Low reticulocyte levels were < 5.175 × 10⁴ cells/μL for men and < 4.926 × 10⁴ cells/μL for women.

CIMT carotid intima-media thickness, GFR glomerular filtration rate.

### Table 2 Odds ratios (ORs) and 95% confidence intervals (CIs) for advanced periodontitis in relation to human T cell leukemia virus-1 (HTLV-1) infections

| HTLV-1 infection | p    |
|------------------|------|
| No of subjects   | 682  |
| No of case (%)   | 147 (21.6) | 34 (24.3) |
| Model 1          | 1.00 | 1.15 (0.75, 1.77) | 0.523 |
| Model 2          | 1.00 | 1.18 (0.76, 1.83) | 0.458 |
| Model 3          | 1.00 | 1.18 (0.76, 1.83) | 0.462 |

Advanced periodontitis is defined as periodontal pocket ≥ 6.0 mm. Model 1: adjusted only for sex and age. Model 2: adjusted further for smoking status (never, former, current), remained number of teeth, status of decayed teeth (presence, absence), carotid intima-media thickness (CIMT), and hemoglobin. Model 3: adjusted further for glomerular filtration rate (GFR).
hematopoietic activity can help to maintain endothelial function even if the magnitude of endothelial damage is severe [17, 18]. In other words, elderly participants with lower hematopoietic activity would have more difficulty in maintaining endothelial function, regardless of the severity of age-related endothelial dysfunction. We previously reported an ambivalent association of reticulocytes in hypertension and atherosclerosis (endothelial dysfunction), in which hypertension and atherosclerosis showed a significantly positive association; however, hypertension showed a significantly positive association with reticulocytes while atherosclerosis showed a significantly inverse association with reticulocytes [18]. The present study partly supports the abovementioned mechanism by indicating that the reticulocyte level serves as an indicator of not only endothelial damage but also endothelial repair activity. Furthermore, this study suggests that individuals with high levels of reticulocytes should have a higher capacity of endothelial maintenance compared to those with low levels of reticulocytes.

In addition, HTLV-1 infection is reported to be positively associated with atherosclerosis [7]. Therefore, compared to participants without HTLV-1 infection, HTLV-1-infected participants might have a greater degree of endothelial dysfunction, which can increase the risk of periodontitis [9–13]. This relationship between hematopoietic activity and endothelial function explains the significantly positive association between HTLV-1 infection and advanced periodontitis only in participants with lower hematopoietic activity (lower endothelial repair activity).

Furthermore, a significant interaction was observed between HTLV-1 status and hematopoietic activity level (low vs. high) with respect to advanced periodontitis. These results also support the abovementioned mechanism by indicating that hematopoietic activity levels could act as a determinant of the association between HTLV-1 and advanced periodontitis.

Some potential limitations of this study warrant consideration. Although the age-related endothelial dysfunction should have an influence on the association between HTLV-1 infection and advanced periodontitis, no data on endothelial function were available to directly assess this relationship. Thus, further analyses that include endothelial function-related data such as flow-mediated dilation [39] will be necessary.

Owing to the lack of knowledge regarding the determination of the exact cutoff point, the median values of reticulocytes were used for the present analysis. Further investigation is necessary to clarify whether the balance between endothelial damage and endothelial repair that contributes to endothelial maintenance can help to determine the exact cutoff point for the present associations. In addition, because this was a cross-sectional study, causal relationships could not be established. Nevertheless, the association identified provides new insights into the potential mechanisms mediating the relationship between HTLV-1 and periodontitis, highlighting the importance of maintaining hematopoietic activity for the overall health of elderly participants.

### Conclusions
HTLV-1 infection is positively associated with advanced periodontitis among elderly participants with lower hematopoietic activity (a lower reticulocyte count) but not among elderly participants with higher hematopoietic activity (a higher reticulocyte count). Since elderly participants with higher hematopoietic activity should have higher tissue-repairing activity [40] with a protective effect against periodontitis progression [23], these findings can help to clarify the underlying mechanism contributing to the progression of periodontitis among elderly participants.

### Abbreviations
ANOVA: Analysis of variance; CIMT: Carotid intima-media thickness; CIs: Confidence intervals; CLEIA: Chemiluminescent enzyme immunoassay; HAM: HTLV-1-associated myelopathy; Hb: Hemoglobin; HTLV-1: Human T cell leukemia virus-1; HTLV-1 infection: Human T cell leukemia virus-1 infection; HTLV-1-associated myelopathy: Human T cell leukemia virus-1-associated myelopathy; Hb: Hemoglobin; HTLV-1: Human T cell leukemia virus-1; Hb: Hemoglobin; HTLV-1: Human T cell leukemia virus-1; Hb: Hemoglobin;
leukemia virus 1; ORs: Odds ratios; RBC: Red blood cell; SD: Standard deviation; TSP: Tropical spastic paraparesis

Acknowledgements
We are grateful to the staff of Goto city office for their outstanding support. We would like to thank Editage (www.editage.jp) for English language editing.

Authors’ contributions
YS designed the study, performed the statistical analyses, interpreted the data, and drafted and revised the manuscript. HY, MK, RF, TI, HF, HH, Kok, Kak, SYK, TS, and AK assisted with the study design, were involved in data collection, and checked the manuscript. HY, SYK, and TM participated in the study concept and checked the manuscript. TM was the general coordinator and designed the study. All authors read and approved the final manuscript.

Funding
This study was supported by Grants-in-Aid for Scientific Research from the Japan Society for the Promotion of Science (No. 18H06448, No. 17H03740).

Availability of data and materials
The datasets generated during and/or analyzed during the current study are not publicly available due to ethical consideration but are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
This study was approved by the Ethics Committee of Nagasaki University Graduate School of Biomedical Sciences (project registration number 14051404). All procedures involving human participants were performed in accordance with the ethical standards of the institution research committee and with the 1964 Helsinki Declaration and its later amendments for comparable ethical standards.

Consent for publication
Not applicable

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

Author details
1Department of Community Medicine, Nagasaki University Graduate School of Biomedical Sciences, Nagasaki-shi, Sakamoto 1–12–4, Nagasaki 852-8523, Japan. 2Department of Cardiovascular Disease Prevention, Osaka Center for Cancer and Cardiovascular Diseases Prevention, Osaka, Japan. 3Department of General Medicine, Nagasaki University Hospital, Nagasaki, Japan. 4Department of Oral Health, Nagasaki University Graduate School of Biomedical Sciences, Nagasaki, Japan. 5Department of Community Oral Health, School of Dentistry, Asahi University, Gifu, Japan. 6Department of Community Medicine, Nagasaki University Graduate School of Biomedical Sciences, Nagasaki, Japan.

Received: 11 January 2019 Accepted: 29 May 2019

Published online: 10 June 2019

References
1. Gessain A, Cassar O. Epidemiological aspects and world distribution of HTLV-1 infection. Front Microbiol. 2012;3:388.
2. Tajima K. The 4th nation-wide study of adult T-cell leukemia/lymphoma (ATL) in Japan: estimates of risk of ATL and its geographical and clinical features. The T- and B-cell Malignancy Study Group. Int J Cancer. 1990;45(2):237–43.
3. Yamaguchi K, Watanabe T. Human T lymphotropic virus type-I and adult T-cell leukemia in Japan. Int J Hematol. 2002;76(Suppl 2):240–5.
4. Murphy EL, Hanchard B, Figueroa JP, Gibbs WN, Loferski WS, Campbell M, et al. Modelling the risk of adult T-cell leukaemia/lymphoma in persons infected with human T-lymphotropic virus type I. Int J Cancer. 1989;43(2):250–3.
5. Kaplan JE, Osame M, Kubota H, Iga T, Nishitani H, Maea Y, et al. The risk of development of HTLV-I-associated myelopathy/tropical spastic paraparesis among persons infected with HTLV-I. J Acquir Immune Defic Syndr. 1990;3(11):1096–101.
6. Maloney EM, Cleggern HR, Morgan OS, Rodgers-Johnson P, Cranston B, Jack N, et al. Incidence of HTLV-I-associated myelopathy/tropical spastic paraparesis (HAM/TSP) in Jamaica and Trinidad. J Acquir Immune Defic Syndr Hum Retrovirol. 1998;17(2):167–70.
7. Yamanashi H, Koyamatsu J, Nagayoshi M, Shimizu Y, Kawashiri SY, Kondo H, et al. Human T-cell leukemia virus-1 infection is associated with atherosclerosis as measured by carotid intima-media thickness in Japanese community-dwelling older people. Clin Infect Dis. 2018;67(2):291–4.
8. Endemmann DH, Schifflin EL. Endothelial dysfunction. J Am Soc Nephrol. 2004;15(8):1983–92.
9. Hayashida H, Salto T, Kawasaki K, Kitamura M, Furugen R, Iwasaki T, et al. Association of periodontitis with carotid artery intima-media thickness and arterial stiffness in community-dwelling people in Japan: the Nagasaki Islands study. Atherosclerosis. 2013;229(1):186–91.
10. Gurav AN. The implication of periodontitis in vascular endothelial dysfunction. Eur J Clin Invest. 2014;44(10):1000–9.
11. Hoffrider R, Hampen K, Glaser S, Lorbëer R, Völzke H, Ewert R, et al. Periodontitis is associated with endothelial dysfunction in a general population: a cross-sectional study. PLoS One. 2013;8(12):e84603.
12. Lira-Junior R, Figueiredo CM, Boukela E, Fischer RG. Severe chronic periodontitis is associated with endothelial and microvascular dysfunctions: a pilot study. J Periodontol. 2014;85(12):1548–57.
13. Boillot A, Bouchard P, Moss K, Offenbacher S, Czemichow S. Periodontitis and retinal microcirculation in the Atherosclerosis Risk in Communities study. J Clin Periodontol. 2015;42(4):342–9.
14. Ong G. Periodontal reasons for tooth loss in an Asian population. J Clin Periodontol. 1996;23(4):307–9.
15. Takakura N, Watanabe T, Suebou S, Yamada Y, Noda T, Ito Y, et al. A role for hematopoietic stem cells in promoting angiogenesis. Cell. 2000;102(2):199–209.
16. Yamada Y, Takakura N. Physiological pathway of differentiation of hematopoietic stem cell population into mural cells. J Exp Med. 2006;203(4):1055–65.
17. Shi Q, Raffi S, Wu MH, Wijelath ES, Yu C, Ishida A, et al. Evidence for circulating bone marrow-derived endothelial cells. Blood. 1998;92(2):362–7.
18. Shimizu Y, Kawashiri SY, Yamanashi H, Koyamatsu J, Fukui S, Kondo H, et al. Reticulocty levels have an ambivalent association with hypertension and atherosclerosis in the elderly: a cross-sectional study. Clin Interv Aging. 2019;14:849–57.
19. Brunsnah SK, McGuire TR, Jackson JD, Lane JT, Garvin KL, O’Kane BJ, et al. Human blood and marrow side population stem cell and Stro-1 positive bone marrow stromal cell numbers decline with age, with an increase in quality of surviving stem cells: correlation with cytokines. Mech Ageing Dev. 2010;131(11–12):719–22.
20. Garvin K, Feschuk C, Sharp JG, Berger A. Does the number or quality of pluripotent bone marrow stem cells decrease with age? Clin Orthop Relat Res. 2007;465:202–7.
21. Cooper BR. The origins of bone marrow as the seedbed of our blood: from antiquity to the time of Osler. Proc (Bayl Univ Med Cent). 2011;24(2):115–8.
22. Anand PS, Sakar DK, Ashok S, Kamath KP. Association of aggressive periodontitis with reduced erythrocyte counts and reduced hemoglobin levels. J Periodontal Res. 2014;49(6):719–28.
23. Hutter JW, van der Velden U, Varoufaki A, Huffels RA, Hoek FJ, Loos BG. Lower numbers of erythrocytes and lower levels of hemoglobin in periodontitis patients compared to control subjects. J Clin Periodontol. 2001;28(10):930–6.
24. Chaturvedi AK, Wilson M, Sanders-Lewis KA, Kati H, Urquhart N, Walters MA, et al. Hematologic and biochemical changes associated with human T lymphotropic virus type 1 infection in Jamaica: a report from the population-based donors study. Clin Infect Dis. 2007;45(8):975–82.
25. Imai E, Hono M, Watanabe T, Iseki K, Yamagata K, Hara S, et al. Prevalence of chronic kidney disease in the Japanese general population. Clin Exp Nephrol. 2009;13(6):621–30.
26. Hara T, Takamura N, Akashi S, Nakazato M, Maeda T, Wada M, et al. Evaluation of clinical markers of atherosclerosis in young and elderly Japanese adults. Clin Chem Lab Med. 2006;44(7):824–9.
27. Brown LJ, Brunelle JA, Kingman A. Periodontal status in the United States, 1988–1991: prevalence, extent, and demographic variation. J Dent Res. 1996;75 Spec5:72–83.
28. Nociti FH Jr, Casati MZ, Duarte PM. Current perspective of the impact of smoking on the progression and treatment of periodontitis. Periodontol 2000. 2015;67(1):187–210.

29. Saotome Y, Tada A, Hanada N, Yoshihara A, Uematsu H, Miyazaki H, et al. Relationship of cariogenic bacteria levels with periodontal status and root surface caries in elderly Japanese. Gerodontology. 2006;23(4):219–25.

30. Frenzen M, Schühler N, Nolden R. Correlation between caries prevalence (DMFS) and periodontal condition (CPITN) in more than 2000 patients. Int Dent J. 1990;40(5):313–8.

31. Shimizu Y, Nakazato M, Sekita T, Kadota K, Yamashita H, Takamura N, et al. Association between hemoglobin levels and arterial stiffness for general Japanese population in relation to body mass index status: the Nagasaki Islands study. Geriatr Gerontol Int. 2014;14(4):811–8.

32. Shimizu Y, Sato S, Koyamatsu J, Yamanashi H, Nagayoshi M, Kadota K, et al. Associations between renal impairment and anemia in older, rural Japanese men: the Nagasaki Island study. J Physiol Anthropol. 2014;33:7.

33. Shimizu Y, Sato S, Koyamatsu J, Yamanashi H, Nagayoshi M, Kawashiri SY, et al. Hepatocyte growth factor and carotid intima-media thickness in relation to circulating CD34-positive cell levels. Environ Health Prev Med. 2018;23(1):16.

34. Shimizu Y, Yamanashi H, Noguchi Y, Koyamatsu J, Nagayoshi M, Kiyoura K, et al. Association between chronic kidney disease and carotid intima-media thickness in relation to circulating CD34-positive cell count among community-dwelling elderly Japanese men. Atherosclerosis. 2019;283(5):89–91.

35. Shimizu Y, Yamanashi H, Noguchi Y, Koyamatsu J, Nagayoshi M, Kiyoura K, et al. Cardio-ankle vascular index and circulating CD34-positive cell levels as indicators of endothelial repair activity in older Japanese men. Geriatr Gerontol Int. 2019. In press.

36. Garlet GP, Giozza SP, Silveira EM, Claudino M, Santos SB, Avila-Campos MJ, et al. Association of human T lymphotropic virus 1 amplification of periodontitis severity with altered cytokine expression in response to a standard periodontopathogen infection. Clin Infect Dis. 2010;50(3):e11–8.

37. Paneni F, Osto E, Costantino S, Mateescu B, Briand S, Coppolino G, et al. Deletion of the activated protein-1 transcription factor JunD induces oxidative stress and accelerates age-related endothelial dysfunction. Circulation. 2013;127(11):1229–40 e1–21.

38. Andriollo-Sanchez M, Hininger-Favier I, Meunier N, Verneire E, O’Connor JM, Maiani G, et al. Age-related oxidative stress and antioxidant parameters in middle-aged and older European subjects: the ZENITH study. Eur J Clin Nutr. 2005;59(Suppl 2):S58–62.

39. Rodriguez-Miguelez P, Seigler N, Harris RA. Ultrasound assessment of endothelial function: a technical guideline of the flow-mediated dilation test. J Vis Exp. 2016;110.

40. Gafter-Gvili A, Zingerman B, Rozen-Zvi B, Ori Y, Green H, Lubin I, et al. Oxidative stress-induced DNA damage and repair in human peripheral blood mononuclear cells: protective role of hemoglobin. PLoS One. 2013; 8(7):e68341.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.