

Association of BMI, smoking and alcohol with multiple myeloma mortality in Asians: a pooled analysis of more than 800,000 participants in the Asia Cohort Consortium

Tomotaka Ugai¹⁾, Hidemi Ito²⁾³⁾, Isao Oze¹⁾, Eiko Saito⁴⁾, Md Shafiur Rahman⁵⁾, Paolo Boffetta⁶⁾⁷⁾, Prakash C. Gupta⁸⁾, Norie Sawada⁹⁾, Akiko Tamakoshi¹⁰⁾, Xiao Ou Shu¹¹⁾, Woon-Puay Koh^{12) 13)}, Yu-Tang Gao¹⁴⁾, Atsuko Sadakane¹⁵⁾, Ichiro Tsuji¹⁶⁾, Sue K.

Park¹⁷⁾, Chisato Nagata¹⁸⁾, San-Lin You¹⁹⁾, Mangesh S. Pednekar⁸⁾, Shoichiro Tsugane⁹⁾,

Hui Cai¹¹⁾, Jian-Min Yuan^{20) 21)}, Yong-Bing Xiang²²⁾, Kotaro Ozasa¹⁵⁾, Yasutake

Tomata¹⁶⁾, Seiki Kanemura¹⁶⁾, Yumi Sugawara¹⁶⁾, Keiko Wada¹⁸⁾, Chien-Jen Chen²³⁾,

Keun-Young Yoo²⁴⁾, Kee Seng Chia¹³⁾, Habibul Ahsan²⁵⁾, Wei Zheng¹¹⁾, Manami Inoue⁹⁾,

Daehee Kang¹⁷⁾, John Potter²⁶⁾, Keitaro Matsuo¹⁾²⁷⁾

1) Division of Cancer Epidemiology and Prevention, Department of Preventive Medicine,

Aichi Cancer Center Research Institute, Nagoya, Japan

2) Division of Cancer Information and Control, Department of Preventive Medicine,

Aichi Cancer Center Research Institute, Nagoya, Japan

3) Division of Descriptive Cancer Epidemiology, Nagoya University Graduate School of

Medicine, Nagoya, Japan

- 20 4) Division of Cancer Statistics, Integration Center for Cancer Control & Information
21 Services, National Cancer Center, Tokyo, Japan
- 22 5) Department of Global Health Policy, Graduate School of Medicine, The University of
23 Tokyo, Tokyo, Japan
- 24 6) Icahn School of Medicine at Mount Sinai, New York, NY, USA
- 25 7) Department of Medical and Surgical Sciences, University of Bologna, Bologna, Italy
- 26 8) Healis Sekhsaria Institute for Public Health, Navi Mumbai, India
- 27 9) Epidemiology and Prevention Group, Center for Public Health Sciences, National Cancer
28 Center, Tokyo, Japan
- 29 10) Department of Public Health, Hokkaido University Graduate School of Medicine,
30 Sapporo, Japan
- 31 11) Division of Epidemiology, Vanderbilt-Ingram Cancer Center, Vanderbilt Epidemiology
32 Center, Nashville, TN, USA
- 33 12) Health Services and Systems Research, Duke-NUS Medical School, Singapore,
34 Singapore
- 35 13) Saw Swee Hock School of Public Health, National University of Singapore, Singapore,
36 Singapore.
- 37 14) Department of Epidemiology, Shanghai Cancer Institute, Shanghai, China.
- 38 15) Department of Epidemiology, Radiation Effects Research Foundation, Hiroshima, Japan

- 39 16) Division of Epidemiology, Department of Public Health and Forensic Medicine, Tohoku
40 University Graduate School of Medicine, Sendai, Japan
- 41 17) Department of Preventive Medicine, Seoul National University College of Medicine,
42 Seoul, Korea.
- 43 18) Department of Epidemiology and Preventive Medicine, Gifu University Graduate School
44 of Medicine, Gifu, Japan
- 45 19) School of Medicine & Big Data Research Center, Fu Jen Catholic University, Taiwan
- 46 120) Department of Epidemiology, Graduate School of Public Health, University of
47 Pittsburgh, Pittsburgh, Pennsylvania, USA.
- 48 21) Division of Cancer Control and Population Sciences, UPMC Hillman Cancer Center,
49 University of Pittsburgh, Pittsburgh, Pennsylvania, USA
- 50 22) Department of Epidemiology, Shanghai Cancer Institute, Shanghai, China.
- 51 23) Genomics Research Center, Academia Sinica, Taipei, Taiwan
- 52 24) Department of Preventive Medicine, Seoul National University, Seoul, Korea.
- 53 25) Department of Public Health Sciences, University of Chicago, Chicago, IL, USA.
- 54 26) Fred Hutchinson Cancer Research Center, Seattle, WA, USA.
- 55 27) Division of Cancer Epidemiology, Nagoya University Graduate School of Medicine
- 56

57 **Key words:** body mass index; smoking; alcohol; multiple myeloma; Asia

58

59 The authors declare no potential conflicts of interest.

60

61 **Corresponding author:**

62 Keitaro Matsuo, M.D., Ph.D., M.Sc.

63 Division of Cancer Epidemiology and Prevention, Department of Preventive Medicine, Aichi

64 Cancer Center Research Institute, Nagoya, Japan

65 1-1 Kanokoden, Chikusa-ku, Nagoya 464-8681, Japan., E-mail: kmatsuo@aichi-cc.jp

66 TEL: +81-52-764-2982 FAX: +81-52-763-5233

67

68

69

70

71

72

73

74

75

76

77 **Abstract:**

78 **Background.**

79 To date, few epidemiological studies have been conducted to elucidate lifestyle-related risk
80 factors for multiple myeloma (MM) in Asia. We investigated the association of body mass
81 index (BMI), smoking, and alcohol intake with the risk of MM mortality through a pooled
82 analysis of more than 800,000 participants in the Asia Cohort Consortium.

83 **Methods.**

84 The analysis included 805,309 participants contributing 10,221,623 person-years of
85 accumulated follow-up across Asia Cohort Consortium cohorts. Hazard ratios (HRs) and 95%
86 confidence intervals (95% CIs) for the association between BMI, smoking and alcohol at
87 baseline and the risk of MM mortality were assessed using a Cox proportional hazards model
88 with shared frailty.

89 **Results.**

90 We observed a statistically significant dose-dependent association between BMI categories
91 and the risk of MM mortality (<18.5 kg/m²: HR=0.80, 95% CI: 0.52–1.24; 18.5 to 24.9
92 kg/m²: reference; 25.0 to 29.9 kg/m²: HR=1.17, 0.94–1.47; ≥30 kg/m²: HR=1.61, 0.99-2.64, p
93 for trend=0.014). By sex, this association was more apparent in women than in men (P for
94 heterogeneity between sexes=0.150). We observed no significant associations between

95 smoking or alcohol consumption and risk of MM mortality.

96 **Conclusion.**

97 This study showed that excess body mass is associated with an increased risk of MM
98 mortality among Asian populations. In contrast, our results do not support an association
99 between smoking or alcohol consumption and the risk of MM mortality in Asian populations.

100 **Impact.**

101 This study provides important evidence on the association of BMI, smoking and alcohol with
102 the risk of MM mortality in Asian populations.

103

104

105

106

107

108

109

110

111

112

113

114

115 **Introduction**

116 Multiple myeloma (MM) is the second most common hematologic malignancy and is
117 characterized by the neoplastic proliferation of plasma cells producing monoclonal
118 immunoglobulins (1). Despite recent progress in treatment, MM is still an incurable disease
119 associated with substantial mortality. MM occurs in people of all races and from all
120 geographic locations but its incidence varies greatly across regions and countries (2). In
121 general, Western countries show a higher incidence than Asian countries. For example, in the
122 US, the age-standardized rates (ASRs) were 4.3 and 3.0 per 100,000 for males and females,
123 respectively. In contrast, the ASRs in Asia were reported to be 1.0 and 0.7 per 100,000 for
124 males and females, respectively (3). The difference in incidence suggests a substantial
125 difference in risk-factor exposure between Western and Asian populations.

126 To date, several epidemiological studies have been conducted to elucidate lifestyle-related
127 risk factors for MM. Body mass index (BMI), smoking, and alcohol intake are the most
128 intensively examined factors. However, most epidemiological evidence on these factors has
129 been obtained from studies in Western populations (4-6), whereas evidence from Asian
130 populations is very limited. Because there are large differences in the prevalence of obesity,
131 smoking, and drinking habits (7), as well as of MM incidence between the Western and Asian
132 populations (3), large prospective studies of the associations between these factors and MM

133 risk in Asian populations are, therefore, needed. Given the limited statistical power due to the
134 small number of MM cases in each cohort in Asia, a pooled analysis of the existing cohorts
135 from the Asian Cohort Consortium is one of the ideal approaches to evaluate these
136 relationships.

137 Here, we investigated the association of body mass index, smoking, and alcohol intake
138 with the risk of MM mortality through a pooled analysis of more than 800,000 participants
139 from the Asian Cohort Consortium.

140

141 **Materials and Methods**

142 *Study population*

143 Details of the Asian Cohort Consortium have been described elsewhere (8,9). Briefly, it is a
144 consortium of cohorts in Asian countries developed to explore the association between
145 genetics, environmental exposure, and the etiology of disease, with sufficient statistical
146 power. Among the cohorts participating in the Asian Cohort Consortium, 16 provided
147 information on MM-related deaths during follow-up, as well as data on BMI, smoking,
148 alcohol intake, and potential confounders (sex, age, and education) at baseline. We excluded
149 one cohort with missing data on vital status. Thus, we included 15 cohorts (9 in Japan, 2 in
150 China, 1 in Taiwan, 1 in Korea, 1 in India, and 1 in Singapore) in this pooled analysis. We
151 excluded participants who met any of the following criteria: 1) invalid or missing data on

152 vital status (n=1,565); 2) missing data on age or sex (n=3,163); 3) invalid or missing data on
153 height or weight (n=2,290); or 4) BMI <15 kg/m² or >40 kg/m² (n=16,006). We included a
154 total of 805,309 participants (384,927 men and 420,382 women) in this analysis. The Asian
155 Cohort Consortium coordinating center at the National Cancer Center Japan obtained
156 de-identified individual participant data from all participating cohorts and harmonized it for
157 the statistical analysis.

158 Pooled analysis of the Asian Cohort Consortium cohorts was approved by the ethical
159 committee of the National Cancer Center Japan (number 2014-041) and each study was
160 approved by respective ethic committees overseeing the participating studies. This analysis
161 was also approved by the Institutional Review Board of Aichi Cancer Center Research
162 Institute.

163

164 *Exposure data and study outcome*

165 Height and weight at baseline were directly measured in 5 cohorts and obtained via self-
166 report in 10 cohorts. Information on smoking, alcohol intake, and potential confounders was
167 obtained through baseline questionnaires. BMI was calculated as weight [kg]/(height [m])².
168 We categorized BMI according to the guidelines of the World Health Organization (10) as
169 follows : <18.5 (underweight); 18.5–24.9 (normal weight); 25.0–29.9 (overweight); and ≥30
170 kg/m² (obese). We also applied a five-category BMI classification [<20.0 , $20.0–22.4$, $22.5–$

171 24.9 (reference), 25.0–27.4, and ≥ 27.5 kg/m²] as used in our previous report of a pooled
172 analysis of BMI and overall mortality in the Asian Cohort Consortium (9). Regarding
173 smoking, participants were classified as never, former, or current smokers, as well as by the
174 cumulative exposure to smoking in pack-years (never smokers; <20 pack-years; and ≥ 20
175 pack-years). Alcohol intake was calculated as grams per week to unify data on alcohol intake
176 across the cohorts and then the participants were classified into the following three groups:
177 nondrinkers; intake of 1–149 g/week of ethanol; and intake of ≥ 150 g/week of ethanol. The
178 exposure period for alcohol intake was the year prior to baseline for SMHS, SWHS,
179 Takayama, KMCC and SCHS, and it was not specified for other cohorts.

180 Study outcome was defined as death due to MM (ICD-9: 203 and ICD-10: C90) during
181 follow-up; cause of death was extracted from death certificates.

182

183 *Statistical analysis*

184 To determine the relative risk of MM mortality associated with BMI, smoking, and alcohol
185 intake, we calculated the hazard ratios (HRs) and 95% confidence intervals (CIs) by a Cox
186 proportional hazards model with shared frailty (STATA command *stcox, shared*) (11) using
187 pooled individual participant data. An individual cohort was considered as the shared frailty
188 variable to account for between-study heterogeneity. The details of this statistical model were
189 described in previous articles (11, 12). We estimated two types of HR: model 1, which was

190 adjusted for age at baseline (continuous) and sex (men or women); and model 2, which was
191 adjusted for age at baseline, sex, education (none, primary, secondary, trade or technical,
192 university, post university, and missing) and exposures of interest including body mass index
193 (<18.5 , $18.5\text{--}24.9$, $25.0\text{--}29.9$, ≥ 30 kg/m²); smoking (never-smoker, <20 pack-years, ≥ 20
194 pack-years); and alcohol intake (non-drinker, <150 g/week of ethanol, ≥ 150 g/week of
195 ethanol). Missing values for covariates were treated as dummy variables in the models. We
196 calculated P values for trend using ordinal variables across each exposure category. We also
197 performed stratified analyses based on geographic regions (East Asia: cohorts from China,
198 Japan, Korea, Singapore, and Taiwan; South Asia: cohorts from India) and countries.
199 Likelihood-based methods were used to test for heterogeneity: between sexes; by smoking
200 status; and across geographic regions and countries. The proportional hazards assumptions
201 were tested using scaled Schoenfeld residuals and were found to be justified. All statistical
202 analyses were performed using Stata version 14.1 software (Stata Corp., College Station, TX,
203 USA), and $P < 0.05$ was considered to be statistically significant.

204

205 **Results**

206 Table 1 shows the main characteristics of the participating cohorts in the Asian Cohort
207 Consortium. The final analysis included 805,309 participants accounting for 10,221,623
208 person-years of accumulated follow-up. Mean age at baseline was approximately 54 years in

209 both men and women. Mean BMI ranged from 21.8 to 24.0 kg/m² in men and from 21.9 to
210 24.1 in women. The proportion of ever smokers was 63.7% in men and 6.8% in women,
211 respectively. Mean alcohol intake was 130 g/week in men and 9 g/week in women. During
212 the follow-up period (12.7 years on average), we identified a total of 428 MM-related deaths,
213 accounting for 0.33% of deaths from all causes.

214 Table 2 shows the adjusted HRs for MM mortality based on BMI. We observed a
215 statistically significant dose-dependent association between BMI categories and the risk of
216 MM mortality (<18.5 kg/m²: HR=0.80, 95% CI: 0.52-1.24; 18.5 to 24.9 kg/ m²: reference;
217 25.0 to 29.9 kg/m²: HR=1.17, 0.94-1.47; ≥30 kg/m²: HR=1.61, 0.99-2.64, p for trend=0.014,
218 per 1 kg/m²: HR=1.04, 1.01-1.07, model 2). By sex, we also observed this significant
219 association among women, but observed no clear association among men (P for heterogeneity
220 between sexes=0.150, Supplementary Table 1). When we applied a five-category BMI
221 classification, we observed similar results (Table 1 and Supplementary Table 1).

222 Table 3 shows results of the stratified analysis by region and country. Similar findings
223 were observed when the analysis was restricted to East Asia, but evaluation for data from
224 South Asia was difficult because the number of MM-related deaths was low. We did not
225 observe significant heterogeneity of association across countries. To evaluate whether
226 smoking modified the association between BMI and the risk of MM mortality, we performed
227 stratified analysis by smoking status. The association between BMI and MM mortality was

228 more apparent in never-smokers than ever-smokers, although no formal evidence of
229 heterogeneity was observed by smoking status (Supplementary Table 2).

230 Supplementary Table 3 shows the adjusted HRs for MM mortality based on smoking and
231 alcohol intake categories. We observed no association between smoking or alcohol intake and
232 risk of MM mortality. However, these associations were difficult to evaluate in women
233 because the proportions of female smokers and drinkers were low and the number of the
234 female cases who were smokers and drinkers was small.

235 Finally, we performed several sensitivity analyses as follows: 1) excluding the first 3 and
236 5 years of follow-up, 2) excluding participants with a past history of cancer to avoid the
237 possibility of reverse causality and 3) excluding female participants in the analysis of
238 smoking and alcohol. These analyses did not change our main results substantially
239 (Supplementary Table 4, 5 and 6). We also conducted another sensitivity analysis by
240 excluding one cohort at a time and ensured that our finding was not driven by any single
241 cohort (Supplementary Table 7).

242

243 **Discussion**

244 In this pooled analysis of more than 800,000 Asian participants, we found a statistically
245 significant dose-dependent association between BMI and MM mortality among Asian
246 populations. By sex, this association this association was more apparent in women than in

247 men. We observed no significant association between smoking or alcohol intake and the risk
248 of MM mortality.

249 A positive association between higher BMI and the risk of MM incidence and mortality
250 has been previously reported (4,12). The 2016 IARC update on body fatness and cancer
251 concluded that excess BMI is a risk factor for MM (13). However, although there is sufficient
252 evidence in Western populations, only a few studies have been conducted in Asia. A
253 prospective cohort study, which involved 781,283 Korean men and 103 MM cases, did not
254 show a significant association between higher BMI and MM incidence, among both sexes
255 combined, with an HR of 0.98 (95% CI: 0.30–3.32) for the BMI category of 27.0–29.9 kg/m²
256 relative to the reference category of 18.5–22.9 kg/m² (14). The JPHC study in Japan, which is
257 one of the 15 cohort studies in this pooled analysis, also did not report a statistically
258 significant association between BMI and MM incidence among both sexes combined (23.0–
259 29.9 kg/m²: reference; 25.0–29.9 kg/m²: HR=0.79, 0.45–1.38; ≥30 kg/m²: HR=0.76, 0.45–
260 1.38) (15). Parr et al. conducted a pooled analysis of 424,519 participants in the Asia-Pacific
261 Cohort Collaboration and did not observe a statistically significant association between
262 obesity and MM mortality for both sexes combined, with an HR of 1.20 (95% CI: 0.59–2.43)
263 for the BMI category of ≥30 kg/m² relative to the reference category of 18.5–22.9 kg/m² (16).
264 In contrast, the JACC study in Japan, another cohort study included in this pooled analysis,
265 showed a statistically significant association between obesity and MM mortality only among

266 women (18.5–25.0 kg/m²: reference; ≥ 30 kg/m²: HR=4.11, 1.45–11.64) (17). The current
267 study observed a statistically significant dose-dependent association between BMI and MM
268 mortality also only among women. A recent meta-analysis suggests no sex difference in the
269 association between BMI and MM risk in mainly Western populations (4). This discrepancy
270 could be explained by the differences in body fat distribution (18) and metabolic profiles or
271 in genetic susceptibility to obesity (19) between Western and Asian populations. The possible
272 sex difference in Asian populations should be elucidated in future studies.

273 Different mechanistic pathways for the effect of excess BMI on the development of MM
274 have been proposed (20). Adiponectin, an adipocyte-derived cytokine that is inversely
275 correlated with BMI, has been shown to inhibit proliferation of MM cells and reduce
276 tumorigenic angiogenesis (21,22). In support of this hypothesis, Hofmann et al reported that
277 low levels of circulating adiponectin were associated with MM risk in overweight and obese
278 individuals (23). They also reported that adiponectin levels were significantly lower among
279 MM patients than among patients with monoclonal gammopathy of undetermined
280 significance (MGUS) – the MM precursor – suggesting that reduced expression of
281 adiponectin may be associated with progression from MGUS to MM (24). A recent analysis
282 showed that there is a large variation in adiponectin levels between the sexes and different
283 ethnic groups (25-28); in general, women have higher levels than men and Western
284 populations have higher levels than Asian populations. This difference may explain the sex

285 differences in BMI-associated MM risk, that we report here.

286 BMI is most commonly used to determine adiposity. However, in recent years, it has been
287 reported to be an imprecise measure of body composition, including visceral and
288 subcutaneous adiposity and muscle mass (29). This misclassification might mask true
289 associations. Future researches with direct measures of body composition by computed
290 tomography or dual energy x-ray absorptiometry could better characterize the association
291 between adiposity and subsequent MM incidence and mortality.

292 Most studies have not found an association between smoking and MM risk and a recent
293 meta-analysis of 40 observational studies confirmed this (6). The International Multiple
294 Myeloma Consortium conducted a pooled analysis of nine case-control studies, including
295 2,670 cases and 11,913 controls, and also did not observe an association with smoking (30).
296 Two studies in Japan were similarly null (31,32). Consistent with these studies, our results do
297 not support an association between smoking and MM risk in Asian populations.

298 A pooled analysis of 59 case-control studies, including 1,567 cases and 7,296 controls,
299 reported that ever-drinking was associated with reduced risk of MM (men: OR=0.72, 95%
300 CI: 0.59–0.89; women: OR=0.81, 95% CI: 0.68–0.95) (33). A recent meta-analysis of 26
301 observational studies reported similar findings (pooled relative risk [RR] =0.88, 95% CI:
302 0.79–0.99) (34). However, most of this evidence has been accumulated in Western
303 populations, whereas evidence in Asian populations remains inconclusive. The JACC study

304 (35) and the JPHC study (36), which are participating Japanese cohorts in the Asian Cohort
305 Consortium, did not find an association with alcohol. In the pooled analysis reported here, we
306 found no evidence that alcohol consumption was associated with the risk of MM mortality.

307 The present study has several strengths, most importantly the analysis of
308 individual-level data from a large multi-site, multi-country cohort, allowing better detection
309 of possible associations and the calculation of more precise estimates. Further, the
310 prospective design is less susceptible to recall bias than case-control studies. Several
311 limitations also warrant consideration. First, some cohorts collected anthropometric data that
312 were self-reported, although the validation of the self-reported height, weight, and BMI was
313 high among these cohorts (37,38). Second, as our analyses were conducted using information
314 at a single time point (baseline), we were unable to consider subsequent changes in BMI,
315 smoking habit, and alcohol intake over time. Third, we were unable to consider the effect of
316 other potential confounding factors, including physical activity, MGUS status and family
317 history of hematological malignancies. Fourth, the study outcome of this analysis was
318 mortality, rather than incidence, and we could not distinguish the possible difference in
319 associations with incidence and survival. However, recent analyses have reported that a
320 higher BMI was associated with longer survival among MM patients (39,40), suggesting that
321 the strength of the association we report here with higher BMI is more likely to be under-
322 than over-estimated. In addition, MM was a highly fatal disease during the follow-up period

323 of this study; therefore, the observed associations with MM mortality are a fair representation
324 of the association with MM incidence. Finally, another potential limitation is the accuracy of
325 diagnosis based on death-certificate data. Some fatal MM cases might not have been reported
326 correctly due to the lack of diagnostic precision in some areas; nonetheless, it seems probable
327 that any misclassification occurred independently of body size, smoking and alcohol intake.

328 In conclusion, this study showed that excess body mass is associated with an increased
329 risk of MM mortality among Asian populations. In contrast, our results do not support an
330 association between smoking or alcohol consumption and the risk of MM mortality in Asian
331 populations.

332

333 **Acknowledgements**

334 This work were supported by the following grants: Shanghai Men's Health Study (SMHS),
335 the US National Cancer Institute R01 CA082729 and UM1 CA173640 (Principal
336 Investigator: Xiao-Ou Shu); Shanghai Women's Health Study (SWHS), the US National
337 Cancer Institute (grant numbers: R37 CA070867 and UM1 CA182910 (Principal
338 Investigator: Wei Zheng); Mumbai Cohort Study, International Agency for Research on
339 Cancer, Lyon, France; Clinical Trials Service Unit, Oxford, UK; World Health Organisation,
340 Geneva, Switzerland (Principal Investigator: Prakash C. Gupta); Japan Public Health
341 Center-based prospective Study (JPHC Study) 1 and 2, National Cancer Center Research and

342 Development Fund (23-A-31[toku] and 26-A-2) (since 2011) and a Grant-in-Aid for Cancer
343 Research from the Ministry of Health, Labour and Welfare of Japan (from 1989 to 2010)
344 (Principal Investigator: Shoichiro Tsugane); Japan Collaborative Cohort Study (JACC),
345 National Cancer Center Research and Development Fund, A Grant-in-Aid for Cancer
346 Research; Grant for Health Services and Grant for Comprehensive Research on
347 Cardiovascular and Life-Style Related Diseases from the Ministry of Health, Labour and
348 Welfare, Japan; Grant for the Scientific Research from the Ministry of Education, Culture,
349 Sports, Science and Technology, Japan (Principal Investigator: Akiko Tamakoshi); Miyagi
350 Cohort Study, National Cancer Center Research and Development Fund (Principal
351 Investigator: Ichiro Tsuji); Ohsaki Cohort Study, National Cancer Center Research and
352 Development Fund (Principal Investigator: Ichiro Tsuji); Radiation Effects Research
353 Foundation, The Japanese Ministry of Health, Labour and Welfare and the US Department of
354 Energy (Principal Investigator: Atsuko Sadakane); Takayama Study, National Cancer Center
355 Research and Development Fund (Principal Investigator: Chisato Nagata); 3 Prefecture
356 Miyagi Study, National Cancer Center Research and Development Fund (Principal
357 Investigator: Ichiro Tsuji); 3 Prefecture Aichi Study, The Japanese Ministry of the
358 Environment (former Environment Agency) (Principal Investigator: Keitaro Matsuo);
359 Singapore Chinese Health Study, the US National Cancer Institutes R01CA144034 and
360 UM1CA182876 (Principal Investigator: Jian-Min Yuan); Community-based Cancer

361 Screening Project (CBCSP), Ministry of Health and Welfare and Ministry of Science and
362 Technology, Taiwan (Principal Investigator: San-Lin You); and ACC Coordinating Center,
363 National Cancer Center Research and Development Fund (30-A-15) (Principal Investigator:
364 Manami Inoue) .

365

366 **References**

- 367 1. Palumbo A, Anderson K. Multiple myeloma. *N Engl J Med* **2011**;364(11):1046-60
- 368 2. Cowan AJ, Allen C, Barac A, Basaleem H, Bensenor I, Curado MP, *et al.* Global
369 Burden of Multiple Myeloma: A Systematic Analysis for the Global Burden of
370 Disease Study 2016. *JAMA Oncol* **2018**;4(9):1221-7
- 371 3. Ferlay J, Soerjomataram I, Dikshit R, Eser S, Mathers C, Rebelo M, *et al.* Cancer
372 incidence and mortality worldwide: sources, methods and major patterns in
373 GLOBOCAN 2012. *Int J Cancer* **2015**;136(5):E359-86
- 374 4. Wallin A, Larsson SC. Body mass index and risk of multiple myeloma: a
375 meta-analysis of prospective studies. *Eur J Cancer* **2011**;47(11):1606-15
- 376 5. Rota M, Porta L, Pelucchi C, Negri E, Bagnardi V, Bellocco R, *et al.* Alcohol drinking
377 and multiple myeloma risk--a systematic review and meta-analysis of the dose-risk
378 relationship. *Eur J Cancer Prev* **2014**;23(2):113-21
- 379 6. Psaltopoulou T, Sergentanis TN, Kanellias N, Kanavidis P, Terpos E, Dimopoulos MA.
380 Tobacco smoking and risk of multiple myeloma: a meta-analysis of 40 observational
381 studies. *Int J Cancer* **2013**;132(10):2413-31
- 382 7. Mendis S. Global status report on noncommunicable diseases 2014. World health
383 organization; 2014.
- 384 8. Rolland B, Smith BR, Potter JD. Coordinating centers in cancer epidemiology
385 research: the Asia Cohort Consortium coordinating center. *Cancer Epidemiol*
386 *Biomarkers Prev* **2011**;20(10):2115-9
- 387 9. Zheng W, McLerran DF, Rolland B, Zhang X, Inoue M, Matsuo K, *et al.* Association
388 between body-mass index and risk of death in more than 1 million Asians. *N Engl J*
389 *Med* **2011**;364(8):719-29
- 390 10. Organization WH. Physical status: The use of and interpretation of anthropometry,
391 Report of a WHO Expert Committee. **1995**.
- 392 11. O'Quigley J, Stare J. Proportional hazards models with frailties and random effects.

- 393 Stat Med **2002**;21(21):3219-33
- 394 12. Teras LR, Kitahara CM, Birmann BM, Hartge PA, Wang SS, Robien K, *et al.* Body
395 size and multiple myeloma mortality: a pooled analysis of 20 prospective studies. Br J
396 Haematol **2014**;166(5):667-76
- 397 13. Lauby-Secretan B, Scoccianti C, Loomis D, Grosse Y, Bianchini F, Straif K. Body
398 Fatness and Cancer--Viewpoint of the IARC Working Group. N Engl J Med
399 **2016**;375(8):794-8
- 400 14. Oh SW, Yoon YS, Shin SA. Effects of excess weight on cancer incidences depending
401 on cancer sites and histologic findings among men: Korea National Health Insurance
402 Corporation Study. J Clin Oncol **2005**;23(21):4742-54
- 403 15. Kanda J, Matsuo K, Inoue M, Iwasaki M, Sawada N, Shimazu T, *et al.* Association of
404 anthropometric characteristics with the risk of malignant lymphoma and plasma cell
405 myeloma in a Japanese population: a population-based cohort study. Cancer
406 Epidemiol Biomarkers Prev **2010**;19(6):1623-31
- 407 16. Parr CL, Batty GD, Lam TH, Barzi F, Fang X, Ho SC, *et al.* Body-mass index and
408 cancer mortality in the Asia-Pacific Cohort Studies Collaboration: pooled analyses of
409 424,519 participants. Lancet Oncol **2010**;11(8):741-52
- 410 17. Khan MM, Mori M, Sakauchi F, Matsuo K, Ozasa K, Tamakoshi A. Risk factors for
411 multiple myeloma: evidence from the Japan Collaborative Cohort (JACC) study.
412 Asian Pac J Cancer Prev **2006**;7(4):575-81.
- 413 18. Wulan SN, Westerterp KR, Plasqui G. Ethnic differences in body composition and the
414 associated metabolic profile: a comparative study between Asians and Caucasians.
415 Maturitas **2010**;65(4):315-9
- 416 19. Akiyama M, Okada Y, Kanai M, Takahashi A, Momozawa Y, Ikeda M, *et al.*
417 Genome-wide association study identifies 112 new loci for body mass index in the
418 Japanese population. Nat Genet **2017**;49(10):1458-67
- 419 20. Fairfield H, Falank C, Avery L, Reagan MR. Multiple myeloma in the marrow:
420 pathogenesis and treatments. Ann N Y Acad Sci **2016**;1364:32-51
- 421 21. Brakenhielm E, Veitonmaki N, Cao R, Kihara S, Matsuzawa Y, Zhivotovsky B, *et al.*
422 Adiponectin-induced antiangiogenesis and antitumor activity involve
423 caspase-mediated endothelial cell apoptosis. Proc Natl Acad Sci U S A
424 **2004**;101(8):2476-81
- 425 22. Dalamaga M, Diakopoulos KN, Mantzoros CS. The role of adiponectin in cancer: a
426 review of current evidence. Endocr Rev **2012**;33(4):547-94
- 427 23. Hofmann JN, Birmann BM, Teras LR, Pfeiffer RM, Wang Y, Albanes D, *et al.* Low
428 Levels of Circulating Adiponectin Are Associated with Multiple Myeloma Risk in
429 Overweight and Obese Individuals. Cancer Res **2016**;76(7):1935-41
- 430 24. Hofmann JN, Mailankody S, Korde N, Wang Y, Tajeja N, Costello R, *et al.*

- 431 Circulating Adiponectin Levels Differ Between Patients with Multiple Myeloma and
432 its Precursor Disease. *Obesity (Silver Spring)* **2017**;25(8):1317-20
- 433 25. Song HJ, Oh S, Quan S, Ryu OH, Jeong JY, Hong KS, *et al.* Gender differences in
434 adiponectin levels and body composition in older adults: Hallym aging study. *BMC*
435 *Geriatr* **2014**;14:8
- 436 26. Boyne MS, Bennett NR, Cooper RS, Royal-Thomas TY, Bennett FI, Luke A, *et al.*
437 Sex-differences in adiponectin levels and body fat distribution: longitudinal
438 observations in Afro-Jamaicans. *Diabetes Res Clin Pract* **2010**;90(2):e33-6
- 439 27. Nakano Y, Tajima S, Yoshimi A, Akiyama H, Tsushima M, Tanioka T, *et al.* A novel
440 enzyme-linked immunosorbent assay specific for high-molecular-weight adiponectin.
441 *J Lipid Res* **2006**;47(7):1572-82
- 442 28. Mente A, Razak F, Blankenberg S, Vuksan V, Davis AD, Miller R, *et al.* Ethnic
443 variation in adiponectin and leptin levels and their association with adiposity and
444 insulin resistance. *Diabetes Care* **2010**;33(7):1629-34
- 445 29. Caan BJ, Cespedes Feliciano EM, Kroenke CH. The Importance of Body
446 Composition in Explaining the Overweight Paradox in Cancer-Counterpoint. *Cancer*
447 *Res* **2018**;78(8):1906-12
- 448 30. Andreotti G, Birmann BM, Cozen W, De Roos AJ, Chiu BC, Costas L, *et al.* A pooled
449 analysis of cigarette smoking and risk of multiple myeloma from the international
450 multiple myeloma consortium. *Cancer Epidemiol Biomarkers Prev* **2015**;24(3):631-4
- 451 31. Ozasa K. Smoking and mortality in the Japan Collaborative Cohort Study for
452 Evaluation of Cancer (JACC). *Asian Pac J Cancer Prev* **2007**;8 Suppl:89-96
- 453 32. Sonoda T, Ishida T, Mori M, Sakai H, Noguchi M, Imai K. A case-control study of
454 multiple myeloma in Japan: association with occupational factors. *Asian Pac J Cancer*
455 *Prev* **2005**;6(1):33-6.
- 456 33. Andreotti G, Birmann B, De Roos AJ, Spinelli J, Cozen W, Camp NJ, *et al.* A pooled
457 analysis of alcohol consumption and risk of multiple myeloma in the international
458 multiple myeloma consortium. *Cancer Epidemiol Biomarkers Prev* **2013**;22(9):1620-7
- 459 34. Psaltopoulou T, Sergentanis TN, Sergentanis IN, Karadimitris A, Terpos E,
460 Dimopoulos MA. Alcohol intake, alcoholic beverage type and multiple myeloma risk:
461 a meta-analysis of 26 observational studies. *Leuk Lymphoma* **2015**;56(5):1484-501
- 462 35. Ozasa K. Alcohol use and mortality in the Japan Collaborative Cohort Study for
463 Evaluation of Cancer (JACC). *Asian Pac J Cancer Prev* **2007**;8 Suppl:81-8
- 464 36. Kanda J, Matsuo K, Inoue M, Iwasaki M, Sawada N, Shimazu T, *et al.* Association of
465 alcohol intake with the risk of malignant lymphoma and plasma cell myeloma in
466 Japanese: a population-based cohort study (Japan Public Health Center-based
467 Prospective Study). *Cancer Epidemiol Biomarkers Prev* **2010**;19(2):429-34
- 468 37. Inoue M, Sobue T, Tsugane S. Impact of body mass index on the risk of total cancer

- 469 incidence and mortality among middle-aged Japanese: data from a large-scale
470 population-based cohort study--the JPHC study. *Cancer Causes Control*
471 **2004**;15(7):671-80
- 472 38. Shimizu N, Nagata C, Shimizu H, Kametani M, Takeyama N, Ohnuma T, *et al.*
473 Height, weight, and alcohol consumption in relation to the risk of colorectal cancer in
474 Japan: a prospective study. *Br J Cancer* **2003**;88(7):1038-43
- 475 39. Beason TS, Chang SH, Sanfilippo KM, Luo S, Colditz GA, Vij R, *et al.* Influence of
476 body mass index on survival in veterans with multiple myeloma. *Oncologist*
477 **2013**;18(10):1074-9
- 478 40. Jung SH, Yang DH, Ahn JS, Lee SS, Ahn SY, Kim YK, *et al.* Decreased body mass
479 index is associated with poor prognosis in patients with multiple myeloma. *Ann*
480 *Hematol* **2014**;93(5):835-40
- 481

Table 1. Characteristics of the cohort studies in the present pooled analysis.

Country and Study	No. of subjects	Enrollment period	Mean Follow-up, years (SD)	Person-Years	Mean Age at Baseline, years (SD)		Mean BMI at Baseline (SD)		Method of Height and Weight Ascertained	% of Ever Smokers		Mean Alcohol Intake at Baseline, g/week (SD)		No. of Deaths	No. of myeloma Deaths	% of myeloma Deaths
					Men	Women	Men	Women		Men	Women	Men	Women			
					China											
SMHS	61,426	2001-2006	9.5 (1.8)	581,041	55.4 (9.7)	NA	23.7 (3.1)	NA	DM	69.6	NA	82 (175)	NA	5,423	22	0.41%
SWHS	74,862	1997-2000	14.9 (2.3)	1,115,384	NA	52.6 (9.1)	NA	24.0 (3.4)	DM	NA	2.8	NA	1 (17)	7,618	35	0.46%
India																
Mumbai	145,093	1991-1997	5.2 (1.5)	755,039	52.6 (10.9)	48.0 (11.0)	22.1 (3.7)	22.8 (4.5)	DM	31.3	0.4	NA	NA	12,456	10	0.08%
Japan																
3pref. Aichi	32,142	1985	11.5 (5.1)	372,220	55.6 (11.0)	56.6 (11.4)	22.3 (2.8)	22.0 (3.0)	SA	82.6	15.2	NA	NA	5,404	14	0.26%
JPHC1	42,728	1990-1992	21.0 (4.3)	897,432	49.5 (5.9)	49.7 (5.9)	23.6 (2.8)	23.6 (3.1)	SA	75.7	7.5	237 (314)	16 (101)	7,392	37	0.50%
JPHC2	55,675	1992-1995	17.7 (4.2)	986,710	54.0 (8.8)	54.4 (8.8)	23.5 (2.9)	23.4 (3.2)	SA	75.5	7.7	211 (289)	15 (79)	12,517	66	0.53%
JACC	86,566	1988-1990	12.7 (3.4)	1,097,249	57.6 (10.2)	57.6 (9.9)	22.6 (2.8)	22.9 (3.1)	SA	76.2	6.0	NA	NA	12,851	72	0.56%
Miyagi	44,842	1990	16.2 (3.7)	725,882	51.7 (7.6)	52.2 (7.4)	23.5 (2.8)	23.7 (3.1)	SA	79.6	8.3	186 (185)	13 (53)	5,233	26	0.50%
Ohsaki	47,607	1995	10.8 (4.3)	513,397	59.5 (10.6)	60.7 (9.9)	23.3 (2.9)	23.7 (3.2)	SA	77.5	8.8	187 (186)	16 (68)	7,993	39	0.49%
RERF	49,424	1963-1993	21.9 (10.3)	1,084,701	52.4 (11.1)	51.9 (15.0)	21.8 (3.0)	21.9 (3.3)	SA	85.4	14.7	204 (262)	15 (63)	25,530	34	0.13%
3pref. Miyagi	29,443	1984	11.6 (5.0)	340,376	56.5 (11.0)	52.2 (7.4)	23.0 (2.9)	23.3 (3.4)	SA	59.3	8.7	NA	NA	5,848	8	0.14%
Takayama	29,640	1992	13.7 (4.0)	405,102	54.9 (12.2)	55.8 (13.0)	22.5 (2.8)	22.0 (2.9)	SA	81.8	15.9	288 (287)	54 (118)	5,465	17	0.31%
Korea																
KMCC	18,962	1994-2004	13.8 (4.7)	261,165	53.5 (14.5)	53.9 (14.3)	23.1 (3.0)	23.9 (3.4)	DM	78.3	8.3	12 (40)	3 (5)	3,477	8	0.23%
Singapore																
SCHS	63,147	1993-1999	11.5 (3.0)	723,862	56.7 (8.0)	56.3 (8.0)	23.0 (3.1)	23.2 (3.2)	SA	58.0	8.8	25 (80)	3 (17)	10,657	33	0.31%
Taiwan																
CBCSP	23,752	1991-1992	15.2 (2.6)	362,062	48.0 (10.2)	46.6 (9.8)	24.0 (3.2)	24.1 (3.5)	DM	56.2	0.9	NA	NA	2,755	7	0.25%
Total	805,309		12.7 (6.3)	10,221,623	54.3 (10.4)	53.6 (10.9)	23.0 (3.2)	23.2 (3.5)		63.7	6.8	130 (222)	9 (55)	130,619	428	0.33%

Abbreviations: BMI, body mass index; SMHS, Shanghai Men's Health Study; SWHS, Shanghai Women's Health Study; Mumbai, Mumbai Cohort Study; 3-Pref Aichi, Three-Prefecture Association for Cancer Research.

Cohort Study Aichi; JPHC, Japan Public Health Center-based prospective Study (1 and 2) ; JACC, The Japan Collaborative Cohort Study; Miyagi, The Miyagi Cohort Study; Ohsaki: Ohsaki Cohort Study; RERF, Radiation Effects Research Foundation; 3-Pref Miyagi, Three-Prefecture Cohort Study; Miyagi; Takayama, Takayama Cohort Study; KMCC, Korea Multi-Center Cancer Cohort; SCHS; Singapore Chinese Health Study; CBCSP, Community-Based Cancer Screening Project; NA, not available.

Table 2. Risk of multiple myeloma mortality according to body mass index

	Body Mass Index					P trend ¹	Per 1 kg/m ²
	<18.5	18.5–24.9	25.0–29.9	≥30			
Person-years	619,504	6,961,719	2,356,839	283,561			
No. of cases	22	280	109	17			
HR (model 1)	0.81 (0.52-1.25)	1.00 (Reference)	1.17 (0.93-1.46)	1.60 (0.98-2.61)		0.016	1.04 (1.01-1.07)
HR (model 2)	0.80 (0.52-1.24)	1.00 (Reference)	1.17 (0.94-1.47)	1.61 (0.99-2.64)		0.014	1.04 (1.01-1.07)
	<20.0	20.0–22.4	22.5–24.9	25.0–27.4	≥27.5		
Person-years	1,590,661	2,894,330	3,096,232	1,706,921	933,479		
No. of cases	58	121	123	76	50		
HR (model 1)	0.89 (0.65-1.22)	1.06 (0.82-1.36)	1.00 (Reference)	1.14 (0.85-1.51)	1.41 (1.01-1.96)	0.035	
HR (model 2)	0.88 (0.64-1.21)	1.06 (0.82-1.36)	1.00 (Reference)	1.14 (0.85-1.51)	1.42 (1.02-1.98)	0.030	

model 1: HRs are adjusted for age and sex.

model 2: HRs are adjusted for age, sex, smoking, alcohol intake and education.

¹ P-values for trends were calculated by assigning scores for categories of body mass index.

Table 3. Risk of multiple myeloma mortality according to body mass index by region and country

	Body Mass Index				P trend ¹	Per 1 kg/m ²	P for heterogeneity ²
	<18.5	18.5–24.9	25.0–29.9	≥30			
East Asia							
Person-years	490,945	6,521,799	2,203,889	249,951			
No. of cases	21	274	106	17			
HR (model 1)	0.86 (0.55-1.35)	1.00 (Reference)	1.14 (0.91-1.43)	1.59 (0.97-2.61)	0.039	1.03 (1.00-1.06)	0.544
HR (model 2)	0.86 (0.55-1.34)	1.00 (Reference)	1.15 (0.92-1.45)	1.64 (1.00-2.67)	0.028	1.03 (1.00-1.07)	0.545
South Asia (India)							
Person-years	128,559	439,921	152,950	33,610			
No. of cases	1	6	3	0			
HR (model 1)	0.49 (0.06-4.10)	1.00 (Reference)	1.59 (0.40-6.40)	NA	0.551	1.06 (0.92-1.23)	
HR (model 2)	0.54 (0.07-4.63)	1.00 (Reference)	1.54 (0.38-6.21)	NA	0.639	1.05 (0.90-1.23)	
China							
Person-years	61,477	1,053,353	512,047	69,548			
No. of cases	1	24	24	8			
HR (model 1)	0.70 (0.09-5.14)	1.00 (Reference)	1.71 (0.97-3.02)	3.85 (1.71-8.69)	0.001	1.14 (1.06-1.22)	0.206
HR (model 2)	0.68 (0.09-5.04)	1.00 (Reference)	1.72 (0.97-3.05)	3.85 (1.69-8.74)	0.001	1.14 (1.06-1.22)	0.196
Japan							
Person-years	362,488	4,559,859	1,367,655	133,067			
No. of cases	18	217	70	8			
HR (model 1)	0.88 (0.54-1.44)	1.00 (Reference)	1.05 (0.80-1.37)	1.20 (0.59-2.43)	0.451	1.01 (0.97-1.05)	
HR (model 2)	0.89 (0.55-1.45)	1.00 (Reference)	1.05 (0.80-1.38)	1.23 (0.61-2.49)	0.422	1.02 (0.98-1.05)	
Korea							
Person-years	11,414	168,680	72,043	9,028			
No. of cases	0	7	1	0			
HR (model 1)	NA	1.00 (Reference)	0.36 (0.04-2.90)	NA	0.473	0.90 (0.71-1.13)	
HR (model 2)	NA	1.00 (Reference)	0.38 (0.05-3.12)	NA	0.561	0.91 (0.72-1.16)	

Singapore

Person-years	44,396	518,521	139,586	21,360		
No. of cases	2	21	10	0		
HR (model 1)	1.06 (0.25-4.54)	1.00 (Reference)	1.89 (0.89-4.02)	NA	0.482	1.05 (0.95-1.17)
HR (model 2)	1.06 (0.25-4.54)	1.00 (Reference)	1.91 (0.90-4.06)	NA	0.480	1.05 (0.95-1.17)

Taiwan

Person-years	11,170	221,386	112,558	16,949		
No. of cases	0	5	1	1		
HR (model 1)	NA	1.00 (Reference)	0.33 (0.04-2.82)	2.22 (0.26-19.0)	0.994	1.04 (0.84-1.30)
HR (model 2)	NA	1.00 (Reference)	0.33 (0.04-2.84)	2.25 (0.26-19.6)	0.985	1.04 (0.84-1.30)

model 1: HRs are adjusted for age and sex.

model 2: HRs are adjusted for age, sex, smoking, alcohol intake and education.

1 P-values for trends were calculated by assigning scores for categories of body mass index, with 1 for <18.5, 2 for 18.5-24.9, 3 for 25.0–29.9, 4 for ≥30 kg/m².

2 Heterogeneity for trend among regions or among countries.

Abbreviations: NA, not available.

BLOOD CANCER DISCOVERY

Association of BMI, smoking and alcohol with multiple myeloma mortality in Asians: a pooled analysis of more than 800,000 participants in the Asia Cohort Consortium

Tomotaka Ugai, Hidemi Ito, Isao Oze, et al.

Cancer Epidemiol Biomarkers Prev Published OnlineFirst August 9, 2019.

Updated version	Access the most recent version of this article at: doi: 10.1158/1055-9965.EPI-19-0389
Supplementary Material	Access the most recent supplemental material at: http://cebp.aacrjournals.org/content/suppl/2019/08/09/1055-9965.EPI-19-0389.DC1
Author Manuscript	Author manuscripts have been peer reviewed and accepted for publication but have not yet been edited.

E-mail alerts	Sign up to receive free email-alerts related to this article or journal.
Reprints and Subscriptions	To order reprints of this article or to subscribe to the journal, contact the AACR Publications Department at pubs@aacr.org .
Permissions	To request permission to re-use all or part of this article, use this link http://cebp.aacrjournals.org/content/early/2019/08/09/1055-9965.EPI-19-0389 . Click on "Request Permissions" which will take you to the Copyright Clearance Center's (CCC) Rightslink site.