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The association between macrovascular complications and intensive care admission, invasive mechanical ventilation, and mortality in people with diabetes hospitalized for coronavirus disease-2019 (COVID-19)

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1 **Title: The association between macrovascular complications and intensive care**
2 **admission, invasive mechanical ventilation, and mortality in people with diabetes**
3 **hospitalized for coronavirus disease-2019 (COVID-19)**

4
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66 **Abstract**

67 **Background:** It is not clear whether pre-existing macrovascular complications (ischemic heart
68 disease, stroke or peripheral artery disease) are associated with health outcomes in people with
69 diabetes mellitus hospitalized for COVID-19.

70 **Methods:** We conducted cohort studies of adults with pre-existing diabetes hospitalized for
71 COVID-19 infection in the UK, France, and Spain during the early phase of the pandemic
72 (between March 2020 - October 2020). Logistic regression models adjusted for demographic
73 factors and other comorbidities were used to determine associations between previous
74 macrovascular disease and relevant clinical outcomes: mortality, intensive care unit (ICU)
75 admission and use of invasive mechanical ventilation (IMV) during the hospitalization. Output
76 from individual logistic regression models for each cohort was combined in a meta-analysis.

77 **Results:** Complete data were available for 4,106 (60.4%) individuals. Of these, 1,652 (40.2%)
78 had any prior macrovascular disease of whom 28.5% of patients died. Mortality was higher for
79 people with compared to those without previous macrovascular disease (37.7% vs 22.4%). The
80 combined crude odds ratio (OR) for previous macrovascular disease and mortality for all four
81 cohorts was 2.12 (95%CI 1.83-2.45 with an I^2 of 60%, reduced after adjustments for age, sex,
82 type of diabetes, hypertension, microvascular disease, ethnicity, and BMI to adjusted OR 1.53
83 [95%CI 1.29-1.81]) for the three cohorts. Further analysis revealed that ischemic heart disease
84 and cerebrovascular disease were the main contributors of adverse outcomes. However,
85 proportions of people admitted to ICU (adjOR 0.48 [95%CI 0.31-0.75], I^2 60%) and the use of
86 IMV during hospitalization (adjOR 0.52 [95%CI 0.40-0.68], I^2 37%) were significantly lower for
87 people with previous macrovascular disease.

88 **Conclusions:** This large multinational study of people with diabetes mellitus hospitalized for
89 COVID-19 demonstrates that previous macrovascular disease is associated with higher
90 mortality and lower proportions admitted to ICU and treated with IMV during hospitalization
91 suggesting selective admission criteria. Our findings highlight the importance correctly assess
92 the prognosis and intensive monitoring in this high-risk group of patients and emphasize the
93 need to design specific public health programs aimed to prevent SARS-CoV-2 infection in this
94 subgroup.

95

96

97 **Keywords:** diabetes, macrovascular disease, mortality, COVID-19

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99

100 **1. Background**

101

102 Coronavirus disease 2019 (COVID-19) is caused by the severe acute respiratory syndrome
103 coronavirus 2 (SARS-CoV-2) and is characterized by a variable clinical presentation that ranges
104 from asymptomatic infection to fatal multi-organ damage and mortality [1,2]. Since the
105 emergence of SARS-CoV-2 in December 2019, cases of COVID-19 have rapidly increased
106 worldwide. The updated WHO estimates on August 19th, 2022, reported 590,659,276 confirmed
107 cases, including 6,440,163 deaths worldwide (<https://covid19.who.int>). The case fatality for
108 COVID-19 has been estimated to be 0.5-1.0% [3,4]. Nevertheless, certain characteristics,
109 including increasing age, male sex, ethnicity, socio-economic deprivation, and comorbidities,
110 have been associated with a higher risk of severe COVID-19 or death [5–7].

111

112 COVID-19 pandemic has had a large negative impact on both diabetes management [8] and
113 diabetes-related mortality [9]. As well, pre-existing diabetes mellitus has been considered a risk
114 factor for increased COVID-19 severity and worse outcomes, including higher mortality,
115 irrespective of age and comorbidity status [7]. The estimates of diabetes prevalence in those
116 who have died of COVID-19 range from 20% to 30% [10,11]. A recent meta-analysis showed
117 that people with diabetes were at higher risk of COVID-19-related mortality in comparison to
118 people without diabetes [11]. In addition, diabetes is associated with more than double the risk
119 for ICU admission and a more than triple the risk of death compared to people without diabetes
120 [12]. Therefore, identifying which clinical factors are associated with greater morbidity and
121 mortality would be useful for prevention and management of high-risk groups during future
122 waves of the pandemic. In that sense, few studies have examined the possibility that micro- and
123 macrovascular complications contribute to susceptibility to acute organ injury [13,14] but with
124 contradictory results [15].

125

126 Our study aimed to assess whether the presence of macrovascular complications (ischemic
127 heart disease, stroke, or peripheral artery disease) prior to hospital admission is associated with
128 intensive care unit admission, mechanical ventilation, and mortality in people with diabetes
129 mellitus hospitalized for COVID-19 in four European cohorts.

130

2.Methods

131 2.1 Study design and participants

132 Retrospective data from hospitalized adults with pre-existing diabetes and concomitant COVID-
133 19 infection were collected in the UK, France, and Spain. Adults with hyperglycaemia at
134 admission but not pre-existing or subsequent diagnosis of diabetes (based on WHO criteria)
135 were excluded from the analysis [16]. COVID-19 was defined as a SARS-CoV-2 infection
136 confirmed by quantitative PCR (qPCR) performed on nasopharyngeal samples obtained by
137 trained personnel and/or by fulfilling clinical and radiological diagnostic criteria at hospital
138 admission. Further descriptions of each dataset have been published previously [17].

139

140 2.2 United Kingdom: association of British clinical diabetologists (ABCD) COVID-19 audit

141 The NHS supports audits with clear guidance for the contributing centers on using routine
142 clinical practice data submitted anonymously via the secure NHS network [18]. Clinicians
143 participating in the ABCD COVID-19 audit submitted data for adults with pre-existing type 1 and
144 type 2 diabetes admitted with COVID-19 from hospitals across the UK. The audit is registered
145 with Oxford University Hospitals NHS Foundation Trust (OUH), a Data Protection Impact
146 Assessment was carried out and the audit was approved by the OUH Caldicott Guardian and
147 the Public Benefit and Privacy Panel in Scotland (reference 2021-0111).

148

149 2.3 France: CORONADO (CORONAVirus-SARS-CoV-2 and diabetes outcomes)

150 The CORONADO study described the phenotypic characteristics and prognosis of people with
151 diabetes admitted with COVID-19 between March 10 and April 10, 2020 [13,19]. CORONADO
152 is a cohort study from French hospitals volunteering to share data on hospitalized COVID-19
153 patients with diabetes. The study was sponsored by the Nantes University Hospital and
154 designed in accordance with the Declaration of Helsinki. It obtained all regulatory approvals.

155

156 2.4 Spain – HM Hospitales cohort

157 The six hospitals in the HM Hospitales group collected anonymized observational data for
158 people infected with COVID-19 during the first wave of the pandemics. This dataset is made
159 available to researchers via “Covid Data Save Lives” [20]. The electronic hospital health records

160 were collected for admitted persons, including pre-existing disease status, medication use,
161 demographic, and outcome. A subset of people with pre-existing diabetes from this cross-
162 sectional database was used in this study. Before access was granted, a formal petition,
163 specific study protocol, and ethics committee approval were obtained. The study was approved
164 by the Ethics Committee of the Primary Health Care University Research Institute (IDIAP) Jordi
165 Gol, Barcelona (approval number: 20/089-PCV).

166

167 **2.5 Spain – Barcelona cohort**

168 An observational cohort study was conducted at the Hospital del Mar and Hospital de la Santa
169 Creu i Sant Pau, two tertiary hospitals in Barcelona providing healthcare to 800,000 people. The
170 two hospitals from Barcelona (Catalonia) collected anonymized observational data for people
171 infected with COVID-19 during the first wave. Demographic, clinical, epidemiological, and
172 whole-episode (laboratory workup, vital signs, treatment) data were extracted from electronic
173 medical records using a standardized data collection method. All patients with type 2 diabetes
174 mellitus admitted for COVID-19 between March and April 2020 were included. The Hospital del
175 Mar Institutional Ethics Committee (CEIm-2020/9352) and the Hospital de la Santa Creu I Sant
176 Pau Ethics Committee (HSCSP-20/117) approved the study and waived the informed consent
177 need due to the study's nature.

178

179 **2.6 Data collection: definitions and outcomes**

180 Demographic data included: age, sex, and type of diabetes. UK and France collected ethnicity
181 data (White/Europid, Black/African, Asian/Asian, Other/Middle East and North African (MENA)).
182 Medication use at the point of admission was collected with particular focus on those
183 medications associated with diabetes or diabetes-related comorbidities. Microvascular disease
184 (including retinopathy, neuropathy, and nephropathy) was collected for the UK, French and
185 Spanish (Barcelona cohort) cohorts. The Spanish cohort (HM Hospitales) collected data on the
186 presence of chronic kidney disease (CKD) alone based on clinical coding records. CKD was
187 defined by eGFR <60 ml/min or the presence of macroalbuminuria (urinary albumin-to-
188 creatinine ratio ≥ 300 mg/g) [21].

189

190 **2.7 Definition of macrovascular complications**

191 History of macrovascular disease was collected for all datasets. The presence of macrovascular
192 complications was defined according to the presence of a previous history of ischemic heart
193 disease (including a history of myocardial infarction and/or coronary artery revascularization or
194 heart failure), cerebrovascular disease (including history of stroke or transient ischemic attack –
195 TIA-) and/or peripheral artery disease (amputation owing to ischemic disease and/or lower limb
196 artery revascularization). Data were obtained based on the information recorded in medical
197 records or according international ICD10 classification. Of the 6,795 people included, 4,106 had
198 complete data for macrovascular complications (all four cohorts) and/or the rest of the variables
199 included (French, UK and Spanish HM Hospitales cohorts). **The flowchart of the study is**
200 **summarized in Figure 1.** The descriptive analysis compared the characteristics of people with
201 and without complete data for macrovascular disease. The comparison of the clinical
202 characteristics of people with complete data compared with those with missing data for
203 macrovascular complications is shown in **Table 1.**

204

205 **2.8 Outcomes**

206 The primary outcome was all-cause mortality (French data were collected to day 28 after
207 admission, and Spanish and UK data included mortality during the whole hospital episode). The
208 secondary outcomes were intensive care unit (ICU) admission for all four cohorts and use of
209 invasive mechanical ventilation (IMV) during the hospitalization for the French and the Spanish
210 HM Hospitales cohorts for which IMV data were available.

211

212 **2.9 Statistical analysis**

213 All quantitative data were tested for normality. Clinical characteristics were expressed as the
214 number (percentage) of participants for categorical variables, mean +/- standard deviation (SD)
215 for normally distributed continuous variables, or median (25th-75th percentile) for non-normally
216 distributed continuous variables. **Multivariable logistic regression models were used to obtain**
217 **odds ratio for the primary (death) and secondary outcomes (ICU admission and IMV). The main**
218 **exposure was the presence of macrovascular complications. Regulatory issues prevented us**
219 **from sharing and combining individual level data from each contributing country, so,**

220 multivariable logistic regression models were used to analyse the association between the
221 presence of macrovascular disease (main exposure) and primary (death) and secondary
222 outcomes (ICU admission and IMV) in each country's data separately. Logistic regression
223 models were then used to adjust for potential confounders: age and sex (model 1), age, sex,
224 type of diabetes, arterial hypertension, and the presence of microvascular disease (model 2)
225 and age, sex, type of diabetes, arterial hypertension, microvascular disease, ethnicity, and BMI
226 (model 3 – data not available for HM cohort). Logistic regressions were performed using R in
227 each contributing country; country-specific odds ratios were then pooled in both common effect
228 and random effect meta-analysis (as needed according to I^2 statistic) and using the inverse
229 variance method. Heterogeneity across studies was evaluated using the I^2 statistic. Results
230 were expressed as odds ratio (OR) and 95% confidence interval (95% CI) and p-values <0.05
231 were considered statistically significant. Statistical analyses were performed with R statistical
232 software version 3.6.1 (<https://www.r-project.org/>).

233

234 3. Results

235 The UK ABCD COVID-19 audit collected data on 3,179 people with diabetes from over 40
236 hospitals between March and October 2020. Of these, 1,846 (58.1%) had complete data
237 required for this study and were included in the analysis. CORONADO investigators collected
238 data on 2,843 people with diabetes from 68 hospitals, with 1,510 (53.1%) having complete data.
239 Spanish investigators from the HM Hospitales collected data on 2,310 individuals at six
240 hospitals. There was complete data for 406 individuals (100.0%) with pre-existing diabetes.
241 Finally, the Spanish investigators from Hospital del Mar and Hospital de la Santa Creu i Sant
242 Pau collected data on 367 individuals, with 344 (93.7%) having complete data. A comparison of
243 the complete dataset to that with missing data within all the countries (**Table 1**). Ethnicity data
244 were not available for the Spain – HM Hospitales cohort.

245

246 The baseline characteristics of the cohort of patients in each country are summarized in **Table**
247 **2**. Data related to macrovascular disease status was available for 4,106 people. Of these, 1,652
248 (40.2%) had any prior macrovascular disease, 1,339 (32.6%) had a previous history of ischemic
249 heart disease, 520 (12.7%) had previous cerebrovascular disease, and 457 (11.1%) had

250 previous peripheral artery disease. In the four included cohorts, people with a history of
251 macrovascular disease were older, had a higher percentage of men and had a higher
252 prevalence of hypertension, dyslipidemia, and microvascular complications than the group with
253 no history of macrovascular disease.

254

255 In total, 1,172 (28.5%, range 19.2-39.0%) people died. Mortality was higher for people with
256 compared to without previous macrovascular disease (37.7% vs 22.4%). The combined crude
257 odds ratio (OR) for previous macrovascular disease and mortality was 2.12 (95%CI 1.83-2.45)
258 (**Figure 2, panel A**), with moderate heterogeneity (I^2 60%). In the multivariable analyses, the
259 results were attenuated after adjusting for age and sex (model 1: OR 1.39 [95%CI 0.86-2.26])
260 (**Figure 2, panel B**) and age, sex, type of diabetes, arterial hypertension, and the presence of
261 microvascular disease (model 2: OR 1.38 [95%CI 0.93-2.04]) (**Figure 2, panel C**). The final
262 model and additional adjustment for ethnicity and BMI (in a subset including 3 of the 4 cohorts)
263 showed similar results (OR 1.53 [95%CI 1.29-1.81]) (**Figure 2, panel D**). Further, each
264 component of macrovascular complications was analysed separately, to know which of the 3
265 diseases contributed the most. Both ischemic heart disease (unadjusted OR 1.78 [95%CI 1.20-
266 2.63]), cerebrovascular disease (unadjusted OR 1.91 [95%CI 1.57-2.34]) and peripheral artery
267 disease (unadjusted OR 1.70 [95%CI 1.38-2.10]) were associated with higher mortality
268 (**Supplemental Figure 1-3**). The results were maintained significant after further adjustments
269 except for peripheral artery disease (**Supplemental Figure 1-3**).

270

271 Regarding the secondary outcomes, 699 people (15.7%, range 8.6-22.3%) were admitted to
272 ICU, and 306 (13.6%, range 9.1-17.8%) required IMV during hospitalization. The proportions
273 admitted to ICU or treated with IMV during hospitalization were lower for those patients with
274 previous macrovascular disease (11.5% vs 20.7% and 11.9% vs 18.5%, respectively). The
275 overall odds ratio (OR) for previous macrovascular disease and ICU admission was 0.48
276 (95%CI 0.31-0.75) in the unadjusted analyses (**Figure 3, panel A**), with moderate heterogeneity
277 (I^2 68%). In the multivariable analyses, the estimates were similar after adjusting for age and
278 sex (model 1: OR 0.61 [95%CI 0.49-0.77]) (**Figure 3, panel B**); age, sex, type of diabetes,
279 arterial hypertension, and the presence of microvascular disease (model 2: OR 0.58 [95%CI

280 0.47-0.72]) (**Figure 3, panel C**); and age, sex, type of diabetes, arterial hypertension,
281 microvascular disease, ethnicity, and BMI (model 3: OR 0.57 [95%CI 0.44-0.74]) (**Figure 3,**
282 **panel D**). Both ischemic heart disease (unadjusted OR 0.53 [95%CI 0.44-0.64]),
283 cerebrovascular disease (unadjusted OR 0.32 [95%CI 0.12-0.84]) and peripheral artery disease
284 (unadjusted OR 0.48 [95%CI 0.34—0.66]) were associated with lower mortality (**Supplemental**
285 **Figure 1-3**). The results were maintained significant after further adjustments in all cases
286 (**Supplemental Figure 1-3**). Finally, the overall odds ratio (OR) for previous macrovascular
287 disease and use of IMV during hospitalization was 0.52 (95%CI 0.40-0.68) in the unadjusted
288 analyses (**Figure 4, panel A**), with little evidence of heterogeneity (I^2 37%). In the multivariable
289 analyses, the results were similar after adjusting for age and sex (model 1: OR 0.63 [95%CI
290 0.47-0.85]) (**Figure 4, panel B**) and age, sex, type of diabetes, arterial hypertension, and the
291 presence of microvascular disease (model 2: OR 0.61 [95%CI 0.45-0.83]) (**Figure 4, panel C**).
292 Ischemic heart disease (unadjusted OR 0.55 [95%CI 0.43-0.71]) was associated with lower
293 mortality (**Supplemental Figure 1-3**). The results were maintained significant after further
294 adjustments in all cases (**Supplemental Figure 1-3**). The association between cerebrovascular
295 disease and peripheral artery disease and use of IMV during hospitalization was not tested for
296 insufficient number of events.

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4. Discussion

300 This is the first European retrospective study to specifically investigate the association between
301 previous macrovascular disease and severe outcomes of people with diabetes mellitus
302 hospitalized for COVID-19. The present study demonstrates that presence of macrovascular
303 complications (ischemic heart disease, stroke, peripheral artery disease) was associated with
304 higher mortality. These findings remain significant after further adjustments for age, sex, type of
305 diabetes, arterial hypertension, microvascular disease, ethnicity, and BMI. However, the
306 proportions admitted to ICU and treated with IMV during hospitalization were lower for patients
307 with previous macrovascular disease reflecting clinical decisions around ICU admission.

308 COVID-19 is known to be associated with poorer outcomes for those with long-term conditions
309 such as diabetes, and several potential mechanisms have been proposed [22]. Moreover, age,
310 sex, ethnicity, deprivation, and other comorbidities such as obesity, hypertension and
311 cardiovascular disease contribute to the increased risk [23]. The underlying mechanisms
312 resulting in adverse outcomes in subjects with diabetes hospitalized for COVID-19 are therefore
313 complex and unknown. In that sense, it has been suggested that both impaired glucose
314 regulation and hyperglycemia [24] and the visceral adipose tissue expansion (and its associated
315 ectopic fat depots) that characterize patients with diabetes and/or obesity activate the
316 inflammatory cascade, increasing the production of interleukin-6, which has been proposed as a
317 determinant factor of the “cytokine storm” associated with severe forms of COVID-19 [25].

318 A recent meta-analysis, including 158 observational studies with a total of 270,212 participants
319 of whom 57,801 had diabetes, reported that people with diabetes were at higher risk of COVID-
320 19-related mortality with an OR 1.87 (95%CI 1.61-2.17) and higher risk of ICU admission (1.59
321 [95%CI 1.15-2.18]) and ventilation requirements (1.44 [95%CI 1.20-1.73]) in comparison to
322 subjects without diabetes [11]. In addition, cardiovascular disease is a common comorbidity
323 observed in patients with COVID-19, associated with increased severity and mortality [22]. In
324 that sense, it has been reported that patients with COVID-19 who have either hypertension or
325 cardiovascular disease have an approximately 3-4-fold higher risk of developing a severe
326 disease [26]. By contrast, recent observational studies have demonstrated a significant

327 association between statins (with anti-inflammatory and vasculo-protective effects) and reduced
328 mortality in patients hospitalized with COVID-19, especially those with diabetes [27].

329 Moreover, an adverse effect on outcomes of diabetic complications in patients with diabetes
330 during the COVID-19 pandemic has been recently identified. In a national population-based
331 study in Scotland, associations with fatal or critical care unit treated COVID-19 among people
332 with diabetes adjusted for age, sex, diabetes duration, and type of diabetes were reported for 35
333 factors, including heart disease (OR 2.43 [95%CI 2.14-2.75]), history of hospital admission with
334 diabetic ketoacidosis (OR 2.87 [95%CI 1.85–4.46]), microalbuminuria (OR 1.35 [95%CI 1.16–
335 1.58]), macroalbuminuria 1.92 [95%CI 1.52–2.43]) and severe retinopathy (OR 1.92 [95%CI
336 1.52–2.43]) [28]. The CORONADO study found that both microvascular (OR 2.14 [95%CI:1.16–
337 3.94]) and macrovascular (OR 2.54 [95%CI: 1.44–4.50]) complications were independently
338 associated with the risk of death on day seven of admission after adjusting for age, sex,
339 comorbidities/complications, and glucose lowering and anti-hypertensive treatment [13,14,19].
340 By contrast, no association was found between mortality (death by day seven of admission) and
341 micro- or macrovascular complications in the ACCREDIT Study [15]. Nevertheless, both cohorts
342 differ in several aspects such as mean age (69.8 in the CORONADO study vs. 74.1 for the
343 ACCREDIT study cohort), median BMI (28.4 kg/m² vs. 27.6 kg/m²), the mean HbA1c (8.1% vs.
344 7.7%), which may partially explain the different results regarding the outcomes [13–15,19]. Our
345 analysis suggests that participants living with diabetes hospitalized for COVID-19 with previous
346 macrovascular complications (ischemic heart disease, stroke, peripheral artery disease) have
347 an approximately 50% higher risk of mortality compared to people with no history of
348 macrovascular disease after adjusting for all available confounding factors and that ischemic
349 heart disease and stroke are the main contributors to this higher risk. However, proportions
350 admitted to ICU or treated with IMV during hospitalization were lower for people with previous
351 macrovascular disease. These results suggest unmeasured differences that may explain these
352 opposing relationships, such as (1) the severity of the disease, (2) early mortality leading to
353 potentially fewer patients admitted to ICU or meeting intubation criteria in the group of people
354 with macrovascular disease or (3) the criteria used for ICU admission or IMV. In addition, it
355 should be emphasized that our analysis revealed moderate heterogeneity in mortality and ICU
356 admission for people with diabetes and previous cardiovascular disease with differences in the

357 strength of the relationship between cohorts. While the higher mortality among people with
358 history of macrovascular disease was consistently higher across the four cohorts, there were
359 more marked differences regarding ICU admission and the use of IMV, being lower compared
360 to the Spanish cohorts in the UK (ICU admission) and French cohorts (both ICU admission and
361 use of IMV).

362

363 Our study is limited by the heterogeneity of data collection methods across the nations due to
364 the use of databases that were designed separately and not specifically to answer the study
365 question and, also by missing data. The meta-analysis combines the individual datasets to
366 increase power but masks heterogeneity across nations. The relatively large proportion of
367 people with missing data on one or more variables in UK and French cohorts has occurred as a
368 consequence of using routinely collected data from clinical practice and could introduce bias if
369 data are not missing at random. As shown in Table 1, summary measures of distribution of most
370 variables used in model 3 (age, sex, type of diabetes, arterial hypertension, BMI, microvascular
371 disease, macrovascular disease, death and ICU admission) were similar or had only modest
372 differences between people with and without missing data in both UK and French cohorts.
373 People of non-white ethnicity were over-represented in the missing data group in the UK cohort
374 but not in the French cohort. Clinical data of the whole-episode, such as vital signs or arterial
375 gasometry parameters, clinical severity scores (i.e. MEWS or CURB-65 score) or markers of
376 inflammation (PCR, IL-6, serum ferritin) were not collected, as well as other potential
377 confounding factors. In addition, the small sample size of patients with type 1 diabetes included
378 made impossible to analyze both groups separately to evaluate the potential differences
379 between both. Lastly, we focused on people hospitalized for COVID-19; thus, our results cannot
380 be generalized to all people with diabetes and COVID-19, especially those with less severe
381 forms of the disease.

382 **Conclusions**

383 In conclusion, this large multinational study of people with diabetes mellitus hospitalized for
384 COVID-19 demonstrates significant associations between previous macrovascular disease and

385 higher mortality and with lower ICU admission and the use of IMV during hospitalization. This
386 study is the first specifically designed to evaluate the association of macrovascular
387 complications (ischemic heart disease, stroke, or peripheral artery disease) as main exposure
388 with mortality, intensive care unit admission and mechanical ventilation in people with diabetes
389 mellitus hospitalized for COVID-19 in Europe. Our findings highlight the importance correctly
390 assess the prognosis and intensive monitoring in this high-risk group of patients and emphasize
391 the need to design specific public health programs aimed to prevent SARS-CoV-2 infection in
392 this subgroup (i.e. reinforcing vaccination campaigns). Nevertheless, further studies are
393 required to confirm and extend these findings in these and other populations.

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415 **List of abbreviations**

416 CKD: chronic kidney disease

417 COVID-19: Coronavirus disease 2019

418 ICU: intensive care unit

419 IMV: invasive mechanical ventilation

420 OR: odds ratio

421 SARS-CoV-2: Severe acute respiratory syndrome coronavirus 2

422 SD: standard deviation

423 95% confidence interval (95% CI)

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445 **Figure Legends**

446 **Figure 1. Inclusion and exclusion criteria**

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448 **Figure 2.** Odds ratio for the association between mortality and the presence of previous
449 macrovascular disease in each of the four cohorts and overall (panel A). I^2 indicates
450 heterogeneity in the estimates. Odds ratio estimates adjusted for: model 1: age and sex (panel
451 B); model 2: model 1 + type of diabetes, arterial hypertension, and the presence of
452 microvascular disease (panel C); model 3: model 2 + ethnicity and BMI (panel D). HM cohort is
453 excluded from model 3 due to lack of ethnicity data.

454

455 **Figure 3.** Odds ratio for the association between intensive care unit admission and the
456 presence of previous macrovascular disease in each of the four cohorts and overall (panel A). I^2
457 indicates heterogeneity in the estimates. Odds ratio estimates adjusted for: model 1: age and
458 sex (panel B); model 2: model 1 + type of diabetes, arterial hypertension, and the presence of
459 microvascular disease (panel C); model 3: model 2 + ethnicity and BMI (panel D). HM cohort is
460 excluded from model 3 due to lack of ethnicity data.

461

462 **Figure 4.** Odds ratio for the association between use of invasive mechanical ventilation during
463 the hospitalization and the presence previous of macrovascular disease in each of the two
464 cohorts and overall (panel A). I^2 indicates heterogeneity in the estimates. Odds ratio estimates
465 adjusted for: model 1: age and sex (panel B); model 2: model 1 + type of diabetes, arterial
466 hypertension, and the presence of microvascular disease (panel C).

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475 **Supplementary Material**

476 **Supplemental Figure 1.** Odds ratio for the association between mortality (panels 1A-1C),
477 intensive care unit admission (panels 2A-2C) and use of invasive mechanical ventilation during
478 the hospitalization (panels 3A-C) and ischemic heart disease in each of the four cohorts and
479 overall. I^2 indicates heterogeneity in the estimates. Odds ratio estimates adjusted for: model 1:
480 age and sex (panel B); model 2: model 1 + type of diabetes, arterial hypertension, and the
481 presence of microvascular disease (panel C).

482

483 **Supplemental Figure 2.** Odds ratio for the association between mortality (panels 1A-1C),
484 intensive care unit admission (panels 2A-2C) and and stroke in each of the four cohorts and
485 overall. I^2 indicates heterogeneity in the estimates. Odds ratio estimates adjusted for: model 1:
486 age and sex (panel B); model 2: model 1 + type of diabetes, arterial hypertension, and the
487 presence of microvascular disease (panel C).

488

489 **Supplemental Figure 3.** Odds ratio for the association between mortality (panels 1A-1C),
490 intensive care unit admission (panels 2A-2C) and peripheral artery disease in each of the four
491 cohorts and overall. I^2 indicates heterogeneity in the estimates. Odds ratio estimates adjusted
492 for: model 1: age and sex (panel B); model 2: model 1 + type of diabetes, arterial hypertension,
493 and the presence of microvascular disease (panel C).

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505 **Declarations**

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507 **Ethics approval and consent to participate**

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509 **United Kingdom: association of British clinical diabetologists (ABCD) COVID-19 audit:**

510 The audit is registered with Oxford University Hospitals NHS Foundation Trust (OUH), a Data

511 Protection Impact Assessment was carried out and the audit was approved by the OUH

512 Caldicott Guardian and the Public Benefit and Privacy Panel in Scotland (reference 2021-0111).

513 **France: CORONADO:** The study was sponsored by the Nantes University Hospital and

514 designed in accordance with the Declaration of Helsinki. It obtained all regulatory approvals.

515 **Spain – HM Hospitales cohort:** The study was approved by the Ethics Committee of the

516 Primary Health Care University Research Institute (IDIAP) Jordi Gol, Barcelona (approval

517 number: 20/089-PCV).

518 **Spain – Barcelona cohort:** The Hospital del Mar Institutional Ethics Committee (CEIm-

519 2020/9352) and the Hospital de la Santa Creu I Sant Pau Ethics Committee (HSCSP-20/117)

520 approved the study and waived the informed consent need due to the study's nature.

521

522 **Consent for publication:** All authors approved the final manuscript and give consent for the

523 publication.

524

525 **Availability of data and materials:** Data are available on request from corresponding author

526 and/or national study leads with appropriate data governance permission.

527

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529

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549

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551 GL, PD, DM collected the Hospital del Mar-Hospital de la Santa Creu I Sant Pau cohort and BV
552 and JFN collected the cohort from HM Hospitales. GL, BV and DM drafted the first version of
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554 conducted the CORONADO study. RR, SHW, YR and KK (with additional colleagues) designed
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