



HAL
open science

Prenatal maternal vitamin D deficiency sex-dependently programs adipose tissue metabolism and energy homeostasis in offspring

Eva Seipelt, Franck Tourniaire, Charlène Couturier, Julien Astier, Béatrice B. Loriod, Hortense Vachon, Michel Puceat, Lourdes Mounien, Jean-Francois Landrier

► To cite this version:

Eva Seipelt, Franck Tourniaire, Charlène Couturier, Julien Astier, Béatrice B. Loriod, et al.. Prenatal maternal vitamin D deficiency sex-dependently programs adipose tissue metabolism and energy homeostasis in offspring. *FASEB Journal*, 2020, 34 (11), pp.14905-14919. 10.1096/fj.201902924RR . hal-03156949

HAL Id: hal-03156949

<https://hal.inrae.fr/hal-03156949>

Submitted on 20 May 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

1 **Prenatal maternal vitamin D deficiency sex-dependently programs adipose tissue**
2 **metabolism and energy homeostasis in offspring**

3
4

5 Eva M. Seipelt^{1,4}, Franck Tourniaire^{1,2}, Charlène Couturier¹, Julien Astier¹, Béatrice Loriod³,
6 Hortense Vachon³, Michel Pucéat^{4*}, Lourdes Mounien^{1*}, Jean-François Landrier^{1,2*}

7
8

9 1 Aix-Marseille Université, C2VN, INRAE, INSERM, 13000, Marseille, France

10 2 CriBioM, Criblage Biologique Marseille, Faculté de médecine de la Timone, Marseille,
11 France.

12 3 Aix-Marseille Université, TGML, TAGC, INSERM, 13000 Marseille, France

13 4 Aix-Marseille Université, MMG, INSERM U1251, Marseille, France.

14
15

* joint last authors.

16
17

Short running title : maternal vitamin D programs offspring metabolism

18
19

Corresponding author and person to whom reprint requests should be addressed:

20 Jean-François Landrier, C2VN ; UMR 1260 INRAE/1263 INSERM/Université d'Aix-
21 Marseille, 27 Bd Jean Moulin, 13385 Marseille cedex 05, France. Phone: +33 4 91 29 42 75;
22 E-mail: jean-francois.landrier@univ-amu.fr

23
24

25 **List of nonstandard abbreviations :**

26

27 25(OH)D: 25-hydroxyvitamin D

28 AGRP: Agouti related-peptide

29 AUC: Area under the curve

30 CTRL: Control

31 EE: Energy expenditure

32 GOX: Carbohydrate oxidation

33 HF: High fat

34 HOMA-IR: Homeostatic model assessment of insulin resistance

35 IPA: Ingenuity pathway analysis

36 ITT: Insulin tolerance test

37 LF: Low fat

38 LOX: Lipid oxidation

39 MC4R: Melanocortin receptor type 4

40 NPY: Neuropeptide Y

41 RQ: Respiratory quotient

42 VDD: Vitamin D deficiency

43

44 **Abstract**

45 *In utero* environment is crucial to ensure normal development of the foetus and to program
46 metabolic health throughout the life. Beside macronutrients, the role of micronutrients,
47 including vitamin D, begins to be explore. The aim of this study was to decipher the impact of
48 maternal vitamin D deficiency (VDD), in normal and high fat (HF) diet context, on adipose
49 tissue metabolism and energy homeostasis in offspring, considering sex-specific responses.
50 Body weight, energy expenditure and spontaneous activity was differential impacted in
51 juvenile male and female offspring born from VDD mice. In adulthood, a HF diet combined
52 with maternal VDD disrupted glucose homeostasis and adiposity in male offspring but not in
53 females. Such phenotypes were associated to different transcriptomic profiles in adipose
54 tissue, that could be related to differential modulation of plasma 17 β -estradiol concentrations.
55 Thus, maternal VDD sex-dependently modulated metabolic fate of the offspring, especially
56 when associated with HF diet in adulthood.

57

58 **Keywords:** Vitamin D, Maternal vitamin D deficiency, offspring, adipose tissue, metabolism.

59

60 **Introduction**

61

62 Vitamin D is an essential micronutrient that is suspected to display an important role on
63 normal foetal development, and throughout the life of the child and future adult. Its role in
64 calcium and phosphate metabolism is well-established and it also appears to be involved in
65 many other physiological processes¹, including control of adipose tissue biology²⁻⁹.
66 Consequently, vitamin D insufficiency is considered as a risk factor in several pathologies,
67 including auto-immune diseases, musculoskeletal defects¹⁰, cardiovascular and metabolic
68 diseases. Such insufficiency is characterised by plasma concentration in 25-hydroxyvitamin D
69 (25(OH)D) below the cut-off value of 50nmol/L¹¹. In fact, vitamin D insufficiency has
70 become a large public health issue^{12,13} and is common for women in child bearing age,
71 pregnant and breastfeeding women. Currently, 54% of pregnant women and 75% of new-
72 borns present a 25(OH)D status < 50nmol/L, also 18% of pregnant women and 29% of new-
73 borns present a severe vitamin D deficiency (25(OH)D < 25nmol/L)^{14,15}. It is noteworthy that
74 maternal vitamin D deficiency (VDD) was associated with pregnancy, foetal and neonatal
75 outcomes¹⁵ such as increased risk factor for preeclampsia¹⁶, gestational diabetes mellitus¹⁷,
76 higher risk of small-for-gestational-age, reduced term birth weight, and lower head
77 circumference¹⁸, even if these results are sometimes controverted¹⁹.

78 Interestingly, recent studies linked maternal vitamin D insufficiency to overweight and
79 increased fat mass in offspring. Indeed, in the INMA cohort, including 2358 pregnant women,
80 maternal VDD was associated with increased risk of overweight in foetal and early postnatal
81 (1-year-old) offspring²⁰. Similarly, in the prospective cohort study Southampton Women's
82 survey (977 pregnant women), maternal VDD was associated with lower fat mass at birth and
83 greater fat mass at 4 and 6 years²¹. These results were confirmed in the prospective cohort
84 Rhea (532 pregnant women), where the maternal 25(OH)D concentrations < 37.7 nmol/L
85 were associated with higher body mass index and central adiposity in the 4 and 6-years-old
86 offspring²². Nevertheless, in a long term follow up (20 years), no association between
87 maternal vitamin D status and cardio-metabolic risk factors was highlighted²³. Thus, those
88 associations remain controversial.

89 To gain further insight on the relationship, several preclinical studies have been implemented.
90 Overall, it has been established that maternal VDD as potential programming long-term effect
91 on metabolic health^{24,25}, nevertheless several important points remain pending. Based on the
92 Developmental origins of Health and disease (DOHaD concept), we hypothesized that the
93 response to prenatal maternal VDD may be sex-specific and influenced by the nutritional

94 environment during adulthood. Indeed, the maternal environment during the pre- and peri-
95 conceptional period, especially nutrition, can modify epigenetic marks, leading to long term
96 phenotypic consequences that may vary according to the sex and the environment of the
97 offspring²⁶.

98 Thus, the aim of the present study is to determine the impact of maternal VDD combined with
99 the impact of obesogenic environment, induced by a high-fat diet, on the sex-specific
100 response during adulthood on energy homeostasis and adipose tissue metabolism.

101

102

103 **Material and Methods**

104

105 ***Animal Experiments***

106 The protocol received the agreement of Aix-Marseille University Ethics Committee and the
107 French Ministry of Research (APAFIS#1300-2015072112279135). Eight-week-old female
108 and male C57BL/6J mice were obtained from Janvier Labs (Le Genest-Saint-Isle, France),
109 fed ad libitum with control food (chow diet A04 from Safe-diets, Augy France) during the 1-
110 week acclimation period and with full access to drinking water. The animals were maintained
111 at 22°C under a 12-hour light, 12-hour dark cycle and a 20% humidity level. Female mice (15
112 per group) were randomly assigned into one of the two experimental groups depending on the
113 diet *i.e.* control (AIN-93G with vitamin D3, 1.0 IU/g) or vitamin D-depleted (AIN-93G
114 without vitamin D3, 0.0 IU/g) for eight-weeks (Supplemental Figure 1), and were mated with
115 males. Weight gain was measured once a week and dietary at 3-weeks of pre-mate diet, at 5
116 days and 15 days of gestational stage (Supplemental Figure 2). After delivery, all females
117 were fed with control diet (AIN-93G) until weaning of the offspring. The litter size was
118 adjusted to 6 pups per females. The body weight of the offspring was evaluated weekly from
119 the weaning until the study end, and not prior weaning to avoid maternal cannibalization and
120 perinatal stress. At six-weeks of age both males and females of the offspring were randomly
121 assigned to receive Low Fat diet (AIN-93M Maintenance Purified Diet) or High Fat diet (DIO
122 Rodent Purified Diet w/45% Energy from Fat) for eight weeks. At the end of the protocol,
123 mice were subjected to food restriction overnight and blood was collected by cardiac puncture
124 anesthesia, serum was isolated by centrifugation at 3000 rpm for 15 min at 4°C and was
125 stored at -80°C. Animals were euthanized by cervical dislocation and various tissue (liver,
126 spleen, hypothalamus and various white adipose tissue deposits) were collected, weighted and
127 stored at -80°C. Eight groups of offspring mice (males and females) were designed to study
128 the impact of maternal diet (CTRL vs VDD) and adult diet (LF vs HF).

129

130 ***Biochemical analysis***

131 To confirm the maternal and offspring vitamin D status, 25(OH)D serum concentration were
132 measured using an in vitro diagnostic enzyme immunoassay kit 25-OH Vitamin D (direct)
133 ELISA kit (PromoKine). Insulin, leptin, testosterone and 17 β -estradiol were measured in
134 plasma using an enzyme-linked immuno-sorbent assay ELISA (Insulin ALPCO Diagnostics,
135 New Hampshire, United States; DuoSet mouse leptin, R&D systems, Minneapolis, United
136 States, Testosterone Demeditec Diagnostics GmbH, Germany; ab108667, Abcam, Cambridge,

137 England, respectively). The manufacturer's protocols were followed. For glucose
138 concentration, mice were subjected to food restriction for 5h and glycemia was measured
139 from tail blood (Accu-Check glucometer, Roche). The HOMA-IR index was calculated
140 according to the following formula: fasting insulin (microU/L) x fasting glucose
141 (nmol/L)/22.5.

142

143 ***Histological analysis***

144 Visceral adipose tissue samples were fixed in 10% buffered formalin, embedded in paraffin
145 and sliced to prepare 5 μm tissue sections whose were stained with hematoxylin and eosin
146 (H&E). The images were captured by a light microscope (EZAD, Leica, Germany; 10X
147 magnification). The adipocyte area (μm^2) were determined using (Image J) software.

148

149 ***Indirect calorimetry***

150 At 5 and 12 weeks old the offspring was acclimated and kept for 24h in an indirect
151 calorimetric cage (Physiocage, Bioseb, Vitrolles, France), as previously described⁶. The
152 calorimetric appliance was composed of a gas analyzer (to measure O_2 consumption and CO_2
153 production as VO_2 and VCO_2) and an activity recorder (locomotion and rearing). The
154 temperature of the calorimetric room was set to 22°C. Four were connected to each gas
155 analyzer, but each cage had specific inlets and outlets. A constant inlet flow (5 cm^3/min) was
156 maintained throughout the experiment. Gases were continuously analyzed with the following
157 sequence: 3 min from cage 1, 3 min from cage 2, 3 minutes from cage 3, 3 minutes from cage
158 4 and then 3 min from room air and thus the volume (mL/min) of O_2 consumed (VO_2) and
159 CO_2 produced (VCO_2) were measured for each mouse. Energy expenditure (EE) was
160 calculated as following ($\text{EE} = (16.3 \times \text{VO}_2 + 4.57 \times \text{VCO}_2)/60$ (watt)). Lipid (LOX) and
161 carbohydrate (GOX) oxidation were calculated according to the following equations: $\text{LOX} =$
162 $(1.69 \times \text{VO}_2 - 1.69 \times \text{VCO}_2) \times (9.46 \times 4.186/60)$ (watt) and $\text{GOX} = (4.57 \times \text{VCO}_2 -$
163 $3.23 \times \text{VO}_2) \times (3.74 \times 4.186/60)$ (watt)²⁷. Total activity was evaluated by summing
164 spontaneous activity and rearing activity, measured in indirect calorimetric cages
165 (Physiocage, Bioseb, Vitrolles, France), and was normalized to the control value.

166

167 ***Insulin Tolerance Test (ITT)***

168 One week before the end of the protocol, mice were subjected to ITT. Mice were subjected to
169 food restriction for 5h and stuffed with an insulin solution (0.05 UI/mL) prepared in saline

170 (0.5 UI/ kg body weight). Glycemia was measured from tail blood at 0, 15, 30, 60, 90, 120
171 minutes after injection (Accu-Check glucometer, Roche Diagnostic, Meylan, France).

172

173 *RNA extraction real time PCR and RNA sequencing*

174 Total RNA was extracted from retroperitoneal adipose tissue or hypothalamus using TRIzol
175 reagent (Thermo Fischer Scientific, Les Ulis, France). For the real time PCR, one µg of total
176 RNA from hypothalamus was used to synthesize cDNAs using random primers and Moloney
177 murine leukemia virus reverse transcriptase (Thermo Fischer Scientific, Les Ulis, France).
178 Real-time quantitative PCR analyses were performed using the Mx3005P Real-Time PCR
179 System (Stratagene, La Jolla, USA) as previously described²⁸. For each condition, expression
180 was quantified in duplicate, and 18S rRNA was used as the endogenous control in the
181 comparative cycle threshold (CT) method²⁹.

182 The sequences of the primers used in this study are reported in supplemental data
183 (Supplemental Table 1).

184 For the RNA sequencing, total RNA was isolated from 3 mice per group and was used for the
185 RNA-seq library preparation, using the kit TruSeq Stranded mRNA by Illumina.

186 Libraries were paired-end sequenced on the Illumina NextSeq 500 sequencer. Reads with a
187 phred score lower than 20 and shorter than 25 bp were removed using Sickle (v1,33). Quality
188 of trim reads were checked using multiQC (v1.0). Trim reads were aligned using STAR
189 aligner (v2.7.0d) with arguments “outFilterMismatchNoverLmax” and
190 “outFilterMultimapNmax” set to 0.08 and 1, respectively.

191 Transcripts discovery was performed using Cufflinks (v2.2.1) with the “library-type”
192 argument set to fr-firststrand, and a GTF file obtained from GENCODE (“Comprehensive gene
193 annotation”, vM1) provided as the genomic annotation. The GTF files produced for each
194 sample by Cufflinks were combined using Cuffmerge. The “class code” assigned to each
195 transcript by Cuffmerge was used to defined unknown transcripts (class code“u”). Only de
196 novo transcripts with counts greater than 0 in at least one RNA-seq sample were kept for
197 subsequent analyses. These de novo transcripts were combined with the GENCODE GTF file
198 to produce the final genomic annotation that was provided to FeatureCounts (v1.6.1) for
199 quantification.

200 Differential gene expression was performed using DESEQ2 between conditions. To create
201 bigwig files, reads from Watson and Crick strands were selected using SAMtools (v1.9) and
202 provided to the bam2wig.py script from the RseQC program suite (v2.6.4). RNA-seq profiles
203 were visualized using the IGV genome browser.

204

205 ***Ingenuity Pathway Analysis (IPA)***

206 Differential gene expression (with p-value adjusted < 0.05), obtained from the RNA seq
207 analysis between our conditions, were used in the IPA software to identify the canonical
208 pathways differentially impacted by the maternal diet.

209

210 ***Statistical analysis***

211 Data are expressed as mean \pm SEM. Significant differences were determined by unpaired
212 Student's *t* test or by ANOVA followed by the Fisher's LSD post hoc test using GraphPad
213 Prism. $p < 0.05$ was considered to be statistically significant.

214

215

216 **Results**

217

218 ***Maternal vitamin D deficiency affects post-weaning energy metabolism of offspring.***

219 Maternal VDD diet consumption prior and during gestation led to significant smaller post-
220 weaning (5 weeks old) body weight in male mice offspring (Fig.1A), that was non-significant
221 in female offspring (Fig.1H). The food consumption remained the same between CTRL and
222 VDD, either for males or females (Fig. 1B, 1I). In VDD males, spontaneous activity,
223 respiratory quotient (RQ) and energy expenditure (EE) were increased (Fig. 1C, D, E)
224 compared with the control group ($p < 0.05$). No difference of lipid oxidation (LOX) was
225 noticed in male (Fig. 1F), but carbohydrate oxidation (GOX) was increased in VDD males
226 (Fig. 1G). For female, despite the increased activity and RQ (Fig. 1J, K) of the VDD group,
227 there was no statistical difference for the EE (Fig. 1L). This was associated with decreased
228 LOX and increased GOX (Fig. 1M, N).

229

230 ***Maternal vitamin D deficiency combined with HF diet affects energy metabolism in adult***
231 ***offspring.***

232 To explore the combined effect of maternal VDD and post-weaning nutrition, mice were
233 submitted to low fat (LF) or high fat (HF) diet, leading thus to 8 groups (CTRL-LF, CTRL-
234 HF, VDD-LF and VDD-HF, for both male and female). Firstly, plasma 25(OH)D
235 concentration was measured in 6 weeks old offspring. Only VDD females presented
236 decreased 25(OH)D values ($108.6\text{nmol/L} \pm 2.852$) compared with the CTRL ($128.9\text{nmol/L} \pm$
237 6.186 ; $p < 0.01$) (Fig. 2A, B). During the diet period (from week 6 to 12), no statistical
238 difference between CTRL-LF and VDD-LF males body weight was observed (except at 9
239 weeks old, $p < 0.05$; Fig. 2C). Under HF diet, the body weight of CTRL males was gradually
240 higher than VDD males (significant from week 9 to 12; Fig. 2C). Concerning females, there
241 was non-significant differences between CTRL-LF and VDD-LF (from week 6 to 12). Under
242 HF diet, the CTRL females showed increased body weight compared with the VDD
243 (significant from week 9 to 12; Fig. 2D).

244 The energy metabolism was explored by indirect calorimetry (performed at 12-weeks of age).
245 No modification of the energy metabolism of VDD offspring (*i.e.* body weight, activity, RQ,
246 EE, LOX and GOX) was observed compared with the CTRL offspring when exposed to LF
247 diet in adulthood, either for male and female (Supplemental Figure 3). Under HF diet, energy
248 metabolism of VDD offspring was disturbed when compared with CTRL (Fig. 2). Males
249 exposed to VDD associated-HF diet displayed a smaller body weight than CTRL-HF ($p <$

250 0.05) and increased food consumption, EE and GOX ($p < 0.05$) (Fig. 2E, F, I, J). No
251 modification of activity, RQ and LOX was observed in males. Female exposed to VDD-HF
252 diets also exhibited smaller body weight than CTRL-HF (Fig. 2L). The food intake tended to
253 be increased (non-significant) in the VDD-HF compared with the CTRL-HF, whereas the
254 activity decreased and EE and GOX increased ($p < 0.05$) (Fig. 2M, N, P, R). No modification
255 of RQ and LOX was observed in females.

256 In order to explain the disruption of feeding behaviour in VDD group, we investigated the
257 expression of the genes of hypothalamic melanocortin pathway known to be involved in the
258 regulation of food intake³⁰. For males, exposed to LF diet in adulthood, agouti related-peptide
259 (AGRP) and melanocortin receptor type 4 (MC4R) were under-expressed in the VDD
260 compare to the CTRL (Supplemental Figure 4A). No statistical differences were observed in
261 HF condition (Supplemental Figure 4B). For females in VDD and CTRL-LF there was not
262 statistical difference (Supplemental Figure 4C). Under HF diet, Neuropeptide Y (NPY) and
263 AGRP were overexpressed ($p < 0.05$) and MC4R under-expressed in VDD-HF compare to
264 CTRL HF (Supplemental Figure 4D). There is no difference in POMC expression between
265 the different groups (Supplemental Figure 4A-D).

266 In order to explain energy metabolism sex discrepancies, the 17β -estradiol and testosterone
267 plasma concentrations were evaluated in all groups (Table 1). No significant modification was
268 observed in male groups (CTRL-LF, CTRL-HF, VDD-LF, VDD-HF) for 17β -estradiol and
269 testosterone. In females, no difference was observed in CTRL condition, whereas in VDD
270 condition, 17β -estradiol was strongly increased under HF diet (Table 1). No modification of
271 testosterone was observed.

272

273 ***Maternal VDD and HF diet-associated affect glucose homeostasis sex-specifically***

274 Biochemical analysis of male offspring highlighted that the glycemia tended to be higher in
275 both CTRL-HF and VDD-HF groups when compared with LF groups (performed at 13
276 weeks; Fig. 3A). Similar pattern was observed for plasma insulin, with an increased
277 concentration for the VDD-HF compared with the CTRL-HF ($p < 0.05$) (Fig. 3B). Thus, the
278 VDD-HF males presented the higher values of plasma insulin, glycemia and HOMA-IR when
279 compared with CTRL-HF, and to CTRL-LF and VDD-LF males (Fig. 3C, D).

280 When challenged with an HF diet, the glycemia of the VDD female tended to be decreased
281 (nonsignificant) compared with the CTRL-HF group to reach the same values as LF groups
282 (Fig. 3G), and plasma insulin of VDD female (VDD-HF) decreased compared with the
283 control group (CTRL-HF) ($p < 0.05$) to reach the same values as LF groups (both CTRL-LF

284 and VDD-HF) (Fig. 3H). The CTRL-HF females displayed higher values than the other three
285 groups regarding of combined plasma insulin, glycemia and HOMA-IR (Fig. 3I, J).
286 Insulin tolerance tests were undertaken in CTRL and VDD offspring (performed at 13 weeks).
287 No statistically differences were observed between LF groups, for both male and female (Fig.
288 3E, 3K). In obesogenic condition, the area under the curve (AUC) of VDD-HF was increased
289 compared with the CTRL HF group of males ($p < 0.01$) (Fig. 3F) and the AUC of VDD-HF
290 was decreased compared with the CTRL of females ($p < 0.05$) (Fig. 3L).

291

292 ***Maternal VDD and HF diet-associated modify morphological parameters and adipose***
293 ***tissue cellularity in sex-specific manner.***

294 After euthanasia, CTRL-LF and VDD-LF males showed similar body weight and weight gain;
295 similarly, to CTRL-HF and VDD-HF (Table 1). Significant differences were observed
296 between LF (both CTRL and VDD) and HF groups (both CTRL and VDD). Compare to the
297 three other groups, VDD-HF males presented the highest fat pad weight (*i.e.* perigonadal,
298 retroperitoneal, inguinal fat pad; Table 1), leading to increased adiposity index and leptin
299 plasma level (Fig. 4A, Table 1)). HF diet in CTRL females increased body weight, weight
300 gain (between weaning and the end of the protocol), fat pad weight, adiposity index and leptin
301 plasma level, whereas VDD-HF displayed similar parameters than LF groups (both CTRL-LF
302 and VDD-LF). HF diet (both CTRL and VDD) increased liver weight in male compare to the
303 LF (both CTRL and VDD). For females, only CTRL-HF increased liver weight compared
304 with other groups. No statistical difference of spleen's weight of males, but the females
305 exposed to HF diet showed increased spleen weight compare to the LF (Table 1).

306 The mean adipocyte area of CTRL and VDD males under LF diet were reduced compared
307 with the adipocytes area of males under HF diet (both CTRL and VDD; Fig. 4A, C and E).
308 The mean adipocyte area increased in CTRL-HF females, but was not different in VDD-HF
309 females, compared with LF groups (both CTRL-LF and VDD-LF; Fig. 4B, D and F).

310

311 ***Maternal and adult offspring diet modulate the expression of mRNA and associated***
312 ***canonical pathways in adipose tissue.***

313 To identify the impacts of maternal (CTRL, VDD) and adult diets (LF, HF) on the offspring
314 transcriptome, we performed RNA sequencing on visceral adipose tissue. Two set of data
315 were used to characterize the impact of the maternal diet and the adult diet. The first one to
316 study the impact of the HF diet on the same condition of maternal diet (*i.e.* CTRL-LF vs
317 CTRL-HF and VDD-LF vs VDD-HF). The second one to study maternal VDD on the same

318 condition of adult diet (*i.e.* CTRL-LF vs VDD-LF and CTRL-HF vs VDD-HF). We
319 highlighted the differential expression of transcripts between our conditions as established in
320 the Supplemental Table 2.

321 Using Ingenuity pathway analysis, we put forward canonical pathways with determinant z-
322 score and non-similar between conditions. We highlighted 16 differentially expressed
323 canonical pathways between CTRL-LF and CTRL-HF males. Interestingly, the triacylglycerol
324 biosynthesis, mitochondrial L-carnitine Shuttle Pathway, and fatty acid beta oxidation were
325 differentially decreased in CTRL-HF compared to CTRL-LF. 19 pathways were differentially
326 regulated between the VDD-LF and the VDD-HF males; including the oxidative
327 phosphorylation which was down-regulated in VDD-HF compared to VDD-LF. Also, 6
328 pathways seemed to be common between the offspring exposed to maternal CTRL diet or
329 VDD diet (Fig. 5).

330 When comparing CTRL-LF and VDD-LF males, 1 canonical pathway was expressed, while 3
331 pathways were expressed between CTRL-HF and VDD-HF males.

332 For the female born from CTRL mice, 31 canonical pathways were differentially expressed
333 between LF and HF, including the fatty acid beta oxidation, the oxidative phosphorylation and
334 triacylglycerol biosynthesis which decreased in CTRL-HF compared to CTRL-LF (Fig. 6).
335 No pathway was regulated when comparing VDD-LF to VDD-HF. Thus, when studying the
336 impact of LF diet between CTRL and VDD females, 3 canonical pathways were differentially
337 expressed, while 27 were differentially expressed for females on HF diet, including oxidative
338 phosphorylation, fatty acid beta oxidation, mitochondrial L-carnitine Shuttle pathway that
339 were induced in VDD-HF compared to CTRL-HF. Interestingly, estrogen biosynthesis
340 pathway appeared as induced in VDD-HF compared to CTRL-HF. One pathway was common
341 between LF and HF diets.

342

343

344

345 **Discussion**

346

347 The maternal diet is now well-established as a key player in the foetal development and long-
348 term effects programming in the offspring. In this study, we highlighted the impacts of
349 maternal VDD on offspring metabolism in normal nutritional condition and under HF
350 challenge. We also explored the sexual dimorphism of the metabolic response.

351 In agreement with previously published data, we reported here that maternal VDD diet
352 differentially affected the energy homeostasis and the body weight of the juvenile offspring (5
353 weeks old). Indeed, the maternal VDD was associated with reduction of body weight of the
354 male mice offspring in our experiment, similarly to previous reports in 15 days old mice²⁴ or
355 at weaning³¹. Other studies reported that VDD male offspring displayed higher body weight³²
356 or not differences³³⁻³⁵, but it is noteworthy that body weight evaluation have been evaluated in
357 adult rats^{32,33} or mice^{34,35}, not in juvenile. Interestingly, in our experiment, no discrepancy in
358 body weight was observed in female, demonstrating thus a strong metabolic sex-specific
359 response which has never been highlighted so far.

360 Such body weight modifications are strongly related to energy metabolism as highlight by
361 indirect calorimetry. Indeed, the increased spontaneous activity and energy expenditure
362 (through the carbohydrate oxidation) of the VDD males, together with the absence food intake
363 difference, could explain the smaller body weight of juvenile VDD males. The lack of body
364 weight impact in female may result from the absence of food intake modification in VDD
365 female. The observed increase in VDD female spontaneous activity could explain the increase
366 in carbohydrate oxidation, which was associated to a decrease in lipid oxidation, resulting in
367 overall no modification of energy expenditure. Interestingly, we observed that both male and
368 female VDD of the offspring were characterized by an increase in spontaneous activity.
369 Similar observations have already been reported in males VDD mice³⁶, and it has been
370 reported that in human, VDD during pregnancy is strongly correlated with the risk of attention
371 deficit and hyperactivity disorder in children³⁷. Nevertheless, in juvenile offspring of VDD
372 rats³⁸, no modification of locomotor activity was observed. The origin of this phenotype is
373 presently not well understood. Neurobehavioral development could explain this phenotype³⁹,
374 but this assumption will require further investigations.

375

376 In adulthood, morphological parameters (body weight, fat pad weight, liver weight, adipocyte
377 area) and energy balance were not modified in male and female offspring of VDD or CTRL
378 mice, when exposed to a control diet (VDD-LF and CTRL-LF), in agreement with previous

379 reports³⁵. All altered parameters in juvenile mice were normalized in adulthood, including
380 reduced body weight and energy expenditure, and increased spontaneous activity, for both
381 CTRL and VDD offspring. In offspring of VDD rats, similar results were reported in
382 adulthood, *i.e.* no effect on food intake, spontaneous activity in 14 weeks old male rats³².
383 Nevertheless, this study also reported differences on body weight and others biological
384 parameters (total cholesterol, triglycerides, HDL, blood glucose, that were higher in VDD
385 compared with CTRL offspring)³². The origin of these contradictory observations is not clear
386 but could be due to different compositions of control diets used, that could be sufficient to
387 unveil the metabolic phenotype associated to maternal VDD in offspring. In agreement with
388 this assumption, we highlighted that challenged offspring with an HF diets (45% of energy
389 from lipids from 9 to 12 weeks of age) led to a smaller body weight of VDD offspring
390 compared with the CTRL offspring for both males and females. Such observation is not fully
391 consistent with previous reports³¹, but could be related to the genetic background of the mice
392 which differ between studies. Nevertheless, our results in males are consistent with the fact
393 that the increased energy expenditure (mainly due to the carbohydrate oxidation), could
394 exceed the increased food intake, leading to smaller body weight of the VDD-HF males
395 compared with CTRL-HF males.

396 Female from VDD-HF diets also presented smaller body weight under HF, compared with LF
397 diet, with a tend to food intake increase. This limitation of body weight might be related to the
398 increase of energy expenditure (through the carbohydrate oxidation). An important issue that
399 remain presently unsolved is the drastic decrease of spontaneous activity in VDD-HF female
400 compared with CTRL-HF.

401 An interesting observation was that in VDD group the level of expression of the hypothalamic
402 melanocortin pathway genes, *i.e.* NPY, AgRP and MC4R, is different of the CTRL mice
403 level, and could explain at least in part food intake modulations. This suggest that VD is
404 important for the accurate development of the melanocortin pathway. In accordance with this
405 hypothesis, VD is known to be important for the brain development⁴⁰. However, further
406 investigations are needed to understand the impact of VD on the development of neural
407 pathways involved in feeding behaviour.

408 Beside body weight and energy metabolism, the glucose homeostasis was also investigated in
409 offspring adulthood. In LF condition, both in males and females, no modification of glucose
410 homeostasis was observed, especially glycemia, consistently with other studies^{24,32-34}.
411 Nevertheless, several studies reported that VDD in male rats led to an increase of fasting
412 insulin, HOMA-IR levels and insulin tolerance at 16 weeks³³, and in VDD male mice to an

413 increase of insulinemia³⁴. Such differences could be related to the model used (mice vs rats³³)
414 or duration of the protocol (14 weeks vs 6 months³⁴). Interestingly, in males submitted to HF
415 diet, we observed higher insulin plasma level, glycemia, HOMA IR and insulin resistance, as
416 previously reported in control conditions^{24,32-34}. Surprisingly, in females, no perturbation of
417 the glucose homeostasis was observed. At the opposite, VDD-HF females displayed similar
418 values as CTRL-LF or VDD-LF for several parameters, including insulinemia, glycemia and
419 HOMA-IR. Such observation has never been reported yet, since most of the studies included
420 only males.

421 Since insulin resistance is strongly linked to adiposity⁴¹, we investigated adiposity index of
422 animals, leptin plasma level and adipose tissue cellularity. Interestingly, adiposity index and
423 leptin plasma concentration were not impacted in males by the VDD in LF condition but was
424 strongly induced by the combination of VDD and HF, similarly to Belenchia et al.³¹ and
425 appeared to be strongly correlated with insulin resistance. No additive effect was reported for
426 adipose cellularity. In females, under LF diet, no modification was observed for adiposity
427 index, leptin plasma level and adipocytes area. As expected, the adiposity index, plasma leptin
428 and cellularity were induced by HF diet in offspring of CTRL mice (CTRL-HF), but
429 surprisingly, VDD-HF group displayed similar adiposity index, plasma leptin and adipocyte
430 area compared with LF groups (both CTRL-LF and VDD-LF). Such observation is fully
431 consistent with the improvement of insulin sensitivity observed in the VDD-HF group.
432 Nevertheless, such effect has never been reported yet and could correspond to an adaptative
433 mechanism based on an adequate diet supply of macronutrients and micronutrients, that
434 appears to be highly sex-specific.

435 To investigate the origin of such adiposity discrepancies, RNA-seq experiments were
436 undertaken on retroperitoneal adipose tissue. We hypothesized that differential transcriptomes
437 could be explain the variation of adiposity between group. In agreement, when comparing
438 male offspring from control mice exposed to LF or HF diet in adulthood (*i.e.* CTRL-LF vs
439 CTRL-HF), we observed that the “mitochondrial L-carnitine shuttle pathway” and the “fatty
440 acid beta oxidation” were down-regulated in the CTRL-HF, which was associated to increase
441 of adiposity. In males from VDD mice submitted to LF or HF (*i.e.* VDD-LF vs VDD-HF),
442 those canonical pathways related to lipid catabolism were not regulated, but we observed a
443 strong repression of oxidative phosphorylation which correlated with fat pad accretion, weight
444 gain and a higher adiposity index. Thus, the discrepancy between offspring of CTRL or VDD
445 mice could be related to the ability to induce lipid oxidation/oxidative phosphorylation in
446 adipose tissue. In addition, the transcriptomic response to maternal diet combined to

447 adulthood diet (*i.e.* CTRL-LF vs VDD-LF and CTRL-HF vs VDD-HF) did not results in
448 major differences in term of lipid metabolism pathways, nor major difference in adiposity,
449 which reinforces the putative role of adipose tissue lipid metabolism modulations to drive
450 adiposity phenotype. It is noteworthy that the contribution of white adipose tissue to whole
451 body energy expenditure is considered relatively small. Nevertheless, there are examples of
452 nutritional and pharmacological interventions in animals resulting in obesity resistance
453 associated with increased oxidative capacity in WAT⁴²⁻⁴⁴. In agreement with the modest
454 contribution of white adipose tissue on the whole energy expenditure , we did not lipid
455 oxidation induction by indirect calorimetry, suggesting that such local induction of lipid
456 oxidation may have beneficial effect in terms of adiposity, but not impact the overall lipid
457 oxidation which involve many other organs.

458 Concerning females, CTRL-HF compared with CTRL-LF repressed several canonical
459 pathways linked to lipid metabolism (*i.e.* fatty acid, beta oxidation, oxidative phosphorylation,
460 triacyl glycerol biosynthesis, mitochondrial l-carnitine shuttle pathway). Similarly to males,
461 such pattern corresponded to a signature of body weight gain and adiposity. Importantly,
462 when comparing VDD-HF to VDD-LF, we did not observe any modification of metabolic
463 canonical pathways, nor modification of adiposity. In addition, the transcriptomic response to
464 maternal diet combined to adulthood diet (*i.e.* CTRL-LF vs VDD-LF and CTRL-HF vs VDD-
465 HF) results in the induction of canonical pathways implicated in lipid metabolism (*i.e.* beta
466 oxidation, oxidative phosphorylation, triacyl glycerol biosynthesis, mitochondrial l-carnitine
467 shuttle pathway) in VDD-HF compared to CTRL-HF. This was in agreement with the
468 decreased of adiposity index, and body weight observed in VDD-HF compared to CTRL-HF.
469 Altogether these observations suggested that adiposity is strongly associated to lipid
470 metabolism pathway that may influence adipose tissue accretion. The origin of such gene
471 expression profiles modifications is presently not established but we speculate that epigenetic
472 mechanisms could link maternal VDD to long-term transcriptional modifications in
473 offspring's adipose tissue. Such assumption will require further investigations.

474 It is obvious that the adiposity and gene expression profile are strongly sex-specific, and
475 notably the effect of VDD maternal under HF diet which leads to highly divergent adiposity
476 in males and females. Sex-specific metabolic discrepancies and notably adiposity are strongly
477 related to estradiol status⁴⁵. Indeed, it is well-established that estrogens promotes
478 subcutaneous fat accumulation⁴⁶ and improve glucose homeostasis⁴⁷. Consequently, the
479 decrease of estrogens associated to menopause is linked to an increase in visceral fat and
480 greater risk for the metabolic syndrome in postmenopausal compared with premenopausal

481 women⁴⁵. Thus, we focussed on 17 β -estradiol plasma level to explain observed phenotypes.
482 Interestingly, we noticed that in males, no significant differences of plasma concentrations
483 were observed, suggesting that estradiol by itself is not a driver element. It is important to
484 keep in mind that estradiol plasma level is not the sole element that impact adipose tissue
485 biology, and estrogen receptors are also important⁴⁵ and would deserve attention in further
486 experiments. Nevertheless, in females, when comparing CTRL-LF to CTRL-HF, 17 β -
487 estradiol plasma level tended to increase but did not reach statistical significance, whereas in
488 VDD mice (VDD-LF vs VDD-HF), the plasma level of 17 β -estradiol was strongly and
489 significantly induced. This is an important point since 17 β -estradiol is well-known to induce
490 metabolic catabolism, leading to weight gain limitation and glucose homeostasis
491 improvement^{45,47}, and such increase of 17 β -estradiol could explain by itself the metabolic
492 improvement observed in VDD-HF female mice. The origin of the induction of 17 β -estradiol
493 is presently unknown, even if epigenetic mechanism are suspected to be involved. Further
494 investigations will be mandatory to explain this observation.
495 To conclude, our study brings new informations on the impact of VDD in metabolic
496 disruption in offspring and notably its predisposition to long term metabolic health
497 complications. Importantly it sheds light on the sex-specific adipose tissue / adiposity
498 response, that need to be taking into account in terms of public health.

499

500

501

502

503 **Funding.** The work has been funded by grants from INRA, INSERM, AMU and the
504 Fondation de France. High throughput sequencing was performed at the TGML Platform,
505 supported by grants from Inserm, GIS IBiSA, Aix-Marseille Université, and ANR-10-INBS-
506 0009-10

507

508

509 **Conflict of interest.** No conflict of interest as to be disclose.

510

511 **Author Contributions:** JF Landrier and E Seipelt designed research; JF Landrier, E Seipelt,
512 L Mounien, B Loriod, M Puceat analysed data; E Seipelt, F Tourniaire, C Couturier, J Astier,
513 H Vachon performed research; JF Landrier, E Seipelt, L Mounien wrote the paper.

514

515 **Figures legends**

516

517 **Fig 1. Indirect calorimetry on juvenile offspring during 24hours to compare the**
518 **energetic metabolism of the VDD and the CTRL.** Parameters measured during 24 h for
519 males and females were Body weight (A, H), Food intake (B, I), spontaneous activity (C, J),
520 Respiratory quotient (RQ, D, K), Energy expenditure (EE, E, L), Lipid oxidation (LOX, F, M)
521 and carbohydrate oxidation (GOX, G, N). Values are presented as mean \pm SEM. Bars not
522 sharing the same letter were significantly different in Fisher's LSD post hoc test. $p < 0.05$

523

524 **Fig 2. Morpho-metabolic follow-up of juvenile and 12 weeks-old offspring.** (A, B)
525 25OH(D) concentration of 5-weeks-old offspring male (A) and female (B) before the adult
526 diet induced (Low fat LF or HF, High fat diet). Growing curves of males (C) and females (D)
527 from the beginning of hf diet to indirect calorimetry at 12-weeks-old. Indirect calorimetry of
528 adults (E-R) offspring during 24hours to compare the energetic metabolism of the VDD and
529 the CTRL on HF diet. Parameters measured during 24 h for males and females were Body
530 weight (E, L), Food intake (F, M), activity (G, N), Respiratory quotient (RQ, H, O), Energy
531 expenditure (EE, LP), Lipid oxidation (LOX, J, Q) and carbohydrate oxidation (GOX, K, R).
532 Values are presented as mean \pm SEM. Bars not sharing the same letter were significantly
533 different in Fisher's LSD post hoc test. $p < 0.05$

534

535 **Fig 3. Glucose homeostasis of 13-weeks old offspring.** Measured of glucose from tail blood
536 of males (A) and females (G). Insulinemia of male (B) and females (H), Ratio glucose/insulin
537 for males (C) and females (F), HOMA-IR for males (D) and females (J). Insulin tolerance test
538 for males on LF diet (E) and HF diet (F), also for females on LF diet (K), and HF diet (L).
539 Values are presented as mean \pm SEM. Bars not sharing the same letter were significantly
540 different in Fisher's LSD post hoc test. $p < 0.05$

541

542 **Fig 4. Adiposity index and adipose tissue histology of offspring.** Adiposity index of the
543 offspring has been established for males (A) and females (B). Representative histological
544 images of visceral adipose tissue of males (E) and females (F) adult offspring after eosin-
545 hematoxylin coloration (10X magnification). Mean adipocyte area determined using Image J
546 software for males (C) and females (D).. Values are presented as mean \pm SEM. Bars not
547 sharing the same letter were significantly different in Fisher's LSD post hoc test. $p < 0.05$

548

549 **Fig 5. Overview of canonical pathway differentially** expressed on visceral adipose tissue of
550 the male offspring.

551

552 **Fig 6. Overview of canonical pathway differentially** expressed on visceral adipose tissue of
553 the female offspring.

554

555 **Table 1.** General morphological parameters and 17 β -estradiol plasma concentrations obtain
556 from the offspring at the protocol end. Values are presented as mean \pm SEM. Bars not sharing
557 the same letter were significantly different in Fisher's LSD post hoc test. $p < 0.05$

558

559

560

561

562

563 **References**

564

- 565 1. Bendik I, Friedel A, Roos FF, Weber P, Eggersdorfer M. Vitamin D: A critical and
566 essential micronutrient for human health. *Front Physiol.* 2014;5 JUL(July):1-14.
567 doi:10.3389/fphys.2014.00248
- 568 2. Bonnet L, Karkeni E, Couturier C, et al. Gene Expression Pattern in Response to
569 Cholecalciferol Supplementation Highlights Cubilin as a Major Protein of 25(OH)D
570 Uptake in Adipocytes and Male Mice White Adipose Tissue. *Endocrinology.*
571 2018;159(2):957-966. doi:10.1210/en.2017-00650
- 572 3. Karkeni E, Bonnet L, Marcotorchino J, et al. Vitamin D limits inflammation-linked
573 microRNA expression in adipocytes in vitro and in vivo: A new mechanism for the
574 regulation of inflammation by vitamin D. *Epigenetics.* 2018;13(2):156-162.
575 doi:10.1080/15592294.2016.1276681
- 576 4. Karkeni E, Marcotorchino J, Tourniaire F, et al. Vitamin D Limits Chemokine
577 Expression in Adipocytes and Macrophage Migration In Vitro and in Male Mice.
578 *Endocrinology.* 2015;156(5):1782-1793. doi:10.1210/en.2014-1647
- 579 5. Marcotorchino J, Gouranton E, Romier B, et al. Vitamin D reduces the inflammatory
580 response and restores glucose uptake in adipocytes. *Mol Nutr Food Res.*
581 2012;56(12):1771-1782. doi:10.1002/mnfr.201200383
- 582 6. Marcotorchino J, Tourniaire F, Astier J, et al. Vitamin D protects against diet-induced
583 obesity by enhancing fatty acid oxidation. *J Nutr Biochem.* 2014;25(10):1077-1083.
584 doi:10.1016/j.jnutbio.2014.05.010
- 585 7. Landrier J-F, Karkeni E, Marcotorchino J, Bonnet L, Tourniaire F. Vitamin D
586 modulates adipose tissue biology: possible consequences for obesity? *Proc Nutr Soc.*
587 2015;25(March):1-9. doi:10.1017/S0029665115004164
- 588 8. Bonnet L, Hachemi MA, Karkeni E, et al. Diet induced obesity modifies vitamin D
589 metabolism and adipose tissue storage in mice. *J Steroid Biochem Mol Biol.*
590 2019;185(May 2018):39-46. doi:10.1016/j.jsbmb.2018.07.006
- 591 9. Landrier J, Mounien L, Tourniaire F. Obesity and Vitamin D Metabolism
592 Modifications. *J Bone Miner Res.* 2019;34(7):1383-1383. doi:10.1002/jbmr.3739
- 593 10. Borg SA, Buckley H, Owen R, et al. Early life Vitamin D depletion alters the postnatal
594 response to skeletal loading in growing and mature bone. *PLoS One.* 2018;13(1):1-17.
595 doi:10.1371/journal.pone.0190675
- 596 11. Turck D, Bresson J, Burlingame B, et al. Dietary reference values for vitamin D. *EFSA*

- 597 *J.* 2016;14(10):e04547. doi:10.2903/j.efsa.2016.4547
- 598 12. Palacios C, Gonzalez L. Is vitamin D deficiency a major global public health problem?
599 *J Steroid Biochem Mol Biol.* 2014;144 Pt A(787):138-145.
600 doi:10.1016/j.jsbmb.2013.11.003
- 601 13. Cashman KD, Dowling KG, Gonzalez-Gross M, et al. Vitamin D deficiency in Europe:
602 pandemic? *Am J Clin Nutr.* 2016;(C):1-12. doi:10.3945/ajcn.115.120873.
- 603 14. Saraf R, Morton SMB, Camargo CA, Grant CC. Global summary of maternal and
604 newborn vitamin D status - a systematic review. *Matern Child Nutr.* 2016;12(4):647-
605 668. doi:10.1111/mcn.12210
- 606 15. Miliku K, Blanken LME, Gaillard R, et al. Maternal Vitamin D concentrations during
607 pregnancy, fetal growth patterns, and risks of adverse birth outcomes. *Am J Clin Nutr.*
608 2016;103(6):1514-1522. doi:10.3945/ajcn.115.123752.1514
- 609 16. Achkar M. Vitamin D status in early pregnancy and risk of preeclampsia. *Am J Obstet*
610 *Gynecol.* 2016;212(4):1-14. doi:10.1016/j.ajog.2014.11.009.Vitamin
- 611 17. Amraei M, Mohamadpour S, Sayehmiri K, Mousavi SF, Shirzadpour E, Moayeri A.
612 Effects of vitamin D deficiency on incidence risk of gestational diabetes mellitus: A
613 systematic review and meta-analysis. *Front Endocrinol (Lausanne).* 2018;9(FEB):1-11.
614 doi:10.3389/fendo.2018.00007
- 615 18. Gernand AD, Simhan HN, Klebanoff MA, Bodnar LM. Maternal serum 25-
616 hydroxyvitamin D and measures of newborn and placental weight in a U.S. multicenter
617 cohort study. *J Clin Endocrinol Metab.* 2013;98(1):398-404. doi:10.1210/jc.2012-3275
- 618 19. Wang H, Xiao Y, Zhang L, Gao Q. Journal of Steroid Biochemistry and Molecular
619 Biology Maternal early pregnancy vitamin D status in relation to low birth weight and
620 small-for-gestational-age offspring. *J Steroid Biochem Mol Biol.* 2017;(157):0-1.
621 doi:10.1016/j.jsbmb.2017.09.010
- 622 20. Morales E, Rodriguez A, Valvi D, et al. Deficit of vitamin D in pregnancy and growth
623 and overweight in the offspring. *Int J Obes.* 2015;39(1):61-68.
624 doi:10.1038/ijo.2014.165
- 625 21. Crozier SR, Harvey NC, Inskip HM, Godfrey KM. Maternal vitamin D status in
626 pregnancy is associated with adiposity in the offspring : prospective observational
627 study. *Am J Clin Nutr.* 2012;96(1):57-63. doi:10.3945/ajcn.112.037473.Maternal
- 628 22. Daraki V, Roumeliotaki T, Chalkiadaki G, et al. Low maternal vitamin D status in
629 pregnancy increases the risk of childhood obesity. *Pediatr Obes.* 2018;13(8):467-475.
630 doi:10.1111/ijpo.12267

- 631 23. Rytter D, Bech BH, Halldorsson TI, et al. Maternal Vitamin D status at week 30 of
632 gestation and offspring cardio-metabolic health at 20 years: A prospective cohort study
633 over two decades. *PLoS One*. 2016;11(10):1-12. doi:10.1371/journal.pone.0164758
- 634 24. Reichetzeder C, Chen H, Föller M, et al. Maternal Vitamin D Deficiency and Fetal
635 Programming - Lessons Learned from Humans and Mice. *Kidney Blood Press Res*.
636 2014;39(4):315-329. doi:10.1159/000355809
- 637 25. Ideraabdullah FY, Belenchia AM, Rosenfeld CS, et al. Maternal vitamin D deficiency
638 and developmental origins of health and disease (DOHaD). *J Endocrinol*.
639 2019;241(2):R65-R80. doi:10.1530/joe-18-0541
- 640 26. Goyal D, Limesand SW, Goyal R. Epigenetic responses and the developmental origins
641 of health and disease. *J Endocrinol*. 2019;242(1):T105-T119. doi:10.1530/JOE-19-
642 0009
- 643 27. Even PC, Mokhtarian A, Pele A. Practical aspects of indirect calorimetry in laboratory
644 animals. *Neurosci Biobehav Rev*. 1994;18(3):435-447.
- 645 28. Fenni S, Hammou H, Astier J, et al. Lycopene and tomato powder supplementation
646 similarly inhibit high-fat diet induced obesity, inflammatory response, and associated
647 metabolic disorders. *Mol Nutr Food Res*. 2017;61(9):1601083.
648 doi:10.1002/mnfr.201601083
- 649 29. Livak KJ, Schmittgen TD. Analysis of Relative Gene Expression Data Using Real-
650 Time Quantitative PCR and the $2^{-\Delta\Delta CT}$ Method. *Methods*. 2001;25(4):402-408.
651 doi:10.1006/meth.2001.1262
- 652 30. Derghal A, Djelloul M, Trouslard J, Mounien L. The Role of MicroRNA in the
653 Modulation of the Melanocortinergic System. *Front Neurosci*. 2017;11(APR):1-8.
654 doi:10.3389/fnins.2017.00181
- 655 31. Belenchia AM, Johnson SA, Eilersieck MR, Rosenfeld CS, Peterson CA. In utero
656 vitamin D deficiency predisposes offspring to long-term adverse adipose tissue effects.
657 *J Endocrinol*. 2017;234(3):301-313. doi:10.1530/JOE-17-0015
- 658 32. Wen J, Hong Q, Wang X, et al. The effect of maternal vitamin D deficiency during
659 pregnancy on body fat and adipogenesis in rat offspring. *Sci Rep*. 2018;8(1):365.
660 doi:10.1038/s41598-017-18770-4
- 661 33. Zhang H, Chu X, Huang Y, et al. Maternal vitamin D deficiency during pregnancy
662 results in insulin resistance in rat offspring, which is associated with inflammation and
663 Ikb α methylation. *Diabetologia*. 2014;57(10):2165-2172. doi:10.1007/s00125-014-
664 3316-7

- 665 34. Nascimento FAM, Ceciliano TC, Aguila MB, Mandarim-de-lacerda CA.
666 Transgenerational Effects on the Liver and Pancreas Resulting from Maternal Vitamin
667 D Restriction in Mice. *J Nutr Sci Vitaminol (Tokyo)*. 2013;59(5):367-374.
668 <http://jlc.jst.go.jp/DN/JST.JSTAGE/jnsv/59.367?lang=en&from=CrossRef&type=abstract>.
669 act.
- 670 35. Belenchia AM, Jones KL, Will M, et al. Maternal vitamin D deficiency during
671 pregnancy affects expression of adipogenic-regulating genes peroxisome proliferator-
672 activated receptor gamma (PPAR γ) and vitamin D receptor (VDR) in lean male mice
673 offspring. *Eur J Nutr*. 2018;57(2):723-730. doi:10.1007/s00394-016-1359-x
- 674 36. Fu L, Chen Y-H, Chen X, Xu S, Yu Z, Xu D-X. Vitamin D deficiency impairs
675 neurobehavioral development in male mice. *Physiol Behav*. 2017;179(July):333-339.
676 doi:10.1016/j.physbeh.2017.07.017
- 677 37. Morales E, Julvez J, Torrent M, et al. Vitamin D in Pregnancy and Attention Deficit
678 Hyperactivity Disorder-like Symptoms in Childhood. *Epidemiology*. 2015;26(4):458-
679 465. doi:10.1097/EDE.0000000000000292
- 680 38. Pan P, Jin DHS, Chatterjee-Chakraborty M, et al. The effects of vitamin D3 during
681 pregnancy and lactation on offspring physiology and behavior in Sprague-Dawley rats.
682 *Dev Psychobiol*. 2014;56(1):12-22. doi:10.1002/dev.21086
- 683 39. Hawes JE, Tesic D, Whitehouse AJ, Zosky GR, Smith JT, Wyrwoll CS. Maternal
684 vitamin D deficiency alters fetal brain development in the BALB/c mouse. *Behav Brain*
685 *Res*. 2015;286:192-200. doi:10.1016/j.bbr.2015.03.008
- 686 40. Eyles D, McGrath J. Vitamin D Brain Development and Function. In: *Vitamin D*. Vol
687 1. Fourth Edi. Elsevier; 2018:563-581. doi:10.1016/B978-0-12-809965-0.00033-1
- 688 41. Rosen ED, Spiegelman BM. Adipocytes as regulators of energy balance and glucose
689 homeostasis. *Nature*. 2006;444(7121):847-853. doi:10.1038/nature05483
- 690 42. Kusminski CM, Scherer PE. Mitochondrial dysfunction in white adipose tissue. *Trends*
691 *Endocrinol Metab*. 2012;23(9):435-443. doi:10.1016/j.tem.2012.06.004
- 692 43. Tourniaire F, Musinovic H, Gouranton E, et al. All- trans retinoic acid induces
693 oxidative phosphorylation and mitochondria biogenesis in adipocytes. *J Lipid Res*.
694 2015;56(6):1100-1109. doi:10.1194/jlr.M053652
- 695 44. Flachs P, Rossmeisl M, Kuda O, Kopecky J. Stimulation of mitochondrial oxidative
696 capacity in white fat independent of UCP1: A key to lean phenotype. *Biochim Biophys*
697 *Acta - Mol Cell Biol Lipids*. 2013;1831(5):986-1003. doi:10.1016/j.bbailip.2013.02.003
- 698 45. Brown L, Clegg D. Central effects of estradiol in the regulation of food intake, body

- 699 weight, and adiposity. *J Steroid Biochem Mol Biol*. 2010;122(1-3):65-73.
700 doi:10.1016/j.jsbmb.2009.12.005
- 701 46. Krotkiewski M, Bjorntorp P, Sjostrom L, Smith U. Impact of obesity on metabolism in
702 men and women. Importance of regional adipose tissue distribution. *J Clin Invest*.
703 1983;72(3):1150-1162.
- 704 47. Riant E, Waget A, Cogo H, Arnal J-F, Burcelin R, Gourdy P. Estrogens Protect against
705 High-Fat Diet-Induced Insulin Resistance and Glucose Intolerance in Mice.
706 *Endocrinology*. 2009;150(5):2109-2117. doi:10.1210/en.2008-0971
707
708