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Are Psychotic-like Experiences related to a Discontinuation of Cannabis Consumption in Young Adults?

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84 Declaration of Interest

85 Dr. Banaschewski served in an advisory or consultancy role for Lundbeck, Medice, Neurim Pharmaceuticals,
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92 **ABSTRACT**

93 *Objective:* To assess changes in cannabis use in young adults as a function of psychotic-like
94 experiences.

95 *Method:* Participants were initially recruited at age 14 in high schools for the longitudinal
96 IMAGEN study. All measures presented here were assessed at follow-ups at age 19 and at age
97 22, respectively. Perceived stress was only assessed once at age 22. Ever users of cannabis
98 (N=552) gave qualitative and quantitative information on cannabis use and psychotic-like
99 experiences using the Community Assessment of Psychic Experiences (CAPE). Of those,
100 nearly all n=549 reported to have experienced at least one psychotic experience of any form at
101 age 19.

102 *Results:* Mean cannabis use increased from age 19 to 22 and age of first use of cannabis was
103 positively associated with a change in cannabis use between the two time points. Change in
104 cannabis use was not significantly associated with psychotic-like experiences at age 19 or 22.
105 In exploratory analysis, we observed a positive association between perceived stress and the
106 experience of psychotic experiences at age 22.

107 *Conclusion:* Age of first use of cannabis influenced trajectories of young cannabis users with
108 later onset leading to higher increase, whereas the frequency of psychotic-like experiences
109 was not associated with a change in cannabis use. The observed association between
110 perceived stress and psychotic-like experiences at age 22 emphasizes the importance of stress
111 experiences in developing psychosis independent of cannabis use.

112 *Keywords:* cannabis use; psychotic-like experiences; age of first use; perceived stress;
113 cannabis discontinuation hypothesis

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1. INTRODUCTION

117 Cannabis is the most used illicit drug in Europe, with estimates that 24.7 million adults
118 have used the drug in the last year (EMCDDA, 2019). Cannabis use across adolescence is
119 reported to increase and reach its peak in young adulthood (Patton et al., 2007; Tucker et al.,
120 2019). Herbal cannabis and its extracts contain numerous cannabinoids, most notably
121 tetrahydrocannabinol (THC) and cannabidiol (CBD). Evidence has linked cannabis
122 consumption to psychosis (Moore et al., 2007), specifically THC, which is known for its
123 psychoactive effect and can cause intoxicating effects (Morgan and Curran, 2008). The
124 potency of THC in cannabis has risen in herbal and in resin cannabis (EMCDDA, 2019). The
125 increased levels of THC may put users at a higher risk for developing psychosis (Di Forti et
126 al., 2019).

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128 Longitudinal studies show that regular cannabis use is associated with an increased risk
129 for schizophrenia and for reporting psychotic symptoms (Hall and Degenhardt, 2008). More
130 frequent cannabis use is independently associated with more frequent or intense symptoms on
131 three psychotic dimensions: positive, negative and depressive (Bernardini et al., 2018;
132 Schubart et al., 2011a; Skinner et al., 2011; Verdoux et al., 2003). The negative dimension
133 refers to one of the key symptom domains of schizophrenia, with negative symptoms
134 including anhedonia or apathy (Selten et al., 1998), whereas the depressive dimension partly
135 overlaps with negative symptoms, but additionally covers more cognitive symptoms of
136 depression (e.g. sadness, pessimism, feeling guilty) that discriminate between depression and
137 negative symptoms (Kibel et al., 1993; Stefanis et al., 2002; Stefanis et al., 2004). According
138 to meta-analyses, psychotic experiences and cannabis intake show a dose-response
139 relationship (Marconi et al., 2016; Ragazzi et al., 2018), which suggests that psychosis and

140 psychotic-like experiences (PLEs) share the same risk factors, thus supporting an association
141 between cannabis use and PLEs.

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143 Not only is continuous cannabis consumption related to psychosis, but also the age of first
144 use is predictive of frequency and intensity of psychotic symptoms (Konings et al., 2008;
145 Ragazzi et al., 2018; Schubart et al., 2011b; Skinner et al., 2011). Such an association is also
146 reported for negative psychotic symptoms, but to a lesser degree (Schubart et al., 2011b).
147 Together, these findings support the hypothesis that the impact of cannabis use is age
148 dependent and stronger for positive psychotic symptoms.

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150 Although the association between cannabis consumption and PLEs is well documented, its
151 causality and directionality are still intensely debated (Degenhardt et al., 2018; DeVlyder et
152 al., 2018; Hall and Degenhardt, 2008; Murray and Hall, 2020). Different theories are
153 discussed: First, the psychosis risk might be primarily caused by familial risk for
154 schizophrenia and only appears to be triggered by cannabis consumption. For example, Proal
155 et al. (2014) showed that both cannabis using and non-using relatives of patients with
156 psychosis showed increased familial risk for psychotic-like symptoms compared with their
157 respective non-psychotic control samples. Secondly, co-occurring genetic or environmental
158 risk factors including stress exposure could contribute to both cannabis use and PLEs in
159 adolescents (Shakoor et al., 2015; Arranz et al., 2018). Thirdly, cannabis use disorder also
160 could directly affect the risk for PLEs (Nesvåg et al., 2017). Fourthly, cannabis could be used
161 as self-medication in face of subclinical symptoms of psychosis to reduce distress (Mané et
162 al., 2015).

163

164 It has been reported that a decrease in cannabis use in $n=705$ young adults aged 18–27
165 years was associated with a decrease in psychotic experiences, while increased consumption

166 was linked to positive symptoms at follow-up (Van Gastel et al., 2014). This association
167 between changes in cannabis use and changes in the frequency of PLEs does not prove a
168 causal relationship, but strongly suggests a bidirectional association and a reduction of PLEs
169 after the cessation of cannabis use. Interestingly, the “cannabis discontinuation hypothesis”
170 suggests that in young adolescents, aversive effects of cannabis use including the
171 manifestation of psychotic symptoms may trigger a reduction in cannabis consumption by
172 self-selection, i.e. a self-imposed protection from the risk of developing enduring psychotic
173 disorders (Sami et al., 2019; Van Gastel et al., 2012). Moreover, cessation of cannabis
174 consumption was predicted by more aversive subjective experiences with cannabis and by no
175 increase in the first three years after first use (Seidel et al., 2019), which could partly be
176 mediated by aversive psychotic experiences. Hence, in the present study we sought to
177 investigate the association of change in cannabis use over a period of 3 years with the
178 occurrence of PLEs in a non-clinical sample of young adults, controlling for potentially
179 confounding factors including age of first use of cannabis, other illicit drug use and socio-
180 economic status.

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182 **1.1. Anecdotal evidence from qualitative interviews for hypothesis generation**

183 Qualitative interviews in our study were conducted within the scope of the interdisciplinary
184 research project ERANID, which focuses on use of illicit drugs including cannabis (ERANID,
185 2015). For the purpose of hypothesis generation, interviews were conducted additionally to
186 quantitative data using a mixed-method approach. Detailed information on the ethnographic
187 methods can be found in section 2.1. One topic that emerged in several interviews was the
188 cessation of cannabis consumption after the experience of psychotic experiences, as suggested
189 by the so-called cannabis discontinuation hypothesis (Sami et al., 2019). For exemplification,
190 we here provide a quote of one participant (age 22):

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192 *“I think that definitely a motivation for stopping was every time I got reasonably high,*
193 *I would start to have paranoid thoughts, not in a psychotic way like, people were*
194 *watching me or whatever [...]. So, yeah, I kind of had enough of that. Taking a break*
195 *has stopped that so I think that was a good decision.”*

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197 **1.2. Hypotheses**

198 We tested the hypothesis that (1) cannabis use at age 19 is predictive of cannabis use at
199 age 22; (2) early age of first use of cannabis is predictive of increase in cannabis use from 19
200 to 22; (3) total occurrence of distressful PLEs at age 19 as well as frequency and distress of
201 positive PLEs are associated with reductions in cannabis use between age 19 and 22; and (4)
202 current cannabis use at age 19 or 22 is associated with current PLEs at these time points.
203 Furthermore, we explored the association of stress effects at age 22 with PLEs and cannabis
204 use.

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2. METHODS

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2.1. Sample

209 The sample was drawn from the longitudinal European IMAGEN cohort (Schumann et
210 al., 2010). The IMAGEN study consists of a community sample recruited at the age of 14
211 (N=2214) from 8 sites across Europe. Follow up 1 (FU1) was conducted at age 16 (N=1700).
212 Here we used data from the second follow up at age 19 (FU2; N=1515) and the third follow
213 up (FU3; N=1360) at age 22. In the current study, we included all participants who had
214 reported to have used cannabis at least once in their life at the age of 19 (for assessment see
215 2.2.2.). Recruitment strategies and inclusion criteria can be found elsewhere (Schumann et al.,
216 2010). The anecdotal evidence provided above was obtained in a subsample ($N = 42$) of the
217 IMAGEN cohort within the scope of the research project Imagen Pathways funded by
218 ERANID (ERANID, 2015). Here, ethnographic interviews on the experience of illicit drug
219 use were conducted at age 22, transcribed by independent assistants, and reoccurring topics in
220 relation to cannabis use were extracted by ethnographic researchers.

221 All study participants were provided with a description of the study and written informed
222 consent was obtained before participation. The research protocol was approved by local
223 Ethics Committees and adhered to the Declaration of Helsinki.

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2.2. Measures

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2.2.1. Psychotic-like experiences (PLEs)

227 *Community Assessment of Psychic Experiences (CAPE)*. PLEs were assessed using the
228 CAPE (Stefanis et al., 2002), a self-report questionnaire consisting of 42 items, which has
229 been found to be a reliable and valid instrument for evaluating the presence of lifetime
230 psychotic-like symptoms in the general population in various languages (Mark and

231 Touloupoulou, 2017, 2016; Mossaheb et al., 2012; Schlier et al., 2015; Vermeiden et al., 2019).
232 The CAPE measures 1) frequency and 2) associated distress of psychotic experiences on three
233 symptom dimensions: positive (*Pos*), negative (*Neg*) and depressive (*Dep*) (Konings et al.,
234 2006; Stefanis et al., 2002). PLEs were not queried explicitly in relation to cannabis
235 consumption, hence the CAPE score reflects PLEs induced by cannabis use as well as non-
236 cannabis related PLEs across lifespan. The frequency scale answers comprise the options:
237 never (0); sometimes (1); often (2); and nearly always (3); whereas the distress scale answer
238 options are: not distressed (0); sometimes (1); often (2); and nearly always (3). Items scores
239 were re-coded (range: 1 to 4) and added up to a total score (*CAPE_{Total}*) and to the sum scores
240 for the positive dimension, i.e. the frequency of positive symptoms and the distress associated
241 with them (CAPE - positive frequency: *CAPE_{PosFreq}*; CAPE - positive distress:
242 *CAPE_{PosDis}*). Sum scores were weighted with number of answered items to account for
243 partial non-responders resulting in a value ranging from 1 to 4. In our analysis, the total score
244 and the weighted sum scores were used as continuous measures.

245 **2.2.2. Cannabis use**

246 *European School Survey Project on Alcohol and Drugs (ESPAD)*. The ESPAD (Hibell et
247 al., 1997) was used to measure the frequency of cannabis use in the past year at age 19 and
248 age 22 respectively in an online design by asking the question: “On how many occasions
249 OVER THE LAST 12 MONTHS have you used marijuana (grass, pot) or hashish (hash, hash
250 oil)?”. Answers were scored between 0-6 according to their use frequencies: never (0); once
251 or twice (1); 3-5 times (2); 6-9 times (3); 10-19 times (4); 20-39 times (5); 40 times or more
252 (6). Additionally, age of first use of cannabis was asked at age 19 using the question: “When
253 did you first try marijuana (grass, pot) or hashish (hash, hash oil)?”.

254 The difference in frequency of cannabis use assessed at FU2 versus FU3 was calculated
255 by subtracting frequency at age 22 from frequency at age 19. The difference in frequency of
256 cannabis use was used as main outcome variables in our analysis.

257 **2.2.3. Stress measures**

258 *Perceived Stress Scale.* The perceived stress scale (PSS) is a self-report scale measuring
259 perceived stress with 10 items (Cohen et al., 1994). The degree to which situations are
260 perceived as unpredictable, uncontrollable and overloaded is assessed using a 5-point Likert
261 scale ranging from never (0), almost never (1), sometimes (2), fairly often (3), very often (4).
262 Total scores range from 0 to 40, with higher scores indicating greater perceived stress.

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264 **2.2.4. Covariates**

265 Additional parameters of drug use were assessed at FU2 and FU3 and used as covariates.
266 Apart from gender, age of first use of cannabis (if applicable), the use of other illicit drugs
267 (ever vs. never), nicotine dependence, parental socio-economic status (SES) and psychiatric
268 disorders were introduced as covariate in our analysis (for details of assessment see
269 supplements). Additionally, recruitment site was introduced as covariate in our analysis. As
270 number of inhabitants is related to urbanicity, which has been associated with psychotic-like
271 experiences in children (Karcher et al., 2020) and considered to be a general risk factor for
272 psychosis in adults in developed countries (Heinz et al., 2013), we ranked the recruitment
273 sites in the order of inhabitants of the respective city to account for possible differences in
274 urbanicity.

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276 **2.3. Data analysis**

277 The analyses were carried out with the statistical package for the social sciences (SPSS
278 20.0). Descriptive statistics for the predictor (*CAPETotal*, *CAPEPosFreq* and *CAPEPosDis*),

279 main outcome variables (cannabis use, change in cannabis use) and all covariates (gender
280 identification, recruitment site, age of first use, other illicit drug use, nicotine dependence,
281 SES, and diagnosis of any psychiatric disorders) were estimated as means and standard
282 deviations (*SD*) for continuous variables and as frequencies for all other variables (Table 1).
283 Listwise exclusion was applied for missing values and a quality check was applied for
284 cannabis use: participants who stated never to have used at age 22, while they indicated
285 cannabis use at age 19, were removed from the original sample of 562 participants ($N=10$).
286 First exploratory analyses including *t*-tests for continuous variables and χ^2 test for categorical
287 variables were conducted to compare the 3 groups of change in use (decrease, unchanged,
288 increase) (Table 2).

289 Regressions (ordinal and linear) were carried out according to our hypotheses with either
290 cannabis use or the change in cannabis use as the outcome measure and, respectively,
291 cannabis use, age of first use, *CAPETotal*, *CAPEPosFreq* and *CAPEPosDis* score as
292 predictors. The predictor variables were tested *a priori* to verify there was no violation of the
293 assumption of no multicollinearity (see T1 in supplements). We first investigated model (I)
294 correcting for gender identification and site. In model (II), the other covariates were
295 additionally included. Post-hoc analyses were performed with the changes in cannabis use and
296 the frequency and distress scores of the positive subscales as outcome variables.

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3. RESULTS

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3.1. Sample characteristics

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Of the 1434 subjects who participated in FU2 and FU3 the IMAGEN study, 562 subjects indicated ever use of cannabis at age 19 and provided data for both follow up time points. After inconsistency checks for cannabis use (see 2.3.), 552 subjects who used cannabis at least once were included in our analysis (221 from UK, 88 from France and 243 from Germany). Of those, nearly all ($n=549$) reported to have experienced at least one psychotic experience of any form at age 19. Average age at FU2 was 19.08 years ($SD = .78$), ranging from 17 to 21 years. Average age at FU3 was 22.59 years ($SD = .69$) ranging from 20 to 25 years. The average time span between two timepoints was 3.51 years ($SD = .74$) (Table 1).

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3.2. Changes in cannabis consumption over time

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In this sample of 552 ever users of cannabis, 37.9% of all participants reduced their cannabis use between age 19 and age 22, about a third showed no change (33.5%), and 28.4% increased their cannabis use over the course of 3 years. More participants reported no use of cannabis within the past year at age 22 (31.5%) than at age 19 (23%). Change in cannabis use was normally distributed (Figure 1) and sample characteristics stratified for three groups (decrease, unchanged, increase) are shown in Table 2.

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In line with our hypothesis, cannabis use at age 19 in the ordinal logistic regression analysis was found to predict cannabis use at age 22 in model (I) ($\beta=.536$, $SD=.042$; Wald $\chi^2(1) = 160.050$, $p<.001$) with an estimated odds ratio of 1.7-fold (95% CI, 1.573 to 1.857) for every unit increase of cannabis use at age 19. Also, gender was found to contribute to the model as covariate ($\beta=.643$, $SD=.163$; Wald $\chi^2(1) = 15.52$, $p<.001$) with an estimated odds ratio of nearly 1.9-fold (95% CI, 1.382 to 2.621) for male gender identification. In model (II) age of first use and other illicit drug use also showed a significant association (see T2 in supplements).

324 **3.3. Age of first use and change in cannabis use**

325 Testing whether early age of onset is predictive of an increase in cannabis use from age 19
326 to age 22 in model (I), we found that age of first use was predictive for the observed change in
327 cannabis consumption, with later age increasing the odds for an increase in consumption
328 ($\beta=.180$, $SD=.057$; Wald $\chi^2(1) = 9.92$, $p=.002$). The estimated odds ratio favored a positive
329 relationship of 1.2-fold (95% CI, 1.070 to 1.340) for every year later the first use occurred
330 (Figure 2). Thus, our hypothesis was not confirmed that early age of onset is predictive of a
331 later increase in cannabis use, with results even pointing in a different direction. In model (II),
332 other illicit drug use, nicotine dependence score, SES and psychiatric diagnosis were
333 introduced as covariates, of which other illicit drug use ever significantly contributed to the
334 increase of cannabis use from age 19 to 22 (Table 3).

335 **3.4. Association between PLEs at age 19 and change in cannabis use between** 336 **age 19 and 22**

337 We did not find PLEs at age 19 to be predictive of the change in cannabis use from age 19
338 and 22 in model (I) using gender and site as covariates (Table 4). Also, no significant
339 association was found for any of the CAPE subscales: *CAPETotal*; *CAPEPosFreq*;
340 *CAPEPosDis*. Applying model (II) with age of first use of cannabis, other illicit drug use
341 ever, smoking and SES did not change the predictive value of PLEs (Table 4).

342 We also explored whether PLEs at age 22 are significantly associated with changes in
343 cannabis use from age 19 to 22, and again observed no significant association, neither in
344 model (I) nor in model (II) (see T3 in supplements).

345 **3.5. Association between current PLEs and current cannabis use at age 19 or**

346 **22**

347 We tested whether current cannabis use at age 19 or 22 is associated with current PLEs at
348 age 19 or 22, respectively. In model (I), an association at age 19 was not confirmed, whereas
349 at age 22, we found frequency of cannabis use to be associated with the *CAPETotal* score
350 ($\beta=.700$, $SD=.212$; Wald $\chi^2(1) = 10.812$, $p=.001$) at age 22 in model (I). When including the
351 covariates in the analysis (model II), only psychiatric diagnoses and SES were significantly
352 associated with the *CAPETotal* score (Table 5).

353 **3.6. Association between perceived stress and PLEs and between perceived** 354 **stress and cannabis use**

355 In exploratory analyses, we observed a positive correlation for perceived stress at age 22
356 and the CAPE total score ($r(539) = .48$, $p < .001$), the *CAPEPosFreq* scale ($r(539) = .305$, $p <$
357 $.001$) and the *CAPEPosDis* scale ($r(539) = .308$, $p < .001$), respectively (Figure 3). For
358 perceived stress and current cannabis use at age 22, no significant association was found ($r_{\tau} =$
359 $-.026$, $p = .428$).

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4. DISCUSSION

363 In this longitudinal study in 552 subjects from the general population, we investigated
364 whether cannabis use and its change between age 19 and 22 are associated with PLEs, and we
365 explored whether perceived stress is associated with cannabis use or PLEs. We observed that
366 cannabis use at age 19 was positively associated with cannabis use three years later (age 22).
367 Surprisingly, later first use of cannabis was associated with an increase in cannabis use
368 between age 19 and 22. Regarding the “cannabis discontinuation hypothesis” (Sami et al.,
369 2019; van Gastel et al., 2014), we could not confirm that (distressful) PLEs predict
370 subsequent reductions in cannabis use. Instead, we observed that frequency of cannabis use
371 was positively associated with PLEs at age 22, however, this finding was no longer significant
372 after including presence of psychiatric diagnoses as a covariate. In our exploratory analysis,
373 we observed perceived stress to be associated with PLEs at age 22, but not with cannabis use.
374
375 Regarding our first results, observing that cannabis use at age 19 is associated with cannabis
376 use 3 years later is a plausible finding, which confirms previous study results (Chen et al.,
377 1997; Jones et al., 2016; Patton et al., 2007). The frequency of cannabis use tends to increase
378 in puberty, and on average still continues increasing between age 19 and 22 (Melchior et al.,
379 2008), which was also found in our sample. From age 19 on, different trajectories can be
380 observed in our data, including no change of use as well as increases or decreases in cannabis
381 use. Surprisingly, in our sample the age of first use of cannabis was positively correlated with
382 change in cannabis use from age 19 to 22, indicating that those who initiated use at age 15
383 and later were more likely to increase their use between age 19 and 22 than those who started
384 earlier. While we hypothesized a straightforward association of early first use with higher
385 frequency in cannabis use, some studies indeed suggest more complex trajectories of cannabis
386 use across adolescence and early adulthood (Scholes-Balog et al., 2016; Taylor et al., 2017).

387 According to Scholes-Balog et al. (2016), early-onset cannabis users often start before the age
388 of 15 and usually show persistent use throughout adolescence (1/month), whereas late-onset
389 users usually start after age 15 and tend to use cannabis less often (3-5/year). In our sample,
390 first users at age 15 decreased their use between 19 and 22, which does not support the
391 hypothesis of a rather persistent use of “early-onset” users. Late-onset users in our study
392 increased their use during early adulthood, which raises the concern of persisting harmful use.
393 Given that our sample was followed up 3 times since the age of 14 (Schumann et al., 2010), it
394 is possible that our results partly reflect a selection bias inherent to the longitudinal study
395 design. Dropouts in longitudinal studies are more likely to use substances and tend to report
396 higher mean use of substances at baseline than non-dropouts (Snow et al., 1992), which could
397 affect our final sample at age 22 and contribute to an underestimation of use. Unlike
398 hypothesized (Mullin et al., 2012; Van Gastel et al., 2014), we did not find an association
399 between PLEs at age 19 (or 22) and the change in cannabis use during this observation period.
400 Therefore, the “cannabis discontinuation hypothesis” (Sami et al., 2019; Van Gastel et al.,
401 2012) was not confirmed.

402 Regarding current cannabis use predicting PLEs at the same time point, the occurrence
403 of other psychiatric diagnoses explained the occurrence of PLEs better than cannabis use (or
404 male gender) at both time points. This may reflect the genetic overlap between several mental
405 disorders (Witt et al., 2017) or common environmental factors contributing to both cannabis
406 use disorder and other mental disorders (Heinz et al., 2013; Van Os et al., 2010). Also, the
407 fact that we did not observe an association may be due to the rather low clinical load of our
408 sample. Our PLE score was rather low compared with Barragan et al., (2010: $M = 68.3$, $SD =$
409 13.4) and this restricted variance may limit significant associations with individual differences
410 in cannabis use.

411

412 Finally, the frequency of PLEs was significantly and positively associated with perceived
413 stress. It has been hypothesized that stress exposure contributes to the manifestation of
414 psychotic experiences (Heinz et al., 2020) or that perceived stress levels indicate an increased
415 vulnerability for severe mental disorders (Fusar-Poli et al., 2017). However, our data are only
416 correlational, and the directionality of this interaction needs to be examined in longitudinal
417 studies. On the other hand, we did not find a significant association between perceived stress
418 and cannabis use, rendering it rather unlikely that cannabis was used as self-medication to
419 reduce stress by a majority of the sample (Mané et al., 2015).

420

421 **4.1. Limitations**

422 The major limitation of this study is that selective drop-outs may have occurred during the
423 observation period. This could reduce power to detect effect of increased cannabis use on
424 PLEs. Also, the fact that consumption data were gathered by self-report via online assessment
425 could possibly lead to either over- or underreporting of illegal drug consumption including
426 cannabis use. However, recent studies have shown that web-based questionnaires are a
427 suitable instrument for scientific research and potential biases regarding drug use are unlikely
428 to be systematic (Martin-Willett et al., 2020; Meyerson and Tryon, 2003; Vleeschouwer et al.,
429 2014). Another potential limitation is that the CAPE questionnaire assesses some PLEs that
430 can be hard to distinguish from acute intoxication effects of cannabis. There is, however,
431 some evidence that high CAPE scores associated with acute cannabis intoxication also reflect
432 psychosis proneness (Genetic Risk and Outcome in Psychosis (GROUP) investigators, 2011).

433

434 **4.2. Conclusion**

435 Altogether, we observed a general increase in cannabis use across early adulthood and a
436 positive correlation with (late) age of first use, supporting the notion of diverse trajectories in
437 cannabis use in the general population (Bourque et al., 2017; Patton et al., 2007). We did not

438 find an association between PLEs and subsequent cannabis use, thus not confirming the
439 hypothesis that distressful or other PLEs induce a decline in cannabis use (Van Gastel et al.,
440 2014). Interestingly, perceived stress at age 22 was associated with PLEs (but not with
441 cannabis consumption), emphasizing the importance of perceived stress for psychosis risk
442 (Fusar-Poli et al., 2017). These findings suggest to further explore stress effects on the
443 manifestation of PLEs and vice versa.

444

445

446

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 660 **Table 1.** Sample characteristics of total study sample (n= 552) by gender identification at age
 661 19 and age 22.

Characteristics	Total sample		N available for analyses	p-value ^{a,b}
	female	male		
<i>N</i>	552			
Gender identification	<i>female</i>	<i>male</i>	<i>female/male</i>	
	258	294		
Parental socio-economic status (SES)* (<i>M ± SD</i>)	5.94±.90	5.86±.93	215/244	.382
Ethnicity*			258/294	.351
Central European	236	270		
Black or mixed Black	8	5		
Asian or mixed Asian	7	13		
Other or mixed other	7	6		
Recruitment site (<i>N</i>)			258/294	.534
London	36	38		
Paris	45	43		
Berlin	29	22		
Hamburg	33	47		
Dresden	21	31		
Dublin	25	38		
Nottingham	40	44		
Mannheim	29	31		
Age of onset of cannabis use* (<i>M ± SD</i>)	15.94±1.59	15.84±1.71	143/217	.583
	<i>female</i>	<i>male</i>	<i>female/male</i>	p-value^{a,c}
Total frequency of PLEs (CAPE <i>Total</i>)				.000
Age 19 (<i>M ± SD</i>)	64.83±12.41	62.12±11.74	258/294	
Age 22 (<i>M ± SD</i>)	62.26±11.03	60.21±11.19	254/289	
Frequency of positive PLEs (CAPE <i>PosFreq</i>)				.000
Age 19 (<i>M ± SD</i>)	1.32±.24	1.32±.25	258/294	
Age 22 (<i>M ± SD</i>)	1.25±.22	1.26±.22	254/289	
Distress of positive PLEs (CAPE <i>PosDis</i>)				.000
Age 19 (<i>M ± SD</i>)	1.79±.48	1.59±.50	246/286	
Age 22 (<i>M ± SD</i>)	2.73±.49	2.50±.45	230/272	
Cannabis use within last 12 month*				.000
Age 19 (<i>yes/no</i>)	192/66	233/61	258/294	
Age 22 (<i>yes/no</i>)	153/105	225/69	258/294	
Other illicit drug use ever*				.000
Age 19 (<i>yes/no</i>)	89/169	110/184	258/294	
Age 22 (<i>yes/no</i>)	140/118	185/109	258/294	
Nicotine Dependence*				.030
Age 19 (<i>M ± SD</i>)	.57±1.38	.70±1.40	258/294	
Age 22 (<i>M ± SD</i>)	.39±1.13	.64±1.42	258/294	
Any disorder (clinical rating, DSM-IV)*				.000
Age 19 (<i>yes/no</i>)	68/175	30/245	243/275	
Age 22 (<i>yes/no</i>)	61/134	40/163	195/203	

662 *Annotations:* *N* = sample size; *M* = mean; *SD* = standard deviation; f = female; m = male; *details of assessment
 663 can be found in supplements; ^a According to one-way ANOVA or χ^2 tests to test for possible differences in
 664 ^bgender groups or ^cbetween age 19 and age 22 for the total sample

Table 2. Sample characteristics of total sample (n= 552) stratified by change in cannabis use between age 19 and age 22: decrease, unchanged or increase.

	Group: Change in cannabis use from age 19 to 22						p-value ^a	
	decrease		unchanged		increase			
<i>N</i>	209		185		158			
Gender identification (<i>female/male</i>)	110/99		81/104		67/81			.093
Timepoint of assessment	<i>age 19</i>	<i>age 22</i>	<i>age 19</i>	<i>age 22</i>	<i>age 19</i>	<i>age 22</i>	<i>age 19</i>	<i>age 22</i>
Total frequency of PLEs (<i>CAPETotal</i>) (<i>M ± SD</i>)	63.90±12.83	60.50±10.48	63.39±12.13	62.97±12.53	62.69±11.16	59.66±11.46	.638	.021
Frequency of positive PLEs (<i>CAPEPosFreq</i>) (<i>M ± SD</i>)	1.33±.24	1.23±.19	1.32±.26	1.28±.25	1.31±.23	1.24±.23	.811	.077
Distress of positive PLEs (<i>CAPEPosDis</i>) (<i>M ± SD</i>)	1.75±.52	2.64±.52	1.63±.47	2.59±.47	1.65±.50	2.56±.47	.054	.294
Age of onset of cannabis use* (<i>M ± SD</i>)	15.78±1.61	15.61±1.75	15.14±1.42	15.11±1.64	16.31±1.43	16.36±1.65	.000	.000
Cannabis use within last 12 month* (<i>yes/no</i>)	209/0	95/114	125/60	125/60	91/67	158/0	.000	.000
Other illicit drug use ever* (<i>yes/no</i>)	88/121	88/121	72/113	79/106	39/119	88/60	.002	.002
Nicotine Dependence* (<i>M ± SD</i>)	.69±1.49	.49±1.23	.70±1.37	.56±1.33	.51±1.29	.51±1.27	.356	.885
Socio-economic status* (<i>M ± SD</i>)	5.94±.86	^b	5.91±1.00	^b	5.82±.89	^b	.530	^b
Any disorder (clinical rating, DSM-IV)* (<i>yes/no</i>)	41/158	39/106	35/138	38/98	7/56	22/124	.373	.348

Annotations: *N* = sample size; *M* = mean; *SD* = standard deviation; *details of assessment can be found in supplements; ^a According to one-way ANOVA or χ^2 tests to test for possible differences between groups; ^b parental socio-economic status was assessed at age 14 and used for our analyses

Table 3. Ordinal regression coefficients (β) and p-values for the association between age of first use of cannabis and changes in cannabis use between age 19 and 22 (differences of ESPAD scores) for model (I) and models (II).

<i>Model</i>	<i>Variable</i>	<i>Association with changes in cannabis use</i>	
		<i>β</i>	<i>p-value</i>
<i>Model (I)</i>	Age of first use of cannabis	<i>.180</i>	<i>.002</i>
	Male gender identification	.268	.165
	Recruitment site	- .456 to .19	.05 to .93
<i>Model (II)</i>	Age of first use of cannabis	<i>.195</i>	<i>.011</i>
	Male gender identification	.254	.323
	Recruitment site	- .391 to .617	.179 to .684
	Other illicit drug use ever	<i>.719</i>	<i>.023</i>
	Nicotine dependence	-.073	.400
	Socio-economic status	-.159	.207
	Any disorder (clinical rating, DSM-IV)	.290	.309

Annotations: For model (II), associations between all factors and change in cannabis use are also displayed. β s with a p-value below 0.05 are shown in italic.

Table 4. Ordinal regression coefficients (β) and p-values for the association between CAPE scores at age 19 (*CAPETotal*; *CAPEPosFreq*; *CAPEPosDis*) and changes in cannabis use (differences of ESPAD scores) for model (I) and models (II).

<i>Model</i>	<i>Variable</i>	<i>Association with changes in cannabis use</i>	
Predictor: <i>CAPETotal</i>		<i>β</i>	<i>p</i> -value
<i>Model (I)</i>	<i>CAPETotal</i>	-.003	.668
	Male gender identification	.264	.086
	Recruitment site	-.475 to .285	.10 to .638
<i>Model (II)</i>	<i>CAPETotal</i>	-.001	.910
	Male gender identification	.187	.417
	Recruitment site	-.874 to .188	.038 to .664
	Age of first use of cannabis	.141	.054
	Other illicit drug use ever	-.551	.024
	Nicotine dependence	.024	.740
	Socio-economic status	-.121	.290
	Any disorder (clinical rating, DSM-IV)	-.042	.897
Predictor: <i>CAPEPosFreq</i>			
<i>Model (I)</i>	<i>CAPEPosFreq</i>	-.122	.686
	Male gender identification	.271	.075
	Recruitment site	-.407 to .391	.203 to .737
<i>Model (II)</i>	<i>CAPEPosFreq</i>	-.239	.595
	Male gender identification	.193	.403
	Recruitment site	-.871 to .177	.039 to .682
	Age of first use of cannabis	.140	.055
	Other illicit drug use ever	-.540	.027
	Nicotine dependence	.025	.730
	Socio-economic status	-.124	.277
	Any disorder (clinical rating, DSM-IV)	-.012	.968
Predictor: <i>CAPEPosDis</i>			
<i>Model (I)</i>	<i>CAPEPosDis</i>	-.257	.101
	Male gender identification	.173	.275
	Recruitment site	-.410 to .343	.153 to .269
<i>Model (II)</i>	<i>CAPEPosDis</i>	-.292	.199
	Male gender identification	-.015	.948
	Recruitment site	-.841 to .141	.049 to .747
	Age of first use of cannabis	.143	.059
	Other illicit drug use ever	-.546	.032
	Nicotine dependence	.029	.691
	Socio-economic status	-.129	.267
	Any disorder (clinical rating, DSM-IV)	-.045	.881

Annotations: For model (II), associations between all factors and change in cannabis use are also displayed. β s with a p-value below 0.05 are shown in italic.

Table 5. Regression coefficients (β) and p-values for the association between *CAPETotal*; and current cannabis use for model (I) and models (II) at age 19 and age 22 respectively.

<i>Model</i>	<i>Variable</i>	<i>Association with CAPETotal</i>	
		<i>β</i>	<i>p-value</i>
Predictor: Cannabis use at age 19			
<i>Model (I)</i>	Cannabis use at age 19	.422	.079
	Male gender identification	-3.133	.002
	Recruitment site	-2.490 to .820	.231 to .689
<i>Model (II)</i>			
	Cannabis use at age 19	.055	.887
	Male gender identification	-.779	.576
	Recruitment site	-4.467 to .342	.011 to .951
	Age of first use of cannabis	-.503	.256
	Other illicit drug use ever	1.040	.501
	Nicotine dependence	.351	.426
	Socio-economic status	.467	.509
	Any disorder (clinical rating, DSM-IV)	13.931	.000
Predictor: Cannabis use at age 22			
<i>Model (I)</i>	Cannabis use at age 22	.700	.001
	Male gender identification	-2.728	.006
	Recruitment site	.228-5.241	.009 to .910
<i>Model (II)</i>			
	Cannabis use at age 22	.092	.774
	Male gender identification	-1.085	.439
	Recruitment site	-3.013 to 4.781	.042 to .267
	Age of first use of cannabis	-.275	.502
	Other illicit drug use ever	2.248	.194
	Nicotine dependence	.968	.047
	Socio-economic status	-1.383	.048
	Any disorder (clinical rating, DSM-IV)	13.237	.000

Annotations: For model (II), associations between all factors and change in cannabis use are also displayed. β s with a p-value below 0.05 are shown in italic.

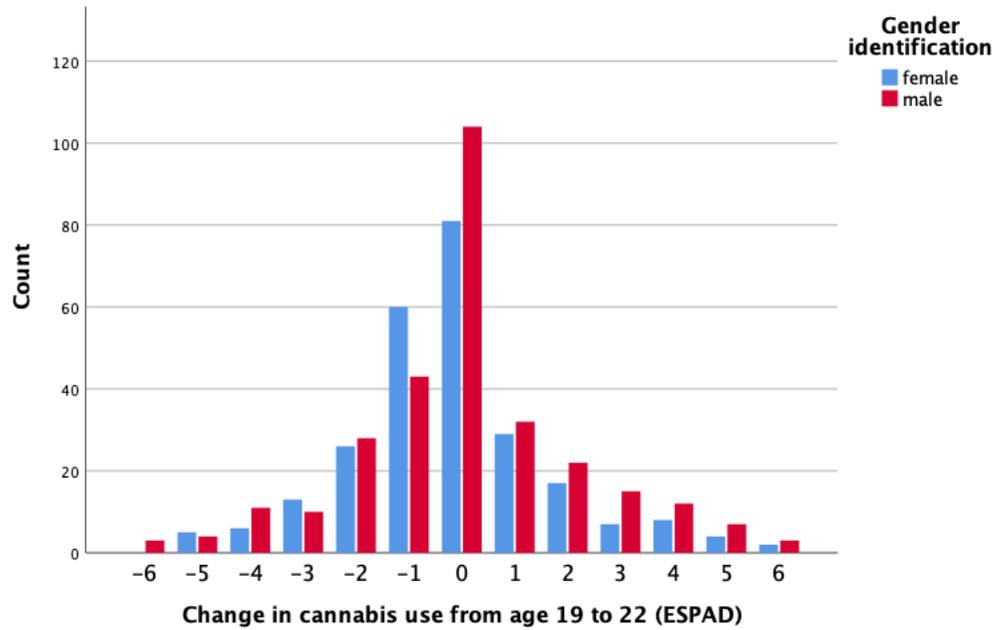


Figure 1. Changes in cannabis use (for last 12 month) from age 19 to age 22 stratified for gender identification. Differences according to ESPAD categories: never (0); once or twice (1); 3-5 times (2); 6-9 times (3); 10-19 times (4); 20-39 times (5); 40 times or more (6).

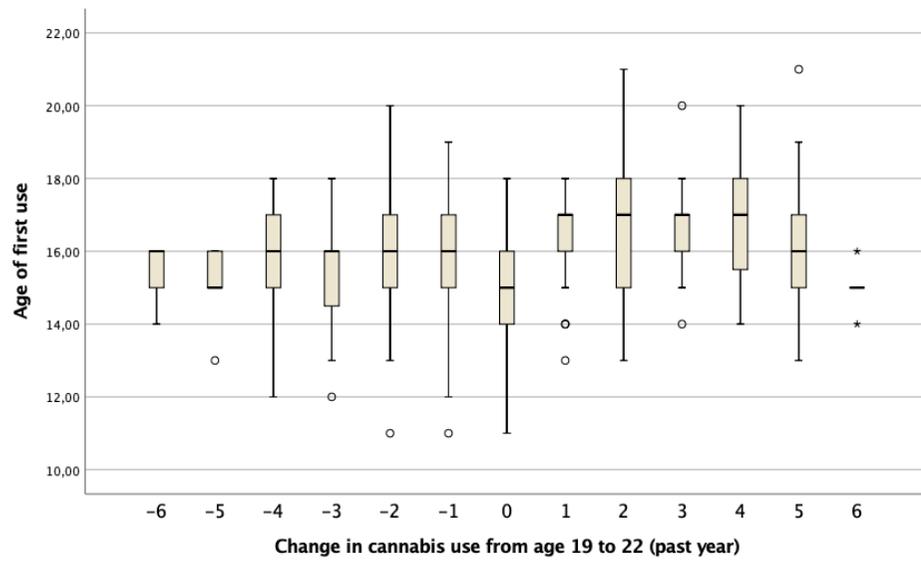


Figure 2. Boxplot for changes in cannabis use within past year and age of first use of cannabis.

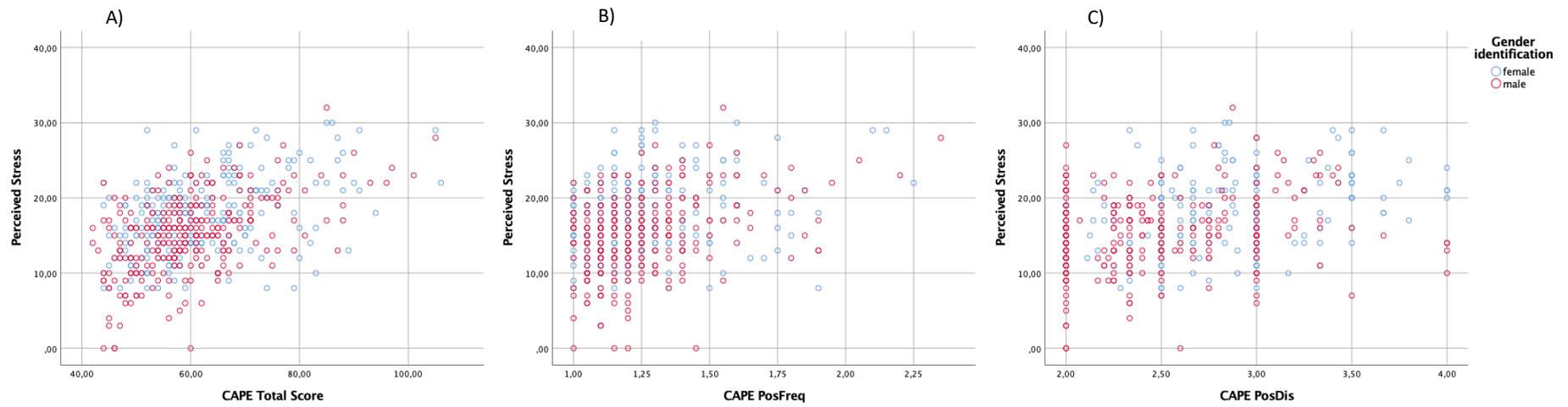


Figure 3. Scatterplot for association of perceived stress (assessed by PSS) and PLEs at age 22 respectively: A) CAPE Total Score B) CAPE *PosFreq*: Frequency of positive dimension C) CAPE *PosDis*: Distress of positive dimension