

1 **Metabolic memory of $\Delta 9$ -tetrahydrocannabinol exposure in pluripotent stem cells**
2 **and primordial germ cells-like cells**

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22 **ABSTRACT**

23 Cannabis, the most consumed psychoactive drug in the world, is increasingly used by pregnant
24 women. However, while cannabinoid receptors are expressed in the early embryo, the impact of
25 phytocannabinoids exposure on early embryonic processes is lacking. Here, we leverage a
26 stepwise in vitro differentiation system that captures early embryonic developmental cascade to
27 investigate the impact of exposure to the most abundant phytocannabinoid, $\Delta 9$ -
28 tetrahydrocannabinol ($\Delta 9$ -THC). We demonstrate that $\Delta 9$ -THC increases the proliferation of naïve
29 mouse embryonic stem cells (ESCs) but not of their primed counterpart. Surprisingly, this
30 increased proliferation, dependent on the CB1 receptor binding, is only associated with moderate
31 transcriptomic changes. Instead, $\Delta 9$ -THC capitalizes on ESCs' metabolic bivalence by increasing
32 their glycolytic rates and anabolic capabilities. A memory of this metabolic rewiring is retained
33 throughout differentiation to Primordial Germ Cell-Like Cells in the absence of direct exposure and
34 is associated with an alteration of their transcriptional profile. These results represent the first in-
35 depth molecular characterization of the impact of $\Delta 9$ -THC exposure on early developmental
36 stages.

37

38 **KEYWORDS**

39 cannabis, $\Delta 9$ -THC, metabolism, embryonic stem cells, primordial germ cells.

40 INTRODUCTION

41 Cannabis is the most widely used psychoactive drug in the world¹. In the United States, an
42 estimated 49.6 million people, roughly 18% of the population, consumed cannabis at least once
43 in 2020, with indications that these numbers will likely increase in the coming years as attitudes
44 and regulations change^{2,3}. In particular, between 7-12% of expecting women report cannabis use,
45 predominantly during the first trimester to alleviate the symptoms of morning sickness⁴⁻⁶. These
46 statistics indicate that a significant number of developing embryos are exposed to cannabis, with
47 limited knowledge of the biological repercussions of such exposure.

48 Among the several hundred unique phytocannabinoids present in *Cannabis sativa*, (-)-
49 trans- Δ^9 -tetrahydrocannabinol (Δ^9 -THC) is chiefly responsible for the psychoactivity of cannabis⁷.
50 As a result, the level of Δ^9 -THC in recreational cannabis has increased over the last 10 years and
51 now commonly accounts for 20% of total compounds⁸. The psychoactive effects of Δ^9 -THC arise
52 from its binding and subsequent activation of the G protein-coupled cannabinoid receptors CB1
53 largely expressed in the central nervous system⁹. In this context, Δ^9 -THC exposure has been
54 shown to durably alter metabolic, transcriptional and epigenetic programs in the brain¹⁰⁻¹³. While
55 over the last decades, significant attention has been paid to Δ^9 -THC's neurological effects, there
56 is also evidence, albeit more limited, of its impact on reproductive functions¹⁴. Data shows CB1
57 expression in the male and the female reproductive tracts, in the pre-implantation embryo and in
58 the placenta^{14,15}. In animal models as well as in humans, exposure to cannabis is associated with
59 reduced fertility, decreased testis weight and sperm count, and impairment of embryo
60 implantation¹⁴. In males, these effects are correlated with an alteration of the sperm transcriptome
61 and epigenome¹⁶⁻¹⁸. Epidemiological evidence also indicates that Δ^9 -THC exposure is associated
62 with long-lasting adverse effects, with exposures in parents affecting the offspring^{13,19}. Despite this
63 accumulating evidence, the molecular impacts and mechanisms of Δ^9 -THC exposure at the
64 earliest stages of development remain to be determined.

65 Progression through states of pluripotency is controlled by metabolic reprogramming in the
66 early mammalian embryo^{20,21}. Accordingly, cultured pluripotent stem cells (PSCs) exhibiting
67 different developmental potentials are marked by specific metabolic signatures, similar to the ones
68 displayed by their in vivo counterparts in the embryo. For instance, mouse embryonic stem cells
69 (ESCs) are naïve PSCs that are functionally equivalent to the inner cell mass (ICM) of the E3.5
70 preimplantation mouse blastocyst²². The extended developmental potential of mouse ESCs is
71 associated with their metabolic bivalence, as these cells rely on both glycolysis and oxidative
72 phosphorylation for energy production. Differentiation of naïve ESCs into primed PSCs such as
73 epiblast-like cells (EpiLCs) is accompanied by an important metabolic shift towards aerobic
74 glycolysis, in link with a highly-proliferative phenotype and a more restricted developmental
75 potential^{20,21,23}. Primordial germ cells (PGCs), the embryonic precursors of gametes in
76 metazoans²⁴, are considered dormant totipotent cells because they possess the unique ability to
77 reacquire totipotency upon fertilization²³. In the mouse embryo, PGCs arise around embryonic day
78 7.5 (E7.5) from a subset of primed PSCs in the epiblast²⁴. Progressive increase in oxidative
79 phosphorylation correlates with the specification and differentiation from epiblast PSCs towards
80 PGCs, a process that can be replicated in vitro by inducing PGC-like cells (PGCLCs) from
81 EpiLCs²⁵. In particular, the extensive metabolic, transcriptional, and epigenetic reprogramming
82 that PGCs undergo during their development has been proposed to be uniquely sensitive to
83 environmental insults, with potential consequences in the offspring²⁶.

84 Here, we deployed this in vitro differentiation system to investigate the impact of $\Delta 9$ -THC
85 exposure on early developmental stages. We demonstrate that exposure of ESCs and EpiLCs to
86 $\Delta 9$ -THC durably alters their metabolome. We reveal that, in the absence of continuous exposure,
87 metabolic memory of $\Delta 9$ -THC is passed onto the PGCLCs stage leading to transcriptional defects
88 in these cells. Together, our findings highlight the role of metabolic reprogramming as a
89 mechanism for early developmental $\Delta 9$ -THC exposure.

90
91 **RESULTS**
92 **$\Delta 9$ -THC induces cellular proliferation of mouse embryonic stem cells but not of mouse**
93 **epiblast-like cells.**

94 To model the impact of early $\Delta 9$ -THC exposure on early embryonic events, we first tested
95 three distinct developmental windows: 1) exposure of ESCs, 2) exposure of EpiLCs and 3)
96 combined ESCs+EpiLCs exposure (Figure 1A). Cells were either exposed to the vehicle (mock)
97 or exposed to $\Delta 9$ -THC in a wide dose range of 10nM-100 μ M, corresponding to the reported
98 physiologically-relevant concentrations of $\Delta 9$ -THC in cannabis users²⁶⁻²⁸.

99 The viability of ESCs exposed to increasing concentrations of $\Delta 9$ -THC for 48h was not
100 significantly altered until the maximal dose of 100 μ M, subsequently serving as a positive control,
101 at which only 13.17% of cells remained alive (Figure 1B, $p < 0.0001$, unpaired T-test). While no
102 significant changes in viability were observed between 10nM and 1 μ M $\Delta 9$ -THC, the number of
103 viable ESCs significantly increased by 1.69, 1.52 and 1.28-fold, respectively, compared to the
104 mock-treated condition (Figure 1C, $p = 0.002$, $p = 0.01$ and $p = 0.03$ for 10nM, 100nM and 1 μ M of $\Delta 9$ -
105 THC, unpaired T-test). To determine whether this increased number of viable cells recovered after
106 $\Delta 9$ -THC exposure was due to higher proliferation, we performed bromodeoxyuridine (BrdU)
107 labeling experiments. Exposed cells were pulsed with BrdU for 30 minutes, and its incorporation
108 in actively dividing cells was measured by flow cytometry. The percentage of BrdU-positive ESCs
109 significantly increased between 10nM and 10 μ M of $\Delta 9$ -THC compared to the mock-treated
110 condition (Figure 1D, $p = 0.01$, $p = 0.001$, $p = 0.05$ and $p = 0.01$ for 10nM, 100nM, 1 μ M and 10 μ M of
111 $\Delta 9$ -THC, unpaired T-test).

112 Next, we derived EpiLCs from unexposed ESCs and performed the same dose-response
113 experiments. Akin to ESCs, $\Delta 9$ -THC exposure in EpiLCs did not significantly alter cellular viability
114 until the dose of 100 μ M (Figure 1E, 11.56% of viable cells, $p < 0.0001$, unpaired T-test). However,
115 contrary to ESCs, $\Delta 9$ -THC exposure in EpiLCs did not significantly increase the number of viable
116 cells nor the percentage of BrdU-positive cells (Figure 1F and Figure 1G). For the dose of 10 μ M
117 of $\Delta 9$ -THC, viable EpiLCs numbers and BrdU-positive EpiLCs decreased compared to the mock-
118 treated condition (Figure 1F, 1.53-fold decrease, $p = 0.0007$, Figure 1G, 1.42-fold decrease,
119 $p = 0.0024$, unpaired T-test).

120 Finally, when continuously exposing ESCs and EpiLCs, cell viability was more significantly
121 and negatively impacted, except at the dose of 100nM of $\Delta 9$ -THC (Figure 1H). Deriving EpiLCs
122 from exposed ESCs and exposing them to $\Delta 9$ -THC for 48h did not significantly affect either their
123 cell number nor their incorporation of BrdU (Figure 1I and Figure 1J), indicating that the increased
124 proliferation observed at the ESCs stage is not carried through the naïve-to-prime transition.

125 Together, the systematic testing of different exposure schemes of $\Delta 9$ -THC in ESCs and
126 EpiLCs revealed that physiologically relevant doses of $\Delta 9$ -THC (10nM-1 μ M) specifically stimulate

127 the proliferation of ESCs, but not of EpiLCs, whether the latter are derived from exposed ESCs or
128 not.

129

130 **Expression of the CB1 receptor does not explain differences in proliferative outcomes.**

131 We next sought to understand the source of variation in proliferative outcomes in response
132 to $\Delta 9$ -THC between naïve mouse embryonic stem cells and primed pluripotent epiblast-like cells.
133 Such differential effects have been previously reported with $\Delta 9$ -THC eliciting the proliferation of
134 neural progenitors²⁹ and of human breast carcinoma cell lines³⁰ but suppressing the proliferation
135 of activated CD4⁺ T cells³¹ and of non-small cell lung cancer cells³². In these studies, the
136 differential expression of cannabinoid receptors at the cell surface was proposed to primarily
137 mediate the variation in cellular outcomes.

138 We therefore first tested whether expression levels of CB1 varied between ESCs and
139 EpiLCs. Western-blot analysis of membrane proteins revealed however that CB1 was expressed
140 at the same levels at the cell surface of both ESCs and EpiLCs (Figure 2A and 2B). We next
141 determined whether the $\Delta 9$ -THC-induced proliferative phenotype in ESCs was due to the
142 engagement of the CB1 cannabinoid receptor. To do so, ESCs were pretreated for 1h with 1 μ M
143 of SR141716 (also known as rimonabant, a specific CB1 blocker³³) then exposed to 100nM or
144 100 μ M of $\Delta 9$ -THC for 48h. Rimonabant pre-treatment did not significantly alter the viability of
145 ESCs compared to conditions exposed to $\Delta 9$ -THC only (Figure 2C) but abolished $\Delta 9$ -THC-
146 induced ESCs increased cell number at 100nM $\Delta 9$ -THC (Figure 2D, 1.53-fold decrease,
147 $p < 0.0001$, when comparing 100nM of $\Delta 9$ -THC +/- 1 μ M of SR141716, unpaired T-test). Notably,
148 SR141716 pre-treatment, while not altering cell viability, reduced cell number compared to control,
149 suggesting a basal role for CB1 in promoting proliferation.

150 Thus, the expression of CB1 at the cell surface does not explain the differential impact of
151 $\Delta 9$ -THC on ESC and EpiLC proliferation even if CB1 engagement is a required event for this effect
152 in ESCs.

153

154 **$\Delta 9$ -THC exposure increases glycolysis in ESCs and EpiLCs.**

155 In the central nervous system, $\Delta 9$ -THC is a known metabolic perturbator which increases
156 bioenergetic metabolism^{12,34}. As mentioned above, the transition of naïve ESCs into the primed
157 state of EpiLCs is accompanied by a switch to glycolysis for energy production^{20,23}. Thus, to
158 capture the impact of $\Delta 9$ -THC at every point of their transition between metabolic states, we used
159 the continuous exposure scheme of ESCs and EpiLCs outlined in Figure 1H-J. Similar to our other
160 exposure schemes, at lower $\Delta 9$ -THC doses, the proliferation of ESCs was observed but not of
161 EpiLCs. We performed these exposures in a wide $\Delta 9$ -THC dose range (10nM-10 μ M) followed by
162 bioenergetics assessment (Figure 3A).

163 First, we assessed the global energy metabolism of exposed cells by measuring the
164 nicotinamide adenine dinucleotide (phosphate) couple ratios (NAD(P)⁺/NAD(P)H) using the WST-
165 1 assay. In ESCs, the ratio of NAD(P)⁺/NAD(P)H significantly increased 1.57, 1.54, 1.29 and 1.38
166 -fold, for 10nM, 100nM, 1 μ M and 10 μ M of $\Delta 9$ -THC, respectively, compared to the mock-treated
167 condition (Figure 3B, $p < 0.0001$, $p < 0.0001$, $p = 0.03$, and $p = 0.0003$ for 10nM, 100nM, 1 μ M and
168 10 μ M of $\Delta 9$ -THC, unpaired T-test). In contrast, no significant increase was observed in
169 NAD(P)⁺/NAD(P)H ratios in exposed EpiLCs (Figure 3B). Consistent with the impact of continuous
170 $\Delta 9$ -THC exposure on EpiLCs viability (Figure 1H), the NAD(P)⁺/NAD(P)H ratios significantly

171 decreased at 10 μ M of Δ 9-THC in EpiLCs (Figure 3B, 59% decrease for 10 μ M of Δ 9-THC
172 compared to the mock-treated condition, $p < 0.0001$, unpaired T-test).

173 Because the elevated NAD(P)⁺/NAD(P)H levels in Δ 9-THC-exposed ESCs could indicate
174 increased mitochondrial activity in the context of oxidative phosphorylation³⁵, we next studied
175 changes in mitochondrial membrane potential of exposed cells using the Mitotracker CMXRos
176 fluorescent dye³⁶. A significant increase in mean fluorescence intensity (MFI) associated with the
177 mitochondrial stain was observed at 100nM of Δ 9-THC in ESCs (Figure 3C, $p = 0.02$, unpaired T-
178 test), indicating that, at this dose, the observed increase in NAD(P)⁺/NAD(P)H could be explained
179 by higher mitochondrial membrane potential. By contrast, no change in EpiLCs mitochondrial
180 activity was detected (Figure 3C), consistent with these cells relying on glycolysis for energy
181 production^{20,23}.

182 Changes in mitochondrial activity in ESCs upon Δ 9-THC exposure, although significant,
183 remained modest and are unlikely to be the sole contributor to the more significant increase in
184 NAD(P)⁺/NAD(P)H upon exposure. Thus, we performed an in-depth analysis of the differential
185 impact of Δ 9-THC on ESCs and EpiLCs bioenergetics by measuring both glycolysis (extracellular
186 acidification rate, ECAR) and mitochondrial respiration (oxygen consumption rate, OCR) using a
187 Seahorse bioanalyzer. At 100nM of Δ 9-THC, the maximal glycolytic capacity of both ESCs and
188 EpiLCs increased significantly (Figure 3D, 15% increase, $p = 0.03$ and 22% increase, $p = 0.03$ for
189 ESCs and EpiLCs, respectively, compared to the mock-treated condition, unpaired T-test). In both
190 cell types, a significant decrease in glycolytic capacity was observed at 10 μ M of Δ 9-THC (Figure
191 3D, 39.8% reduction, $p = 0.0006$ and 44.8% reduction, $p = 0.0001$, for ESCs and EpiLCs,
192 respectively, compared to the mock-treated condition, unpaired T-test). Of note, the maximal
193 glycolytic capacity of EpiLCs in the untreated condition was higher than the one of ESCs, in
194 agreement with their metabolic shift towards aerobic glycolysis (Figure 3D, 7.88% higher ECAR
195 rate in mock-treated EpiLCs compared to mock-treated ESCs, $p = 0.03$, unpaired T-test). As a
196 consequence, Δ 9-THC exposure significantly impacted more glycolysis in EpiLCs than ESCs,
197 both in basal capacity and upon mitochondrial inhibition by oligomycin (Supplementary Figure 1A
198 and Figure 1B). In addition, at 100nM of Δ 9-THC, the maximal respiratory capacity of ESCs was
199 significantly increased compared to the mock-treated condition (Figure 3E, 21.8% increase,
200 $p = 0.03$, unpaired T-test). This increase was observed only for the maximal respiratory capacity of
201 ESCs, but not for basal respiration, nor for ATP-linked respiration (Supplementary Figure 1C),
202 suggesting that Δ 9-THC impact on mitochondrial respiration does not support increased energetic
203 production. In agreement with EpiLCs metabolic shift towards a glycolytic phenotype, increasing
204 doses of Δ 9-THC did not alter their maximal respiratory capacity (Figure 3E), nor their global
205 oxygen consumption rate (Supplementary Figure 1D). In both cell types, a significant decrease in
206 oxygen consumption rate was observed at 10 μ M of Δ 9-THC (Figure 3E and Supplementary Figure
207 1C and Figure 1D).

208 Together, our analysis of cellular bioenergetics following Δ 9-THC exposure showed an
209 increased glycolytic rate in ESCs that was also observed in EpiLCs. However, the increased
210 oxygen consumption and the associated increase in mitochondrial activity were observed only in
211 ESCs following exposure to 100nM of Δ 9-THC, likely for the oxidization of the accumulating
212 pyruvate generated from glycolysis.

213 **Δ9-THC-induced increase in glycolysis supports anabolism and ESCs proliferation**

214 Because our data indicated that the impact of Δ9-THC exposure on stem cells' bioenergetics
215 did not result in greater ATP production, we next sought to characterize the global metabolic
216 impact of Δ9-THC in these cells. ESCs and EpiLCs were continuously exposed to 100nM Δ9-THC
217 and intracellular metabolites were detected and quantified by mass spectrometry (Figure 4A-E).
218 To explore the metabolic signatures in the different samples, we performed a global principal
219 component analysis (PCA) (Figure 4B). All samples clustered in well-defined groups of replicates,
220 both by cell type on the first principal component (accounting for 65.81% of the variation) and by
221 Δ9-THC exposure on the second principal component (accounting for 20.83% of the variation). Of
222 the 126 metabolites detected in ESCs, 39 were significantly upregulated (Figure 4C and
223 Supplementary Figure 2A) and only two metabolites – NADPH and Adenine – were significantly
224 downregulated. Of the 138 metabolites detected in EpiLCs, 95 were significantly upregulated
225 (Figure 4C and Supplementary Figure 2B) and only one metabolite – NADPH – was significantly
226 downregulated. In agreement with the PCA, the overlap of over-expressed metabolites in
227 response to Δ9-THC exposure was important between the two stem cell populations (Figure 4C,
228 accounting for 79.49% and 32.63% of all upregulated metabolites in ESCs and EpiLCs,
229 respectively). The functional interpretation of the significantly upregulated metabolites confirmed
230 the Δ9-THC-associated increase in energy metabolism in the two stem cell populations. Indeed,
231 amongst the 25 metabolic pathways upregulated, pyruvate metabolism and glycolysis were
232 detected in both ESCs and EpiLCs (Figure 4D and Figure 4E, respectively). Increased
233 mitochondrial respiration was also seen in ESCs with the enrichment of (ubi)quinone metabolism,
234 indicating an increased synthesis of ubiquinone that serves as an electron carrier in oxidative
235 phosphorylation. Of note, metabolite measurements showed that the ratio of glutathione in its
236 reduced to oxidated form (GSH/GSSG) was unchanged in both stem cell types in response to Δ9-
237 THC (Supplementary Figure 2C), suggesting that the increased mitochondrial respiration does not
238 cause an overt elevation of oxidative stress. Importantly, and in agreement with the PCA, in both
239 ESCs and EpiLCs, Δ9-THC exposure elicited an increase in metabolic pathways that feed
240 anabolic reactions, in particular contributing to the synthesis of amino acids (tyrosine, tryptophan,
241 arginine, alanine, valine, (iso)leucine, etc.), nucleotides ("Pyrimidine metabolism", "Purine
242 metabolism"), NAD(P)+ ("Nicotinate and nicotinamide metabolism") and fatty acids ("Butanoate
243 metabolism") (Figure 4D and Figure 4E).

244 Extensive metabolic profiling of ESCs and EpiLCs upon Δ9-THC exposure thus indicated
245 that the increased glycolytic rates in both stem cell populations, rather than provoking an increased
246 production of energy under the form of ATP, participated in increased anabolism. Such increased
247 anabolism could explain the proliferation observed in ESCs upon Δ9-THC exposure. To test this
248 hypothesis, we exposed ESCs to 100nM of Δ9-THC for 48h as above but 24h before the harvest,
249 cells were exposed to 10mM of 2-deoxyglucose (2-DG), an inhibitor of glycolysis³⁷. Despite
250 increasing the energy stress (Supplementary Figure 3), inhibition of glycolysis by 2-DG did not
251 significantly impact viability over this shorter time frame and at this concentration (Figure 4F).
252 Importantly, glycolytic inhibition by 2-DG abrogated the Δ9-THC-induced increase in both cell
253 number and NAD(P)+/NAD(P)H levels (Figure 4G and Figure 4H, 1.39-fold reduction and
254 $p < 0.0001$ and 1.68-fold reduction and $p = 0.0064$, respectively, when comparing 100nM of Δ9-THC
255 +/- 10mM 2-DG, unpaired T-test). Thus, exposure to Δ9-THC increases anabolism in both ESCs
256 and EpiLCs, however, this increased anabolism only supports cellular proliferation in ESCs.

257 **Δ 9-THC exposure is associated with the upregulation of genes involved in anabolic**
258 **pathways in ESCs but not in EpiLCs.**

259 Our data shows that Δ 9-THC exposure increases anabolic pathways in both ESCs and
260 EpiLCs and that this causes the proliferation of ESCs but not of EpiLCs. We thus next examined
261 whether this differential impact of Δ 9-THC on ESCs and EpiLCs was mirrored by a change in
262 these cells' transcriptomes. To this aim, we performed RNA-sequencing (RNA-seq) on ESCs and
263 EpiLCs continuously exposed to 100nM Δ 9-THC or to the vehicle control (Figure 5A).

264 Unsupervised exploration of the global transcriptome by PCA revealed that the vast majority
265 of data variation could be attributed to the cell type (PC1, accounting for 98% of the variation)
266 rather than to Δ 9-THC exposure (PC2, accounting for 1% of the variation, Figure 5B). This
267 suggests that Δ 9-THC exposure only moderately impacts ESC and EpiLC transcriptomes. In
268 agreement, we identified a low number of differentially expressed genes (DEGs) in both ESCs
269 and EpiLCs (Figure 5C and Figure 5D, respectively). In ESCs, only 12 genes were significantly
270 upregulated with a $\log_2(\text{fold-change}) > 0.5$ and only 9 were significantly downregulated at the same
271 threshold (Figure 5C, significance corresponds to adjusted $p\text{-value} \leq 0.05$). More genes were
272 differentially expressed when looking at lower fold-changes ($|\log_2(\text{fold-change})| > 0.25$, Figure 5C),
273 confirming that the magnitude of transcriptional effects due to Δ 9-THC exposure is moderate. This
274 low transcriptional impact following Δ 9-THC exposure was also observed in EpiLCs (Figure 5D).
275 Nevertheless, gene ontologies (GO) associated with Δ 9-THC-induced DEGs revealed the
276 biological significance of these low transcriptional changes (Figure 5E). In particular, GO terms
277 associated with metabolic pathways involved in anabolism were significantly over-represented for
278 upregulated genes in ESCs following Δ 9-THC exposure (Figure 5E), such as: "Cellular aromatic
279 compound metabolic process", "Cellular nitrogen compound biosynthetic process",
280 "Organonitrogen compound metabolic process". This suggests that the glycolytic rewiring elicited
281 by Δ 9-THC exposure in ESCs has some transcriptional support. Indeed, when performing joint
282 pathway integration between our transcriptomics data and our targeted metabolomics³⁸, we
283 observed that Δ 9-THC-induced perturbed genes and metabolites were associated with the
284 observed anabolic effects (Figure 5F). In contrast, GO terms associated with metabolism were not
285 found within the upregulated DEGs in EpiLCs. However, several GO terms relating to alterations
286 in cellular components were enriched in EpiLCs (Figure 3E), such as: "Organelle organization",
287 "Cellular component organization or biogenesis", "Microtubule-based process". This indicates that
288 Δ 9-THC exposure significantly upregulated genes in EpiLCs that impact organelles structure,
289 integrity and position, in agreement with several reports in the literature^{39,40}.

290 Together, our analysis of ESCs and EpiLCs transcriptomes reveals a difference in the
291 response of these stem cell populations to Δ 9-THC exposure: the transcriptional alterations
292 observed in ESCs supported their increased anabolism and proliferation, whereas changes in
293 EpiLCs gene expression did not correlate with their metabolic changes.

294

295 **Proliferation of Primordial Germ Cell-Like Cells stemming from prior Δ 9-THC exposure.**

296 PGCs display a distinct transcriptomic and metabolic profile compared to their cellular
297 precursors that are recapitulated in vitro during the differentiation of ESCs into EpiLCs and then
298 of EpiLCs into PGCLCs. Thus, we asked whether the metabolic alterations observed in ESCs and
299 EpiLCs could lead to an altered differentiation program in PGCLCs. To this aim, we continuously
300 exposed ESCs and EpiLCs to a Δ 9-THC dose range of 10nM-1 μ M (or mock control), before

301 changing to a $\Delta 9$ -THC-free media and inducing PGCLCs differentiation (Figure 6A). In particular,
302 we took advantage of ESCs that harbor two fluorescent reporters for germline markers,
303 *Blimp1:mVenus* and *Stella:CFP*⁴¹. Thus, the induction efficiency of PGCLCs within 5-days old
304 embryoid bodies can be detected by monitoring the fluorescence associated with each cell in flow
305 cytometry, allowing for the determination of a double-negative population (DN), a single-positive
306 population (SP) wherein *Blimp1:mVenus* is expressed and a double-positive population (DP)
307 expressing both *Blimp1:mVenus* and *Stella:CFP*, which represents the true specified PGCLC
308 population.

309 We first measured the impact of ESCs + EpiLCs $\Delta 9$ -THC exposure on PGCLC induction
310 efficiency. Flow analyses revealed a dose-dependent increase in the induction efficiency of SP
311 and DP cell populations (Figure 6B). Specifically, at 100nM $\Delta 9$ -THC, a significant decrease in DN
312 was observed, with a corresponding significant increase of 1.14-fold in SP and of 1.64-fold in DP
313 cells (Figure 6C, $p=0.0002$, $p=0.05$, and $p<0.0001$ for 100nM of $\Delta 9$ -THC in DN, SP and DP
314 populations respectively compared to the mock-treated condition, unpaired T-test). To determine
315 if the increased proportion of PGCLCs generated from exposed precursors was due to higher
316 proliferative kinetics, we performed a proliferation tracing assay⁴². The tracing dye was added to
317 the cells on the day of aggregates formation, and fluorescence attenuation due to cell division was
318 measured in each subpopulation on day 5. At 100nM $\Delta 9$ -THC, a smaller proportion of DN cells
319 underwent two or three mitotic divisions compared to the control (Figure 6D, 1.14-fold fewer cells
320 and 1.12-fold fewer cells, $p=0.05$ and $p=0.04$ for 2 divisions and 3 divisions, respectively, unpaired
321 T-test). In parallel, for the same dose, a significantly higher proportion of SP and DP cells
322 underwent three mitotic divisions compared to the control (Figure 6D, 1.24-fold and 1.11-fold,
323 $p=0.03$ and $p=0.0035$, for 3 divisions in SP and DP cells, compared to the control, unpaired T-
324 test). These results, therefore, indicate that the higher number of PGCLCs observed upon $\Delta 9$ -
325 THC exposure originates from their increased proliferation during their specification and
326 differentiation.

327 Thus, $\Delta 9$ -THC causes an alteration of the developmental kinetics that PGCLCs normally
328 undergo, even in the absence of direct exposure.

329

330 **PGCLCs derived from $\Delta 9$ -THC-exposed cells present an altered metabolism and** 331 **transcriptome**

332 Since exposure to $\Delta 9$ -THC prior to their specification increased the number of PGCLCs and
333 ESCs and PGCLCs share similar metabolic programs^{20,23}, we next sought to characterize their
334 associated metabolic and transcriptional changes. We, therefore, assessed the impact of
335 exposure of ESCs + EpiLCs to 100nM $\Delta 9$ -THC on PGCLCs metabolism (Figure 7A).

336 First, NAD(P)⁺/NAD(P)H assessment revealed a modest but significant 1.17-fold increase
337 in NAD(P)⁺/NAD(P)H ratio in whole day 5 embryoid bodies deriving from exposed ESCs + EpiLCs
338 compared to those deriving from mock-treated cells (Figure 7B, $p=0.01$, unpaired T-test). To
339 garner cell type-specific information on whether these metabolic changes were related to
340 mitochondrial activity and the differentiation of PGCLCs, we assessed the mitochondrial
341 membrane potential of each subpopulation in day 5 embryoid bodies. Embryoid bodies were
342 incubated with Mitotracker CMXRos³⁶, and dissociated and analyzed by flow cytometry. The MFI
343 associated with the mitochondrial stain was then measured in each subpopulation (Figure 7C). A
344 significant increase in MFI was observed in DN, SP and DP populations deriving from exposed

345 ESCs +EpiLCs compared to those deriving from mock controls (Figure 7C, 1.17, 1.16, 1.23 -fold,
346 $p=0.006$, $p=0.05$ and $p=0.01$ for DN, SP and DP, respectively, unpaired T-test). These results
347 indicate that the metabolic changes induced by $\Delta 9$ -THC prior to PGCLCs induction and
348 differentiation are not reset during the profound reprogramming that PGCLCs undergo.

349 Because our results indicated a sustained impact of $\Delta 9$ -THC beyond the period of direct
350 exposure, we further examined PGCLCs by performing a transcriptomic analysis. In particular,
351 day 5 embryoid bodies deriving from ESCs + EpiLCs, either exposed to 100nM of $\Delta 9$ -THC or
352 mock-exposed, were sorted and the total RNA of DP subpopulations, representing true PGCLCs,
353 was analyzed by RNA-seq (Figure 7A). Unsupervised analysis of the global transcriptome in DP
354 PGCLCs by PCA delineated a transcriptional signature of prior $\Delta 9$ -THC exposure (Figure 7D, PC1
355 accounting for 59% of the variance and PC2 accounting for 24% of variance). Volcano plot of
356 DEGs between DP PGCLCs deriving from mock- or 100nM $\Delta 9$ -THC-exposed ESCs and EpiLCs
357 revealed that most of the significant transcriptional change was towards downregulation rather
358 than upregulation (Figure 7E, 11 genes were significantly upregulated whereas 97 were
359 significantly downregulated, $|\log_2(\text{fold-change})|>0.25$ and adjusted $p\text{-value}\leq 0.05$). Despite the low
360 number of upregulated DEGs, the functional annotation of their associated GO terms showed that
361 all terms enriched corresponded to metabolic processes involved in oxidative phosphorylation
362 (Figure 7F, “Aerobic electron transport chain”, “Mitochondrial respiratory chain complex I
363 assembly”, “Electron transport couple proton transport”). Thus, our data indicate that the metabolic
364 changes induced by exposure to $\Delta 9$ -THC prior to PGCLCs specification are retained through
365 transcriptional reprogramming. Importantly, while our results show that pre-specification $\Delta 9$ -THC
366 exposure increases PGCLCs number and mitochondrial activity, the functional annotation of GO
367 terms associated with downregulated DEGs suggests degradation of PGCLCs quality. Indeed,
368 and reminiscent of GO terms observed in EpiLCs, several GO terms relating to alterations in
369 structural cellular components (“Anatomical structure morphogenesis”, “Cellular anatomical
370 entity”), and in particular the interface with the extracellular environment (“External encapsulating
371 structure organization”, “Membrane”, “Cell periphery”, “Extracellular region”, “Extracellular space”,
372 “Extracellular matrix structural constituent”) were enriched (Figure 7G). Furthermore, GO terms
373 associated with cell adhesion and junction (“Cell adhesion”, “Cell migration”, “Collagen metabolic
374 process”, “Cell junction”, “Anchoring junction”, “Collagen trimer”) were also enriched in
375 downregulated genes.

376 Together, our data show that $\Delta 9$ -THC exposure in ESCs and EpiLCs durably alters their
377 metabolome and that these changes are carried through PGCLCs specification and differentiation,
378 leading to an alteration of PGCLCs transcriptional program (Figure 8).

379

380 **DISCUSSION**

381 With greater social acceptance and legalization, cannabis use has increased worldwide^{1,3}.
382 Yet, the impact of such heightened use on reproductive functions, and in particular, on the earliest
383 developmental stages are not well understood. Cannabis use directly alters adult male fertility and
384 causes abnormal embryo implantation¹⁴. Using a well-characterized in vitro model of early
385 embryonic differentiation events culminating into the differentiation of PGCLCs, our study is the
386 first to shed light on the impact of $\Delta 9$ -THC at these stages which unfold during the first trimester
387 in humans⁴⁻⁶.

388 Our data revealed the differential effects of $\Delta 9$ -THC on naïve and primed pluripotent stem
389 cells, respectively represented by ESCs and EpiLCs. In particular, exposure to $\Delta 9$ -THC increased
390 ESCs proliferation which was in a similar range to what has been previously reported for human
391 breast carcinoma cell lines (about 30-50% between 10nM and 1 μ M of $\Delta 9$ -THC)³⁰. Differential
392 expression and use of cannabinoid receptors on the surface of exposed cells have been shown
393 to correlate with $\Delta 9$ -THC proliferative phenotypes²⁹⁻³². However, our experiments demonstrated
394 that despite being required for $\Delta 9$ -THC-induced proliferation in ESCs, CB1 expression did not
395 significantly differ at the surface of ESCs and EpiLCs.

396 Because $\Delta 9$ -THC is a known perturber of mitochondrial function as previously described
397 in the central nervous system^{12,34}, we studied the metabolic impact of its exposure in ESCs and
398 EpiLCs. Our data indicate that, at 100nM, $\Delta 9$ -THC exposure increased the glycolytic rate in both
399 ESCs and EpiLCs. Bioenergetics analyses and metabolite measurements showed that this
400 increased glucose metabolism did not support increased energy production in the mitochondria,
401 but rather, that it led to the accumulation of metabolic intermediates used in anabolic reactions for
402 the synthesis of amino acids, nucleotides, and lipids. Thus, the metabolic signatures associated
403 with $\Delta 9$ -THC exposure are reminiscent of those inherently occurring during naïve-to-prime
404 transition, during which increased aerobic glycolytic rates feed anabolic reactions ultimately
405 fueling proliferation²⁴. We verified this model by testing the requirement of increased glycolysis to
406 support proliferation and indeed observed that ESCs proliferation upon $\Delta 9$ -THC exposure is
407 abrogated in the presence of the glycolytic inhibitor 2-DG.

408 Transcriptomic analyses revealed that the metabolic reprogramming induced by $\Delta 9$ -THC
409 exposure in ESCs was transcriptionally encoded, with increased expression of genes involved in
410 anabolic pathways. In contrast, functional annotations of DEGs in EpiLCs did not show such
411 transcriptional control of increased anabolism. Comparing the outputs of the metabolomic and
412 transcriptomic analyses (i.e. PCA plots and volcano plots), the impact of $\Delta 9$ -THC at these early
413 stages seems to be primarily metabolic, although the moderate effects on the transcriptome
414 appear to support the metabolic outcome as revealed by our integrated analysis (Figure 5F).
415 Together, we propose that $\Delta 9$ -THC exposure elicits a reprogramming of ESCs that (1) coaxes
416 them to rely more on aerobic glycolysis, (2) drives anabolic pathways, and therefore (3) leads to
417 their proliferation. In EpiLCs, the impact of $\Delta 9$ -THC exposure is not sufficient to override the
418 cellular and metabolic programs of these already highly proliferative cells that are fully reliant on
419 aerobic glycolysis (Figure 8).

420 Finally, we assessed the impact of $\Delta 9$ -THC exposure in ESCs and EpiLCs on the
421 differentiation of PGCLCs. Our data indicate that at the physiologically relevant dose of 100nM of
422 $\Delta 9$ -THC, a significant increase in PGCLCs was observed. In particular, during PGCLCs
423 differentiation, metabolic reprogramming and increased oxidative phosphorylation play a critical
424 role in the reacquisition of an extended developmental potential^{20,23}. Thus, we investigated
425 whether the metabolic alterations observed in ESCs and EpiLCs upon $\Delta 9$ -THC exposure could
426 be carried through PGCLCs differentiation. Metabolic characterization revealed that PGCLCs
427 arising from exposed ESCs and EpiLCs showed increased mitochondrial respiration. Thus, in the
428 absence of continuous exposure, $\Delta 9$ -THC still has lasting consequences on the metabolome of
429 embryonic germ cells. A recent study in drosophila reported that nutrient stress induces oocytes
430 metabolites remodeling that drives the onset of metabolic diseases in the progeny⁴³. This indicates
431 that non-DNA-associated factors, such as germline metabolites, can act as factors of inheritance.

432 Similarly, we show here that exposure to $\Delta 9$ -THC remodels ESCs and EpiLCs metabolome and
433 that a metabolic memory of this exposure is retained during PGCLCs differentiation (Figure 8). In
434 addition to metabolic remodeling, we show that the PGCLCs transcriptome is also altered. In
435 particular, despite proliferation and a higher number of cells, the number of DEGs that were
436 downregulated in PGCLCs deriving from $\Delta 9$ -THC-exposed ESCs and EpiLCs suggests a general
437 degradation of PGCLCs' homeostasis. Functional annotation further indicated that these
438 downregulated genes are related to structural cellular components, to the interaction with the
439 extracellular environment and, specifically, to cell adhesion and junction. During the development
440 of the central nervous system, perinatal $\Delta 9$ -THC exposure has also been associated with
441 alteration in cell adhesion, with an impact on neuronal interactions and morphology⁴⁴⁻⁴⁶. Cell-cell
442 adhesion is crucial in PGCs' formation both in cell culture systems⁴⁷ as well as in vivo where it
443 controls PGCs motility during their migration towards the developing somatic gonad⁴⁸. Our results
444 thus suggest that exposure to $\Delta 9$ -THC prior to specification affects embryonic germ cells'
445 transcriptome and metabolome, with potentially adverse consequences on cell-cell adhesion that
446 could impact their normal development in vivo.

447 Together, our studies reveal a moderate but significant impact of $\Delta 9$ -THC exposure on early
448 embryonic processes. Our work also highlights the importance of the metabolic remodeling
449 induced by $\Delta 9$ -THC and its potential role as a driver of exposure memory through differentiation
450 stages.

451 **METHODS**

452

453 **Data availability**

454 The RNA sequencing data from this study is made available at the Gene Expression
455 Omnibus (GEO) under the following accession number GSE226955. All other data are available
456 upon request.

457

458 **Cell culture and PGCLCs model**

459 Mouse ESCs containing the two fluorescent reporters *Blimp1::mVenus* and *Stella::ECFP*
460 (BVSC cells) were described previously⁴¹. The female BVSC clone H18 was kindly provided by
461 Mitinori Saitou and cells were seeded on coated plates (Poly-L-ornithine [0.001%; A-004-C;
462 Sigma-Aldrich] and laminin [300ng/mL; L2020; Sigma-Aldrich]) in 2i+LIF culture medium (N2B27
463 Media, CHIR99021 [30 μ M; NC9785126; Thermo Fisher], PD0325901 [10 μ M; NC9753132;
464 Thermo Fisher], ESGRO[®] Leukemia Inhibitory Factor (LIF) [1,000 U/mL, ESG1106; Sigma-
465 Aldrich]) for 48h. Differentiation of ESCs to EpiLCs was performed by seeding the cells on Human
466 Plasma Fibronectin (HPF)-coated plates [16.7 μ g/mL; 33016015; Thermo Fisher] in the presence
467 of EpiLC induction medium (N2B27 medium containing activin A [20ng/mL; 50-398-465; Thermo
468 Fisher], basic fibroblast growth factor (bFGF) [12ng/mL; 3139FB025; R&D Systems], and
469 KnockOut Serum Replacement [KSR, 1%; Thermo Fisher]). For PGCLCs induction, 44h EpiLCs
470 were harvested using TrypLE[™] Select (1X) (Thermo Fisher) and seeded either in 96-wells plate
471 (Nunclon Sphera, Thermo Fisher) or in EZsphere plates for large-scale induction (Nacalai) in the
472 presence of GK15 medium (Glasgow's Minimal Essential Medium [GMEM, 11710035, Thermo
473 Fisher] supplemented with 15% KSR, 0.1 mM Minimal Essential Medium Nonessential Amino
474 Acids [MEM-NEAA], 1 mM sodium pyruvate, 0.1 mM 2-mercaptoethanol, 100U/mL penicillin,
475 0.1mg/mL streptomycin, and 2 mM L-glutamine in the presence of bone morphogenetic protein 4
476 [BMP4; 500ng/mL; 5020-BP-010/CF; R&D Systems], LIF, stem cell factor [SCF; 100ng/mL; 50-
477 399-595; R&D Systems], bone morphogenetic protein 8b [BMP8b; 500ng/mL; 7540-BP-025; R&D
478 Systems], and epidermal growth factor [EGF; 50ng/mL; 2028EG200; R&D Systems]. Cells were
479 culture for 5d before collection, dissociation of embryoid bodies and downstream experiments. All
480 cells were cultured in a humidified environment at 37°C under 5% CO₂.

481

482 **Δ 9-THC exposures**

483 To assess the impact of Δ 9-THC exposure on the developmental trajectory of PGCLCs,
484 three exposure schemes were tested: 1) ESCs exposure only, 2) EpiLCs exposure only, and 3)
485 ESCs+EpiLCs exposure. The stock of Δ 9-THC was obtained from the National Institute on Drug
486 Abuse (7370-023 NIDA; Bethesda, MD). The stock was adjusted to a concentration of 200mM
487 diluted in ethanol, aliquoted and stored according to the DEA's recommendations. The dose range
488 of 0-100 μ M was determined based on Δ 9-THC physiological measurements in the blood, plasma,
489 and follicular fluid²⁶⁻²⁸. For each exposure, new aliquots of Δ 9-THC were diluted in ESCs or EpiLCs
490 culture media in coated tubes (Sigmacote, Sigma Aldrich). Exposure was performed for 48h.
491 Solubility tests were performed and ethanol was added to reach the same amount for each Δ 9-
492 THC concentration (0.05% ethanol). Vehicle control corresponded to 0.05% ethanol added to the
493 respective culture media for ESCs or EpiLCs. All experiments performed are authorized under
494 DEA registration number RA0546828.

495

496 **PGCLCs induction efficiency**

497 Changes in PGCLCs induction were calculated by flow cytometry. Practically, d5 aggregates
498 were harvested, dissociated using TrypLE™ Select, and resuspended in fluorescence-activated
499 cell sorting (FACS) buffer (1×Dulbecco's phosphate buffered saline [DPBS], 1% BSA, 1 mM
500 EDTA, 25 mM HEPES). Quantification of subfractions of double-positive PGCLCs
501 (Blimp1::mVenus+ and Stella::ECFP+), single-positive (Blimp1::mVenus+) and double-negative
502 cells was performed on a BD Biosciences LSRII (UCLA BSCRC Flow Cytometry Core). Cells were
503 initially identified by forward- and side-scatter gating, with back-gating used to verify the accuracy
504 by which target cell populations were identified. Cell populations of interest were identified by 2-D
505 plots displaying the parameter of interest, using embryoid bodies cultured in GK15 medium without
506 added cytokines and BMPs as a negative control. Manually defined gates as well as quadrants
507 were used, as indicated. The FlowJo software was used to calculate percent of induction and
508 generate the associated graphs (version 10, FlowJo, LLC).

509

510 **Cell viability and proliferation studies**

511 The viability and viable cell count of ESCs and EpiLCs was calculated using Trypan blue
512 (0.4%, Thermo Fisher) on a Countess II FL Automated Cell Counter (Thermo Fisher). For BrdU
513 incorporation studies, cells were permeabilized, fixed, and stained using the BrdU Flow Kit
514 (PerCP-Cy™5.5 Mouse anti-BrdU, BD Biosciences) before analysis by flow cytometry on a BD
515 Biosciences LSRII (UCLA BSCRC Flow Cytometry Core). Quantification of PGCLCs proliferation
516 was performed using CellTrace™ Yellow (5µM, added at the induction, Thermo Fisher), which
517 binds to intracellular amines after diffusing through cell membranes. The overall fluorescent signal,
518 that gradually decreases as cell division occurs, reflects the number of cell divisions occurring and
519 was measured on a BD Biosciences LSRII (UCLA BSCRC Flow Cytometry Core). The FlowJo
520 software was used to calculate percent of induction, numbers of cell division and generate the
521 associated graphs (version 10, FlowJo, LLC).

522

523 **CB1 antagonist treatment**

524 To block the effects of Δ^9 -THC on the cannabinoid receptor CB1, ESCs were plated on 48-
525 well plate and were pre-treated with 1µM of SR141716/Rimonabant (SML0800, Sigma Aldrich) for
526 1h before being exposed to the dose range of Δ^9 -THC, as above. After 24h incubation, this
527 procedure was repeated and cells were harvested after 48h total incubation. The viability and
528 viable cell count was calculated using Trypan blue (0.4%, Thermo Fisher) on a Countess II FL
529 Automated Cell Counter (Thermo Fisher). The concentration of 1µM of Rimonabant was chosen
530 based on previous experiments³⁹ and did not impact cell viability nor cell number on its own.

531

532 **Western blotting**

533 Membrane proteins were extracted from cell pellets using the Mem-PER™ Plus Membrane
534 Protein Extraction Kit (89842, Thermo Fisher) according to the manufacturer's protocol. Western
535 blotting was performed with 25µg of protein extracts. The immunodetection was assessed using
536 primary antibodies targeting CB1 (101500, Cayman Chemical) or β -actin (3700, Cell Signaling
537 Technology) as loading control. Horseradish peroxidase (HRP)-conjugated secondary antibodies
538 were used for chemiluminescence detection (Amersham).

539

540 **WST-1 assay**

541 The colorimetric assay WST-1 was used according to the manufacturer's instructions
542 (Roche). The tetrazolium salt WST-1 is reduced by mitochondrial dehydrogenases to formazan
543 using NAD(P)H as co-substrates. Thus, the quantity of formazan is directly proportional to
544 NAD(P)⁺.

545

546 **Mitochondrial activity**

547 Staining for mitochondria was performed by incubating cells at 37°C with 250nM MitoTracker
548 CMXRos (M7512, Thermo Fisher) for 30min³⁶. Cells were washed and analyzed by flow cytometry
549 on a BD Biosciences LSR II (UCLA BSCRC Flow Cytometry Core). The FlowJo software (version
550 10, FlowJo, LLC) was used to calculate the mean fluorescence intensity (MFI) corresponding to
551 the average fluorescence intensity of each event of the selected cell population within the chosen
552 fluorescence channel associated to MitoTracker CMXRos.

553

554 **Seahorse experiments**

555 The extracellular acidification rate (ECAR) and the oxygen consumption rate (OCR) are
556 indicative of glycolysis and mitochondrial respiration, respectively. A total of 10×10³ ESCs and
557 8×10³ EpiLCs were seeded on Seahorse XF96 plates (101085-004, Agilent Technologies) and
558 exposed to increasing doses of Δ9-THC for 48h. On the day of the assay, cells were washed with
559 assay medium (unbuffered DMEM assay medium [5030, Sigma Aldrich] supplemented with
560 31.6mM NaCl, 3mg/L phenol red, 5mM HEPES, 5mM glucose, 2mM glutamine and 1mM sodium
561 pyruvate). For OCR measurement, compounds were injected sequentially during the assay
562 resulting in final concentrations of 2μM oligomycin, 0.75μM and 1.35μM FCCP, 1μM rotenone and
563 2μM antimycin. ECAR was measured in parallel. The measured quantities were normalized to the
564 protein content as measured by a BCA quantitation (23227, Thermo Fisher).

565

566 **Mass spectrometry-based metabolomics analysis**

567 To extract intracellular metabolites, cells were rinsed with cold 150mM ammonium acetate
568 (pH7.3) then incubated with 80% ice-cold methanol supplemented with 10 nmol D/L-norvaline for
569 1h. Following resuspension, cells were pelleted by centrifugation (15,000g, 4°C for 15min). The
570 supernatant was transferred into a glass vial and metabolites were dried down under vacuum then
571 resuspended in 70% acetonitrile. Mass spectrometry analysis was performed at the UCLA
572 Metabolomics Center with an UltiMate 3000RSLC (Thermo Scientific) coupled to a Q Exactive
573 mass spectrometer (Thermo Scientific) in polarity-switching mode with positive voltage 3.0 kV and
574 negative voltage 2.25 kV. Separation was achieved using a gradient elution with (A) 5mM NH₄AcO
575 (pH 9.9) and (B) acetonitrile. The gradient ran from 15% (A) to 90% (A) over 18 min, followed by
576 an isocratic step for 9 minutes and re-equilibration for 7 minutes. Metabolites were quantified as
577 area under the curve based on retention times and using accurate mass measurements (≤ 3 ppm)
578 with the TraceFinder 3.1 software (Thermo Scientific). For heatmap depiction, the relative amounts
579 of metabolites were normalized to the mean value across all samples for one same condition and
580 to the number of viable cells harvested in parallel on a control plate. Pathway enrichment for up-
581 and downregulated KEGG metabolites ($|\log_2(\text{fold-change})|=0.25$) was determined using the
582 MetaboAnalyst 5.0 platform (www.metaboanalyst.ca)³⁸.

583 **RNA-sequencing**

584 Total RNA was extracted from ESCs and EpiLCs pellets using the AllPrep DNA/RNA Micro
585 Kit (Qiagen), according to the manufacturer's protocol. For PGCLCs, d5 embryonic bodies were
586 harvested and cells were dissociated using TrypLE™ Select followed by resuspension in
587 fluorescence-activated cell sorting (FACS) buffer (1×Dulbecco's phosphate buffered saline
588 [DPBS], 1% BSA, 1 mM EDTA, 25 mM HEPES) and cell suspension were passed through a cell
589 strainer (70µm). Cells were sorted on a BD Biosciences FACSAria III (UCLA BSCRC Flow
590 Cytometry Core). Practically, cell populations of interest, being double-positive (Blimp1::mVenus+
591 and Stella::ECFP+) were sorted and collected in microtubes containing GK15 medium. Total RNA
592 was extracted from double-positive PGCLCs using the AllPrep DNA/RNA Micro Kit (Qiagen). RNA
593 concentration was measured using a NanoDrop™ 2000 UV spectrophotometer (Thermo Fisher).
594 Libraries were prepared with the KAPA mRNA HyperPrep Kit (BioMek) or with the RNA library
595 prep kit (ABCclonal) following the manufacturers' protocols. Briefly, poly(A) RNA were selected,
596 fragmented and double-stranded cDNA synthesized using a mixture of random and oligo(dT)
597 priming, followed by end repair to generate blunt ends, adaptor ligation, strand selection, and
598 polymerase chain reaction (PCR) amplification to produce the final library. Different index adaptors
599 were used for multiplexing samples in one sequencing lane. Sequencing was performed on an
600 Illumina NovaSeq 6000 sequencers for paired end (PE), 2×150 base pair (bp) runs. Data quality
601 check was performed using Illumina Sequencing Analysis Viewer (SAV) software. Demultiplexing
602 was performed with Illumina Bcl2fastq2 program (version 2.19.1.403; Illumina Inc.).

603

604 **Differential gene expression analysis**

605 The quality of the reads was verified using FastQC⁴⁹ before reads were aligned to the mm10
606 reference genome (GRCm39) using STAR⁵⁰ with the following arguments: --readFilesCommand
607 zcat --outSAMtype BAM SortedByCoordinate --quantMode GeneCounts --
608 outFilterMismatchNmax 5 --outFilterMultimapNmax 1. The quality of the resulting alignments was
609 assessed using QualiMap⁵¹. The Python package HTseq was used for gene counts⁵² using the
610 following arguments: --stranded=no --idattr=gene_id --type=exon --mode=union -r pos --
611 format=bam. Output files were filtered to remove genes with low count (≤ 10) then were used for
612 differential gene expression analysis using DESeq2⁵³. The negative binomial regression model of
613 ComBat-seq was used to correct unwanted batch effects⁵⁴. For a gene to be classified as showing
614 differential expression between treated and untreated cells, a threshold of $|\log_2(\text{fold-change})|=0.5$
615 and Benjamini-Hochberg adjusted p-value ≤ 0.05 had to be met.

616

617 **Gene Ontology (GO) Analysis**

618 Lists of differentially expressed genes were generated from read counts using DESeq2
619 Bioconductor package⁵³. Enrichment of GO terms in lists of up- and downregulated genes
620 ($|\log_2(\text{fold-change})|=0.25$) was determined using g:Profiler⁵⁵. Redundant GO terms were removed
621 using reduce + visualize gene ontology (REVIGO)⁵⁶. Terms were included if the fold enrichment
622 (frequency of DEGs in each GO term to the frequency of total genes in GO terms) was higher than
623 1.5 and if the Benjamini-Hochberg-adjusted p-value was less than 0.05. Plots for GO terms were
624 generated using a custom R script⁵⁷.

625 **Statistical Methods**

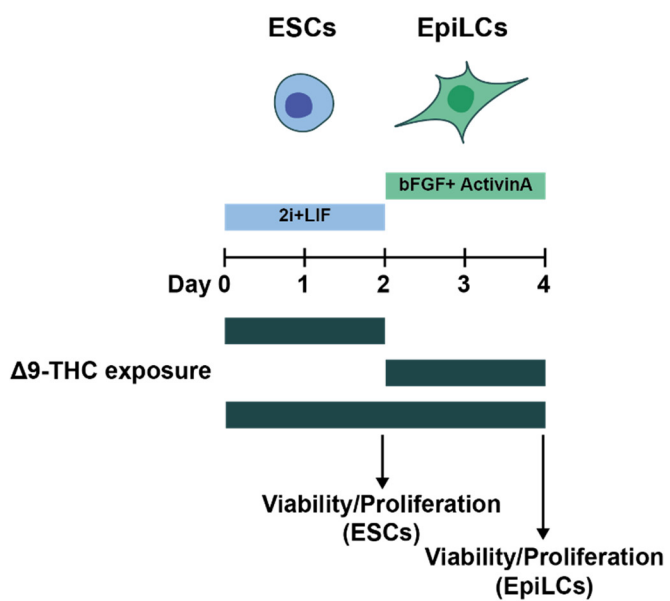
626 Statistical analyses, when not otherwise specified, were performed using GraphPad Prism
627 9 software. For significance testing, two-tailed T-tests were performed on pairwise comparisons.
628 In all cases, significance was determined by p-values less than or equal to 0.05. Each figure
629 corresponds to at least three independent biological repeats with three technical replicates (N=3,
630 n=3), unless otherwise specified. Number of asterisks on plots indicate level of statistical
631 significance: *(p<0.05), **(p<0.01), ***(p<0.001) and ****(p<0.0001).

632 **ACKNOWLEDGMENTS**

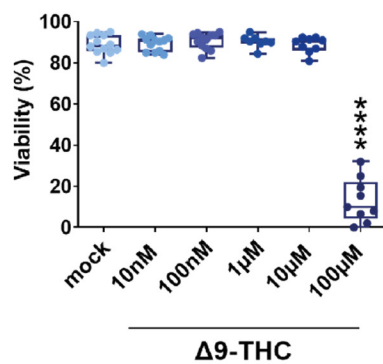
633 We thank Jessica Scholes, Felicia Codrea and Jeffrey Calimlim from the UCLA BSCRC
634 Flow Cytometry Core for their assistance in cell sorting. We thank Linsey Stiles (UCLA
635 Mitochondria and Metabolism Core), Johanna ten Hoeve-Scott and Thomas Graeber (UCLA
636 Metabolomics Center) for their technical support in the metabolomics analyses. We thank Xinmin
637 Li and his team at the UCLA Technology Center for Genomics & Bioinformatics for their technical
638 support in the high-throughput sequencing. We thank Luigi Bellocchio and Giovanni Marsicano for
639 their recommendations in $\Delta 9$ -THC exposure setups. We would like to acknowledge the UCLA
640 Cannabis Research Initiative (UCRI) and Dr Ziva Cooper for their continuous support and
641 encouragements.

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643 and the Iris Cantor-UCLA Women's Health Center and the NCATS UCLA CTSI Grant Number
644 UL1TR001881. RV is a postdoctoral fellow and acknowledges support from the Belgian-American
645 Educational Foundation (BAEF).

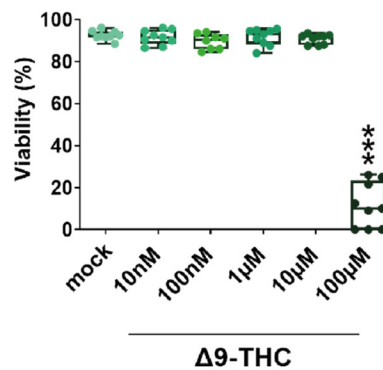
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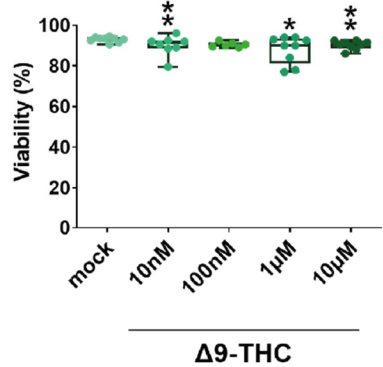
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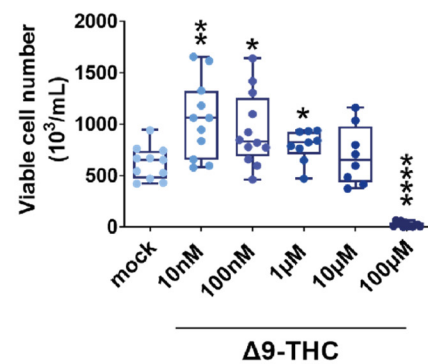
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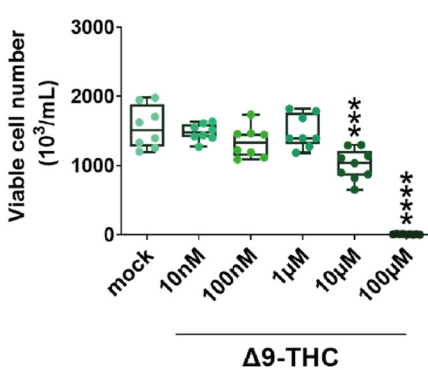
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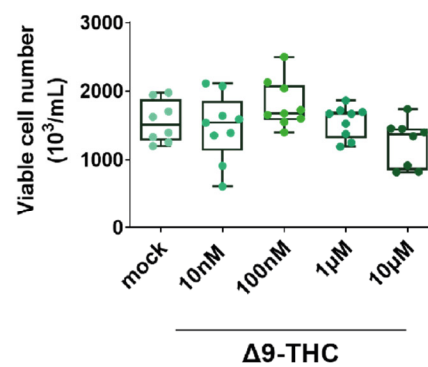
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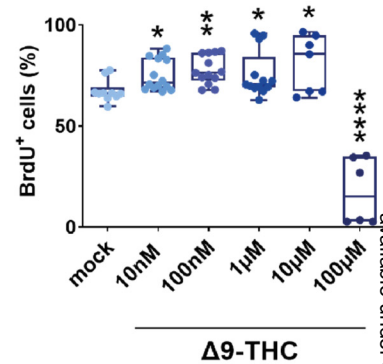
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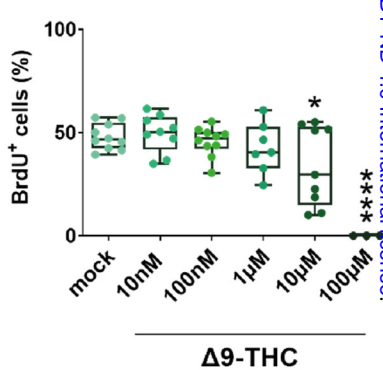
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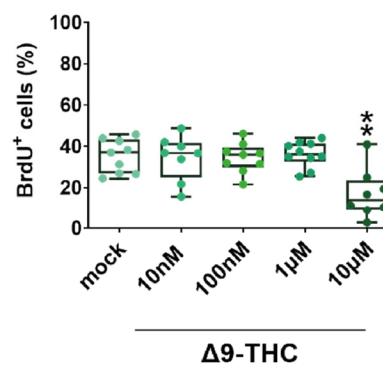
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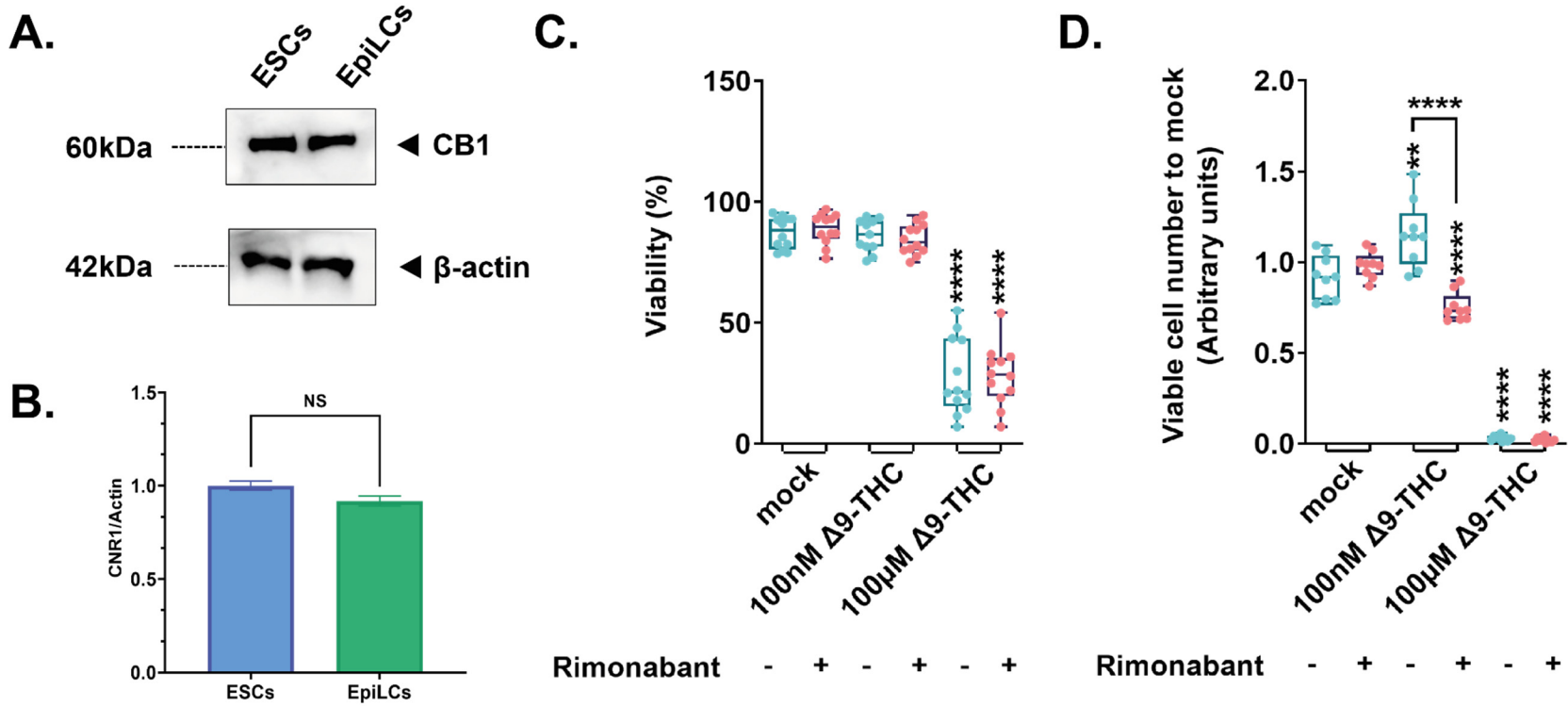


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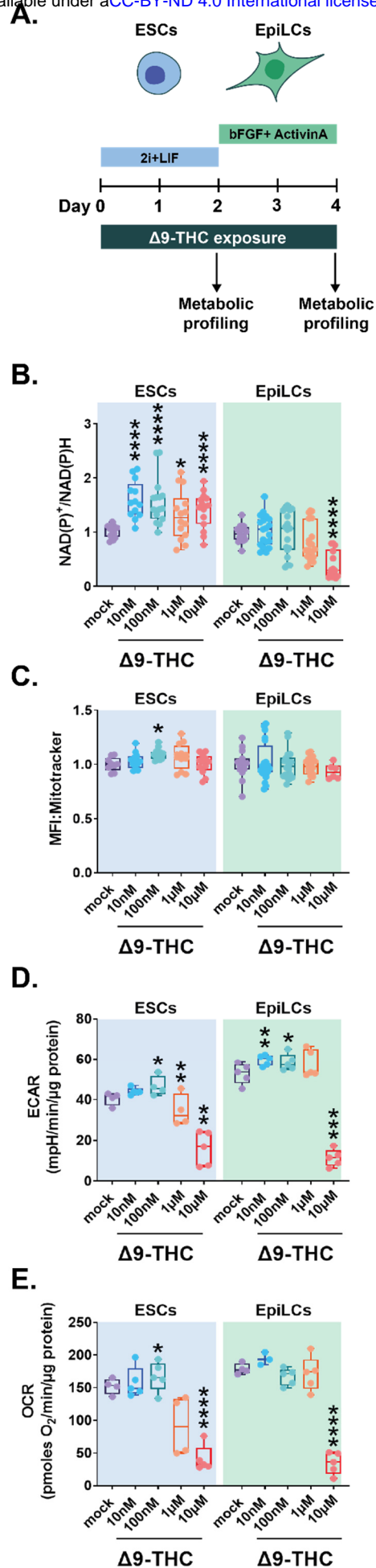


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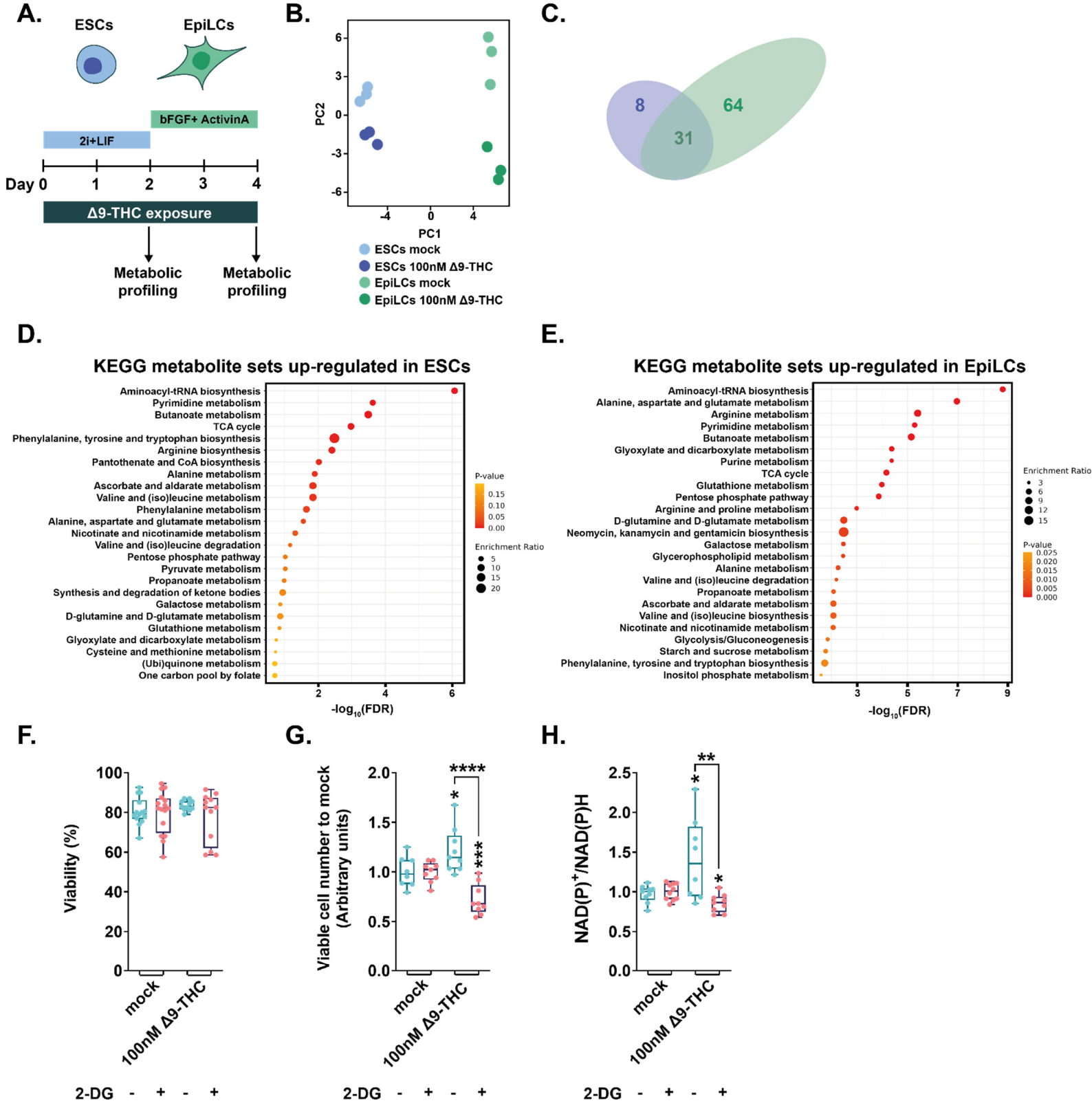


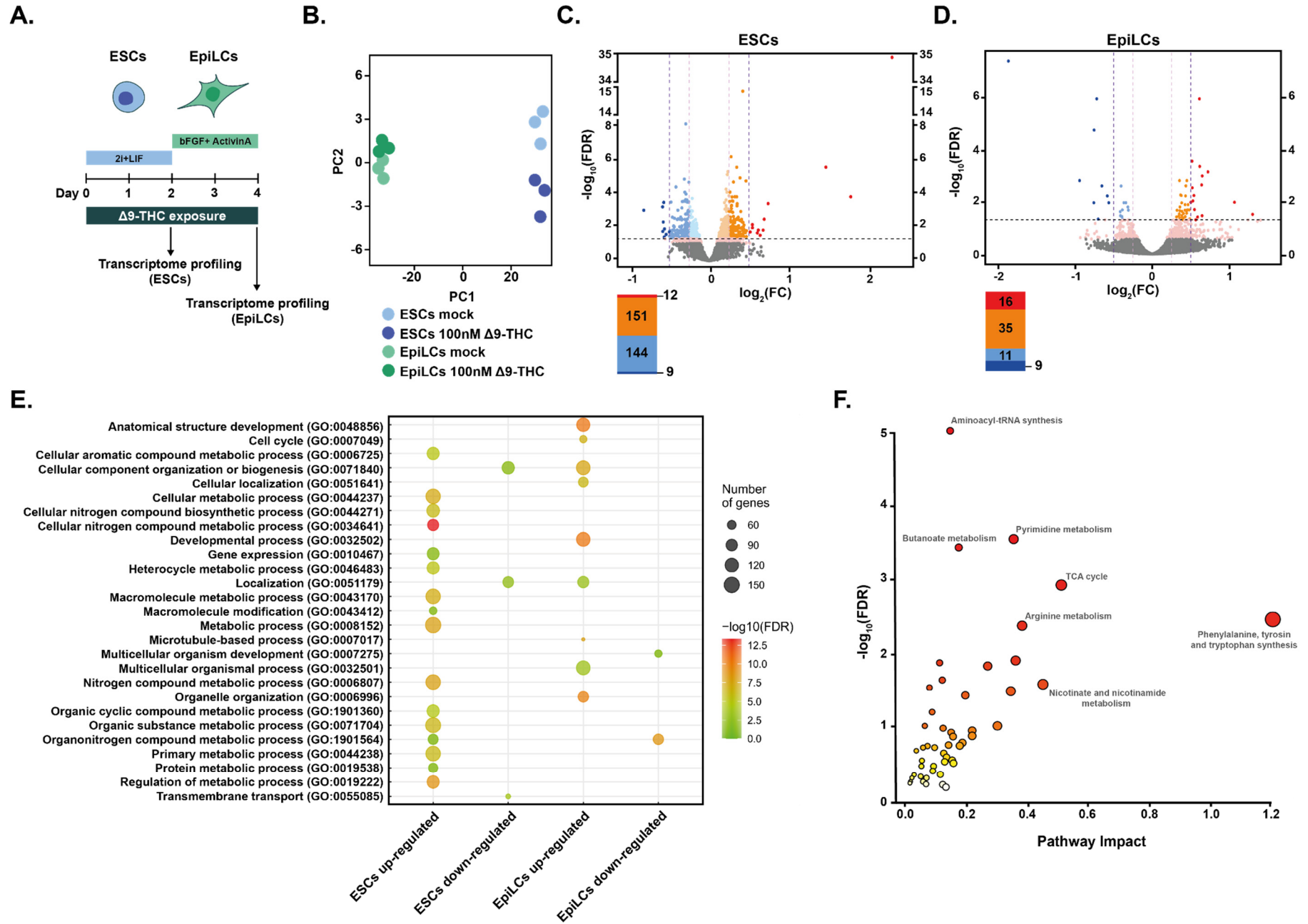


649 **Figure 3**

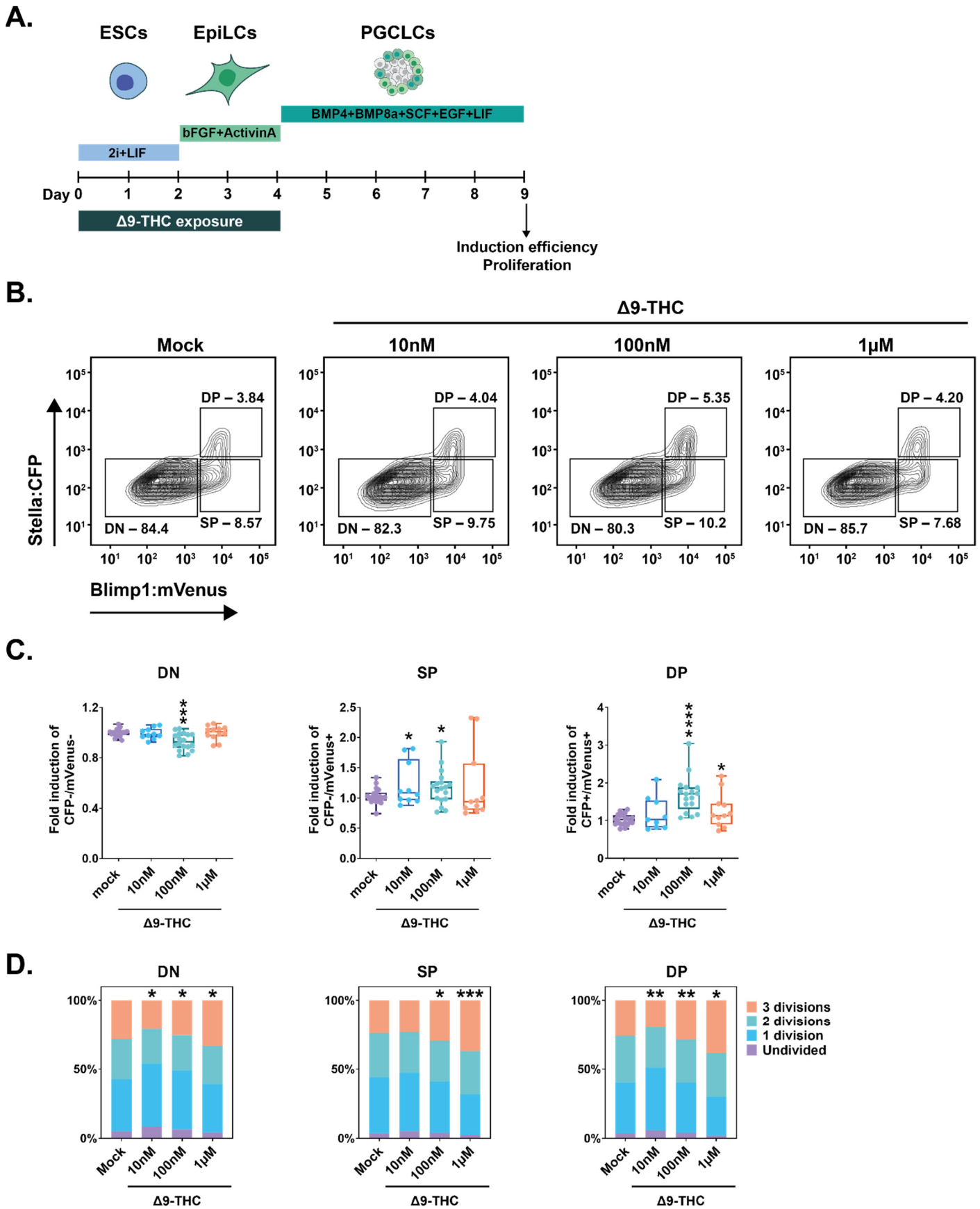


650 **Figure 4**

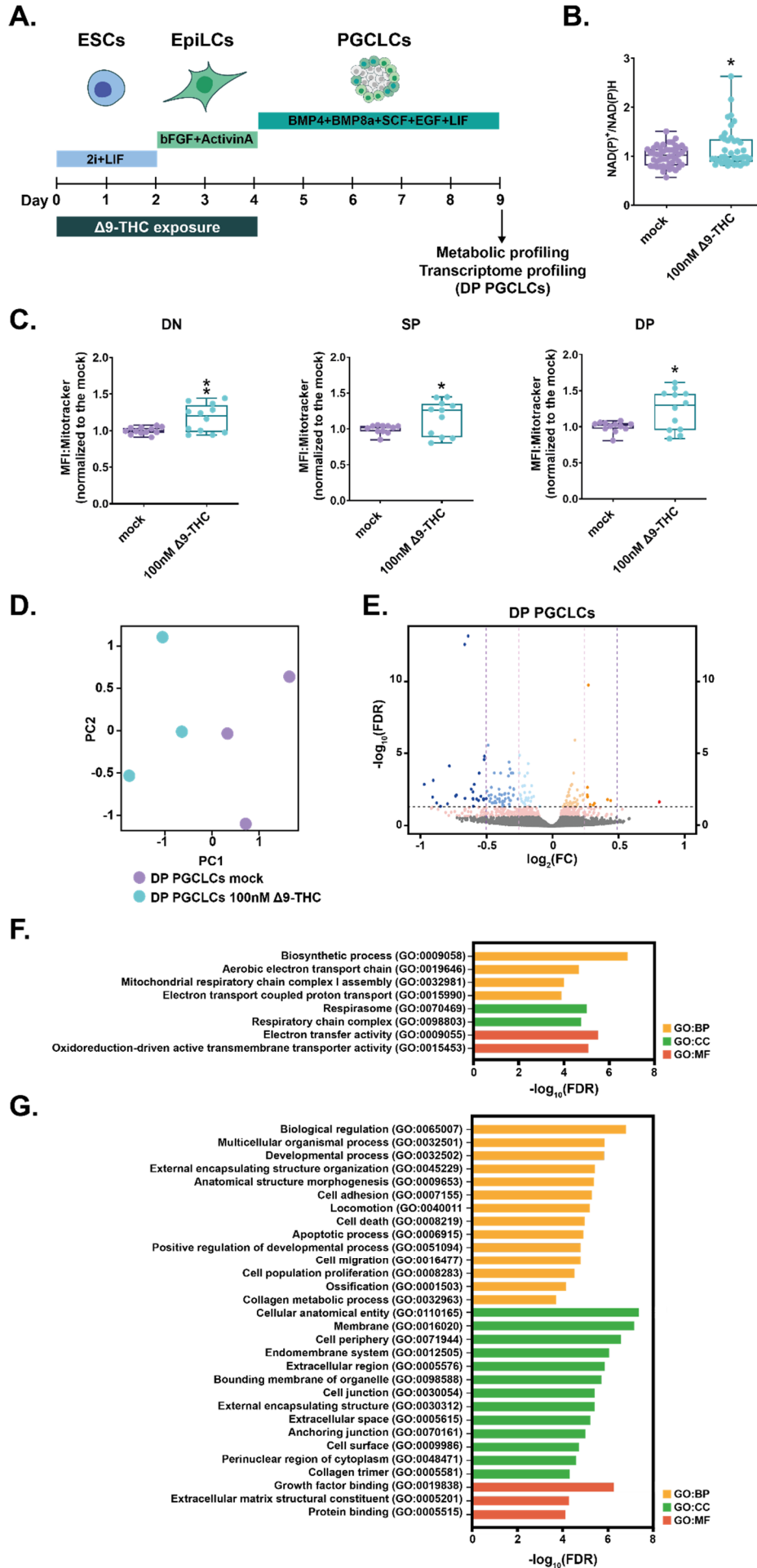


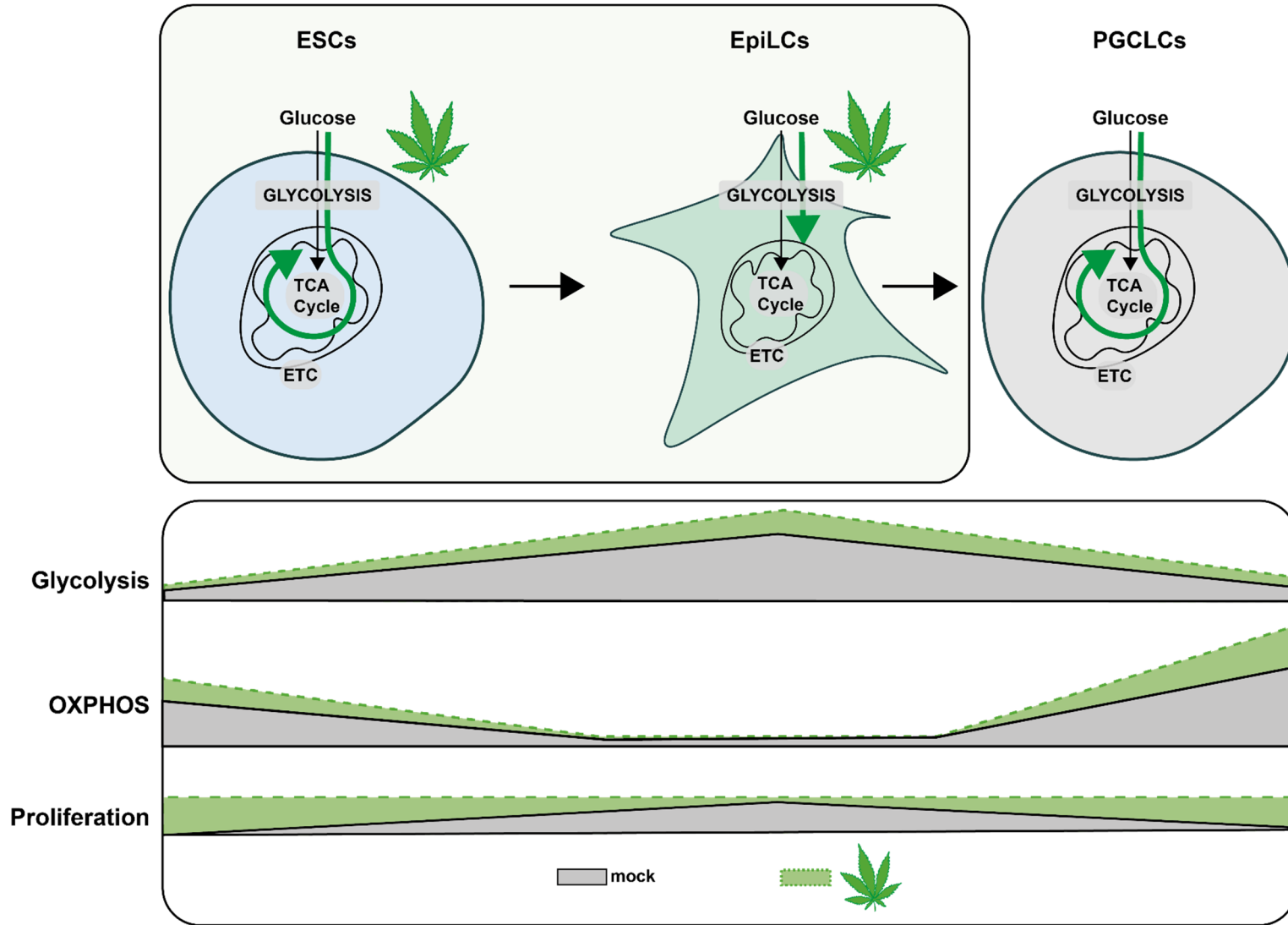


652 **Figure 6**



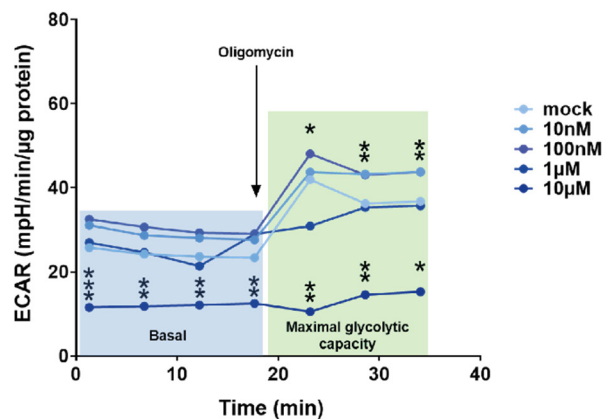
653 Figure 7



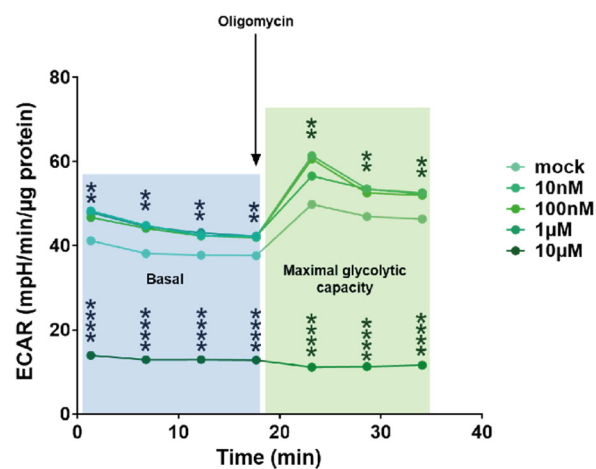


655 SUPPLEMENTARY FIGURES
 656 Supplementary Figure 1

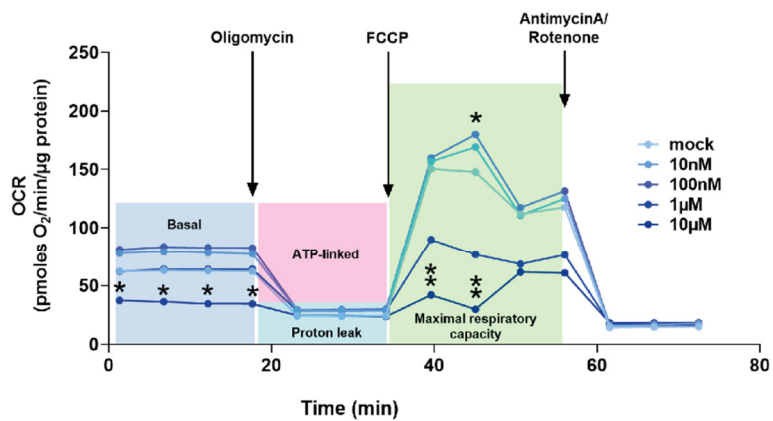
A.



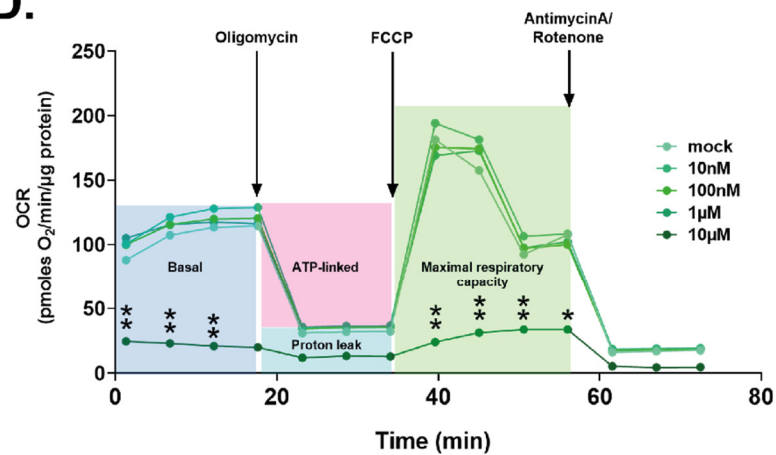
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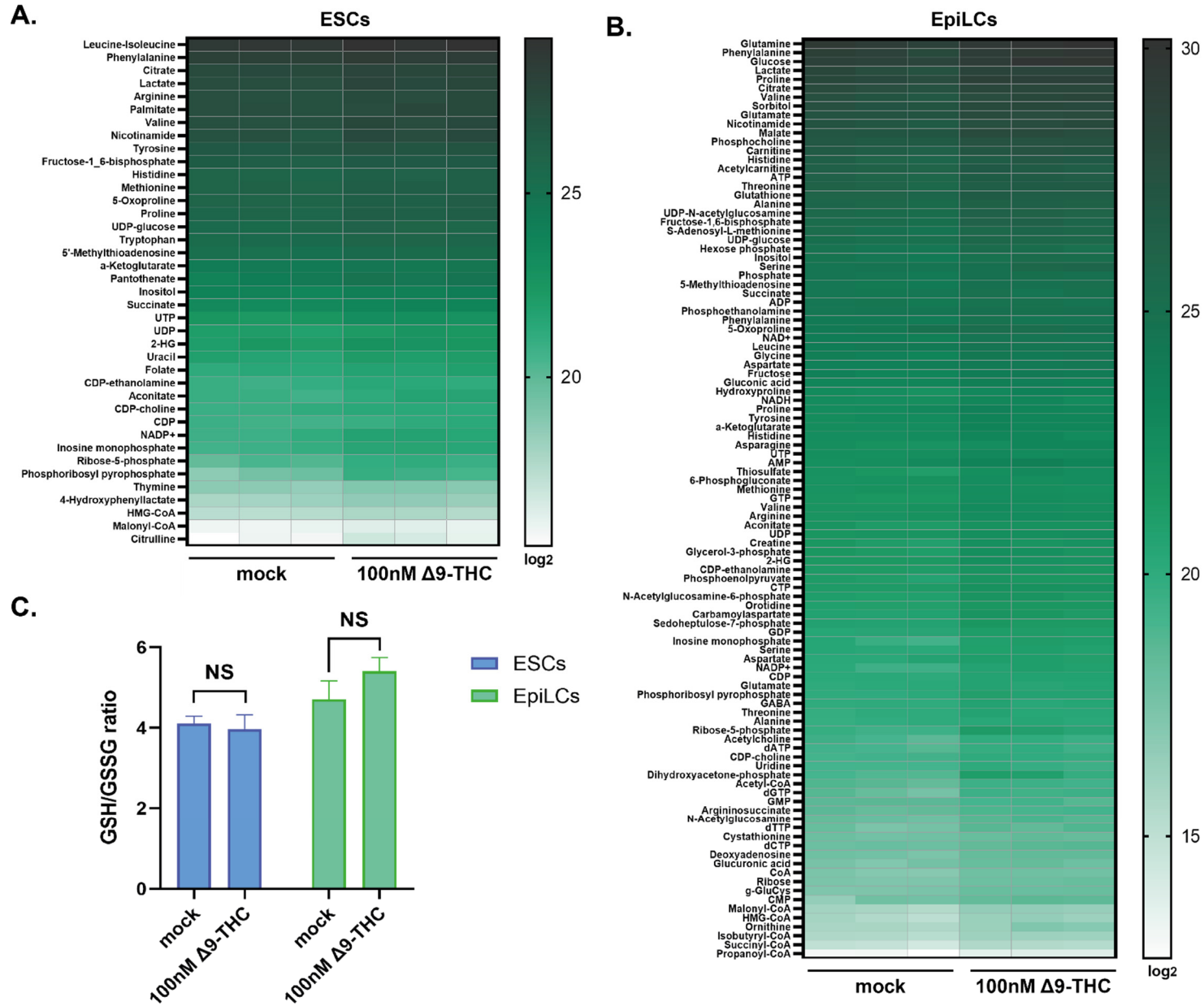


C.

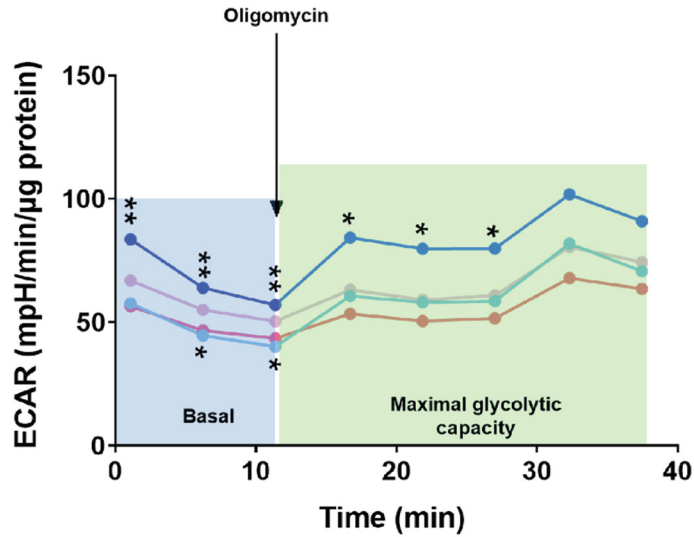


D.



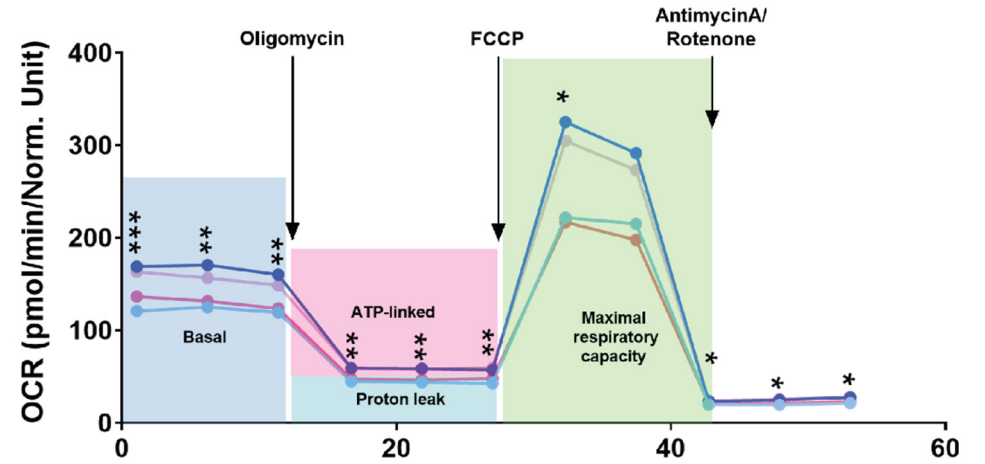


A.



- mock
- 100nM Δ 9-THC
- mock + 10mM 2-DG
- 100nM Δ 9-THC + 10mM 2-DG

B.



- mock
- 100nM Δ 9-THC
- mock + 10mM 2-DG
- 100nM Δ 9-THC + 10mM 2-DG

659 **FIGURE LEGENDS**

660 **Figure 1: $\Delta 9$ -THC exposure provokes the proliferation of ESCs but not EpiLCs**

661 **(A)** Diagram illustrating $\Delta 9$ -THC exposure scheme and experimental strategy. bFGF: basic
662 fibroblast growth factor, ESCs, embryonic stem cells; EpiLCs, epiblast-like cells; LIF, leukemia
663 inhibitory factor. **(B, E, H)** Whisker boxplot indicating the median cellular viability of stem cells
664 exposed to the different $\Delta 9$ -THC doses and associated errors. **(C, F, I)** Whisker boxplot indicating
665 the median number of viable cells exposed to the different $\Delta 9$ -THC doses indicated and
666 associated errors. **(D, G, J)** Whisker boxplot indicating the median percentage of BrdU-stained
667 cells exposed to the different $\Delta 9$ -THC doses and associated errors. ESCs exposed cells are
668 presented in (B, C and D). EpiLCs exposed cells deriving from unexposed ESCs are presented in
669 (E, F and G). EpiLCs exposed cells deriving from exposed ESCs are presented in (H, I and J).

670

671 **Figure 2: Implication of the CB1 receptor in the proliferative phenotype.**

672 **(A)** Western blot analysis of transmembrane protein extracts of ESCs or EpiLCs. Antibodies raised
673 against CB1 or β -actin serving as a loading control were used for immunoblotting. **(B)**
674 Quantification of the gel presented in (A) was done using Image Studio (version 5.2). **(C)** Whisker
675 boxplot indicating the median cellular viability of stem cells exposed to the different $\Delta 9$ -THC and
676 rimonabant doses indicated and their associated errors. **(D)** The median numbers of viable cells
677 exposed to the different $\Delta 9$ -THC and rimonabant doses indicated were normalized to their own
678 control (+/- rimonabant). Median and associated errors were plotted in whisker boxplots.

679

680 **Figure 3: $\Delta 9$ -THC exposure provokes an increase in glycolytic rates in ESCs and EpiLCs.**

681 **(A)** Diagram illustrating $\Delta 9$ -THC exposure scheme and experimental strategy. **(B)** The
682 NAD(P)⁺/NADPH ratio of stem cells exposed to the different $\Delta 9$ -THC doses was normalized to
683 the one measured in the mock-treated condition. Median and associated errors were plotted in
684 whisker boxplots. **(C)** Mean fluorescence intensity (MFI) associated with the Mitotracker CMXRos
685 stain was normalized to the one measured in the mock-treated condition. Median and associated
686 errors were plotted in whisker boxplots. **(D)** Median and associated error of the maximal
687 extracellular acidification rate (ECAR) measured in cells exposed to the different $\Delta 9$ -THC doses
688 and normalized to the protein content was plotted in whisker boxplots. **(E)** Median and associated
689 error of the maximal oxygen consumption rate (OCR) measured in cells exposed to the different
690 $\Delta 9$ -THC doses and normalized to the protein content was plotted in whisker boxplots. For (B and
691 C), 5 technical repeats of 3 biological repeats (n=15) were plotted. One same representative
692 experiment out of three independent experiments was used to plot results in (D and E).

693

694 **Figure 4: $\Delta 9$ -THC-induced glycolysis sustain anabolism and ESCs proliferation**

695 **(A)** Diagram illustrating $\Delta 9$ -THC exposure scheme and experimental strategy. **(B)** PCA of the
696 metabolomics profiling of either ESCs or EpiLCs mock-exposed or exposed to 100nM $\Delta 9$ -THC.
697 **(C)** Venn diagram showing the overlap in upregulated metabolites following $\Delta 9$ -THC exposure in
698 ESCs and EpiLCs. **(D and E)** KEGG metabolite sets enrichment analysis for upregulated
699 metabolites in ESCs and EpiLCs, respectively, performed by MetaboAnalyst³⁸. KEGG, Kyoto
700 Encyclopedia of Genes and Genomes. **(F)** Whisker boxplot indicating the median cellular viability
701 of stem cells exposed to 100nM of $\Delta 9$ -THC and 10mM of 2-DG, as indicated, and their associated
702 errors. **(G)** The median numbers of viable cells exposed to 100nM of $\Delta 9$ -THC and 10mM of 2-DG,

703 as indicated, were normalized to their own control (+/- 2-DG). Median and associated errors were
704 plotted in whisker boxplots. **(H)** The NAD(P)+/NADPH ratio of stem cells exposed to 100nM of Δ 9-
705 THC and 10mM of 2-DG, as indicated, was normalized to the one measured in the mock-treated
706 condition (+/- 2-DG). Median and associated errors were plotted in whisker boxplots.

707

708 **Figure 5: Metabolic changes following Δ 9-THC exposure in ESCs are transcriptionally**
709 **encoded.**

710 **(A)** Diagram illustrating Δ 9-THC exposure scheme and experimental strategy. **(B)** PCA of the
711 transcriptomics profiling of either ESCs or EpiLCs mock-exposed or exposed to 100nM Δ 9-THC.
712 **(C and D)** Volcano plot in ESCs and EpiLCs, respectively, showing significance [expressed in
713 \log_{10} (adjusted p-value or false-discovery rate, FDR)] versus fold-change [expressed in \log_2 (fold-
714 change, FC)]. Thresholds for significance (adjusted p-value \leq 0.05) and gene expression fold-
715 change [$|\log_2(\text{FC})|>0.25$ or $|\log_2(\text{FC})|>0.5$] are shown as dashed lines. Color code is as follows:
716 $\log_2(\text{FC})>0.5$ in red, $\log_2(\text{FC})>0.25$ in orange, $\log_2(\text{FC})>0$ in light orange, $\log_2(\text{FC})<0$ in light blue,
717 $\log_2(\text{FC})>-0.25$ in blue, $\log_2(\text{FC})>0.5$ in dark blue and p-value <0.01 in pink. **(E)** Gene ontology
718 (GO) terms associated with up- and downregulated DEGs [$|\log_2(\text{FC})|>0.25$ and p <0.01] in ESCs
719 and EpiLCs as determined by g:Profiler⁵⁵. **(F)** Joint pathway analysis performed by the multi-omics
720 integration tool of MetaboAnalyst³⁸. The p-values were weighted based on the proportions of
721 genes and metabolites at the individual pathway level.

722

723 **Figure 6: PGCLCs deriving from ESCs and EpiLCs exposed to 100nM of Δ 9-THC**
724 **proliferate.**

725 **(A)** Diagram illustrating Δ 9-THC exposure scheme and experimental strategy. **(B)** Representative
726 flow contour plots showing distribution of live-gated events, gating strategy for Stella:CFP versus
727 Blimp1:mVenus and percentages of cells in each subpopulations for ESCs and EpiLCs exposed
728 to the different doses of Δ 9-THC indicated. DN: double negative, SP: single positive, DP: double
729 positive subpopulations. **(C)** The percentage of events in the gates associated to each
730 subpopulation was normalized to the one measured in the mock-treated condition. Median and
731 associated errors were plotted in whisker boxplots independently for each subpopulation. **(D)**
732 Representative histograms showing CellTrace™ Yellow staining profile of cells arising from ESCs
733 and EpiLCs exposed to the different doses of Δ 9-THC indicated. The Y-axis represents the
734 average percentage of cells in each category of subpopulations undividing (purple), undergoing 1
735 division (blue), 2 divisions (green) or 3 divisions (orange). One representative experiment out of
736 three is represented.

737

738 **Figure 7: Δ 9-THC exposure prior to specification increases mitochondrial respiration in**
739 **PGCLCs.**

740 **(A)** Diagram illustrating Δ 9-THC exposure scheme and experimental strategy. **(B)** The
741 NAD(P)+/NADPH ratio of embryoid bodies arising from ESCs and EpiLCs exposed to 100nM of
742 Δ 9-THC was normalized to the one measured in the mock-treated condition. Median and
743 associated errors were plotted in whisker boxplots. **(C)** Mean fluorescence intensity (MFI)
744 associated with the Mitotracker CMXRos stain in each subpopulation was normalized to the one
745 measured in the mock-treated condition. Median and associated errors were plotted in whisker
746 boxplots. **(D)** PCA of the transcriptomics profiling of DP PGCLCs deriving from ESCs and EpiLCs

747 either mock-exposed or exposed to 100nM Δ 9-THC. **(E)** Volcano plot in DP PGCLCs showing
748 significance [expressed in \log_{10} (adjusted p-value or false-discovery rate, FDR)] versus fold-
749 change [expressed in \log_2 (fold-change, FC)]. Thresholds for significance and different enrichment
750 ratios [$|\log_2(\text{FC})| > 0.25$ or $|\log_2(\text{FC})| > 0.5$] are shown as dashed lines. Color code is as follows:
751 $\log_2(\text{FC}) > 0.5$ in red, $\log_2(\text{FC}) > 0.25$ in orange, $\log_2(\text{FC}) > 0$ in light orange, $\log_2(\text{FC}) < 0$ in light blue,
752 $\log_2(\text{FC}) > -0.25$ in blue, $\log_2(\text{FC}) > 0.5$ in dark blue and p-value < 0.01 in pink. **(F and G)** Gene
753 ontology (GO) terms associated with up- and downregulated DEGs [$|\log_2(\text{FC})| > 0.25$ and
754 p < 0.01], respectively, as determined by g:Profiler⁵⁵.

755

756 **Figure 8: Metabolic impact of Δ 9-THC exposure in pluripotent stem cells and primordial**
757 **germ cells-like cells.**

758 Diagram illustrating the impact of Δ 9-THC exposure on stem cells metabolism.

759

760 **Supplementary Figure 1: Extracellular acidification rates and oxygen consumption rates**
761 **in ESCs and EpiLCs upon Δ 9-THC exposure.**

762 **(A and B)** Traces were plotted for the extracellular acidification rate (ECAR) measurements in
763 ESCs and EpiLCs, respectively, exposed to the different Δ 9-THC doses indicated and normalized
764 to the protein content. The oligomycin injection time is indicated by an arrow and allows to
765 differentiate basal glycolytic rate from maximal glycolytic rate (when mitochondria are inhibited).
766 The datapoints used in the main figure correspond to the first timepoint in the maximal glycolytic
767 capacity section. **(C and D)** Traces were plotted for the oxygen consumption rate (OCR)
768 measurements in ESCs and EpiLCs, respectively, exposed to the different Δ 9-THC doses
769 indicated and normalized to the protein content. The oligomycin, FCCP and AntimycinA/Rotenone
770 injection times are indicated by arrows and allow to differentiate basal respiration from ATP-
771 coupled respiration and maximal respiratory capacity. The datapoints used in the main figure
772 correspond to the second timepoint in the maximal respiratory capacity section. FCCP: Carbonyl
773 cyanide-p-trifluoromethoxyphenylhydrazone.

774

775 **Supplementary Figure 2: Metabolite profiling in ESCs and EpiLCs upon Δ 9-THC exposure.**

776 **(A and B)** Heatmaps showing the \log_2 of the amount of each metabolite upregulated in ESCs and
777 EpiLCs upon exposure to 100nM of Δ 9-THC. The relative amounts of metabolites were normalized
778 to the mean value across all samples for one same condition and to the number of viable cells
779 harvested in parallel on a control plate. **(C)** Histograms showing the ratio of reduced to oxidized
780 glutathione (GSH/GSSG) based on the amounts measured in the metabolomics profiling.

781

782 **Supplementary Figure 3: Extracellular acidification rates and oxygen consumption rates**
783 **in ESCs upon Δ 9-THC and 2-DG exposure.**

784 **(A)** Traces were plotted for the extracellular acidification rate (ECAR) measurements in ESCs
785 exposed to 100nM of Δ 9-THC and 10mM of 2-DG, as indicated, and normalized to the protein
786 content. The oligomycin injection time is indicated by an arrow and allows to differentiate basal
787 glycolytic rate from maximal glycolytic rate (when mitochondria are inhibited). **(B)** Traces were
788 plotted for the oxygen consumption rate (OCR) measurements in ESCs exposed to 100nM of Δ 9-
789 THC and 10mM of 2-DG, as indicated, and normalized to the protein content. The oligomycin,
790 FCCP and AntimycinA/Rotenone injection times are indicated by arrows and allow to differentiate

791 basal respiration from ATP-coupled respiration and maximal respiratory capacity. FCCP: Carbonyl
792 cyanide-p-trifluoromethoxyphenylhydrazone.

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