

1 **Influence of sex, age, pubertal maturation and body mass index on circulating white blood**
2 **cell counts in healthy European adolescents - The HELENA STUDY**

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25

26 **Abstract**

27 Percentiles 10th, 25th, 50th, 75th and 90th are presented for total circulating white blood cells
28 (WBC), neutrophils, lymphocytes, monocytes, eosinophils and basophils in healthy European
29 adolescents (12.5-17.5 years, n=405, 48.9% boys), considering age, sex, puberty and body mass
30 index (BMI). CD3⁺ (mature T cells), CD4⁺ (T helper), CD8⁺ (T cytotoxic), CD16⁺56⁺ (natural
31 killer), CD19⁺ (B cells), CD45RA⁺ (naïve) and CD45RO⁺ (memory) lymphocytes were also
32 analysed by immunophenotyping. Girls presented higher WBC, neutrophil, CD3⁺CD45RO⁺ and
33 CD4⁺CD45RO⁺ cell counts and CD3⁺/CD19⁺ ratio, and lower CD3⁺CD45RA⁺ and
34 CD4⁺CD45RA⁺ counts than boys. Age was associated with higher neutrophil counts and
35 CD3⁺/CD19⁺ ratio and lower CD19⁺ counts; in boys, with lower CD3⁺CD45RA⁺,
36 CD4⁺CD45RA⁺ and CD8⁺CD45RA⁺ counts as well; in girls, with higher WBC, CD3⁺CD45RO⁺,
37 and CD4⁺CD45RO⁺ counts. Pubertal maturation in boys was associated with lower WBC and
38 lymphocyte counts; in girls, with higher basophil, CD3⁺CD45RO⁺ and CD4⁺CD45RO⁺ values.
39 BMI was associated with higher WBC counts; in boys, also with higher lymphocyte counts; in
40 girls, with higher neutrophil, CD4⁺, CD3⁺CD45RO⁺ and CD4⁺CD45RO⁺ counts. *Conclusions:*
41 Our study provides normative values for circulating immune cells in adolescents, highlighting

42 the importance of considering sex, age, pubertal maturation and BMI when establishing
43 reference values for WBC in paediatric populations.

44 **Key words:** adolescents; immune cells; immunophenotyping; sex; puberty; body mass index.

45 **Abbreviations used**

46 BMI – body mass index

47 WBC – white blood cells

48 **What is known?**

49 Reference values for white blood cell counts and immunophenotyping of lymphocyte subsets can
50 constitute useful clinical tools and health indicators in both adult and paediatric populations.

51 Like other health indicators, they can vary according to a number of factors, such as age or sex.

52 **What is new?**

53 Specific data from WBC in European adolescents are limited, and even less is known about
54 changes in lymphocyte subsets during adolescence. This work provides normative values
55 obtained from adolescents from 9 European countries, considering the influence of sex, age,
56 pubertal maturation and BMI and finding that:

- 57 • Girls had higher WBC, neutrophil and CD45RO⁺ (memory) cell values, while boys had
58 higher percentages of lymphocytes, monocytes and eosinophils and CD45RA⁺ (naive)
59 cell counts.
- 60 • Age was associated with higher WBC, neutrophil, CD45RO⁺ and CD3⁺/CD19⁺ values,
61 and lower percentage of total lymphocytes and CD45RA⁺ cell counts.
- 62 • Pubertal maturation was associated, on the contrary, with lower WBC and lymphocytes,
63 but only in boys. In girls, pubertal maturation was linked to higher CD45RO⁺ cell counts.
- 64 • BMI was associated with higher WBC, mainly due to greater lymphocyte counts in boys,
65 and to neutrophil and CD45RO⁺ cell counts in girls.

66 **Introduction**

67 Total white blood cell (WBC) counts and the evaluation of the different subtypes of white blood
68 cells are useful clinical indicators and are frequently used as diagnostic tools for adults as well as
69 for children. Besides providing information on acute inflammatory and infectious states, the
70 immunological status serves as an indicator of many other physiological processes, and immune
71 markers are becoming increasingly used to study alterations other than infections – for example,
72 as early as in the 70s, Friedman and colleagues reported an association between WBC count and
73 myocardial infarction [9]. Immune function is closely related to overall health and nutritional
74 status [31], and during the last decade, WBC count has been related to different metabolic
75 alterations, such as impaired glucose tolerance, type 2 diabetes mellitus, obesity, or the metabolic
76 syndrome, and this has been observed in adults [10, 25] as well as in children and adolescents [5,
77 36]. The analysis of lymphocyte subsets by flow cytometry (known as immunophenotyping) is
78 an ever more widely used tool not only for assessing health status but also specifically in
79 nutritional evaluation, and helps identify subjects at risk of disease.

80 Adequate and reliable reference values from healthy populations become a key point in
81 the clinical use of any biological parameter. Variations can occur as a consequence of
82 physiological, ethnic or environmental factors; therefore, normative values should be specific for
83 a given population group. For example, childhood and adolescence are growth periods in which
84 all the systems in the body are developing physically and functionally. The immune system itself
85 experiences a series of modifications from birth until adulthood; during childhood and
86 adolescence the immune cells vary in both their number and functionality [1, 6, 8, 13, 16, 18, 28,
87 32, 34]. Therefore, normative values for immune cells obtained from adult populations can be
88 misleading when applied to children and adolescents. In addition, sex-related and even ethnicity-
89 related variations in the values of circulating immune cells have been acknowledged [1, 4, 7, 17,
90 28, 29, 34, 38,], highlighting the importance of providing reference values for specific

91 population groups and geographical areas. However, information about WBC counts and specific
92 lymphocyte subsets on healthy adolescents is still scarce [1, 8, 16, 28, 32, 34], as most available
93 data belong to populations with particular immune-related diseases, or the studies, although
94 useful and informative, have usually been conducted on relatively modest sample sizes for this
95 particular age group.

96 The present study aims to provide normative ranges for total and differential WBC counts
97 and for selected lymphocyte subsets in a representative sample of healthy European adolescents,
98 attending to variations due to sex, age, degree of pubertal maturation, and body mass index.

99

100 **METHODS**

101 **Study design and sample selection**

102 A European multicentre cross-sectional study (CSS) was performed with the objective of
103 assessing a “healthy lifestyle in Europe by nutrition in adolescence” (HELENA). The HELENA-
104 CSS aimed to obtain reliable and comparable data on nutrition and other health indicators such
105 as physical activity and fitness, body composition, cardiovascular disease risk factors, vitamin
106 and mineral status, and immunological and genetic markers in European adolescents [21]. The
107 methodology used in this study has been published elsewhere [20]. The study was performed
108 according to the ethical guidelines of the Edinburgh revision of the 1964 Declaration of Helsinki
109 (2000), the International Conferences on Harmonization for Good Clinical Practice and the
110 legislation on clinical research from each of the participating countries. The protocol was
111 approved by the Research Ethics Committees of the participating centres. Written informed
112 consent was obtained from the parents of the adolescents and from the adolescents themselves
113 [2].

114 Briefly, subjects aged 12.5-17.5 years were recruited randomly from schools in ten cities
115 belonging to nine countries across Europe (Athens and Heraklion in Greece, Dortmund in

116 Germany, Ghent in Belgium, Lille in France, Pécs in Hungary, Rome in Italy, Stockholm in
117 Sweden, Vienna in Austria and Zaragoza in Spain). The total eligible HELENA-CSS population
118 consisted of 3,528 adolescents. Blood samples were obtained in one third of participants,
119 resulting in a representative subpopulation of 1,089 adolescents (approximately 100 boys and
120 girls per city). The size of this subpopulation was previously calculated as sufficient to account
121 for the expected variability in blood measurements.

122

123 **Data exclusion**

124 Those subjects with conditions that might interfere with or imply stimulation of normal immune
125 function were excluded from the analysis. Exclusion criteria included: suffering from allergies,
126 suffering from fever on the 24 hours prior to blood sampling, having had a cold or any infection
127 during the week prior to the day of blood sampling, having taken any medication in the previous
128 24 hours or for more than 7 days in the previous 30 days, having taken any vitamin or mineral
129 supplement during the previous month, and having been vaccinated in the two weeks prior to the
130 day of blood sampling. After applying these exclusion criteria, the final study population
131 consisted of 405 subjects (48.9% boys).

132

133 **Measurements of pubertal maturation and body mass index**

134 Evaluation of the degree of pubertal maturation was assessed by a medical doctor, according to
135 the Tanner and Whitehouse classification [33]. Anthropometric data were also collected
136 following harmonized protocol procedures previously described [23]. Body mass index (BMI)
137 was calculated as: body weight (kg) / [height (m)]². Standardized BMI values (z-scores) were
138 calculated and the sample was classified according to quartiles of standardized BMI values.

139

140 **Blood sampling, white blood cell profiling and immunophenotyping**

141 Venous blood samples were collected in EDTA Monovette (Sarstedt, Germany) tubes between
142 8.00 and 10.00 a.m. after a 12-hour overnight fast. WBC counts and percentages were
143 determined in each participating city with automated blood cell counters. For immuno-
144 phenotyping of lymphocyte subsets, blood samples were collected in EDTA-K3E Vacutainer
145 (BD Biosciences) tubes. Blood aliquots were taken into 1.5 ml plastic tubes and diluted 1:1 with
146 Cytochex™ Reagent (Streck Laboratories, Omaha, NE, USA). The samples were all sent to the
147 CSIC group laboratory (Madrid, Spain) within 7 days from collection. The methodology for
148 WBC determination and for collection, preparation and shipping of the blood samples to Madrid
149 was standardized amongst all participating cities [11].

150

151 *Immunophenotyping*

152 Blood aliquots were incubated for 30 minutes at room temperature in the dark with
153 fluorochrome-conjugated monoclonal antibodies (BD Biosciences, San José, CA, USA), to
154 differentially label those cells positive for the surface markers CD45 (the pan-leukocyte marker),
155 CD3 (T mature cells), CD4 (helper T cells), CD8 (cytotoxic T cells), CD19 (B cells), CD16⁺56
156 (natural killer cells), CD45RO (memory cells) and CD45RA (naïve cells). A quadruple
157 immunostaining procedure was performed as follows: CD3/CD8/CD45/CD4,
158 CD45RA/CD45RO/CD8/CD3, CD45RA/CD45RO/CD4/CD3 and CD3/CD16⁺56/CD45/ /CD19.
159 After lysis of red blood cells, lymphocytes were analysed by flow cytometry (FACScan Plus
160 Dual Laser, Becton Dickinson Sunnyvale, CA). The lympho-gate was defined on the forward
161 and side scatter patterns of lymphocytes. The analysis protocol gated on lymphocytes stained
162 with PerCP and/or APC and the selected population was then analysed with the two remaining
163 colours (FITC and PE) to obtain percentages of cell expressing the specific antigens.

164 Percentages of the following lymphocyte subsets were obtained: total mature T cells
165 (CD45⁺CD3⁺, or CD3⁺ for simplicity), helper T cells (CD45⁺CD3⁺CD4⁺, or CD4⁺), cytotoxic T

166 cells (CD45⁺CD3⁺CD8⁺, or CD8⁺), natural killer (NK) cells (CD45⁺CD3⁻16⁺56⁺, or CD16⁺56⁺),
167 B cells (CD45⁺CD3⁻CD19⁺, or CD19⁺), naïve T cells (CD3⁺CD45RA⁺, CD4⁺CD45RA⁺ and
168 CD8⁺CD45RA⁺, or CD3RA⁺, CD4RA⁺ and CD8RA⁺), and memory T cells (CD3⁺CD45RO⁺,
169 CD4⁺CD45RO⁺ and CD8⁺CD45RO⁺, or CD3RO⁺, CD4RO⁺ and CD8RO⁺). Absolute cell counts
170 were calculated from total lymphocyte numbers, which were determined with automated blood
171 cell counters in each participating city. The CD4⁺/CD8⁺ and the CD3⁺/CD19⁺ ratios were also
172 calculated.

173

174 **Statistical analysis**

175 All the analyses conducted on the HELENA-CSS data were adjusted by a weighing factor to
176 balance the studied population according to the age and sex distribution of the theoretical
177 sample. Adolescents were grouped into four age categories: 12.5-13.9 years (from 12.5 years to
178 the day before the 14th birthday), 14-14.9 years (from the 14th birthday to the day before the 15th
179 birthday), 15-15.9 years (from the 15th birthday to the day before the 16th birthday), and 16-17.5
180 years (from the 16th birthday to 17.5 years). The *Chi-squared* (X^2) test was used to compare
181 frequencies of sex, age categories, and Tanner stages between the studied and the excluded
182 groups, as well as to compare frequencies of age categories and Tanner stages between sexes.

183 Absolute counts and percentages of cells are presented as percentiles 10th, 25th, 50th
184 (median), 75th and 90th for description of the population classified by sex and age. Data were
185 then studied according to sex, age, Tanner stage and standardized BMI categories. Normality of
186 variables was checked by the Kolmogorov-Smirnov test, and those not normally distributed were
187 appropriately transformed when necessary.

188 Differences between sexes were analysed with the Mann-Whitney U test. Subsequent
189 tests to assess the influence of age, pubertal maturation and BMI were conducted separately in
190 boys and girls. Associations between cell values and age and between cell values and BMI were

191 assessed with Pearson partial correlation test, controlling for city of origin (centre). Differences
192 in cell values between Tanner stages and BMI categories were assessed by analysis of
193 covariance (ANCOVA), adjusting for centre (and age when comparing Tanner stages). Due to
194 the small number of individuals found within Tanner stages I and II, these two groups were
195 combined to allow statistical analysis.

196 Statistical significance was set at $P < 0.05$. All statistical analyses were performed with
197 IBM® SPSS® Statistics v.19 for Windows.

198

199 **RESULTS**

200 **Characteristics of the population**

201 There were no differences in the proportions of sexes between included and excluded
202 adolescents (48.9 vs. 45.5% boys; $\chi^2 = 1.196$, $P = 0.274$). Similarly, no significant differences
203 were found in the distributions of age categories ($\chi^2 = 1.146$, $P = 0.766$), Tanner stages
204 ($\chi^2 = 3.851$, $P = 0.278$) or BMI categories ($\chi^2 = 7.164$, $P = 0.067$) between the adolescents
205 included in the analysis and their excluded peers.

206 The average age of the studied population was 14.9 ± 1.2 years (range 12.5-17.4 y).
207 There were significantly fewer adolescents between 16 and 17.5 years (22%) than in the other
208 age groups ($\chi^2 = 16.392$, $P < 0.01$). In relation to pubertal maturation, most of the adolescents
209 (78.2%) were found at either the IV or V Tanner stages, as could be expected according to the
210 age range. Boys and girls were similar in age, Tanner stage distributions and mean BMI values
211 in each quartile of BMI z-scores (boys: 17.7 ± 1.1 (Q1), 19.8 ± 0.8 (Q2), 21.8 ± 1.2 (Q3), and
212 27.0 ± 3.2 (Q4) kg/m^2 ; girls: 17.9 ± 1.1 (Q1), 20.0 ± 0.8 (Q2), 22.0 ± 1.1 (Q3), and 26.3 ± 3.0
213 (Q4) kg/m^2).

214

215

216 **Influence of sex on immune cell counts and percentages**

217 Statistical comparison between sexes showed that girls in general presented higher values for
218 total WBC and neutrophil counts (**table 1**). Percentages of neutrophils were also higher in girls,
219 while those of lymphocytes, monocytes and eosinophils were greater in boys (**supplementary**
220 **table 1**).

221 **Table 2** shows the percentiles 10th, 25th, 50th (median), 75th and 90th of the absolute
222 counts of the selected lymphocyte subsets, separately by sex and group of age. Percentiles of the
223 lymphocyte subsets percentages can be found in **supplementary table 2**. Boys showed a
224 tendency to higher counts of CD3⁺CD45RA⁺ and CD4⁺CD45RA⁺ naïve cells (differences
225 significant at 14-14.9 y), while girls presented higher counts of CD3⁺CD45RO⁺ (differences
226 being significant at 14-14.9 and 16-17.5 y) and CD4⁺CD45RO⁺ memory cells (significant at 16-
227 17.5 y). The ratio CD3⁺/CD19⁺ was also higher in girls (difference being significant at 15-15.9
228 years) (**table 2**). Similarly, boys had higher percentages of CD3⁺CD45RA⁺ (significant at 14-
229 14.9 y and 16-17.5 y), CD4⁺CD45RA⁺ and CD8⁺CD45RA⁺ (differences significant at 14-14.9 y);
230 girls in turn had higher percentages of CD3⁺ and CD4⁺ (differences significant at 15-15.9 y),
231 CD3⁺CD45RO⁺ (differences significant at 14-14.9 and 16-17.5 y), CD4⁺CD45RO⁺ and
232 CD8⁺CD45RO⁺ (differences significant at 14-14.9 y) (**supplementary table 2**).

233

234 **Influence of age on immune cell counts and percentages**

235 Total WBC and neutrophil counts increased with age in the total population ($r = 0.121$, $P = 0.015$
236 and $r = 0.153$, $P = 0.003$, respectively; and **table 1**). Neutrophil percentages also increased with
237 age, while the percentage of lymphocytes decreased, both in the total population ($r = 0.167$ and
238 $r = -0.173$, respectively, $P = 0.001$) and in boys (**supplementary table 1**).

239 Age was also associated with increased counts of CD4⁺CD45RO⁺ ($r = 0.179$, $P = 0.001$)
240 and the ratio CD3⁺/CD19⁺ ($r = 0.240$, $P < 0.001$), and decreased counts of CD19⁺ ($r = -0.250$,

241 $P < 0.001$), $CD3^+CD45RA^+$ ($r = -0.118$, $P = 0.036$), and $CD4^+CD45RA^+$ ($r = -0.124$,
242 $P = 0.027$). The percentages of these subsets showed the same relationships with age ($\%CD19^+$:
243 $r = -0.255$; $\%CD3^+CD45RA^+$: $r = -0.184$; $\%CD4^+CD45RA^+$: $r = -0.238$; $\%CD3^+CD45RO^+$:
244 $r = 0.206$; $\%CD4^+CD45RO^+$: $r = 0.235$; all $P < 0.01$). In boys alone, age was inversely
245 correlated with counts of $CD19^+$, $CD8^+$ and naïve cells ($CD3^+CD45RA^+$, $CD4^+CD45RA^+$ and
246 $CD8^+CD45RA^+$), and positively with the ratio $CD3^+/CD19^+$ (**table 2**). In girls alone, older age
247 was associated as well with lower $CD19^+$ and higher $CD3^+/CD19^+$, but also with increased
248 counts of $CD3^+CD45RO^+$ and $CD4^+CD45RO^+$ (**table 2**). As a result, in both boys and girls there
249 was a trend to decreased percentages of naïve cells and increased of memory cells with age
250 (**supplementary table 2**).

251

252 **Influence of pubertal maturation on immune cell counts and percentages**

253 In boys, pubertal maturation (Tanner stage) was associated with lower WBC and lymphocyte
254 counts, independently of age (**table 3**); lymphocyte percentages were also lower in more
255 developed Tanner stages (**supplementary table 3**). This decrease was reflected in most
256 lymphocyte subsets: $CD3^+$, $CD4^+$, $CD8^+$, $CD19^+$, $CD3^+CD45RA^+$, $CD3^+CD45RO^+$,
257 $CD4^+CD45RO^+$, $CD8^+CD45RA^+$ and $CD8^+CD45RO^+$ cell counts were all lower in more
258 advanced Tanner stages (**table 4**).

259 In girls, higher basophil counts (**table 3**) and percentages (**supplementary table 3**), and
260 $CD3^+CD45RO^+$ and $CD4^+CD45RO^+$ cell counts were higher in more advanced pubertal
261 maturation stages (**table 4**).

262

263 **Influence of BMI on immune cell counts and percentages**

264 In boys, counts of total WBC, neutrophils and lymphocytes were higher with increasing BMI
265 (**table 5**). The increase in lymphocyte counts was reflected in most subsets ($CD3^+$, $CD4^+$, $CD8^+$,

266 natural killer, CD19⁺, CD3⁺CD45RO⁺, CD4⁺CD45RO⁺ and CD8⁺CD45RO⁺), whereas the ratio
267 CD4⁺/CD8⁺ (**table 6**) and the percentages of naïve cells (CD3⁺CD45RA⁺ and CD4⁺CD45RA⁺)
268 decreased (**supplementary table 6**).

269 In girls, BMI showed a positive relationship with total WBC and neutrophil counts (**table**
270 **5**), neutrophil percentages (**supplementary table 5**), and memory cell counts (CD3⁺CD45RO⁺
271 and CD4⁺CD45RO⁺) (**table 6**), and a negative one with percentages of CD8⁺CD45RA⁺ cells
272 (**supplementary table 6**).

273

274 **DISCUSSION**

275 The current study describes the immune cell profile of a representative sample of healthy
276 European adolescents. Our results show that sex, age, pubertal maturation and BMI are factors
277 that influence normal circulating counts and percentages of immune cells in adolescents.

278 There were clear sex-related differences in the percentages and absolute counts of
279 immune cells in our population. In general, girls had higher total WBC, neutrophil and memory
280 T cells values (in particular, CD3⁺CD45RO⁺ and CD4⁺CD45RO⁺), while boys had higher
281 percentages of lymphocytes and eosinophils, and higher counts of naïve T cells (all three subsets,
282 CD3⁺CD45RA⁺, CD4⁺CD45RA⁺ and CD8⁺CD45RA⁺). Similar trends for differences in WBC
283 between sexes were observed in a previous study in Spanish adolescents, although the authors
284 did not find statistical significance [27]. With regards to lymphocyte subsets, our observations
285 are also in agreement with previous studies [1, 28, 34]. On the contrary, other authors have found
286 no effect of sex on WBC counts [38] or lymphocyte subsets [32] in adolescents. Sex dimorphism
287 in WBC counts and percentages can be explained by differences in sex hormones, as suggested
288 by Rudy and colleagues [28]. Sex hormones have been shown to modulate immune function at
289 various levels, and a greater immune responsiveness has been observed in girls in general [12,
290 35].

291 A trend was observed towards higher WBC counts with age, in agreement with the
292 findings reported by Bartlett and colleagues [1]. In particular, neutrophil counts were elevated in
293 older boys and girls in our study; this led to increasing percentages of this cell type and
294 decreasing percentages of lymphocytes in boys, while no significant changes in cell percentages
295 with age were observed in girls. In contrast, age was associated with decreasing values for B
296 cells (CD19⁺) leading to a higher CD3⁺/CD19⁺ ratio, and with a significant shift towards lower
297 naïve/memory cells ratios. Lower B cell numbers with age were also observed by Bartlett [1].
298 Likewise, the changes in naïve and memory T cells are in agreement with previous studies [16,
299 28], and coherent with an age-related maturation process of the immune system [32]. Age-sex
300 interactions were observed: boys presented higher counts of naïve cells than girls, but the
301 difference was ameliorated by age; in contrast, the counts of memory cells were higher in girls,
302 and the gap increased with age. As a consequence, the changes in percentages of naïve and
303 memory T cells, which were similar in both boys and girls, were the result of decreased naïve
304 cell counts in boys, and of increased memory cell counts in girls. This difference between boys
305 and girls could suggest a more mature or experienced immune system in the girls. As our
306 adolescents were age-matched, the cause must be related to other parameters of growth rather
307 than age, like pubertal maturation.

308 For that reason, pubertal maturation was considered for the analysis of immune cell
309 profiles in our study. Similarly to age, the effect of pubertal maturation on WBC counts and
310 percentages showed a distinct sex-related pattern. In boys, pubertal maturation was associated
311 with lower WBC counts, lymphocyte counts and percentages; the decrease in lymphocyte counts
312 was reflected in most subsets (CD3⁺, CD4⁺, CD8⁺, CD19⁺, CD3⁺CD45RA⁺, CD3⁺CD45RO⁺,
313 CD8⁺CD45RA⁺ and surprisingly, also CD4⁺CD45RO⁺ and CD8⁺CD45RO⁺ cells). This could be
314 explained by the physiological increase in androgen levels, mainly testosterone, since studies in
315 men have reported negative associations between testosterone levels and WBC counts [3, 30]. In

316 girls, pubertal maturation was associated with increases in basophil counts and percentages and
317 higher memory T cell counts. The lack of other significant relationships between pubertal
318 maturation and immune cell counts could be related to the fact that the degree of pubertal
319 maturation of the female sample in this study was high, with 82% of them being in the IV and V
320 Tanner stages (in boys this proportion was lower, 73.7%).

321 Finally, BMI z-scores were also significantly associated with values of circulating
322 immune cells in the studied population, and again the relationship was sex-specific. Higher BMI
323 was in general associated with higher counts of total WBC and memory T cells (all three subsets
324 in boys and $CD3^+CD45RO^+$ and $CD4^+CD45RO^+$ in girls). In boys, both neutrophils and
325 lymphocytes were elevated, but the increase in lymphocytes was proportionally greater. In this
326 line, in addition to memory cells, higher BMI z-scores in boys were also associated to elevated
327 cell counts in most subsets ($CD3^+$, $CD4^+$, $CD8^+$, NK, $CD19^+$, and $CD8^+CD45RA^+$) and with a
328 lower $CD4^+/CD8^+$ ratio. In girls, BMI was mainly linked to counts of neutrophils and memory
329 cells. Alterations in immune parameters have been associated with both insufficient and
330 excessive body weight [reviewed in 19 and 26]. In adults, positive associations were reported
331 between BMI and circulating WBC, neutrophils, lymphocytes and monocytes [15]. Later, other
332 authors have found similar age-independent, positive associations between WBC counts and
333 BMI in children and adolescents [14, 36], in line with our observations. Furthermore, obesity in
334 children and adolescents has been linked to elevated counts of neutrophils, monocytes, total T
335 cells and helper T cells [37]. Our results, like those by Hsieh [14] and Wu [36], show that this
336 increase is not merely the result of obesity, but it takes place in a linear manner with increasing
337 BMI. It is worth highlighting this correlation, as WBC count has been related to features of the
338 metabolic syndrome in both adult [10, 22] and paediatric populations [5, 14, 36], and a follow-up
339 study in Japanese adults concluded that higher WBC counts were associated with higher risk of
340 developing metabolic syndrome in the future [24]. The explanation for the relationship between

341 BMI and memory T cells or its potential implications is less straightforward. On the one hand, it
342 could suggest that increasing BMI provides a more favorable environment for immune system
343 development, similarly to age and pubertal maturation. On the other hand, we could instead be
344 facing similar outcomes with different origins, and BMI-related changes could indicate abnormal
345 or excessive activation of the immune system. This hypothesis would be supported by the
346 relationships between circulating WBC and the metabolic syndrome mentioned above. However,
347 the underlying processes linking nutritional status, BMI and immune function in children and
348 adolescents clearly requires further research.

349 In clinical practice to date, sex and age have been routinely taken into account for the
350 establishment of normative ranges, since they constitute easy-to-obtain information. Other
351 physiological features like pubertal maturation or BMI, however, are not frequently available or
352 assessed when analysing blood variables. In light of ours and other authors' results, we
353 recommend considering these characteristics when setting reference values for or performing
354 blood measurements in paediatric populations.

355 Finally, two considerations should be made in relation to the present study. On the one
356 hand, the number of subjects suitable for statistical analysis was not balanced in relation to the
357 actual population size of each city. On the other, the sample studied included adolescents from
358 different ethnic origins, and as it was mentioned above, ethnicity can be an influential factor for
359 differences in WBC counts [1, 17, 38]. Despite these caveats, our work contributes to the
360 development of a database of haematological reference values in healthy European adolescents.
361 This is particularly strengthened by the use of standardized protocols and methods across all
362 centres participating in the study.

363 In conclusion, the present work provides data on normal values for white blood cell
364 counts and percentages from healthy European adolescents, and highlights the importance of
365 taking into account the influence of sex, age, the degree of pubertal maturation and BMI when

366 comparing or using white blood cell counts for clinical and research purposes in paediatric
367 populations.

368

369 **Acknowledgements**

370 Authors wished to dedicate this work to the memory of Dr. Javier Romeo, and to acknowledge
371 his personal and professional contribution to our group.

372 The present study was performed as part of the HELENA study (www.helenastudy.com).

373 Authors gratefully acknowledge the participation of all HELENA Study members (listed in the
374 Appendix), as well as the financial support of the European Community 6th RTD Framework
375 Programme (contract FOOD-CT-2005-007034). F.P.d.H. acknowledges funding from a JAE-
376 Doc contract cofounded by the Spanish National Research Council (CSIC) and the European
377 Social Fund. K.V. acknowledges funding from the Research Foundation Flanders (Brussels,
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382 final content. All authors read and approved the final manuscript.

383

384 **Conflict of interest**

385 None of the authors had a personal or financial conflict of interest.

386

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Table 1. White blood cell (WBC) counts (cell/ μ l) in European adolescents, according to age categories and stratified for sex.

Age range (years)	Boys					Girls					
	12.5-13.9	14-14.9	15-15.9	16-17.5	R	12.5-13.9	14-14.9	15-15.9	16-17.5	R	
N	48	55	46	49		56	48	63	40		
WBC	10 th	4,106	4,382	3,787 ^{§§}	4,630	0.107	4,493	4,368	4,934^{§§}	5,230	0.154*
	25 th	4,854	5,011	5,214 ^{§§}	5,055		5,151	5,202	5,670^{§§}	5,682	
	50 th	5,721	5,706	6,100 ^{§§}	6,105		6,010	6,404	6,741^{§§}	6,520	
	75 th	6,572	6,579	6,804 ^{§§}	6,810		6,929	7,823	8,011^{§§}	7,702	
	90 th	7,605	8,000	7,568 ^{§§}	7,490		8,457	9,573	8,753^{§§}	8,350	
Neutrophils	10 th	1,786[§]	1,906^{§§}	1,787[§]	2,197^{§§}	0.184*	2,310[§]	1,870^{§§}	2,138[§]	2,700^{§§}	0.162*
	25 th	2,117[§]	2,328^{§§}	2,371[§]	2,576^{§§}		2,834[§]	2,587^{§§}	2,860[§]	3,153^{§§}	
	50 th	2,940[§]	2,681^{§§}	3,310[§]	3,190^{§§}		3,227[§]	3,555^{§§}	3,990[§]	3,835^{§§}	
	75 th	3,448[§]	3,641^{§§}	3,984[§]	3,635^{§§}		3,898[§]	4,873^{§§}	4,900[§]	4,370^{§§}	
	90 th	4,771[§]	4,458^{§§}	4,737[§]	4,700^{§§}		5,158[§]	6,357^{§§}	5,591[§]	4,930^{§§}	
Lymphocytes	10 th	1,688	1,539	1,450	1,460	-0.130	1,483	1,418	1,570	1,700	0.054
	25 th	1,895	1,757	1,700	1,918		1,800	1,691	1,870	1,846	
	50 th	2,200	2,241	2,040	2,120		2,100	2,141	2,140	1,960	
	75 th	2,411	2,561	2,334	2,283		2,408	2,535	2,500	2,488	
	90 th	3,106	2,887	2,768	2,960		2,663	3,088	3,003	2,940	
Monocytes	10 th	295	300	330	300	0.074	324	280	350	280	0.092
	25 th	360	370	376	399		395	340	400	376	
	50 th	420	461	445	460		440	430	500	510	
	75 th	522	600	543	510		514	551	570	578	
	90 th	700	769	648	620		700	743	714	610	
Eosinophils	10 th	80	80	40	60	-0.056	57	49	64	60	-0.074
	25 th	100	100	100	100		100	80	100	70	
	50 th	119	160	140	135		137	130	125	110	
	75 th	200	224	232	191		230	200	200	186	
	90 th	300	400	455	310		424	255	334	280	
Basophils	10 th	0	0	0	0	0.029	0	1	0	0	0.090
	25 th	0	10	8	20		0	10	10	20	
	50 th	20	20	20	30		20	30	30	20	
	75 th	40	45	30	31		33	40	50	40	
	90 th	79	92	64	60		58	76	100	50	

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Data are presented as percentiles 10th, 25th, 50th (median), 75th, and 90th. [§]Significant differences between boys and girls for a given age category, as assessed by the Mann-Whitney U test, [§] $P < 0.05$, ^{§§} $P < 0.01$. R is the partial correlation coefficient between cell counts and age, controlling for centre; bold rows indicate significant correlations, * $P < 0.05$, ** $P < 0.01$.

Table 2. Estimated cell counts (cell/ μ l) of selected lymphocyte subsets in adolescents, according to age categories and stratified for sex.

		Boys					Girls				
Age range (years)		12.5-13.9	14-14.9	15-15.9	16-17.5		12.5-13.9	14-14.9	15-15.9	16-17.5	
N		36	49	41	45	R	39	42	50	26	R
CD3 ⁺	10 th	1,147	1,005	952	943	-0.149	1,041	922	1,117	1,151	0.081
	25 th	1,239	1,208	1,139	1,145		1,278	1,138	1,291	1,275	
	50 th	1,477	1,524	1,355	1,464		1,384	1,498	1,536	1,374	
	75 th	1,758	1,737	1,695	1,598		1,573	1,772	1,951	1,819	
	90 th	2,330	1,974	2,002	2,127		1,889	2,124	2,180	1,930	
CD4 ⁺	10 th	580	535	491	487	-0.139	573	459	616	567	0.109
	25 th	652	660	628	581		651	612	706	687	
	50 th	806	824	771	791		779	789	842	785	
	75 th	1,055	962	902	972		888	950	1,054	1,095	
	90 th	1,154	1,150	1,044	1,070		1,057	1,181	1,292	1,212	
CD8 ⁺	10 th	372	349	343	309	-0.154*	393	333	392	383	0.009
	25 th	490	443	441	358		442	425	463	472	
	50 th	541	601	531	539		581	558	584	515	
	75 th	770	724	704	683		672	735	725	633	
	90 th	1,016	915	850	888		757	970	983	725	
CD3-CD16 ⁺ 56 ⁺	10 th	164	119	145	165	-0.030	148	172	168	138	0.022
	25 th	248	201	222	217		191	242	238	180	
	50 th	333	298	350	376		275	312	312	322	
	75 th	410	417	447	447		434	418	388	400	
	90 th	754	647	562	542		579	531	507	784	
CD3-CD19 ⁺	10 th	160	157	153	131	-0.263**	181	140	150	99	-0.266**
	25 th	221	198	195	174		231	196	174	127	
	50 th	286	283	251	219		267	244	238	241	
	75 th	333	377	311	279		385	304	299	285	
	90 th	514	514	439	375		501	409	354	321	
CD3 ⁺ CD45RA ⁺	10 th	637	643[§]	485	461	-0.199*	570	510 [§]	609	535	-0.016
	25 th	770	735[§]	711	612		718	660 [§]	678	673	
	50 th	958	1,000[§]	809	860		870	815 [§]	887	824	
	75 th	1,113	1,138[§]	1,016	1,001		1,024	1,015 [§]	1,149	911	
	90 th	1,448	1,218[§]	1,213	1,195		1,236	1,386 [§]	1,408	1,193	
CD3 ⁺ CD45RO ⁺	10 th	346	332 [§]	317	385 [§]	-0.016	347	322[§]	437	517[§]	0.237**
	25 th	451	437 [§]	392	463 [§]		440	475[§]	530	572[§]	
	50 th	526	482 [§]	564	575 [§]		520	630[§]	625	660[§]	
	75 th	735	674 [§]	705	672 [§]		619	778[§]	749	791[§]	

CD4+CD45RA+	90 th	867	807 [§]	847	832 [§]	-0.196*	818	897[§]	928	942[§]	-0.042
	10 th	306	311[§]	248	201		306	232 [§]	287	197	
	25 th	382	381[§]	322	304		364	288 [§]	343	347	
	50 th	497	524[§]	416	418		479	422 [§]	448	451	
	75 th	626	629[§]	482	596		570	590 [§]	643	512	
CD4+CD45RO+	90 th	752	709[§]	640	708	0.006	716	694 [§]	744	590	0.344**
	10 th	227	194	194	256 [§]		205	187	279	261[§]	
	25 th	245	251	234	292 [§]		252	269	313	338[§]	
	50 th	313	304	338	337 [§]		291	358	371	389[§]	
	75 th	414	430	442	406 [§]		343	428	483	566[§]	
CD8+CD45RA+	90 th	487	556	508	466 [§]	-0.186*	401	522	540	644[§]	0.021
	10 th	268	249	221	206		200	203	255	230	
	25 th	311	298	285	231		282	271	299	301	
	50 th	384	414	362	367		380	342	391	334	
	75 th	489	523	505	446		471	450	543	441	
CD8+CD45RO+	90 th	645	663	619	603	-0.036	574	558	678	497	0.032
	10 th	90	65	79	78		84	85	92	94	
	25 th	107	98	102	110		118	130	128	139	
	50 th	164	159	152	155		146	182	180	173	
	75 th	224	225	199	224		217	290	231	213	
CD4+/CD8+	90 th	264	290	249	328	0.051	313	354	300	240	0.104
	10 th	1.04	0.85	0.98	0.87		1.03	0.89	1.00	1.14	
	25 th	1.16	1.21	1.16	1.13		1.22	1.11	1.19	1.24	
	50 th	1.29	1.49	1.36	1.62		1.44	1.46	1.55	1.56	
	75 th	1.58	1.86	1.78	1.97		1.64	1.64	1.78	1.87	
CD3+/CD19+	90 th	2.18	2.23	2.22	2.36	0.189*	1.98	2.02	2.11	2.86	0.311**
	10 th	3.08	3.10	3.44[§]	3.90		3.04	3.62	3.99[§]	4.74	
	25 th	4.32	3.75	4.02[§]	5.24		3.87	4.06	5.27[§]	6.10	
	50 th	5.95	5.25	5.82[§]	6.11		4.75	5.69	6.61[§]	6.89	
	75 th	7.10	7.14	6.90[§]	7.74		6.68	7.14	8.31[§]	9.93	
90 th	8.71	9.24	8.58[§]	10.76	7.87	11.00	12.04[§]	13.79			

599 Data are presented as percentiles 10th, 25th, 50th (median), 75th, and 90th. Lymphocyte populations are designated by their cell markers. [§]Significant differences
600 between boys and girls for a given age category, as assessed by the Mann-Whitney U test; [§]*P*<0.05, ^{§§}*P*<0.01. R is the partial correlation coefficient between
601 cell counts and age, controlling for centre; bold rows indicate significant correlations, **P*<0.05, ***P*<0.01.
602

603 **Table 3.** White blood cell (WBC) counts (cell/ μ l) in European adolescents, according to Tanner stages and stratified for sex.

Tanner stage N	Boys					Girls					
		I+II 12	III 34	IV 77	V 55	P	I+II 7	III 26	IV 97	V 60	P
WBC	10 th	4,073	4,803	4,160	4,357	0.037	3,920	4,353	4,696	4,995	0.404
	25 th	4,751	5,469	4,940	4,946		4,423	5,030	5,481	5,357	
	50 th	6,050	5,965	5,706	5,938		5,091	6,065	6,541	6,500	
	75 th	7,271	6,769	6,581	6,811		6,878	8,179	7,612	8,003	
	90 th	9,129	8,273	7,454	7,300		-	9,581	8,693	8,400	
Neutrophils	10 th	1,243	2,129	1,766	2,021	0.068	1,290	1,908	2,153	2,444	0.089
	25 th	2,158	2,833	2,169	2,447		1,543	2,361	2,864	2,975	
	50 th	3,095	3,206	2,891	3,210		2,259	3,515	3,599	3,662	
	75 th	4,057	3,636	3,504	4,000		3,601	5,056	4,486	4,699	
	90 th	4,794	5,097	4,197	4,710		-	7,111	5,335	5,476	
Lymphocytes	10 th	1,803	1,650	1,537	1,298	0.002	1,550	1,385	1,566	1,500	0.981
	25 th	1,898	1,880	1,903	1,575		1,917	1,658	1,850	1,828	
	50 th	2,247	2,138	2,155	1,935		2,122	2,097	2,120	2,140	
	75 th	2,984	2,567	2,397	2,310		2,339	2,468	2,510	2,500	
	90 th	3,845	3,275	2,870	2,728		-	2,790	2,991	2,893	
Monocytes	10 th	350	299	290	316	0.825	340	279	331	291	0.873
	25 th	367	360	350	382		346	362	400	370	
	50 th	420	408	420	469		392	452	494	439	
	75 th	540	540	521	510		702	570	570	526	
	90 th	661	720	618	632		-	735	671	627	
Eosinophils	10 th	85	54	50	69	0.922	51	61	60	74	0.206
	25 th	111	100	100	100		98	94	80	100	
	50 th	164	150	130	130		130	158	110	140	
	75 th	219	220	223	203		165	263	200	200	
	90 th	428	426	311	358		-	513	280	354	
Basophils	10 th	5	0	4	0	0.061	10	1	0	0	0.011
	25 th	22	16	10	9		15	10	10	20	
	50 th	38	20	20	20		30	29	20	30	
	75 th	41	40	30	40		40	32	40	60	
	90 th	60	74	46	60		-	72	50	100	

604 Data are presented as percentiles 10th, 25th, 50th (median), 75th, and 90th. Bold rows indicate significant differences between Tanner stages, as
605 assessed by analysis of covariance (ANCOVA), controlling for centre and age, $P < 0.05$.

606 **Table 4.** Estimated cell counts (cell/ μ l) of selected lymphocyte subsets in European adolescents, according to Tanner stages and stratified for sex.

		Boys					Girls				
Tanner stage		I+II	III	IV	V	<i>P</i>	I+II	III	IV	V	<i>P</i>
N		9	26	67	52		7	19	65	55	
CD3 ⁺	10 th	1,157	1,039	1,066	876	0.002	1,135	819	1,112	1,089	0.362
	25 th	1,259	1,401	1,237	1,007		1,295	990	1,236	1,202	
	50 th	1,694	1,517	1,467	1,316		1,366	1,364	1,470	1,549	
	75 th	2,095	1,903	1,672	1,526		1,466	1,575	1,835	1,858	
	90 th	-	2,254	1,991	1,909		-	1,938	2,205	2,051	
CD4 ⁺	10 th	584	542	551	479	0.010	632	477	585	572	0.308
	25 th	644	735	681	575		727	593	684	652	
	50 th	780	894	835	737		815	689	827	834	
	75 th	1,084	960	994	882		881	862	1,080	1,015	
	90 th	1,457	1,168	1,143	1,014		-	1,069	1,252	1,223	
CD8 ⁺	10 th	492	383	347	279	0.012	330	291	382	392	0.325
	25 th	510	454	430	358		412	388	447	455	
	50 th	671	602	554	464		528	478	559	590	
	75 th	822	843	692	653		606	632	676	733	
	90 th	1,026	1,038	780	846		-	815	883	938	
CD3-CD16+56 ⁺	10 th	247	161	136	149	0.067	119	127	168	170	0.536
	25 th	276	249	210	210		185	183	241	221	
	50 th	319	357	342	323		326	304	339	286	
	75 th	506	430	446	455		496	360	440	375	
	90 th	-	687	540	590		-	540	554	571	
CD3-CD19 ⁺	10 th	162	182	159	116	0.049	144	152	137	128	0.078
	25 th	210	209	200	172		169	185	217	171	
	50 th	282	276	256	225		342	240	273	232	
	75 th	337	325	359	287		418	293	321	289	
	90 th	-	586	460	398		-	466	417	355	
CD3+CD45RA ⁺	10 th	737	649	591	476	0.024	745	515	528	533	0.698
	25 th	814	754	732	582		832	631	666	674	
	50 th	1,005	980	939	777		901	727	846	901	
	75 th	1,405	1,189	1,094	975		1,016	1,039	1,056	1,076	
	90 th	-	1,294	1,193	1,129		-	1,329	1,427	1,297	
CD3+CD45RO ⁺	10 th	354	394	356	297	<0.001	304	284	422	465	0.021
	25 th	477	486	459	373		357	354	474	528	
	50 th	549	592	559	467		454	479	634	636	
	75 th	722	775	682	585		536	684	784	753	

CD4+CD45RA+	90 th	-	907	777	776	0.102	-	813	918	942	0.824
	10 th	318	331	227	238		388	263	245	235	
	25 th	408	355	370	296		472	332	363	332	
	50 th	487	449	491	384		538	390	475	426	
	75 th	639	667	629	506		565	608	598	580	
CD4+CD45RO+	90 th	783	769	701	646	-	720	740	729	0.025	
	10 th	213	198	227	191	186	182	241	270		
	25 th	235	293	280	233	239	203	289	314		
	50 th	303	400	327	303	281	283	363	387		
	75 th	352	457	419	361	306	312	445	493		
CD8+CD45RA+	90 th	851	487	505	485	-	398	558	619	0.473	
	10 th	336	282	222	219	193	201	228	257		
	25 th	367	316	297	245	247	256	301	288		
	50 th	458	413	383	325	407	325	363	412		
	75 th	622	540	451	446	509	440	469	499		
CD8+CD45RO+	90 th	-	684	593	624	-	560	631	649	0.069	
	10 th	107	88	79	49	61	60	105	94		
	25 th	120	117	109	89	82	103	128	138		
	50 th	159	160	164	130	109	150	171	186		
	75 th	211	242	199	197	166	236	229	268		
CD4+/CD8+	90 th	-	366	261	246	-	320	303	332	0.478	
	10 th	1.09	0.98	0.99	0.92	1.04	0.90	1.16	0.94		
	25 th	1.13	1.12	1.22	1.14	1.26	1.21	1.27	1.14		
	50 th	1.20	1.28	1.55	1.43	1.78	1.52	1.50	1.48		
	75 th	1.39	1.72	1.93	1.91	2.14	1.79	1.76	1.77		
CD3+/CD19+	90 th	1.48	2.07	2.15	2.47	-	2.03	2.11	1.99	0.417	
	10 th	3.09	3.40	3.28	3.54	2.99	3.71	3.64	3.86		
	25 th	5.34	4.81	4.37	4.13	3.22	4.04	4.73	4.97		
	50 th	6.38	5.38	5.79	5.79	4.47	5.12	5.56	6.77		
	75 th	7.28	7.28	6.97	7.68	7.09	7.35	7.32	9.68		
90 th	-	8.08	8.11	10.96	-	10.56	10.54	11.78			

607 Data are presented as percentiles 10th, 25th, 50th (median), 75th, and 90th. Lymphocyte populations are designated by their cell markers. Bold rows indicate significant
608 differences between Tanner stages, as assessed by analysis of covariance (ANCOVA), controlling for centre and age, $P < 0.05$.
609

610 **Table 5.** White blood cell (WBC) counts (cell/ μ l) in European adolescents, according to BMI z-scores and stratified for sex.

		Boys						Girls					
BMI z-scores		Q1	Q2	Q3	Q4			Q1	Q2	Q3	Q4		
N		43	42	42	43	<i>P</i>	<i>R</i>	41	41	38	39	<i>P</i>	<i>R</i>
WBC	10 th	4,629	4,180	4,160	5,175	0.001	0.195**	4,392	4,484	5,208	5,221	0.005	0.208**
	25 th	5,223	4,700	4,909	5,775			5,054	5,101	5,716	5,772		
	50 th	6,079	5,463	5,711	6,403			6,260	6,300	6,804	6,574		
	75 th	6,512	6,504	6,781	7,200			7,161	7,751	7,938	8,321		
	90 th	7,709	7,150	7,487	8,825			8,221	8,480	9,068	9,836		
Neutrophils	10 th	2,032	1,769	1,893	2,211	0.020	0.115	1,799	2,225	2,529	2,644	0.003	0.224**
	25 th	2,381	2,275	2,111	2,800			2,693	2,636	3,000	3,170		
	50 th	3,084	2,799	2,910	3,269			3,190	3,512	3,806	4,146		
	75 th	3,791	3,539	3,537	4,009			4,200	4,384	4,902	5,070		
	90 th	4,658	4,387	4,308	5,042			4,666	5,646	6,021	6,287		
Lymphocytes	10 th	1,373	1,456	1,551	1,879	<0.001	0.331**	1,590	1,497	1,488	1,367	0.806	0.029
	25 th	1,703	1,671	1,729	2,170			1,829	1,694	1,856	1,870		
	50 th	2,060	1,992	2,089	2,380			2,093	2,035	2,195	2,122		
	75 th	2,378	2,190	2,331	2,817			2,482	2,456	2,439	2,552		
	90 th	2,676	2,553	2,862	3,157			2,834	2,940	2,802	3,092		
Monocytes	10 th	300	300	319	300	0.261	0.115	294	284	349	300	0.224	0.071
	25 th	363	388	370	380			390	363	413	370		
	50 th	420	420	460	500			476	432	500	463		
	75 th	511	500	583	580			559	553	555	600		
	90 th	672	613	660	709			665	654	707	733		
Eosinophils	10 th	80	90	50	50	0.066	-0.095	70	60	50	50	0.742	0.034
	25 th	100	100	100	100			90	95	81	100		
	50 th	139	130	190	120			120	105	143	100		
	75 th	209	190	277	180			200	200	220	200		
	90 th	307	320	516	246			316	272	300	398		
Basophils	10 th	0	0	0	0	0.506	-0.126	0	0	0	0	0.527	-0.045
	25 th	10	10	10	10			11	10	10	0		
	50 th	30	20	20	20			30	21	21	20		
	75 th	50	30	40	30			47	35	50	40		
	90 th	70	45	60	60			100	50	80	100		

611 Data are presented as percentiles 10th, 25th, 50th (median), 75th, and 90th. Bold rows indicate significant differences between quartiles of standardized body mass index values
612 (BMI z-scores), as assessed by analysis of covariance (ANCOVA), controlling for centre, *P*<0.05. *R* is the partial correlation coefficient between cell counts and BMI z-scores,
613 controlling for centre, **P*<0.05, ***P*<0.01.

Table 6. Estimated cell counts (cell/ μ l) of lymphocyte subsets in European adolescents, according to BMI z-scores and stratified for sex.

		Boys					Girls						
BMI z-scores		Q1	Q2	Q3	Q4			Q1	Q2	Q3	Q4		
N		43	42	42	43	<i>P</i>	<i>R</i>	41	41	38	39	<i>P</i>	<i>R</i>
CD3 ⁺	10 th	850	994	958	1,244	0.014	0.238**	1,084	1,017	946	1,126	0.301	0.122
	25 th	1,083	1,088	1,162	1,411			1,158	1,183	1,269	1,314		
	50 th	1,457	1,238	1,476	1,696			1,382	1,364	1,523	1,517		
	75 th	1,657	1,590	1,622	1,978			1,578	1,756	1,741	1,881		
	90 th	2,038	1,740	1,979	2,269			1,986	2,165	1,930	2,301		
CD4 ⁺	10 th	443	529	551	591	0.085	0.174*	579	522	572	621	0.265	0.164*
	25 th	632	600	617	749			620	649	671	723		
	50 th	808	748	788	942			711	783	871	845		
	75 th	957	858	895	1,086			903	960	1,020	1,075		
	90 th	1,139	1,053	982	1,178			1,248	1,224	1,145	1,237		
CD8 ⁺	10 th	280	300	346	425	0.009	0.266**	385	359	381	373	0.634	0.067
	25 th	386	380	413	537			413	455	434	463		
	50 th	507	492	554	659			526	552	557	578		
	75 th	649	643	684	864			700	681	711	713		
	90 th	852	758	809	1,013			863	787	782	1,046		
CD3 ⁺ CD16 ⁺ 56 ⁺	10 th	149	149	137	192	0.042	0.192*	119	150	154	181	0.766	0.064
	25 th	222	204	198	303			191	216	199	240		
	50 th	307	323	282	389			320	275	307	320		
	75 th	374	422	437	558			464	361	400	403		
	90 th	463	491	579	704			565	496	531	570		
CD3 ⁺ CD19 ⁺	10 th	131	153	152	186	0.013	0.221**	145	105	153	132	0.700	-0.001
	25 th	185	169	202	224			185	167	202	188		
	50 th	239	219	286	287			257	239	266	241		
	75 th	291	281	352	433			316	288	327	310		
	90 th	417	336	482	502			430	431	397	422		
CD3 ⁺ CD45RA ⁺	10 th	469	505	513	642	0.413	0.133	577	513	522	556	0.962	0.036
	25 th	699	649	700	789			664	633	717	695		
	50 th	928	808	905	1,015			805	880	878	857		
	75 th	1,113	976	1,040	1,215			1,003	1,097	1,031	1,073		
	90 th	1,192	1,112	1,193	1,365			1,282	1,455	1,267	1,429		
CD3 ⁺ CD45RO ⁺	10 th	292	356	339	479	0.003	0.266**	367	426	393	456	0.036	0.171*
	25 th	375	401	441	572			454	486	456	556		
	50 th	475	499	514	690			556	569	576	656		
	75 th	636	589	649	780			696	731	745	775		

CD4 ⁺ CD45RA ⁺	90 th	819	654	874	914	0.932	0.045	880	903	861	940	0.917	0.034
	10 th	216	263	259	234			223	247	244	266		
	25 th	357	314	328	368			336	330	362	348		
	50 th	483	391	453	475			428	449	488	424		
	75 th	613	501	543	637			567	596	577	598		
CD4 ⁺ CD45RO ⁺	90 th	718	705	667	719	0.002	0.261**	655	828	709	690	0.012	0.254**
	10 th	191	204	196	278			225	225	219	261		
	25 th	227	256	254	308			274	286	289	306		
	50 th	291	302	328	408			313	362	341	398		
	75 th	410	373	399	487			387	406	438	493		
CD8 ⁺ CD45RA ⁺	90 th	487	436	466	556	0.131	0.191*	533	485	612	620	0.854	0.033
	10 th	220	227	222	291			248	199	230	199		
	25 th	246	252	287	348			279	292	298	297		
	50 th	382	343	370	431			341	373	380	373		
	75 th	471	446	450	609			474	462	486	472		
CD8 ⁺ CD45RO ⁺	90 th	549	489	638	713	0.019	0.248**	570	568	635	697	0.060	0.095
	10 th	51	74	89	110			84	113	88	92		
	25 th	85	97	107	150			127	135	110	155		
	50 th	144	133	150	206			158	172	155	209		
	75 th	190	196	197	236			229	233	226	266		
CD4 ⁺ /CD8 ⁺	90 th	242	244	312	300	0.286	-0.161*	290	334	290	336	0.713	0.100
	10 th	0.91	1.07	1.08	0.91			0.89	1.03	1.15	0.99		
	25 th	1.16	1.20	1.21	1.05			1.09	1.23	1.24	1.20		
	50 th	1.54	1.41	1.36	1.30			1.43	1.43	1.55	1.51		
	75 th	2.03	1.91	1.70	1.71			1.84	1.72	1.75	1.83		
CD3 ⁺ /CD19 ⁺	90 th	2.48	2.36	2.04	2.19	0.250	-0.057	2.05	2.10	1.95	2.15	0.336	0.094
	10 th	3.58	3.74	3.12	3.44			3.74	3.71	3.71	3.70		
	25 th	4.02	5.34	3.74	4.56			4.03	4.60	4.75	5.44		
	50 th	5.89	6.07	4.73	5.46			5.33	6.31	5.57	6.77		
	75 th	7.85	7.12	6.74	7.11			7.87	8.54	6.68	7.63		
90 th	9.90	7.67	10.24	8.61	10.90	13.48	9.29	11.50					

615 Data are presented as percentiles 10th, 25th, 50th (median), 75th, and 90th. Lymphocyte populations are designated by their cell membrane markers. Bold rows indicate
616 significant differences between quartiles of standardized body mass index (BMI z-scores), as assessed by analysis of covariance (ANCOVA), controlling for centre,
617 $P < 0.05$. R is the partial correlation coefficient between cell counts and BMI z-scores, controlling for centre; * $P < 0.05$, ** $P < 0.01$.
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